National Institute for Health and Care Excellence

Draft for consultation

Type 2 diabetes in adults: management (medicines update)

[F1.1] Evidence reviews for subsequent pharmacological management of type 2 diabetes: 1.1.1 to 1.1.6

NICE guideline GID-NG10336

Evidence reviews underpinning recommendations 1.8.6-1.8.32, 1.8.34, 1.8.38-1.8.60 and recommendations for research in the NICE guideline

August 2025

Draft for Consultation

This evidence review was developed by NICE



Disclaimer

The recommendations in this guideline represent the view of NICE, arrived at after careful consideration of the evidence available. When exercising their judgement, professionals are expected to take this guideline fully into account, alongside the individual needs, preferences and values of their patients or service users. The recommendations in this guideline are not mandatory and the guideline does not override the responsibility of healthcare professionals to make decisions appropriate to the circumstances of the individual patient, in consultation with the patient and/or their carer or guardian.

Local commissioners and/or providers have a responsibility to enable the guideline to be applied when individual health professionals and their patients or service users wish to use it. They should do so in the context of local and national priorities for funding and developing services, and in light of their duties to have due regard to the need to eliminate unlawful discrimination, to advance equality of opportunity and to reduce health inequalities. Nothing in this guideline should be interpreted in a way that would be inconsistent with compliance with those duties.

NICE guidelines cover health and care in England. Decisions on how they apply in other UK countries are made by ministers in the <u>Welsh Government</u>, <u>Scottish Government</u>, and <u>Northern Ireland Executive</u>. All NICE guidance is subject to regular review and may be updated or withdrawn.

Copyright

© NICE 2025. All rights reserved. Subject to Notice of Rights.

ISBN:

Contents

| 1. | Sub | sequen | t pharmacological management | 5 |
|----|------|--------|---|-----|
| | 1.1. | Revie | w question | 5 |
| | | 1.1.1. | Introduction | 5 |
| | | 1.1.2. | Summary of the protocol | 5 |
| | | 1.1.3. | Methods and process | 8 |
| | | 1.1.4. | Effectiveness evidence | 9 |
| | | 1.1.5. | Summary of studies included in the effectiveness evidence | 17 |
| | | 1.1.6. | Summary of the effectiveness evidence (network meta-analysis) – combined strategies | 261 |
| | | 117 | References | 266 |

1. Subsequent pharmacological management

1.1. Review question

- 4 For different population subgroups, which individual and/or combinations of pharmacological
- 5 therapies are most clinically and cost effective as subsequent treatment for the management
- 6 of type 2 diabetes?

1

2

3

7

1.1.1. Introduction

- 8 Type 2 diabetes is a chronic metabolic condition characterised by insulin resistance (that is,
- 9 the body's inability to effectively use insulin) and insufficient pancreatic insulin production,
- 10 resulting in high blood glucose levels (hyperglycaemia). The consequences of this include
- 11 macrovascular complications (such as myocardial infarction, stroke and heart failure),
- microvascular complications (such as chronic kidney disease, retinopathy, neuropathy and
- 13 sexual problems), acute complications (such as hyper- and hypoglycaemia, diabetic
- 14 ketoacidosis and hyperosmolar hyperglycaemic state) and other complications (such as gum
- disease, increased risk of pancreatitis, cancer, polycystic ovary syndrome and other
- 16 conditions). There are approximately 5.6 million people living with diabetes in the UK, 90% of
- 17 those having type 2 diabetes and the incidence rises each year. The condition accounts for
- 18 10% of NHS annual budget with almost 80% of that being spent on managing the
- 19 complications of type 2 diabetes.
- 20 The NICE guideline on Type 2 diabetes in adults: management was last updated in 2022
- 21 (NG28) where the focus was on cardiovascular impact. In this update we examine the holistic
- benefits of pharmacological therapy for type 2 diabetes to understand the effects of
- treatments on a range of factors including quality of life, cardiovascular and renal protection,
- 24 weight management, other adverse effects (such as arrhythmias, falls and liver disease) and
- 25 glycaemic control. This considers a wide range of trials and focusses on specific
- subpopulations of interest within the population with type 2 diabetes: people with type 2
- 27 diabetes and heart failure, people with type 2 diabetes and atherosclerotic cardiovascular
- disease, people with type 2 diabetes and chronic kidney disease and people with type 2
- diabetes and different levels of risk of developing cardiovascular disease in the future. This
- 30 will allow for clinical and cost-effectiveness evidence to be identified, considered and
- 31 modelled to allow a comprehensive assessment of the effects of these treatments. The
- 32 subsequent treatment review considers trials where previous medication has been provided
- to all people and any medication is either being added to this treatment or switched with this
- 34 treatment.

35

36

1.1.2. Summary of the protocol

Table 1: PICO characteristics of review question

| Population | Adults (age ≥18 years) with type 2 diabetes mellitus who are currently receiving antidiabetic medication |
|------------|--|
| | The population will be stratified into different groups for the analysis, these include: |
| | People with type 2 diabetes mellitus and heart failure |

- People with type 2 diabetes mellitus and atherosclerotic cardiovascular disease
- People with type 2 diabetes mellitus and chronic kidney disease

- People with type 2 diabetes mellitus and low cardiovascular risk with no other comorbidities
- People with type 2 diabetes mellitus and high cardiovascular risk (or mixed/unclear cardiovascular risk) with no other comorbidities

A stratum where all groups were analysed together was not included as the committee agreed this would not add any value.

Exclusion:

- Children and young people (age <18 years) with type 2 diabetes mellitus
- Pregnant people with type 2 diabetes mellitus
- People with type 1 diabetes mellitus
- People with type 2 diabetes mellitus who are hyperglycaemic and require rescue treatment

Interventions

Pharmacological therapies for people with type 2 diabetes.

- All therapies will be examined on an individual drug level (rather than a class level).
- All doses will be pooled together.

Different strategies to optimise treatment (stratify trials by the strategy used in the trial):

- · Adding a new treatment
- Stopping a previous treatment
- · Switching to a different treatment
- Biguanides
 - Metformin hydrochloride standard release
 - o Metformin hydrochloride slow release
- DPP-4 inhibitors
 - Alogliptin (Vipidia)
 - Linagliptin (Trajenta)
 - Saxagliptin (Onglyza)
 - Sitagliptin (Januvia)
 - o Vildagliptin (Galvus)
- GLP-1 receptor agonist
 - o Dulaglutide (Trulicity)
 - Exenatide (Byetta)
 - Liraglutide (Victoza)
 - o Lixisenatide (Lyxumia)
 - o Semaglutide (Rybelsus, Ozempic)
- Dual GIP/GLP-1 receptor co-agonists
 - Tirzepatide (Mounjaro)
- SGLT2 inhibitors
 - Canagliflozin (Invokana)
 - o Dapagliflozin (Forxiga)
 - o Empagliflozin (Jardiance)
 - o Ertugliflozin (Steglatro)
- Sulfonylureas

| | o Gliclazideo Glimepiride |
|-------------|--|
| | ∘ Glipizide |
| | ⊙ Tolbutamide |
| | Thiazolidinediones |
| | o Pioglitazone |
| | Combinations of therapies listed above (combinations may include medicines being given separately or combination products) |
| Comparisons | Different strategies to optimise treatment |
| | Different pharmacological therapies listed in the intervention section to each other |
| | An oral formulation compared with an injectable formulation of the same medication |
| | Insulin (all types and doses pooled together in the same drug class) |
| | Placebo |
| Outcomes | Outcomes will be extracted in this review for inclusion in the review. The final time point (end point of the trial) reported will be extracted and used in the analysis where possible. |
| | Health-related quality of life (continuous outcomes): |
| | All-cause mortality (time-to-event/dichotomous outcome) |
| | Cardiovascular mortality (time-to-event/dichotomous outcome) |
| | Major Cardiovascular Events (MACE) (where multiple MACE values are reported [for example: 3-item MACE and 4-item MACE], the highest number MACE value will be prioritised) (time-to-event/dichotomous outcome) |
| | o 3-item MACE |
| | o 4-item MACE |
| | ○ 5-item MACE |
| | Events making up MACE (not previously stated) (time-to-event/dichotomous outcomes): |
| | ○ Non-fatal stroke |
| | o Non-fatal myocardial infarction |
| | o Unstable angina |
| | Hospitalisation for heart failure |
| | Renal events (time-to-event/dichotomous outcome): |
| | Acute kidney injury |
| | Persistent signs of worsening kidney disease (including doubling of serum creatinine) |
| | Development of end stage kidney disease (including need for renal replacement therapy and transplant) |
| | ○ Death from renal cause |
| | Serious adverse events (time-to-event/dichotomous outcome): |
| | ○ Cardiac arrhythmia (including atrial fibrillation) |
| | o Diabetic ketoacidosis |
| | o Falls requiring hospitalisation |

| | Progression of liver disease (to non-alcoholic fatty liver disease, to fibrosis, to cirrhosis, to end stage liver disease) (time-to-event/dichotomous outcome) |
|--------------|--|
| | Remission (time-to-event/dichotomous outcome) |
| | Acute diabetic complications (time-to-event/dichotomous outcome): |
| | Hypoglycaemia episodes |
| | At night hypoglycaemic episodes |
| | Severe hypoglycaemic episodes |
| | Continuous outcomes: |
| | HbA1c change (absolute change scores prioritised over percentage change scores) |
| | ○ Weight change |
| | ○ BMI change |
| Study design | Systematic reviews of randomised-controlled trials and randomised-controlled |
| | trials Published network meta-analyses and individual patient data analyses were considered for inclusion. |

1 For full details see the review protocol in report F2, appendix A.

1.1.3. Methods and process

- This evidence review was developed using the methods and process described in Developing NICE guidelines: the manual. Methods specific to this review question are described in the review protocol in report F2, appendix A and the methods document.
- 6 Declarations of interest were recorded according to NICE's conflicts of interest policy.

2

3

4

5

1.1.4. Effectiveness evidence

1.1.4.1. Included studies

1

2

- 3 Fifteen studies^{17, 50, 57, 136, 164, 232, 235, 236, 249, 288, 293, 334, 391, 396, 423} were included in the evidence for
- 4 population model 1 (people with type 2 diabetes and heart failure). All of these trials
- 5 compared an 'adding' strategy, examining the effect of adding an intervention drug to other
- 6 glucose-lowering drugs. Fourteen comparisons were identified. For all the models, studies
- 7 were only included if they reported outcomes at follow-up times of 24 weeks or longer.
- 8 Where multiple timepoints were reported, the longest timepoint was extracted. For population
- 9 model 1, the extracted timepoints ranged between 6 and 50 months.
- Twenty-five studies^{3, 17, 50, 52, 57, 78, 129, 132, 136, 164, 212, 213, 215, 232, 272, 275, 293, 296, 333, 345, 375, 391, 392, 396, 423}
- were included in the evidence for population model 2 (people with type 2 diabetes and
- 12 atherosclerotic CVD). All of these trials compared an 'adding' strategy, examining the effect
- of adding an intervention drug to other glucose-lowering drugs. Twenty-one comparisons
- were identified, and the extracted follow-up time ranged between 6 and 76 months.
- Twenty-seven studies^{29, 50, 70, 102, 118, 138, 140, 161, 199, 201, 205, 248, 254, 257, 274, 288, 289, 299, 308, 334, 358, 368, 380,}
- 16 387, 396, 404, 423 were included in the evidence for population model 3 (people with type 2
- diabetes and CKD). Most of these trials compared an 'adding' strategy, examining the effect
- of adding an intervention drug to other glucose-lowering drugs. No trials were identified that
- 19 examined a 'stopping' strategy. Twenty-one comparisons were identified, and the extracted
- 20 follow-up time ranged between 5.5 and 60 months.
- 21 No evidence was identified for population model 5 (people with type 2 diabetes and lower
- 22 risk for cardiovascular disease).
- 23 Three hundred and seventystudies^{1, 2, 4-28, 30-49, 51, 53-56, 58-69, 71-77, 79-101, 103-117, 119-123, 126-131, 133-135,}
- 24 137, 139, 141-160, 162-198, 200, 202-204, 206-211, 214, 216-247, 250-253, 255, 256, 287, 336, 364, 407 124, 125, 258-271, 273, 276-286, 290-
- 25 292, 294, 295, 297, 298, 300-307, 309-335, 337-357, 359-363, 365-367, 369-374, 376-386, 388-390, 393-406, 408-423 were included in
- the evidence for population model 5 (people with type 2 diabetes and higher risk for
- 27 cardiovascular disease. Almost all of these trials compared an 'adding' strategy, examining
- the effect of adding an intervention drug to other glucose-lowering drugs. Eleven
- comparisons included a 'switching' strategy and none compared a 'stopping' strategy. One
- 30 hundred and twenty-one comparisons were identified, and the follow-up times ranged
- 31 between 5.5 and 76 months.

32

Table 2: Summary of comparisons present in each population group in the protocol

| Comparat or class 1 | Compar ator 1 | Comparat or class 2 | Compar ator 2 | Hea rt fail ure | Atheroscl erotic cardiova scular disease | Chro nic kidn ey dise ase | Lower cardiova scular risk | Higher cardiova scular risk |
|------------------------|-----------------------------|------------------------|--------------------------------------|--------------------------|--|--|-------------------------------------|--------------------------------------|
| Biguanide | Metform in | Placebo | Placebo | No | No | No | No | Yes |
| Biguanide | Metform in modified release | Biguanide | Metformi n standard release | No | No | No | No | Yes |
| Biguanide | Metform in | Insulin | Insulin | No | No | No | No | Yes |
| DPP-4 inhibitor | Aloglipti n | Placebo | Placebo | Yes | Yes | No | No | Yes |

| | | | | | 1 | | • | 1 |
|---------------------|-------------------|------------------------------|------------------|-----|-----|-----|----|-----|
| DPP-4 inhibitor | Linaglipt in | Placebo | Placebo | Yes | No | Yes | No | Yes |
| DPP-4 inhibitor | Linaglipt in | Biguanide | Metformi n | No | No | No | No | Yes |
| DPP-4 inhibitor | Saxagli ptin | Placebo | Placebo | No | Yes | Yes | No | Yes |
| DPP-4 inhibitor | Sitaglipti n | Placebo | Placebo | Yes | Yes | No | No | Yes |
| DPP-4 inhibitor | Sitaglipti n | Biguanide | Metformi n | No | No | No | No | Yes |
| DPP-4 inhibitor | Sitaglipti n | DPP-4 inhibitor | Linaglipti n | No | No | No | No | No |
| DPP-4 inhibitor | Sitaglipti n | Insulin | Insulin | Yes | Yes | No | No | Yes |
| DPP-4 inhibitor | Vildagli ptin | Placebo | Placebo | Yes | No | No | No | No |
| DPP-4 inhibitor | Vildagli ptin | Biguanide | Metformi n | No | No | No | No | Yes |
| DPP-4 inhibitor | Vildagli ptin | Insulin | Insulin | No | No | No | No | Yes |
| DPP-4 inhibitor | Vildagli ptin | DPP-4 inhibitor | Aloglipti n | No | No | No | No | Yes |
| DPP-4 inhibitor | Vildagli ptin | DPP-4 inhibitor | Saxaglip tin | No | No | No | No | Yes |
| DPP-4 inhibitor | Vildagli ptin | DPP-4 inhibitor | Sitaglipti n | No | No | Yes | No | No |
| SGLT-2 inhibitor | Canaglif lozin | Placebo | Placebo | Yes | Yes | Yes | No | Yes |
| SGLT-2 inhibitor | Canaglif lozin | DPP-4 inhibitor | Sitaglipti n | No | No | No | No | Yes |
| SGLT-2 inhibitor | Canaglif lozin | GLP-1 receptor agonist | Liragluti de | No | No | No | No | Yes |
| SGLT-2 inhibitor | Canaglif lozin | GLP-1 receptor agonist | Semaglu tide | No | No | No | No | Yes |
| SGLT-2 inhibitor | Dapaglif lozin | Placebo | Placebo | Yes | Yes | Yes | No | Yes |
| SGLT-2 inhibitor | Dapaglif lozin | DPP-4 inhibitor | Saxaglip tin | No | No | No | No | Yes |
| SGLT-2 inhibitor | Dapaglif lozin | DPP-4 inhibitor | Sitaglipti n | No | No | No | No | Yes |
| SGLT-2 inhibitor | Dapaglif lozin | DPP-4 inhibitor | Vildaglip tin | No | Yes | No | No | Yes |

| SGLT-2 inhibitor | Dapaglif lozin | GLP-1 receptor agonist | Liragluti de | No | No | No | No | Yes |
|------------------------------|-------------------|------------------------------|------------------|-----|-----|-----|----|-----|
| SGLT-2 inhibitor | Dapaglif lozin | GLP-1 receptor agonist | Exenatid e | No | No | No | No | Yes |
| SGLT-2 inhibitor | Empagli flozin | Placebo | Placebo | Yes | Yes | Yes | No | Yes |
| SGLT-2 inhibitor | Empagli flozin | DPP-4 inhibitor | Linaglipti n | No | No | Yes | No | Yes |
| SGLT-2 inhibitor | Empagli flozin | DPP-4 inhibitor | Sitaglipti n | No | Yes | No | No | Yes |
| SGLT-2 inhibitor | Empagli flozin | DPP-4 inhibitor | Vildaglip tin | No | No | No | No | Yes |
| SGLT-2 inhibitor | Empagli flozin | GLP-1 receptor agonist | Liragluti de | No | No | No | No | Yes |
| SGLT-2 inhibitor | Empagli flozin | GLP-1 receptor agonist | Semaglu tide | No | No | No | No | Yes |
| SGLT-2 inhibitor | Empagli flozin | Insulin | Insulin | No | No | No | No | Yes |
| SGLT-2 inhibitor | Ertuglifl ozin | Placebo | Placebo | Yes | Yes | Yes | No | Yes |
| SGLT-2 inhibitor | Ertuglifl ozin | DPP-4 inhibitor | Sitaglipti n | No | No | No | No | Yes |
| GLP-1 receptor agonist | Dulaglut ide | Placebo | Placebo | No | Yes | No | No | Yes |
| GLP-1 receptor agonist | Dulaglut ide | DPP-4 inhibitor | Sitaglipti n | No | No | No | No | Yes |
| GLP-1 receptor agonist | Dulaglut ide | GLP-1 receptor agonist | Exenatid e | No | No | No | No | Yes |
| GLP-1 receptor agonist | Dulaglut ide | Insulin | Insulin | No | No | Yes | No | Yes |
| GLP-1 receptor agonist | Exenati de | Placebo | Placebo | Yes | Yes | No | No | Yes |
| GLP-1 receptor agonist | Exenati de | DPP-4 inhibitor | Sitaglipti n | No | No | No | No | Yes |
| GLP-1 receptor agonist | Exenati de | GLP-1 receptor agonist | Liragluti de | No | No | No | No | Yes |

| | 1 | 1 | | | 1 | 1 | 1 | 1 |
|------------------------------|------------------|------------------------------|------------------|-----|-----|-----|----|-----|
| GLP-1 receptor agonist | Exenati de | Insulin | Insulin | Yes | Yes | Yes | No | Yes |
| GLP-1 receptor agonist | Liragluti de | Placebo | Placebo | Yes | No | Yes | No | Yes |
| GLP-1 receptor agonist | Liragluti de | DPP-4 inhibitor | Linaglipti n | No | No | Yes | No | No |
| GLP-1 receptor agonist | Liragluti de | DPP-4 inhibitor | Saxaglip tin | No | No | No | No | Yes |
| GLP-1 receptor agonist | Liragluti de | DPP-4 inhibitor | Sitaglipti n | No | Yes | Yes | No | Yes |
| GLP-1 receptor agonist | Liragluti de | DPP-4 inhibitor | Vildaglip tin | No | No | No | No | Yes |
| GLP-1 receptor agonist | Liragluti de | GLP-1 receptor agonist | Dulagluti de | No | No | No | No | Yes |
| GLP-1 receptor agonist | Liragluti de | Insulin | Insulin | Yes | Yes | No | No | Yes |
| GLP-1 receptor agonist | Lixisena tide | Placebo | Placebo | Yes | Yes | No | No | Yes |
| GLP-1 receptor agonist | Lixisena tide | DPP-4 inhibitor | Sitaglipti n | No | No | No | No | Yes |
| GLP-1 receptor agonist | Lixisena tide | GLP-1 receptor agonist | Exenatid e | No | No | No | No | Yes |
| GLP-1 receptor agonist | Lixisena tide | GLP-1 receptor agonist | Liragluti de | No | No | No | No | Yes |
| GLP-1 receptor agonist | Lixisena tide | Insulin | Insulin | No | No | No | No | Yes |
| GLP-1 receptor agonist | Semagl utide | Placebo | Placebo | Yes | No | Yes | No | Yes |
| GLP-1 receptor agonist | Semagl utide | GLP-1 receptor agonist | Dulagluti de | No | No | Yes | No | Yes |
| GLP-1 receptor agonist | Semagl utide | DPP-4 inhibitor | Sitaglipti n | No | No | No | No | Yes |

| GLP-1 receptor agonist | Semagl utide | GLP-1 receptor agonist | Exenatid e | No | No | No | No | Yes |
|----------------------------------|--------------------------|----------------------------------|--------------------------------------|----|-----|----|----|-----|
| GLP-1 receptor agonist | Semagl utide | GLP-1 receptor agonist | Liragluti de | No | No | No | No | Yes |
| GLP-1 receptor agonist | Semagl utide; oral | GLP-1 receptor agonist | Semaglu tide; subcuta neous | No | No | No | No | Yes |
| GLP-1 receptor agonist | Semagl utide | Insulin | Insulin | No | No | No | No | Yes |
| GIP/GLP-1 receptor agonist | Tirzepat ide | Placebo | Placebo | No | No | No | No | Yes |
| GIP/GLP-1 receptor agonist | Tirzepat ide | GIP/GLP-1 receptor agonist | Dulagluti de | No | No | No | No | Yes |
| GIP/GLP-1 receptor agonist | Tirzepat ide | GIP/GLP-1 receptor agonist | Semaglu tide | No | No | No | No | Yes |
| GIP/GLP-1 receptor agonist | Tirzepat ide | Insulin | Insulin | No | Yes | No | No | Yes |
| Sulfonylur ea | Gliclazid e | DPP-4 inhibitor | Vildaglip tin | No | No | No | No | Yes |
| Sulfonylur ea | Glimepir ide | Placebo | Placebo | No | No | No | No | Yes |
| Sulfonylur ea | Glimepir ide | Biguanide | Metformi n | No | No | No | No | Yes |
| Sulfonylur ea | Glimepir ide | DPP-4 inhibitor | Linaglipti n | No | Yes | No | No | Yes |
| Sulfonylur ea | Glimepir ide | DPP-4 inhibitor | Saxaglip tin | No | No | No | No | Yes |
| Sulfonylur ea | Glimepir ide | DPP-4 inhibitor | Vildaglip tin | No | No | No | No | Yes |
| Sulfonylur ea | Glimepir ide | SGLT-2 inhibitor | Canaglifl ozin | No | No | No | No | Yes |
| Sulfonylur ea | Glimepir ide | SGLT-2 inhibitor | Dapaglifl ozin | No | No | No | No | Yes |
| Sulfonylur ea | Glimepir ide | SGLT-2 inhibitor | Empaglif lozin | No | No | No | No | Yes |
| Sulfonylur ea | Glimepir ide | SGLT-2 inhibitor | Ertugliflo zin | No | No | No | No | Yes |

| Sulfonylur ea | Glimepir ide | GLP-1 receptor agonist | Exenatid e | No | No | No | No | Yes |
|--------------------|------------------|------------------------------|-------------------|----|-----|-----|----|-----|
| Sulfonylur ea | Glimepir ide | GLP-1 receptor agonist | Liragluti de | No | No | No | No | Yes |
| Sulfonylur ea | Glimepir ide | Sulfonylur ea | Gliclazid e | No | No | No | No | Yes |
| Sulfonylur ea | Glimepir ide | Thiazolidin edione | Pioglitaz one | No | Yes | No | No | No |
| Sulfonylur ea | Glimepir ide | Insulin | Insulin | No | Yes | Yes | No | Yes |
| Sulfonylur ea | Glipizid e | Biguanide | Metformi n | No | No | No | No | Yes |
| Sulfonylur ea | Glipizid e | Placebo | Placebo | No | No | No | No | Yes |
| Sulfonylur ea | Glipizid e | DPP-4 inhibitor | Aloglipti n | No | No | No | No | Yes |
| Sulfonylur ea | Glipizid e | DPP-4 inhibitor | Saxaglip tin | No | No | No | No | Yes |
| Sulfonylur ea | Glipizid e | DPP-4 inhibitor | Sitaglipti n | No | No | No | No | Yes |
| Sulfonylur ea | Glipizid e | SGLT-2 inhibitor | Dapaglifl ozin | No | No | No | No | Yes |
| Thiazolidin edione | Pioglitaz one | Placebo | Placebo | No | Yes | Yes | No | Yes |
| Thiazolidin edione | Pioglitaz one | Biguanide | Metformi n | No | No | No | No | Yes |
| Thiazolidin edione | Pioglitaz one | DPP-4 inhibitor | Sitaglipti n | No | No | No | No | Yes |
| Thiazolidin edione | Pioglitaz one | DPP-4 inhibitor | Vildaglip tin | No | No | No | No | Yes |
| Thiazolidin edione | Pioglitaz one | SGLT-2 inhibitor | Dapaglifl ozin | No | No | No | No | Yes |
| Thiazolidin edione | Pioglitaz one | SGLT-2 inhibitor | Empaglif lozin | No | No | No | No | Yes |
| Thiazolidin edione | Pioglitaz one | GLP-1 receptor agonist | Exenatid e | No | No | No | No | Yes |
| Thiazolidin edione | Pioglitaz one | Sulfonylur ea | Gliclazid e | No | No | No | No | Yes |
| Thiazolidin edione | Pioglitaz one | Sulfonylur ea | Glimepiri de | No | No | No | No | Yes |

| | 1 | | | ı | | 1 | ı | |
|--------------------|--|------------------------------|-------------------|----|----|-----|----|-----|
| Thiazolidin edione | Pioglitaz one | Sulfonylur ea | Glipizide | No | No | No | No | Yes |
| Thiazolidin edione | Pioglitaz one | Insulin | Insulin | No | No | No | No | Yes |
| Combinati on | Dapaglif lozin + Saxagli ptin | Placebo | Placebo | No | No | Yes | No | No |
| Combinati on | Dapaglif lozin + Saxagli ptin | SGLT-2 inhibitor | Dapaglifl ozin | No | No | Yes | No | Yes |
| Combinati on | Dapaglif lozin + Saxagli ptin | DPP-4 inhibitor | Saxaglip tin | No | No | No | No | Yes |
| Combinati on | Dapaglif lozin + Saxagli ptin | DPP-4 inhibitor | Sitaglipti n | No | No | No | No | Yes |
| Combinati on | Dapaglif lozin + Saxagli ptin | Sulfonylur ea | Glimepiri de | No | No | No | No | Yes |
| Combinati on | Dapaglif lozin + Saxagli ptin | Insulin | Insulin | No | No | No | No | Yes |
| Combinati on | Dapaglif lozin + Exenati de | SGLT-2 inhibitor | Dapaglifl ozin | No | No | No | No | Yes |
| Combinati on | Dapaglif lozin + Exenati de | GLP-1 receptor agonist | Exenatid e | No | No | No | No | Yes |
| Combinati on | Empagli flozin + Liragluti de | GLP-1 receptor agonist | Liragluti de | No | No | No | No | Yes |
| Combinati on | Empagli flozin + Liragluti de | SGLT-2 inhibitor | Empaglif lozin | No | No | No | No | Yes |
| Combinati on | Empagli flozin + Liragluti de | Insulin | Insulin | No | No | No | No | Yes |
| Combinati on | Ertuglifl ozin + | DPP-4 inhibitor | Sitaglitpi n | No | No | No | No | Yes |

| | Sitaglipti n | | | | | | | |
|----------------------------|---|------------------------------|---|----|----|----|----|-----|
| Combinati on | Ertuglifl ozin + Sitaglipti n | SGLT-2 inhibitor | Ertugliflo zin | No | No | No | No | Yes |
| Combinati on | Liragluti de + Metform in | Biguanide | Metformi n | No | No | No | No | Yes |
| Combinati on | Glimepir ide + Metform in | Biguanide | Metformi n | No | No | No | No | Yes |
| Combinati on | Glimepir ide + Metform in modified release | Combinati on | Glimepiri de + Metformi n standard release | No | No | No | No | Yes |
| Combinati on | Pioglitaz one + Metform in | Sulfonylur ea | Glimepiri de | No | No | No | No | Yes |
| Combinati on | Pioglitaz one + Metform in | Thiazolidin edione | Pioglitaz one | No | No | No | No | Yes |
| Combinati on | Pioglitaz one + Aloglipi n | Thiazolidin edione | Pioglitaz one | No | No | No | No | Yes |
| Combinati on | Pioglitaz one + Exenati de | Thiazolidin edione | Pioglitaz one | No | No | No | No | Yes |
| Combinati on | Pioglitaz one + Exenati de | Insulin | Insulin | No | No | No | No | Yes |
| Insulin combinatio n | IDegLira | Placebo | Placebo | No | No | No | No | Yes |
| Insulin combinatio n | IDegLira | GLP-1 receptor agonist | Liragluti de | No | No | No | No | Yes |
| Insulin combinatio n | IDegLira | Insulin | Insulin | No | No | No | No | Yes |

| Insulin combinatio n | IGlarLixi | GLP-1 receptor agonist | Lixisenat ide | No | No | No | No | Yes |
|----------------------------|-----------|------------------------------|------------------|----|----|----|----|-----|
| Insulin combinatio n | IGlarLixi | Insulin | Insulin | No | No | No | No | Yes |

- 1 See also the study selection flow chart in report F2 (appendix C) and study evidence tables
- 2 in report F2 (appendix D). The forest plots and GRADE tables can be found in reports F3-7
- 3 (forest plots in Appendix F3-6, GRADE tables in Appendix F3-5 and F7).

4 1.1.4.2. Excluded studies

6

5 See the excluded studies list in report F8, appendix O.

1.1.5. Summary of studies included in the effectiveness evidence

7 Table 3: Summary of studies included in the evidence review

| | | Intervention and | | |
|---|--|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| Abdul-Ghani 2017 Qatar | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease: T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 52 (1) years Time since type 2 diabetes diagnosed: 10.7 (0.5) months | Strategy: Adding N = 231 Pioglitazone + exenatide (n=123) Insulin (n=108) Concomitant therapy: Metformin + Sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 60% | All-cause mortality, Non- fatal stroke, Hospitalisation for heart failure, Hypoglycaemia episodes, Severe hypoglycaemic episodes, Weight change Follow up: 36 months | Study location: Qatar Sources of funding: Supported by a Qatar Foundation grant NPRP 5- 273-3-079. One author's salary is paid in part by the South Texas Veterans Health Care System. |
| Abreu 2019 SIMPLE (NCT019669 78) | Model 5: People with type 2 diabetes at higher risk of cardiovascular | Strategy: Adding N = 120 Liraglutide (n=59) Insulin (n=61) | Health-related quality of life, Hypoglycaemia episodes, Severe hypoglycaemic | Study location: United States of America. Sources of |

| | | Intervention and | | |
|-----------|---|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 47.4 (9.5216) years Time since type 2 diabetes diagnosed: Not stated/unclear | Concomitant therapy: Insulin determir and metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 67.5% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 75.80% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | episodes, HbA1c change, Weight change Follow up: 6 months | funding: Funded by a Novo Nordisk Investigator Initiated Study Grant. |
| Adel 2022 | Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 106 Empagliflozin (n=52) Placebo (n=54) Concomitant therapy: Insulin Antihyperglycaemic treatment received: No additional information available. | Cardiovascular mortality, Unstable angina, HbA1c change, Weight change Follow up: 6 months | Study location: 2 centres in Iran. Sources of funding: Medication provided free of charge by Abidi Pharmaceutical Company, Iran. Funded by the Vice Chancellor for Research of Ahvaz Jundishapur University, Iran. |

| | | latera e Company | | |
|-----------------------------------|---|--|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| · | Mean age (SD): Not stated/unclear Time since type 2 diabetes diagnosed: Not stated/unclear | | | |
| Ahmann 2015 NN2211- 3917 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.4 (10.1944) years Time since type 2 diabetes diagnosed: 12.1 (6.9516) years | Strategy: Adding N = 450 Liraglutide (n=225) Placebo (n=225) Concomitant therapy: Basal insulin +/- metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 92.60% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 67.40% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Hypoglycaemia episodes, HbA1c change, Weight change, BMI change Follow up: 6 months | Study location: Multicenter trial - Argentina, Canada, Finland, Germany, India, Mexico, the Netherlands, Serbia and the United States of America. Sources of funding: Funded by Novo Nordisk. |
| Ahmann 2018 SUSTAIN 3 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney | Strategy: Adding N = 813 Semaglutide (n=406) Exenatide (n=407) Concomitant therapy: Metformin, thiazolidinedione +/-sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 96.60% DPP-4 inhibitors: | Health-related quality of life, All-cause mortality, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 12 months | Study location: Multicenter Sources of funding: Funded by Novo Nordisk A/S. |

| | | Intervention and | | |
|-------------------------|--|--|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.55 years Time since type 2 diabetes diagnosed: 9.2 years | Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 48.1% | | |
| Ahren 2004 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.8 (10.4881) years Time since type 2 diabetes diagnosed: 5.55 (3.9698) years | Strategy: Adding N = 107 Vildagliptin (n=56) Placebo (n=51) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality Follow up: 12 months | Study location: Multicenter. Sources of funding: Support from the Swedish Research Council. |
| Ahren 2013 GetGoal-M | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease | Strategy: Adding N = 680 Lixisenatide AM (n=255) Lixisenatide PM (n=255) Placebo AM and PM | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c | Study location: Multicenter trial - Australia, Canada, Chile, Czech Republic, Germany, Croatia, Mexico, Morocco, the |

| | | Intervention and | | |
|-------------------------|--|---|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.7667 (9.7156) years Time since type 2 diabetes diagnosed: 6.1 (5.1955) years | (n=170) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | change, Weight change Follow up: 5.5 months | Philippines, Romania, Russian Federation, South Africa, Spain, Ukraine, United States of America and Venezuela. Sources of funding: Funded by Sanofi. |
| Ahren 2014 HARMONY 3 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.775 (9.9708) years Time since type 2 diabetes diagnosed: 6.125 (4.87) years | Strategy: Adding N = 1012 Albiglutide (n=302) Sitagliptin (n=302) Glimepiride (n=307) Placebo (n=101) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, Severe hypoglycaemic episodes Follow up: 24 months | Study location: Multicenter Sources of funding: Funded by GlaxoSmithKline. |

| | _ | Intervention and | | _ |
|-------------------------|---|--|--|---|
| Study | Population | comparison | Outcomes | Comments |
| Ahren 2017 SUSTAIN 2 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.1333 (10.0097) years Time since type 2 diabetes diagnosed: 6.5667 (5.1465) years | Strategy: Adding N = 1231 Semaglutide 0.5 mg (n=410) Semaglutide 1.0 mg (n=410) Sitagliptin (n=411) Concomitant therapy: Metformin ± thiazolidinedione Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 99.7% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 0.7% | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 12 months | Study location: Multicenter trial - Bulgaria, Czech Republic, Hungary, Norway, Portugal, Romania, Spain, Sweden, Turkey, Ukraine, Argentina, Hong Kong, India, Japan, Mexico, Russia, South Africa and Thialand. Sources of funding: Novo Nordisk A/S. |
| Ando 2021 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Switching N = 40 Canagliflozin (n=20) Liraglutide (n=20) Concomitant therapy: Basal insulin +/- biguanides, alphaglucosidase inhibitors or glinides Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 3.00% Biguanides: 29.4% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear | Health-related quality of life, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 5.5 months | Study location: Japan. Sources of funding: Supported by the Initiative for Realizing Diversity in the Research Environment 2016. |

| | | Intervention and | | |
|-------------|---|---|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | Mean age (SD): 57.05 (12.2729) years Time since type 2 diabetes diagnosed: 9.1 (6.9502) years | SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Araki 2015A | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 61.0333 (9.8358) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 336 Empagliflozin 10mg (n=136) Empagliflozin 25mg (n=137) Metformin (n=63) Concomitant therapy: Monotherapy with sulfonylurea Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Japan (86 centres). Sources of funding: Funded by Boehringer Ingelheim and Eli Lilly and Company. |
| Araki 2015B | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People | Strategy: Adding N = 361 Dulaglutide (0.75 mg) (n=181) Insulin glargine (n=180) Concomitant therapy: Biguanide and/or sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not | All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Nonfatal stroke, Falls requiring hospitalisation, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c | Study location: 35 sites in Japan Sources of funding: Sponsored by Eli Lilly K. K. Japan. Multiple authors declare funding and honoraria from numerous pharmaceutical companies |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|---|--|--|---|---|
| | without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.8 (10.9062) years Time since type 2 diabetes diagnosed: 8.85 (6.4079) years | stated/unclear Biguanides: 36% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 18.5% | change, Weight change Follow up: 6 months | |
| Arechavaleta 2011 Sitagliptin Protocol 803 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.25 (9.9026) years Time since type 2 diabetes diagnosed: 6.75 (4.7014) years | Strategy: Adding N = 1035 Sitagliptin (n=516) Glimepiride (n=519) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 7 months | Study location: Multicenter trial. Sources of funding: Funded by Merck Sharp & Dohme Corp. |
| Aroda 2016B LixiLan-L | Model 5: People with type 2 diabetes at higher risk of | Strategy: Adding N = 736 Insulin glargine + | All-cause mortality, Cardiovascular mortality, | Study location: Australia, Canada Chile, Czech |

| | | Intervention and | | |
|-------------------------|--|--|--|--|
| Study | Population | comparison | Outcomes | Comments |
| Arada 2017 | cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.95 (9.0558) years Time since type 2 diabetes diagnosed: 12.05 (6.7521) years | lixisenatide (iGlarLix) once daily (n=367) Insulin glargine once daily (n=369) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 180.5% DPP-4 inhibitors: 4% GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 15% | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 7 months | Republic, Denma rk, Estonia, Hungary, Lithuania, Mexico Netherlands, Pol and, Romania, Russia ,Slovakia, Spain, Sweden, Ukraine Sources of funding: Sanofi |
| Aroda 2017 SUSTAIN 4 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.4667 (10.4345) years Time since type 2 | Strategy: Adding N = 1089 Semaglutide 0.5mg (n=362) Semaglutide 1.0mg (n=362) Insulin glargine (n=365) Concomitant therapy: Metformin ± sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 48.3% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear | All-cause mortality, Cardiovascular mortality, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 7 months | Study location: Multicenter trial. Sources of funding: Funded by Novo Nordisk A/S. |

| | | Intervention and | | |
|--------------------------|---|--|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | diabetes diagnosed: 8.5667 (6.2556) years | Sulfonylureas: Not stated/unclear | | |
| Aroda 2019A DUAL VIII | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.6 (10.0501) years Time since type 2 diabetes diagnosed: 10.1 (6.1502) years | Strategy: Adding N = 1012 Insulin degludec/liraglutide (n=506) Insulin glargine (n=506) Concomitant therapy: Metformin, sulfonylurea or pioglitazone Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 0.5% Biguanides: 98% DPP-4 inhibitors: 31.5% GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 64.5% | All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Nonfatal stroke, Unstable angina, Hospitalisation for heart failure, Development of end stage kidney disease, Cardiac arrhythmia, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 24 months | Study location: Multicenter trial. Sources of funding: Funded by Novo Nordisk. |
| Arturi 2017 | Model 1: People with type 2 diabetes and heart failure Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease T2DM and heart failure: People with heart failure: T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases | Strategy: Adding N = 32 Liraglutide (n=10) Sitagliptin (n=10) Glargine (n=12) Concomitant therapy: Metformin +/- sulfonylurea Antihyperglycaemic treatment received: No additional information available. | Hospitalisation for heart failure, Severe hypoglycaemic episodes, HbA1c change Follow up: 12 months | Study location: Italy Sources of funding: No funding from any specific grant from any funding agency in the public, commercial, or not-for profit sector. |

| | | Intomiontion and | | |
|-----------------------|--|---|--|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 60 (8.9693) years Time since type 2 diabetes diagnosed: Not stated/unclear | | | |
| Aschner 2012 EASIE | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 53.6 (8.7976) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 515 Sitagliptin (n=265) Insulin glargine (n=250) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 0.2% Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Non-fatal myocardial infarction, Unstable angina, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multicenter Sources of funding: Sanofi |
| Attaran 2023 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease | Strategy: Adding N = 73 Pioglitazone 30 mg daily (n=36) Empagliflozin 10 mg daily (n=37) Concomitant | Non-fatal stroke, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, BMI change | Study location: Iran Sources of funding: Supported by the Iran University of Medical Sciences No. |
| | | | | |

| | | Intervention and | | |
|----------------------|--|--|---|---|
| Study | Population | comparison | Outcomes | Comments |
| Avilée Conto | failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 52 (7) years Time since type 2 diabetes diagnosed: 7.95 (5.7196) years | Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: 51.40% GLP-1 receptor agonists: Not stated/unclear Insulin: 14.3% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 51.40% | Follow up: 5.5 months | IR.IUMS.REC.13 98.1408. Medication provided by Abidi Pharmaceutrical company |
| Avilés-Santa 1999 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 53.85 (8.6177) years Time since type 2 diabetes | Strategy: Adding N = 43 Metformin (n=21) Placebo (n=22) Concomitant therapy: Insulin Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Texas, US Sources of funding: Partly by Bristol-Myers Squibb |

| | | Intervention and | | |
|------------|--|--|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | diagnosed: 9.65 (5.5942) years | | | |
| Ba 2017 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57 (9.4005) years Time since type 2 diabetes diagnosed: 7 (5.1561) years | Strategy: Adding N = 498 Sitagliptin (n=249) Placebo (n=249) Concomitant therapy: Sulfonylurea ± metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 53.8% | All-cause mortality, Cardiovascular mortality, Nonfatal stroke, Unstable angina, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: multicenter; 32 centers in China Sources of funding: Merck & Co. Inc. FW, LX, MEH, SSE, and RRS are all current or former employees of Merck Sharp & Dohme Corp., a subsidiary of Merck & Co. Inc. (Kenilworth, NJ, USA) and may own stock or stock options in the company. FW also reports employment at Novartis Pharmaceuticals. |
| Babar 2021 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing | Strategy: Adding N = 240 Empagliflozin (n=120) Placebo (n=120) Concomitant therapy: Metformin + sitagliptin Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Pakistan Sources of funding: NR |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|-------------|---|--|---|--|
| , | cardiovascular disease Mean age (SD): 52.965 (8.8535) years Time since type 2 diabetes diagnosed: Not stated/unclear | | | |
| Bae 2021 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 61.855 (9.9295) years Time since type 2 diabetes diagnosed: 13.89 (7.8705) years | Strategy: Adding N = 119 Empagliflozin (n=60) Pioglitazone (n=59) Concomitant therapy: metformin + sulfonylurea + DPP-4 inhibitor Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, BMI change Follow up: 5.5 months | Study location: Multicentre, South Korea Sources of funding: Supported by research grants from Yuhan Corporation (Pharmaceutical company) |
| Bailey 2010 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular | Strategy: Adding N = 546 Dapagliflozin 2.5mg (n=137) Dapagliflozin 5 mg (n=137) Dapagliflozin 10 mg (n=135) Placebo (n=137) Concomitant therapy: Metformin Antihyperglycaemic | All-cause mortality, Cardiovascular mortality, Persistent signs of worsening kidney disease, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change | Study location: 80 sites across USA, Canada, Argentina, Mexico and Brazil Sources of funding: Bristol- Myers Squibb and AstraZeneca. The authors also declare |

| Study | Donulation | Intervention and | Outcomes | Comments |
|--------------------------------|---|--|---|---|
| Study | disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 53.925 (9.7327) years Time since type 2 diabetes diagnosed: 19.825 (5.6411) years | treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | Follow up: 23.7 months | numerous grants and honoraria from multiple pharmaceutical companies |
| Bailey 2016 LIRA- SWITCH | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.4 (10.1577) years Time since type 2 diabetes diagnosed: 7.75 (5.9565) years | Strategy: Switching N = 406 Liraglutide (n=202) Sitagliptin (n=204) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multicenter trial. Sources of funding: Funded by Novo Nordisk. |
| Bajaj 2014 | Model 5: People with type 2 diabetes at higher risk of cardiovascular | Strategy: Adding N = 272 Linagliptin (n=183) Placebo (n=89) | All-cause mortality, Cardiovascular mortality, Non- fatal stroke, | Study location: 52 trial centres in Asia, Europe and North America |

| | | Intervention and | | |
|--------------|---|--|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.15 (9.2963) years Time since type 2 diabetes diagnosed: Not stated/unclear | Concomitant therapy: Metformin + pioglitazone Antihyperglycaemic treatment received: No additional information available. | Unstable angina, Hospitalisation for heart failure, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Sources of funding: Boehringer Ingelheim. The funders participated in the study design, data collection and data analysis. |
| Barnett 2012 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.25 (9.3773) years Time since type 2 | Strategy: Adding N = 455 Saxagliptin (n=304) Placebo (n=151) Concomitant therapy: Insulin ± metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 69.00% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 100% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 5.5 months | Study location: Multicenter Sources of funding: Funding was provided by Bristol-Myers Squibb and AstraZeneca. Au thors declare numerous grants and honoraria for multiple pharmaceutical companies. |

| | | lutom continue and | | |
|---|--|---|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| , | diabetes diagnosed: 12 (7.0695) years | · | | |
| Barnett 2013 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 74.9 (4.3357) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 241 Linagliptin (n=162) Placebo (n=79) Concomitant therapy: Metformin ± sulfonylurea ± basal insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 0.3% Biguanides: 85.8% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 20.60% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 57.00% | All-cause mortality, Cardiovascular mortality, Nonfatal stroke, Unstable angina, Cardiac arrhythmia, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: Sponsored by Boehringer Ingelheim |
| Barnett 2014 EMPA-REG RENAL - CKD2 | Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 704 Stage 2 CKD - Placebo (n=97) Stage 2 CKD - Empagliflozin 10 mg (n=98) Stage 2 CKD - Empagliflozin 25 mg (n=97) Stage 3 CKD - Placebo (n=187) Stage 3 CKD - Empagliflozin 25 mg (n=188) Stage 4 CKD - Placebo (n=37) Stage 4 CKD - Empagliflozin 25 mg (n=37) Concomitant | All-cause mortality, Cardiac arrhythmia, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Multicenter Sources of funding: Boehringer Ingelheim, Eli Lilly |

| | | Intervention and | | |
|----------------------------------|---|---|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | Includes results for a subgroup for people with different stages of chronic kidney disease. Mean age (SD): 63.225 (6.823) years Time since type 2 diabetes diagnosed: Not stated/unclear | therapy: Antidiabetes treatment excluding SGLT-2 inhibitors Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 7.7% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 27.2% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 4.8% | | |
| Bergenstal 2009 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 52.15 (10.7609) years Time since type 2 diabetes diagnosed: 8.5 (6.1033) years | Strategy: Adding N = 248 Biphasic insulin aspart once daily (n=124) Concomitant therapy: Metformin + sulfonylurea Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: 102 sites in the USA Sources of funding: The study was supported by Novo Nordisk. |
| Bergenstal 2010 DURATION-2 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease | Strategy: Adding N = 491 Exenatide (n=160) Sitagliptin (n=166) Pioglitazone (n=165) | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Unstable angina, | Study location: USA, India and Mexico. Sources of funding: Amylin |

| | | Intervention and | | |
|--------------------|---|--|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 52.3333 (10.3489) years Time since type 2 diabetes diagnosed: 5.6667 (4.6858) years | Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Acute kidney injury, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Pharmaceuticals and Eli Lilly |
| Berndt-Zipfel 2013 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.5 (8.0623) years Time since type 2 diabetes | Strategy: Adding N = 44 Vildagliptin (n=22) Glimepiride (n=22) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Not stated Sources of funding: Not stated |

| | | Intervention and | | |
|----------------------------------|---|--|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | diagnosed: 7.25 (7.0838) years | | | |
| Billings 2018 DUAL VII | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.3 (8.8015) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 506 Insulin degludec and liraglutide fixed-ratio combination (n=252) Basal-bolus insulin (Insulin glargine and insulin aspart) (n=254) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Non-fatal myocardial infarction, Unstable angina, Hospitalisation for heart failure, Cardiac arrhythmia, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: Trial funded by Novo Nordisk |
| Bizino 2019 MAGNA VICTORIA | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Mixed population | Strategy: Adding N = 49 Liraglutide (n=23) Placebo (n=26) Concomitant therapy: Metformin with or without sulfonylurea and/or insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 65% | HbA1c change, Weight change Follow up: 6 months | Study location: Trial conducted at Leiden University Medical Centre, Leiden, Netherlands Sources of funding: Novo Nordisk funded the study. |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|------------------------------------|---|---|--|--|
| | T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.5 (6.5509) years Time since type 2 diabetes diagnosed: 11 (6.5509) years | SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 28.5% | | |
| Blonde 2015 AWARD-4 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.3667 (9.2374) years Time since type 2 diabetes diagnosed: 12.7333 (6.9687) | Strategy: Adding N = 884 Duaglutide 1.5 mg (n=295) Dulaglutide 0.75 mg (n=293) Insulin glargine (n=296) Concomitant therapy: Insulin lispro with or without metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 1.1% Biguanides: 72.3% DPP-4 inhibitors: 49% GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 29% | All-cause mortality, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Multicenter Sources of funding: Eli Lilly and Company |
| Blonde 2020 LIRA- ADD2SGLT2i | years Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and | Strategy: Adding N = 303 Liraglutide (n=203) Placebo (n=100) Concomitant therapy: SGLT2i +/- metformin Antihyperglycaemic treatment received: | All-cause mortality, Cardiovascular mortality, Acute kidney injury, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change | Study location: Multicentre, multinational trial at 74 sites in Brazil, India, Israel, Mexico, the Russian Federation and the United States. |

| | | Intervention and | | |
|-----------|--|---|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.35 (10.0347) years Time since type 2 diabetes diagnosed: 9.85 (7.0395) years | Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 94.60% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: 50.60% Sulfonylureas: Not stated/unclear | Follow up: 6 months | funding: Novo Nordisk |
| Bode 2013 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 63.6333 (6.2386) years Time since type 2 diabetes diagnosed: 11.6667 (7.4404) years | Strategy: Adding N = 714 Canagliflozin 100 mg (n=241) Canagliflozin 300 mg (n=236) Placebo (n=237) Concomitant therapy: None or monotherapy/combination therapy (including metformin, sulfonylurea, DPP-4 inhibitor, alpha glucosidase inhibitor, GLP-1 agonist, or insulin) Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 85.3% DPP-4 inhibitors: 9.1% GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 48.7% | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 24 months | Study location: Multicenter Sources of funding: Study sponsored by Janssen Research & Development |

| | | 1.4 | | |
|---------------|---|--|---|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| Bolinder 2012 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Mixed population T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Mean age (SD): 60.7 (7.5706) yrs years Time since type 2 diabetes diagnosed: 5.75 (4.9208) years | Strategy: Adding N = 180 Dapagliflozin + Metformin (n=89) Placebo + Metformin (n=91) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Health-related quality of life, All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 24 months | Study location: Conducted at 40 sites in Bulgaria, Czech Republic, Hungary, Poland, and Sweden Sources of funding: AstraZeneca and Bristol-Myers Squibb |
| Bolli 2008 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher | Strategy: Adding N = 576 Vildagliptin 100 mg (n=295) Pioglitazone 30mg (n=281) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Non-fatal stroke, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Worldwide. Study conducted at 118 centers; Germany (26), UK (25), USA (24), Spain (16), Italy (12), Switzerland (5), Austria (4), South Africa (3) andAustralia (3). Sources of funding: Novartis Pharmaceuticals Corporation |

| | | lusta magastica a surel | | |
|--------------------------|---|--|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | cardiovascular risk: Not stated/unclear Mean age (SD): 56.65 (9.4972) years Time since type 2 diabetes diagnosed: 6.4 (5.0486) years | | | |
| Bolli 2014 GetGoal-F1 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Mean age (SD): 56 (9.2085) years Time since type 2 diabetes diagnosed: 6 (4.4138) years (mean) | Strategy: Adding N = 482 Lixisenatide onestep dose increase (n=161) Lixisenatide twostep dose increase (n=161) Placebo (n=160) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 17.5 months | Study location: Multicenter Sources of funding: Sanofi |
| Bosi 2007 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart | Strategy: Adding N = 544 Vildagliptin 50 mg (n=177) Vildagliptin 100 mg (n=185) Placebo (n=182) Concomitant therapy: Metformin | All-cause mortality, Cardiovascular mortality, Non- fatal stroke, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change | Study location: East Hanover, New Jersey, United States. Study was conducted at 109 centers in the U.S, France, Italy and Sweden. Sources of |

| Otrod | Domaile Com | Intervention and | Outoo | 0 |
|------------------------|---|---|--|--|
| Study | Population failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Noinformation stated/unclear Mean age (SD): 54.2 (9.8386) years Time since type 2 diabetes | Antihyperglycaemic treatment received: No additional information available. | Outcomes Follow up: 5.5 months | Comments funding: Novartis Pharmaceuticals Corporation |
| Brown 2020 DAPA-LVH | diagnosed: 6.2667 (5.1722) years Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 544 Dapagliflozin (n=32) Placebo (n=34) Concomitant therapy: Metformin with or without other antihyperglycaemic drugs Antihyperglycaemic treatment received: No additional information available. | Diabetic ketoacidosis, Hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 12 months | Study location: Scotland Sources of funding: Novartis Pharmaceuticals Corporation |

| | | Intervention and | | |
|--------------------------------|---|--|---|---|
| Study | Population | comparison | Outcomes | Comments |
| Bunck 2009 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.4 (1.3) years Time since type 2 diabetes diagnosed (SD): 4.9 (1.1) years | Strategy: Adding N = 69 Exenatide (n=36) Insulin glargine (n=33) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Multicenter. Sweden, Finland and the Netherlands. Sources of funding: Amylin Pharmaceuticals and Eli Lilly and Company |
| Buse 2004 Exenatide- 113 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 377 Exenatide 10 mcg twice daily (n=129) Exenatide 5 mcg twice daily (n=125) Placebo (n=123) Concomitant therapy: Sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear | Non-fatal myocardial infarction, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 7 months | Study location: USA (101 sites) Sources of funding: Amylin Pharmaceuticals and Eli Lilly. |

| | | Intervention and | | |
|---------------------|--|--|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | 55.3333 (10.6788) years Time since type 2 diabetes diagnosed: 6.2 (5.5749) years | SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 45% | | |
| Buse 2009 LEAD 6 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.4 (1.3) years Time since type 2 diabetes diagnosed (SD): 4.9 (1.1) years | Strategy: Adding N = 464 Liraglutide (n=233) Exenatide (n=231) Concomitant therapy: Metformin and sulfonylureas Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Multicenter. Sources of funding: Novo Nordisk A/S. |
| Buse 2011 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease | Strategy: Adding N = 261 Exenatide 10 mcg twice daily (n=138) Placebo (n=123) Concomitant therapy: Insulin glargine with or without metformin or pioglitazone or metformin + pioglitazone Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 70.5% | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 7 months | Study location: Multicenter Sources of funding: Sponsored and funded by the Alliance of Eli Lilly and Company and Amylin Pharmaceuticals. |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|-------------------------|--|---|---|--|
| | T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59 (9.4842) years Time since type 2 diabetes diagnosed: 12 (7) years | DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Buse 2013 DURATION 6 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease: T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57 (9.5) years Time since type 2 diabetes diagnosed (SD): 8.5 (6.0) years | Strategy: Adding N = 911 Liraglutide (n=461) Exenatide (n=450) Concomitant therapy: No additional information available. Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter. Sources of funding: Novo Nordisk A/S. |
| Buse 2014 DUAL II | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not | Strategy: Adding N = 413 IDegLira (n=207) Insulin degludec (n=206) Concomitant therapy: No additional information available. Antihyperglycaemic | Non-fatal myocardial infarction, Non- fatal stroke, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change | Study location: Multicenter. Sources of funding: Novo Nordisk. |

| | | Intervention and | | |
|---------------------------|--|--|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.5 (10.1) years Time since type 2 diabetes diagnosed (SD): 10.5 (6.5) years | treatment received: No additional information available. | Follow up: 6 months | |
| Camerini- Davalos 1994 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: People without chronic kidney disease: People without chronic kidney disease: T2DM and chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): Not stated/unclear Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 70 Glipizide 5 mg daily (n=40) Placebo (n=30) Concomitant therapy: Insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 61.20% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Follow up: 36 months | Study location: Metropolitan Hospital Center Diabetes CLinic, New York, NY, USA Sources of funding: Supported in part by Diabetes Research Fund, New York, NY; the Michael J. Bilotto Research Fund of HOPE for Diabetes Foundation, New York, NY; the Veterans Administration Research Fund, Washington, DC; Roerig-Pfizer Pharmaceuticals, New York, NY. |
| Cannon 2020 VERTIS CV | Model 1: People with type 2 diabetes and heart failure | Strategy: Adding N = 8246 Ertugliflozin (n=5499) | All-cause mortality, Cardiovascular mortality, 3-point MACE, 4-point | Study location: Multicenter Sources of funding: Merck |

| | | Intervention and | | |
|--------------------------------|---|--|---|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Mixed population T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Mixed population T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Includes results for a subgroup for people with or without heart failure and with or without chronic kidney disease. Mean age (SD): 64.4 (8.0668) years Time since type 2 diabetes diagnosed: 13 (8.3334) years | Placebo (n=2747) Concomitant therapy: Monotherapy or combination therapy of any approved agent Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 76.60% DPP-4 inhibitors: 11.00% GLP-1 receptor agonists: 3.3% Insulin: 47.7% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 41% | MACE, Non-fatal myocardial infarction, Non-fatal stroke, Unstable angina, Hospitalisation for heart failure, Acute kidney injury, Persistent signs of worsening kidney disease, Development of end stage kidney disease, Death from renal causes, Cardiac arrhythmia, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 36 months | Sharp & Dohme and Pfizer |
| Capehorn 2020 SUSTAIN 10 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease | Strategy: Adding N = 577 Semaglutide 1.0 mg weekly (n=290) Liraglutide 1.2 mg daily (n=287) Concomitant | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c | Study location: Multicenter Sources of funding: Funded by Novo Nordisk |

| | | Intervention and | | |
|-------------------------------|---|---|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.5 (10.2544) years Time since type 2 diabetes diagnosed: 9.25 (5.9044) years | therapy: Metformin, sulfonylurea or SGLT2 inhibitor monotherapy or combination therapy Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 94.8% DPP-4 inhibitors: 0.20% GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: 24.6% Sulfonylureas: 46.8% | change, Weight change, BMI change Follow up: 7 months | |
| Cefalu 2013 CANTATA- SU | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.1667 (9.2354) years Time since type 2 diabetes diagnosed: 6.6 (5.3385) years | Strategy: Adding N = 1452 Glimepiride 6/8 mg daily (n=484) Canagliflozin 100 mg daily (n=483) Canagliflozin 300 mg daily (n=485) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Death from renal causes, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Multicenter Sources of funding: Funded by Janssen Research and Development, LLC |

| | | Intervention and | | |
|--|--|--|---|---|
| Study | Population | comparison | Outcomes | Comments |
| Cefalu 2015 | Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 62.9 (7.3599) years Time since type 2 diabetes diagnosed: 12.45 (8.4548) years | Strategy: Adding N = 914 Dapagliflozin (n=455) Placebo (n=459) Concomitant therapy: Drugs excluding rosiglitazone Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 16.80% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Unstable angina, Hospitalisation for heart failure, Acute kidney injury, Persistent signs of worsening kidney disease, Development of end stage kidney disease, Progression of liver disease, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Multicenter trial. Sources of funding: Supported by Bristol-Myers Squibb and AstraZeneca. An author was supported in part by a grant from the National Institute of General Medical Sciences of the National Institutes of Health (1-U54-GM-104940). |
| Charbonnel 2006 Sitagliptin 020 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not | Strategy: Adding N = 701 Sitagliptin 100 mg daily (n=464) Placebo (n=237) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, HbA1c change Follow up: 5.5 months | Study location: Multicenter Sources of funding: Funded by Merck Research Laboratories |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|------------------|--|--|--|---|
| | stated/unclear Mean age (SD): 54.55 (10.1691) years Time since type 2 diabetes diagnosed: 6.3 (5.1742) years | | | |
| Charbonnel 2013 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.25 (10.4083) years Time since type 2 diabetes diagnosed: 7.9 (5.5454) years | Strategy: Adding N = 653 Sitagliptin 100 mg daily (n=326) Liraglutide 1.2 mg daily (n=327) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: Sponsored by Merck Sharp & Dohme Corp, subsidiary of Merck & Co., Inc. |
| Charpentier 2009 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not | Strategy: Adding N = 299 Pioglitazone 30 mg daily (n=145) Placebo (n=154) Concomitant therapy: Metformin + sulfonylurea or metiglinide Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not | All-cause mortality, Cardiovascular mortality, Hospitalisation for heart failure, Hypoglycaemia episodes, HbA1c change Follow up: 7 months | Study location: France (52 hospitals, diabetology or internal medical services and 16 diabetes specialists) Sources of funding: Sponsored by Takeda France |

| | | Intervention and | | |
|-----------|--|--|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.7 (9.4466) years Time since type 2 diabetes diagnosed: 12.3 (8.4845) years | stated/unclear Biguanides: 0.7% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Chen 2016 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 62.895 (6.5404) years Time since type 2 diabetes diagnosed: 6.975 (2.3216) years | Strategy: Adding N = 73 Vildagliptin 100 mg daily (n=37) Saxagliptin 5 mg daily (n=36) Concomitant therapy: Metformin and Gliclazide Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hospitalisation for heart failure, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, BMI change Follow up: 5.5 months | Study location: Guangzhou, China Sources of funding: Reports study not funded. |
| Chen 2017 | Model 1: People with type 2 diabetes and heart failure Model 2: People with type 2 diabetes and atherosclerotic | Strategy: Adding Exenatide (n=14) Insulin (n=12) Concomitant therapy: Metformin +/- sulfonylurea | All-cause mortality, HbA1c change, BMI change Follow up: 6 months | Study location: The Netherlands. Sources of funding: Supported by Eli Lilly which had a partnership with |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|----------|---|--|--|--|
| | cardiovascular disease T2DM and heart failure: People with heart failure T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Antihyperglycaemic treatment received: No additional information available. | Cutcomes | Amylin, the manufacturer of exenatide at the time the trial was designed and data was collected. |
| SUPER | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.1 (8.0507) years Time since type 2 diabetes diagnosed: 13.35 | Strategy: Adding N = 462 Saxagliptin + insulin (n=232) Placebo + insulin (n=230) Concomitant therapy: Insulin with or without metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 5.5 months | Study location: China Sources of funding: Industry funding - AstraZeneca |
| Cho 2019 | (6.8667) years Model 5: People with type 2 diabetes at higher risk of | Strategy: Switching N = 71 Dapagliflozin 5mg | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c | Study location: Japan Sources of |

| | | late man of the second | | |
|-------------|---|---|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 63.35 (10.099) years Time since type 2 diabetes diagnosed: Not stated/unclear | (n=36) Pioglitazone 15 - 30mg (n=35) Concomitant therapy: Other hypoglycemic agents Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 15.5% Biguanides: 65.00% DPP-4 inhibitors: 52.3% GLP-1 receptor agonists: 7% Insulin: 14.00% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 26.80% | change, Weight change Follow up: 5.5 months | funding: There was no financial support for this trial. |
| Civera 2008 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 61.7 (9.7346) years Time since type 2 | Strategy: Adding N = 25 Metformin (n=12) NPH insulin (n=13) Concomitant therapy: Metformin + insulin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, HbA1c change, Weight change Follow up: 5.5 months | Study location: Spain Sources of funding: No additional information |

| | | 1.4 | | |
|---------------|--|--|--|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| , | diabetes diagnosed: 9.05 (5.3506) years | | | |
| Cusi 2019 | Model 5: People with type 2 diabetes at higher risk of | Strategy: Adding N = 56 Canagliflozin 300 | HbA1c change, Weight change Follow up: 5.5 | Study location: USA Sources of |
| | cardiovascular disease T2DM and heart | mg daily (n=26) Placebo daily (n=30) Concomitant | months | funding: Funding by Janssen Research & Development |
| | failure: Not stated/unclear T2DM and atherosclerotic | therapy: Metformin ± DPP-4 inhibitor Antihyperglycaemic | | |
| | cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not | treatment received: No additional information available. | | |
| | Mean age (SD): 58 (9.5501) years Time since type 2 diabetes | | | |
| | diagnosed: Not stated/unclear | | | |
| da Silva 2016 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease | Strategy: Adding N = 35 Sitagliptin (n=18) NPH Insulin (n=17) Concomitant | HbA1c change, Weight change, BMI change Follow up: 12 months | Study location: Brazil Sources of funding: São Paulo Research Foundation |
| | T2DM and heart failure: People without heart | therapy: Metformin + sulfonylurea (glyburide) | | Todridation |
| | failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and | Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: | | |
| | chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not | Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear | | |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|--|--|---|---|---|
| | stated/unclear Mean age (SD): 56.75 (6.7977) years Time since type 2 diabetes diagnosed: 10.9 (6.6785) years | SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 100% | | |
| Dagogo-Jack 2018 VERTIS SITA2 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.75 (6.7977) years Time since type 2 diabetes diagnosed: 10.9 (6.6785) years | Strategy: Adding N = 35 Ertugliflozin 15mg (n=153) Ertugliflozin 5mg (n=156) Placebo (n=153) Concomitant therapy: Metformin + a DPP-4 inhibitor or a sulfonylurea. A small number of people were on triple therapy. Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Multicenter Sources of funding: Merck & Co |
| Dahl 2022 SURPASS-5 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear | Strategy: Adding N = 475 Tirzepatide 15 mg once weekly (n=120) Tirzepatide 10 mg once weekly (n=119) Tirzepatide 5 mg once weekly (n=116) Placebo once weekly (n=120) Concomitant therapy: Insulin +/- | All-cause mortality, Cardiovascular mortality, 4-point MACE, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 11 months | Study location: Multicenter Sources of funding: Eli Lilly and Company |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|----------------------------|---|---|--|--|
| Study | T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 60.75 (10) years Time since type 2 diabetes diagnosed: 13.325 (7.3278) years | metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 98.5% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 118.80% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | Outcomes | Comments |
| D'Alessio 2015 EAGLE | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.25 (8.8499) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strated/unclear Strategy: Adding N = 944 Liraglutide 0.6 mg - 1.8 mg once daily (n=470) Insulin glargine (n=474) Concomitant therapy: Metformin ± sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 1% Biguanides: 470.5% DPP-4 inhibitors: 100% GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 320.5% | Non-fatal stroke, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multicenter Sources of funding: Sanofi |
| Davies 2009 HEELA | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease | Strategy: Adding N = 235 Exenatide (n=118) Insulin glargine (n=117) | Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c | Study location: Multicentre at 36 centres in the UK Sources of funding: NR. A. K. and C. N. are |

| | | lutement to the | | |
|-------------|---|--|--|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Mean age (SD): 56.5 (9.1277) years Time since type 2 diabetes diagnosed: 8.7 (4.5015) years | Concomitant therapy: 2-3 oral drugs (metformin, sulfonylurea, thiazolidinedione) Antihyperglycaemic treatment received: No additional information available. | change, Weight change Follow up: 6 months | employees of Eli Lilly and Company |
| Davies 2013 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease | Strategy: Adding N = 216 Exenatide 2mg once weekly (n=111) Insulin detemir titrated (2.0 IU/day to 62.0 IU/day) (n=105) Concomitant therapy: Metformin ± sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 108% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, At night hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 5.5 months | Study location: UK Sources of funding: Eli Lilly and Company and Amylin Pharmaceutical, LLC |

| | | Intervention and | | |
|----------------------------------|---|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | Mean age (SD): Not stated/unclear Time since type 2 diabetes diagnosed: Not stated/unclear | SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 77% | | |
| Davies 2015 SCALE Diabetes | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.8667 (10.5332) year years Time since type 2 diabetes diagnosed: 7.2 (5.4135) years | Strategy: Adding N = 846 Liraglutide 3.0mg daily (n=423) Liraglutide 1.8 mg daily (n=211) Placebo daily (n=212) Concomitant therapy: 1 to 3 drugs (metformin, thiazolidinedione, sulfonylurea) Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 158% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 3.7% | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Development of end stage kidney disease, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 13 months | Study location: Multicenter Sources of funding: Novo Nordisk |
| Davies 2016 LIRA-RENAL | Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not | Strategy: Adding N = 279 Liraglutide 1.8 mg once daily (n=140) Placebo (n=139) Concomitant therapy: Monotherapy or dual therapy combinations with metformin, | All-cause mortality, Cardiovascular mortality, Non-fatal stroke, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI | Study location: Multicenter Sources of funding: Sponsored by Novo Nordisk A/S. |

| | | Intervention and | | |
|-------------|---|--|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 67.15 (8.1519) years Time since type 2 diabetes diagnosed: 15.05 (8.2323) years | sulfonylurea, and/or pioglitazone; or monotherapy with basal or premix insulin, or any combination of basal or premix insulin with metformin and/or pioglitazone Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 9.4% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SUlfonylureas: 12.3% | change Follow up: 6 months | |
| Davies 2017 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.95 (8.4073) years Time since type 2 diabetes | Strategy: Adding N = 421 Placebo (n=71) Semaglutide 2.5mg (n=70) Semaglutide 5mg (n=70) Semaglutide 10mg (n=69) Semaglutide 20mg (n=70) Semaglutide 40mg standard (n=71) Semaglutide 40mg slow (n=70) Semaglutide 40mg fast (n=70) Semaglutide SC 1mg (n=69) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 84.9% DPP-4 inhibitors: Not stated/unclear | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Nonfatal stroke, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: Editorial support funded by Novo Nordisk |

| | | Intervention and | | |
|-----------------------|--|---|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | diagnosed: 5.975 (4.2113) years | GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Davies 2021 STEP 2 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.3333 (10.6773) years Time since type 2 diabetes diagnosed: 8.0333 (6.1017) years | Strategy: Adding N = 1210 Semaglutide 2.4mg (n=404) Semaglutide 1.0mg (n=403) Placebo (n=403) Concomitant therapy: Background oral antidiabetic drugs Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 0.1% Biguanides: 91.8% DPP-4 inhibitors: 0.5% GLP-1 receptor agonists: 0.1% Insulin: Not stated/unclear SGLT-2 inhibitors: 24.8% Sulfonylureas: 25.5% | Health-related quality of life, All-cause mortality, Acute kidney injury, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 15.7 months | Study location: Multicenter Sources of funding: Novo Nordisk |
| DeFronzo 2005 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear | Strategy: Adding N = 336 Exenatide 20mcg daily (n=113) Exenatide 10mcg daily (n=110) Placebo (n=113) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: 82 sites in the U.S. Sources of funding: Supported by Amylin Pharmaceuticals, San Diego, California, and Eli Lilly, Indianapolis, Indiana |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|--|--|---|--|---|
| | T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 53 (10.3705) years Time since type 2 diabetes diagnosed: 5.9 (5.5981) years | stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| DeFronzo 2009 Saxagliptin 014 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.6 (9.9983) years Time since type 2 diabetes diagnosed: 6.525 (5.1043) years | Strategy: Adding N = 743 Placebo (n=179) Saxagliptin 2.5mg (n=192) Saxagliptin 5mg (n=191) Saxagliptin 10mg (n=181) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 5.5 months | Study location: US Sources of funding: sponsored and monitored by Bristol-Myers Squibb and AstraZeneca |
| DeFronzo 2012 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease | Strategy: Adding N = 1168 Pioglitazone (n=388) Alogliptin 12.5mg + Pioglitazone (n=390) Alogliptin 25mg + Pioglitazone (n=390) | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change | Study location: Multicenter Sources of funding: Supported by Takeda Global Research & Development |

| | | Intervention and | | |
|------------------|---|---|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.5333 (9.4927) years Time since type 2 diabetes diagnosed: 6.3567 (5.4731) years | Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Follow up: 6 months | Center, Takeda Pharmaceuticals North America, Inc. |
| DeFronzo 2015 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.225 (9.2193) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 674 Empagliflozin 2g/linagliptin 5mg (n=134) Empagliflozin 10 mg/linagliptin 5 mg (n=135) Empagliflozin 25 mg (n=140) Empagliflozin 10 mg (n=137) Linagliptin 5 mg (n=128) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 134.8% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Hospitalisation for heart failure, Hypoglycaemia episodes, HbA1c change Follow up: 12 months | Study location: Multicenter Sources of funding: Boehringer Ingelheim and Eli Lilly and Company |

| | | Intervention and | | |
|--------------------------------|---|---|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | | Sulfonylureas: Not stated/unclear | | |
| Del Prato 2014 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.3667 (9.6709) years Time since type 2 diabetes diagnosed: 5.5333 (4.9827) years | Strategy: Adding N = 2639 Alogliptin 12.5 mg (n=880) Alogliptin 25 mg (n=885) Glipizide (n=874) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, 3-point MACE, Non-fatal myocardial infarction, Non-fatal stroke, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 24 months | Study location: 310 study sites in North and South America, Europe, Asia, South Africa and Australia/New Zealand Sources of funding: Takeda Pharmaceuticals International, Inc. |
| Del Prato 2021 SURPASS-4 | Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear | Strategy: Adding N = 2002 Tirzepatide (n=997) Insulin (n=1005) Concomitant therapy: Metformin +/- sulfonylurea +/- SGLT-2 inhibitors Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 94.5% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not | All-cause mortality, Cardiovascular mortality, 4-point MACE, Non-fatal myocardial infarction, Non-fatal stroke, Unstable angina, Hospitalisation for heart failure, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 24 months | Study location: Multicenter Sources of funding: Funded by Eli Lilly and Company. Authors received grants and honoraria from a variety of pharmaceutical companies. |

| | | Intervention and | | |
|-----------------|--|--|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 63.6 (8.5499) years Time since type 2 diabetes diagnosed: Not stated/unclear | stated/unclear SGLT-2 inhibitors: 25.5% Sulfonylureas: 54.5% | | |
| DePaoli 2014 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.55 (10.655) years Time since type 2 diabetes diagnosed: 8.55 (6.1) years | Strategy: Adding N = 121 Pioglitazone (n=60) Placebo (n=61) Concomitant therapy: SU +/- metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 80.20% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 100% | Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Not available Sources of funding: Study was funded by InteKrin Therapeutics, Inc. |
| Derosa 2010A | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure | Strategy: Adding N = 168 Pioglitazone + vildagliptin (n=83) Glimepiride + vildagliptin (n=85) Concomitant therapy: Vildagliptin Antihyperglycaemic | Hypoglycaemia episodes, HbA1c change, Weight change, BMI change Follow up: 12 months | Study location: Multicentre study in Italy Sources of funding: NR |

| | | Intervention and | | |
|-----------------|---|---|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.5 (5.5167) years Time since type 2 diabetes diagnosed: 6.5 (2.5436) years | treatment received: No additional information available. | | |
| Derosa 2010B | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.5 (5.526) years Time since type 2 diabetes diagnosed: 5.5 (2.5528) years | Strategy: Adding N = 151 Sitagliptin (n=75) Metformin (n=76) Concomitant therapy: Pioglitazone Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, HbA1c change, Weight change, BMI change Follow up: 12 months | Study location: Multicentre study in Italy Sources of funding: NR |
| Derosa 2011B | Model 5: People with type 2 diabetes at higher risk of | Strategy: Adding N = 111 Exenatide (n=57) | Hypoglycaemia episodes, HbA1c change, Weight change, BMI | Study location: Multicentre trial in Italy |

| | | lutement | | |
|-----------------|---|--|---|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| Study | cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.5 (6.5329) years Time since type 2 diabetes diagnosed: Not stated/unclear | Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Follow up: 12 months | Sources of funding: NR |
| Derosa 2012A | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.35 (8.3723) | Strategy: Adding N = 178 Sitagliptin (n=91) Placebo (n=87) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Multicentre study in Italy Sources of funding: University of PaviaSigma-Tau |

| | | Intervention and | | |
|-----------------|--|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | years Time since type 2 diabetes diagnosed: 5.6 (2.458) months | | | |
| Derosa 2012B | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 53.3 (7.727) years Time since type 2 diabetes diagnosed: 6.2 (3.8007) months | Strategy: Adding N = 167 Vildagliptin (n=84) Placebo (n=83) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, HbA1c change, Weight change, BMI change Follow up: 12 months | Study location: Multicentre study conducted in Italy Sources of funding: NR |
| Derosa 2012C | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher | Strategy: Adding N = 171 Exenatide (n=86) Placebo (n=85) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, HbA1c change, Weight change, BMI change Follow up: 12 months | Study location: Multicentre study in Italy Sources of funding: None |

| | | Intomorphism and | | |
|-----------------|--|---|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | cardiovascular risk: Not stated/unclear Mean age (SD): 57 (7.5038) years Time since type 2 diabetes diagnosed: 7.7 (2.9529) months | | | |
| Derosa 2014A | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.5 (9.4743) years Time since type 2 diabetes diagnosed: 6.85 (4.2028) months | Strategy: Adding N = 167 Glimepiride (n=81) Vildagliptin (n=86) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | HbA1c change, Weight change, BMI change Follow up: 5.5 months | Study location: Italy Sources of funding: NR |
| Derosa 2014B | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease | Strategy: Adding Sitagliptin (n=102) Placebo (n=103) Concomitant therapy: Oral antidiabetic drugs | HbA1c change, Weight change, BMI change Follow up: 24 months | Study location: Multicentre trial in Italy Sources of funding: NR |
| | T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular | Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 100% Biguanides: Not stated/unclear | | |

| Canal- | Domulation | Intervention and | Outoo | Comments |
|-------------------------------|---|---|--|---|
| Study | disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): Not stated/unclear Time since type 2 diabetes diagnosed: Not stated/unclear | comparison DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 57.7% | Outcomes | Comments |
| Diamant 2010 DURATION-3 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58 (9.5241) years Time since type 2 diabetes diagnosed: 7.9 (6) years | Strategy: Adding N = 456 Exenatide (n=233) Insulin glargine (n=223) Concomitant therapy: Metformin ± sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 70% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Health-related quality of life, All-cause mortality, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 36 months | Study location: Multicentre trial. 72 sites across the USE, Puerto Rico, the European Union, Russia, Australia, Korea, Taiwan and Mexico Sources of funding: Amlyn Pharmaceuticals Inc and Eli Lilly and Company. Authors have received grants and honoraria from multiple pharmaceutical companies. |
| Diamant 2014 4B | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not | Strategy: Adding N = 627 Exenatide (n=315) Insulin lispro (n=312) Concomitant therapy: Metformin + insulin glargine | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Acute kidney injury, Hypoglycaemia episodes, At | Study location: Multicenter Sources of funding: Study was part of the Eli Lilly and Company / Amlyn Pharmaceuticals |

| | | Intervention and | | |
|-----------------|--|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.45 (9.4519) years Time since type 2 diabetes diagnosed: 11 years | Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 100% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 7 months | Alliance and the Bristol-Myers Squibb / AstraZeneca Alliance. Authors received grants and honoraria from a number of different pharmaceutical companies. |
| Dobs 2013 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.6 (9.0496) years Time since type 2 diabetes diagnosed: 9.35 (6.2278) years | Strategy: Adding N = 278 Sitagliptin (n=181) Placebo (n=97) Concomitant therapy: Metformin + PPAR gamma agonist, metformin + sulfonylurea, sulfonylurea + PPAR gamma agonist Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12.4 months | Study location: 41 sites across North and South America, Europe and Asia Sources of funding: Study sponsored by Merck Sharp and Dohme Corp. Numerous authors are current or former employees of Merk Sharp and Dohme Corp. |
| Dorkhan 2009 | Model 5: People with type 2 | Strategy: Adding N = 30 | HbA1c change, BMI change | Study location: NR |
| | 71 | | J - | |

| | | 1.4 | | |
|------------|--|---|--|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 61.15 (7.6697) years Time since type 2 diabetes diagnosed: 10.3 (6.7915) years | Pioglitazone (n=15) Insulin glargine (n=15) Concomitant therapy: Metformin with sulfonylurea or meglitinide Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 100% | Follow up: 6 months | Sources of funding: Study was in part financially supported by grants from Sanofi-Aventis, The Crafoord Foundation, and The Swedish Heart and Lung Association. Authors declare various honoraria's with Eli Lilly and Sanofi-Aventis. One author owns shares and stock options in NovoNordisk A/S. |
| Douek 2005 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 183 Metformin (n=92) Placebo (n=91) Concomitant therapy: Insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 100% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Five hospitals in southwest England Sources of funding: Supported by the Special Trustees for the United Bristol Hospitals and the NHS Executive Southwest. Lipha Pharmaceuticals donated trial medication. |

| | | Intonvention and | | |
|------------------------|--|--|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | 58 (8.325) years Time since type 2 diabetes diagnosed: 9.5 (5.2) years | | | |
| Dungan 2014 AWARD-6 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.65 (9.6052) years Time since type 2 diabetes diagnosed: 7.2 | Strategy: Adding N = 599 Dulaglutide (n=299) Liraglutide (n=300) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Falls requiring hospitalisation, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: Sponsored by Eli Lilly and Company. Authors state numerous grants and honoraria from multiple pharmaceutical companies. |
| Dungan 2016 AWARD-8 | (5.4) years Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear | Strategy: Adding N = 300 Dulaglutide (n=240) Placebo (n=60) Concomitant therapy: Glimepiride Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not | All-cause mortality, Non-fatal stroke, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: NR Sources of funding: Funded by Eli Lilly and Company. First author declares funding and honoraria from multiple pharmaceutical companies |

| | | lusta magastica a surel | | |
|-------------------|--|--|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| · | T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.95 (9.71) years Time since type 2 diabetes diagnosis: 7.3 (5.0386) years | stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 100% | | |
| Ferdinand 2019 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Mean age (SD): 56.85 (9.3) years Time since type 2 diabetes diagnosed: 9.3 (7.0668) years | Strategy: Adding N = 150 Empagliflozin (n=78) Placebo (n=72) Concomitant therapy: None or stable oral hypoglycemic drug Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Persistent signs of worsening kidney disease, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: 92 centers in the United States Sources of funding: Boehringer Ingelheim |
| Fernandez 2008 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease | Strategy: Adding N = 20 Pioglitazone (n=10) Placebo (n=10) Concomitant therapy: insulin | Hypoglycaemia episodes, HbA1c change Follow up: 8.3 months | Study location: Texas, the US Sources of funding: American Diabetes Association Take |

| | | Intervention and | | |
|--------------------|--|--|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): Not stated/unclear Time since type 2 diabetes diagnosed: Not stated/unclear | Antihyperglycaemic treatment received: No additional information available. | | da Pharmaceuticals |
| Ferrannini 2009 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: People without chronic kidney disease: T2DM and chronic kidney disease: T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Mean age (SD): | Strategy: Adding N = 2789 Vildagliptin (n=1396) Glimepiride (n=1393) Concomitant therapy: metformin + placebo Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Cardiac arrhythmia, Hypoglycaemia episodes, HbA1c change Follow up: 12 months | Study location: Germany, United States Sources of funding: Novartis Pharmaceuticals |

| | | Intervention and | | |
|---------------|--|---|---|---|
| Study | Population | comparison | Outcomes | Comments |
| · | 57.48 (9.1705) years Time since type 2 diabetes diagnosed: 5.73 (5.1056) years | | | |
| Filozof 2010a | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.45 (10.0483) years Time since type 2 diabetes diagnosed: 6.6 (5.1991) years | Strategy: Adding N = 1007 Vildagliptin (n=513) Gliclazide (n=494) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, HbA1c change, Weight change Follow up: 12 months | Study location: Unclear, but appears to be Switzerland and France Sources of funding: Novartis Pharmaceuticals |
| Filozof 2010b | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear | Strategy: Adding N = 914 Vildagliptin (n=456) metformin (n=458) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Acute kidney injury, Hypoglycaemia episodes, HbA1c change Follow up: 5.5 months | Study location: Germany, United States Sources of funding: Novartis Pharmaceuticals |

| | | 1.4 | | |
|-------------------------|--|---|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.95 (9.8911) years Time since type 2 diabetes diagnosed: 4.65 (4.9251) years | | All-causa | |
| Fioretto 2018 DERIVE | Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): Not stated/unclear Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 321 Dapagliflozin 10 mg once daily (n=160) Placebo (n=161) Concomitant therapy: Stable glucose-lowering therapy (diet, exercise +oral antidiabetic drug [excluding SGLT2-inhibitors] and/or long/intermediate/mi xed insulin) Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 66.7% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 49.80% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 40.8% | All-cause mortality, Cardiovascular mortality, Nonfatal stroke, Hospitalisation for heart failure, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: International (88 sites in Bulgaria, Canada, Czech Republic, Italy, Poland, Spain, Sweden, USA). Sources of funding: Funded by AstraZeneca and supported by grant from National Institutes of Health, Grant/Award Number: UL1TR001111. |
| Fonseca 2007 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and | Strategy: Adding N = 296 vildagliptin (n=144) placebo (n=152) Concomitant therapy: insulin Antihyperglycaemic treatment received: No additional | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change | Study location: Germany (ten), Finland (five), Spain (four) and the USA (49) Sources of funding: Novartis Pharmaceuticals |

| Study | | | Intoniontian | | |
|--|------------|--|---|--|--|
| atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk of cardiovascular disease Fonseca Model 5: People with type 2 diabetes at higher failure: People without heart failure: People without heart failure: T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and higher cardiovascular disease: Not stated/unclear T2DM and higher cardiovascular disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear St | Study | Population | | Outcomes | Comments |
| with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not st | | atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.25 (10.5598) years Time since type 2 diabetes diagnosed: 14.65 | information | | |
| Forst 2005 Model 5: People with type 2 diabetes at higher Strategy: Adding N = 173 HbA1c change Unclear: appears to be Germany | | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.05 (9.0556) years Time since type 2 diabetes diagnosed: Not | N = 313 Sitagliptin (n=157) Placebo (n=156) Concomitant therapy: Pioglitazone + metformin Antihyperglycaemic treatment received: No additional information | mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 | Multicenter Sources of funding: Merck Sharp & Dohme |
| | Forst 2005 | with type 2 diabetes at higher | N = 173 | HbA1c change | Unclear: appears |

| | | Intomion and | | |
|------------|--|---|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 62.6 (7.9304) months years Time since type 2 diabetes diagnosed: 85.75 (86.8345) months | glimepiride (n=84) Concomitant therapy: other antidiabetic medication Antihyperglycaemic treatment received: No additional information available. | Follow up: 5.5 months | Sources of funding: TAKEDA Germany |
| Forst 2014 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.3333 (10.0701) years Time since type 2 | Strategy: Adding N = 342 Canagliflozin 100 mg (n=113) Canagliflozin 300 mg (n=114) Placebo/Sitagliptin (n=115) Concomitant therapy: metformin + pioglitazone Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, HbA1c change, Weight change Follow up: 6 months | Study location: Canada, Finland, France, Germany, Greece, India, Mexico, Spain, Thailand, United Kingdom, United States Sources of funding: Janssen Research & Development, LLC |

| | | Intervention and | | |
|--------------------------|---|---|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | diabetes diagnosed: 10.5333 (6.9493) years | | | |
| Forst 2015 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 66.75 (10.881) years Time since type 2 diabetes diagnosed: 7.95 (5.1347) years | Strategy: Adding N = 161 Vildagliptin (n=82) NPH insulin (n=79) Concomitant therapy: Glimepiride Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 5.5 months | Study location: Germany Sources of funding: Novartis Pharma GmbH |
| Frias 2016 DURATION-8 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular | Strategy: Adding N = 695 Exenatide 2 mg weekly + Dapagliflozin 10 mg daily (n=231) Exenatide 2 mg weekly + Placebo (n=231) Dapagliflozin 10 mg daily + Placebo (n=233) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional | All-cause mortality, Cardiovascular mortality, Acute kidney injury, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 24 months | Study location: Multicenter Sources of funding: AstraZeneca |

| | | Into many Control | | |
|------------|--|---|--|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | risk: Not stated/unclear Mean age (SD): 54.3333 (9.6763) years Time since type 2 diabetes diagnosed: 7.3667 (5.6711) years | information available. | | |
| Frias 2018 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.95 (7.0827) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 318 Tirzepatide 1 mg (n=53) Tirzepatide 5 mg (n=55) Tirzepatide 10 mg (n=52) Tirzepatide 15 mg (n=53) Dulaglutide 1.5 mg (n=54) Placebo (n=51) Concomitant therapy: Current metformin treatment continued Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 6 months | Study location: 47 sites in Poland, Puerto Rico, Slovakia, and USA Sources of funding: Eli Lilly and Company |
| Frias 2020 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular | Strategy: Adding N = 444 Saxagliptin + Dapagliflozin (n=227) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional | All-cause mortality, Hospitalisation for heart failure, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change | Study location: 87 centres in Germany, the Czech Republic, Hungary, Mexico, Poland, Romania, Russia, Sweden, the UK and the United States. Sources of |

| | | Intomiontion and | | |
|-------------------------|--|---|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.1 (9.6707) years Time since type 2 diabetes diagnosed: 7.8 (6.4491) years | information available. | Follow up: 36 months | funding: AstraZeneca |
| Frias 2021 SURPASS-2 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.575 (10.4287) years Time since type 2 diabetes diagnosed: 8.625 (6.455) years | Strategy: Adding N = 1878 Tirzepatide 5 mg (n=470) Tirzepatide 10 mg (n=469) Tirzepatide 15 mg (n=470) Semaglutide 1 mg (n=469) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Cardiac arrhythmia, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 9 months | Study location: 128 sites in the United States, Argentina, Australia, Brazil, Canada, Israel, Mexico, and the United Kingdom Sources of funding: Eili Lilly |
| Frias 2023 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease | Strategy: Adding N = 103 Dulaglutide (n=50) Placebo (n=53) Concomitant | All-cause mortality, Cardiovascular mortality, Cardiac arrhythmia, Diabetic ketoacidosis, | Study location: The US, Hungary, Poland, and Slovakia Sources of |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|--------------|--|---|--|---|
| | T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.55 (9.8458) years Time since type 2 diabetes diagnosed: Not stated/unclear | therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 6 months | funding: Eli Lilly and Company |
| Fujioka 2003 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.3333 years Time since type 2 diabetes diagnosed: 3 years | Strategy: Switching N = 217 Extended release metformin (1000 mg) (n=75) Extended release metformin (1500 mg) (n=71) Immediate release formulin (1000mg) (n=71) Concomitant therapy: None Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes Follow up: 5.5 months | Study location: 42 centres in the United States Sources of funding: NR |

| | | 1.4 | | |
|----------------------------------|---|--|---|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| Gadde 2017 DURATION- NEO-2 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 53.7 (9.4888) years Time since type 2 diabetes diagnosed: 8.3667 (5.6986) years | Strategy: Adding N = 364 Exenatide QWS-AI (n=181) Sitagliptin (100 mg) (n=122) Placebo (n=61) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Health-related quality of life, Non-fatal myocardial infarction, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6.5 months | Study location: 81 centres in the USA Sources of funding: AstraZeneca. Primary author delcares funding from Bristol-Myers Squibb, Eisai and the NIDDK. A second author was an employee of Bristol-Myers Squibb during the conduct of the study and two further authors are employees of AstraZeneca |
| Galindo 2023 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 145 Insulin degludec/liraglutide (n=72) Basal-bolus Insulin (n=73) Concomitant therapy: Oral agents +/or basal insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: US Sources of funding: Grant funding: National Institutes of Health (NIH) and National Institute of Diabetes and Digestive and Kidney DiseaseNovo Nordisk |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|---------------|---|--|---|---|
| | Mean age (SD): 54.15 (9.9006) years Time since type 2 diabetes diagnosed: Not stated/unclear | Insulin: 15.1% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Galle 2012 | Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease: T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 69.25 (8.1689) years Time since type 2 diabetes diagnosed: 13.1 (9.057) years | Strategy: Adding N = 39 Pioglitazone 30 mg once daily (n=20) Placebo (n=19) Concomitant therapy: Insulin Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 6 months | Study location: Germany (12 sites) Sources of funding: Sponsored by TAKEDA Pharma GmbH, Aachen, Germany |
| Gallwitz 2011 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not | Strategy: Adding N = 363 Exenatide (n=182) Insulin aspart 70/30 (n=181) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor | Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: 68 sites in Germany Sources of funding: Two authors are employed by Lilly Deutschland, Germany, a further author is employed by Eli Lilly Austria. |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|-----------------------------|---|---|---|--|
| | stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57 (9.9503) years Time since type 2 diabetes diagnosed: 5 (4.5263) years | agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Gallwitz 2012A | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.8 (9.4) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 1551 Linagliptin (n=776) Glimepiride (n=775) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Nonfatal stroke, Unstable angina, Hospitalisation for heart failure, Falls requiring hospitalisation, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 24 months | Study location: Multicenter Sources of funding: Boehringer Ingelheim. Multiple authors declare funding and honoraria with numerous pharmaceutical companies |
| Gallwitz 2012B EUREXA | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular | Strategy: Adding N = 1029 Exenatide (n=515) Glimegiride (n=514) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear | All-cause mortality, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change | Study location: Multicenter Sources of funding: Eli Lilly and Company; Amlyn Pharmaceuticals. Multiple authors declare funding and honoraria with numerous |

| | | Intervention and | | |
|----------------------------------|--|---|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56 (9.561) years Time since type 2 diabetes diagnosed: 5.65 (4.5571) years | Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | Follow up: 36 months | pharmaceutical companies |
| Gao 2023 SURPASS- AP-Combo | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.125 (11.3268) years Time since type 2 diabetes diagnosed: 7.655 (5.734) years | Strategy: Adding N = 917 Tirzepatide 5mg (n=230) Tirzepatide 10mg (n=228) Tirzepatide 15mg (n=229) Insulin glargine (n=230) Concomitant therapy: Metformin +/- sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 52.5% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, 4-point MACE, Non-fatal myocardial infarction, Non-fatal stroke, Hospitalisation for heart failure, Cardiac arrhythmia, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 9 months | Study location: Multicentre trial (Asia-Pacific region). Sources of funding: Funded by Eli Lilly and Company. |
| Garber 2007 | Model 5: People with type 2 diabetes at higher risk of | Strategy: Adding N = 463 Vildagliptin 50 mg | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c | Study location: USA and Romania |

| | | 1.4 | | |
|-------------|---|---|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | cardiovascular disease T2DM and heart failure: without hear failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.266 (9.333) years Time since type 2 diabetes diagnosis: 4.7 (4.6) years | daily (n=147) Vildagliptin 100 mg daily (n=158) Placebo (n=158) Concomitant therapy: Pioglitazone 45 mg daily Antihyperglycaemic treatment received: No additional information available. | change Follow up: 5.5 months | Sources of funding: Novartis Pharmaceuticals Corporation The findings from this study are included in the pairwise analysis but not the NMA analysis due to this study being identified during quality checks. It was agreed that this was unlikely to change the results of the analysis. |
| Garber 2008 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.2333 (10.7333) years Time since type 2 | Strategy: Adding N = 515 Vildagliptin 50 mg daily (n=170) Vildagliptin 100 mg daily (n=169) Placebo (n=176) Concomitant therapy: Glimepiride 4mg daily Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, hypoglycaemia episodes, severe hypoglycaemia episodes, HbA1c change, weight change Follow up: 5.5 months | Study location: USA, Sweden, Finland, Argentina, and Lithuania Sources of funding: Novartis Pharmaceuticals The findings from this study are included in the pairwise analysis but not the NMA analysis due to this study being identified during quality checks. It was agreed that this was unlikely to change the results of the analysis. |

| | | Intonvoction and | | |
|---------------------------------|---|--|---|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | diabetes diagnosis: 7.133 (5.433) years | | | |
| Garber 2009 LEAD-3 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Mean age (SD): 53.4 (10.9) years Time since type 2 diabetes diagnosed: 5.4 (5.2) years | Strategy: Adding N = 917 Liraglutide 1.2mg (n=251) Liraglutide 1.8mg (n=247) Glimepiride (n=248) Concomitant therapy: No additional information Antihyperglycaemic treatment received: No additional information | HbA1c change Follow up: 12 months | Study location: USA and Mexico. Sources of funding: Novo Nordisk. |
| Garvey 2020 SCALE Insulin | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher | Strategy: Adding N = 396 Liraglutide (n=198) Placebo (n=198) Concomitant therapy: Insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 0.5% Biguanides: 88.60% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 91.9% SGLT-2 inhibitors: | Health-related quality of life, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 13 months | Study location: 53 sites globally Sources of funding: Study was sponsored by Novo Nordisk. The authors declare multiple research grants and honoraria funded by multiple pharmaceutical companies |

| Population cardiovascular risk: Not stated/unclear Mean age (SD): 56.75 (10.8593) years Time since type 2 diabetes diagnosed: 12.1 (6.8502) years Model 5: People | Intervention and comparison 22.2% Sulfonylureas: 35.1% | Outcomes | Comments |
|--|--|---|--|
| risk: Not stated/unclear Mean age (SD): 56.75 (10.8593) years Time since type 2 diabetes diagnosed: 12.1 (6.8502) years Model 5: People | Sulfonylureas: 35.1% | | |
| | | | |
| with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.2 (10.6) years Time since type 2 diabetes diagnosed: 8.5333 (6.5058) years | Strategy: Adding N = 938 Tirzepatide 10mg (n=312) Tirzepatide 15mg (n=311) Placebo (n=315) Concomitant therapy: Biguanide, sulfonylureas, SGLT-2 inhibitors, thiazolidinediones and alpha-glucosidase inhibitors Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 0.8% Biguanides: 88.7% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: 20.4% Sulfonylureas: 26.6% | Health-related quality of life, All-cause mortality, 3-point MACE, Persistent signs of worsening kidney disease, Cardiac arrhythmia, Progression of liver disease, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 18 months | Study location: Multicenter trial. Sources of funding: Funded by Eli Lilly and Company. |
| Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People | Strategy: Adding N = 213 Pioglitazone (n=110) Placebo (n=103) Concomitant therapy: Metformin Antihyperglycaemic | All-cause mortality, Cardiovascular mortality, HbA1c change, Weight change, BMI change Follow up: 5.5 months | Study location: Italy Sources of funding: Takeda Italia SpA, Rome, Italy |
| dT cd sT cris N5T dd8 y N wdricd | iseases '2DM and hronic kidney isease: Not tated/unclear '2DM and higher ardiovascular sk: Not tated/unclear Mean age (SD): 4.2 (10.6) years ime since type 2 iabetes iagnosed: .5333 (6.5058) ears Model 5: People vith type 2 iabetes at higher sk of ardiovascular isease | Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 0.8% Biguanides: 88.7% DPP-4 inhibitors: Not stated/unclear Sk: Not GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: 20.4% Sulfonylureas: iagnosed: .5333 (6.5058) ears Model 5: People vith type 2 iabetes at higher sk of ardiovascular isease Concomitant therapy: Metformin Antihyperglycaemic | Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 0.8% Biguanides: 88.7% DPP-4 inhibitors: Alpha-glucosidase inhibitors: DPP-4 inhibitors: Alpha-glucosidase inhibitors: DPP-4 inhibitors: Alpha-glucosidase inhibitors: DPP-4 inhibitors: Alexander agonists: Not stated/unclear agonists: Not stated/unclear lnsulin: Not stated/unclear SGLT-2 inhibitors: Dean age (SD): Alexander agonists: Dean age (SD): Alexander agonist |

| | | Intervention and | | |
|-----------------------------|---|--|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.4 (8.409) years Time since type 2 diabetes diagnosed: 5.75 (5.0989) years | No additional information available. | | |
| Gerstein 2019A REWIND | Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Mixed population T2DM and chronic kidney disease: Mixed population T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Includes results for a subgroup for | Strategy: Adding N = 9901 Dulaglutide (n=4949) Placebo (n=4952) Concomitant therapy: Antihyperglycemic therapy except DPP-4 inhibitor or GLP-1 receptor agonist Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 81.2% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 23.80% SGLT-2 inhibitors: 0.1% Sulfonylureas: 46% | All-cause mortality, Cardiovascular mortality, 3-point MACE, Non-fatal myocardial infarction, Non-fatal stroke, Unstable angina, Hospitalisation for heart failure, Acute kidney injury, Persistent signs of worsening kidney disease, Development of end stage kidney disease, Cardiac arrhythmia, Progression of liver disease, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 64.8 months | Study location: Multicenter Sources of funding: Eli Lilley & Co. |

| | | 1.4 | | |
|-----------------------------|--|--|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | people with atherosclerotic cardiovascular disease. Mean age (SD): 66.2 (6.5) years Time since type 2 diabetes diagnosed: 10.55 (7.2502) years | | | |
| Giorgino 2015 AWARD-2 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.6667 (9.3503) years Time since type 2 diabetes diagnosed: 9 (6) years | Strategy: Adding N = 807 Dulaglutide 1.5 mg (n=273) Dulaglutide 0.75 mg (n=272) Insulin glargine (n=262) Concomitant therapy: Metformin + glimepiride Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 17.7 months | Study location: NR Sources of funding: Eli Lilly and Company |
| Giugliano 1993 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular | Strategy: Adding N = 50 Metformin (n=27) Placebo (n=23) Concomitant therapy: Insulin Antihyperglycaemic treatment received: No additional | HbA1c change Follow up: 6 months | Study location: Unclear - authors were based in Italy Sources of funding: NR |

| | | lutama attan | | |
|--------------------------|--|--|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 11.6 (1.2468) years Time since type 2 diabetes diagnosed: 11.7 (1.2) years | information available. | | |
| Gohari 2022 EMPA-CARD | Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 62.84 (7.9196) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 95 Empagliflozin 10 mg (n=47) Placebo (n=48) Concomitant therapy: Background oral antidiabetic drugs Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 96.80% DPP-4 inhibitors: 12.60% GLP-1 receptor agonists: Not stated/unclear Insulin: 7.4% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 31.60% | All-cause mortality, Cardiovascular mortality, HbA1c change Follow up: 6 months | Study location: Iran Sources of funding: Dr. Abidi Pharmaceutical company and Zanja University Medical Sciences (Grant Number: 1602001000) |
| Göke 2010 | Model 5: People with type 2 diabetes at higher risk of | Strategy: Adding N = 858 Saxagliptin (n=428) | All-cause mortality, Cardiovascular mortality, | Study location: International, multicentre trial taking place at |

| | | Intervention and | | |
|-----------------|--|--|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.55 (10.3153) years Time since type 2 diabetes diagnosed: 5.45 (4.6013) years | Glipizide (n=430) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 46.80% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 24 months | 130 study sites in Germany, Finland, United Kingdom, Hungary, India, South Korea, Netherlands, Norway, Russia, Slovakia and Vietnam. Sources of funding: Bristol- Myers Squibb and AstraZeneca |
| Goodman 2009 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.8 (10.4914) years Time since type 2 | Strategy: Adding N = 370 Vildagliptin AM (n=125) Vildagliptin PM (n=123) Placebo (n=122) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 5.5 months | Study location: Multicentre trial conducted at 67 centres in the USA and Europe Sources of funding: Novartis Pharmaceutical Corporation |

| | | Intervention and | | |
|--|---|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | diabetes diagnosed: Not stated/unclear | | | |
| Gough 2014 DUAL-I | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55 (9.9266) years Time since type 2 diabetes diagnosed: 6.9333 (5.4147) years | Strategy: Adding N = 1660 Insulin degludec + liraglutide once daily (n=833) Insulin degludec titrated once daily (n=413) Liraglutide 1.8 mg once daily (n=414) Concomitant therapy: Metformin ± pioglitazone Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 457.3% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, 3-point MACE, Non-fatal myocardial infarction, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: Novo Nordisk |
| Gram 2011 South Danish Diabetes Study | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular | Strategy: Adding N = 184 NPH Insulin titrated + placebo twice daily (n=46) NPH insulin titrated + metformin 1000- 2000 mg daily (n=45) Insulin aspart titrated + placebo twice daily (n=48) Insulin aspart titrated + metformin 1000- 2000 mg daily (n=45) Concomitant therapy: Insulin Antihyperglycaemic treatment received: | Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 24 months | Study location: Denmark Sources of funding: No additional information. |

| | | Intervention and | | |
|---------------------|--|--|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | risk: Not stated/unclear Mean age (SD): 56.1 (8.2332) years Time since type 2 diabetes diagnosed: 8.325 (4.6241) years | No additional information available. | | |
| Green 2015 TECOS | Model 1: People with type 2 diabetes and heart failure Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease T2DM and heart failure: People without heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases: T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Includes results for a subgroup for people with or without heart failure. Mean age (SD): 65.45 (7.9502) years Time since type 2 diabetes | Strategy: Adding N = 14671 Sitagliptin (n=7332) Placebo (n=7339) Concomitant therapy: Monotherapy or combination therapy of any approved agent Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 81.6% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 23.2% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 45.3% | All-cause mortality, Cardiovascular mortality, 3-point MACE, 4-point MACE, Non-fatal myocardial infarction, Unstable angina, Hospitalisation for heart failure, Persistent signs of worsening kidney disease, Development of end stage kidney disease, Severe hypoglycaemic episodes, HbA1c change Follow up: 36 months | Study location: Multicenter Sources of funding: Merck Sharp & Dohme; |

| | | Intervention and | | |
|-------------------------------|---|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | diagnosed: 11.6 (8.1) years | | | |
| Grey 2014 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): not stated/unclear Time since type 2 diabetes diagnosed: not stated/unclear | Strategy: Adding Pioglitazone (n=43) Placebo (n=43) Concomitant therapy: Insulin and / or other oral hypoglycaemic Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 20% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | HbA1c change, Weight change Follow up: 12 months | Study location: NR Sources of funding: Grant support from the Health Research Council of New Zealand. Two authors declare funding from multiple pharmaceutical companies |
| Groop 2017 MARLINA- T2D | Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 360 Linagliptin 5mg once daily (n=182) Placebo (n=178) Concomitant therapy: Antidiabetic drugs including insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 34.2% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 31.6% | All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Nonfatal stroke, Hospitalisation for heart failure, Development of end stage kidney disease, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 5.5 months | Study location: International (Canada, Denmark, Finland, France, Germany, Japan, the Philippines, South Korea, Spain, Taiwan, USA and Vietnam) Sources of funding: Supported by the Boehringer Ingelheim and Eli Lilly and Company, Diabetes Alliance. |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|---------------------------------------|---|--|--|---|
| | Mean age (SD): 60.55 (9.6603) years Time since type 2 diabetes diagnosed: Not stated/unclear | SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Group 2022 The GRADE Study | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): Not stated/unclear Mean age (SD): Not stated/unclear Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 5047 Glimepiride (n=1254) Liraglutide (n=1262) Sitagliptin (n=1268) Insulin glargine (n=1263) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Health-related quality of life, All-cause mortality, Cardiovascular mortality, 3-point MACE, Hospitalisation for heart failure, Severe hypoglycaemic episodes, HbA1c change Follow up: 60 months | Study location: 36 clinical centres - it appeared that these were based in the US Sources of funding: National Institute of Diabetes and Digestive and Kidney Diseases and others. The manufacturers contributed trial medications under clinical-trial agreements with the NIDDK but had no role in the design, conduct, or analysis of the trial: donated medications and supplies were from Becton Dickinson, Bristol Myers Squibb, Merck, Novo Nordisk, Roche Diagnostics, and Sanofi. |
| Grunberger 2018 VERTIS RENAL | Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People with chronic | Strategy: Adding N = 467 Ertugliflozin 15 mg once daily (n=155) Ertugliflozin 5 mg once daily (n=158) Placebo (n=154) Concomitant therapy: Antihyperglycemic therapy (monotherapy or combination therapy including sulfonylureas or insulin) | All-cause mortality, Cardiovascular mortality, Hospitalisation for heart failure, Persistent signs of worsening kidney disease, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Multicenter Sources of funding: Merck Sharp & Dohme Corp. subsidiary of Merck & Co., Inc., Kenilworth, NJ, USA and Pfizer Inc. |

| Study Population Ridney disease T2DM and higher cardiovascular risk: Not stated/unclear Siguanides: 24.6% Mean age (SD): 67.2333 (8.5679) years Time since type 2 diabetes at higher risk of cardiovascular disease Concomitant therapy: Metformin a datherosclerotic cardiovascular disease: Not stated/unclear T2DM and a therosclerotic cardiovascular disease: Not stated/unclear T2DM and heart failure: Not stated/unclear risk of cardiovascular disease Not stated/unclear risk of stated/unclear risk of cardiovascular disease Not stated/unclear risk of stated/unclear risk of cardiovascular disease Not stated/unclear rizbM and a therosclerotic cardiovascular disease: Not stated/unclear rizbM and ethoric kidney disease: Not stated/unclear rizbM and higher cardiovascular risk: Not stated/unclear rizbM and ethoric kidney disease: Not stated/unclear rizbM and higher cardiovascular risk: Not stated/unclear rizbM and phane cardiovascular risk: Not stated/unclear rizbM and higher cardiovascular risk: Not stated/unclear rizbM and higher cardiovascular risk: Not stated/unclear rizbM and phane cardiovascular risk: Not stated/unclear risk of cardiovascular risk: Not stated/unclear ri | | | Intervention and | | |
|--|-------|--|---|---|---------------------------------------|
| T2DM and higher cardiovascular risk: Not stated/unclear stated/unclear stated/unclear stated/unclear stated/unclear stated/unclear sulfunith risk of cardiovascular disease (Concomitant therapy: Metfornia and antherosclerotic cardiovascular disease: T2DM and after T2DM and higher cardiovascular disease: T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular disease: Not stated/unclear T2DM and higher data disease: Not stat | Study | | | Outcomes | Comments |
| SPECIFY with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and adherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 53.5 (9.4501) years Guja 2017 DURATION-7 Guja 2017 DURATION-7 Model 5: People with type 2 diabetes at higher risk of cardiovascular disease Concomitant therapy: Metformin, Multicenter Sources of funding: AstraZeneca Sources of funding: AstraZeneca | | T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 67.2333 (8.5679) years Time since type 2 diabetes diagnosed: 14.1667 (8.5452) | treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 24.6% DPP-4 inhibitors: 13.5% GLP-1 receptor agonists: 2.8% Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: | | |
| DURATION-7 with type 2 diabetes at higher risk of cardiovascular disease N = 464 mortality, Cardiovascular mortality, Acute kidney injury, Hypoglycaemia episodes, Severe therapy: Metformin, Multicenter Sources of kidney injury, Hypoglycaemia episodes, Severe hypoglycaemic | | with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 53.5 (9.4501) years Time since type 2 diabetes diagnosed: 5.05 | Strategy: Adding N = 388 Saxagliptin (n=194) Glimepiride (n=194) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information | myocardial infarction, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 11 | 11 sites in China Sources of funding: |
| | | Model 5: People with type 2 diabetes at higher risk of cardiovascular | N = 464 Exenatide (n=233) Placebo (n=231) Concomitant | mortality, Cardiovascular mortality, Acute kidney injury, Hypoglycaemia episodes, Severe | Multicenter Sources of funding: |

| | | Intervention and | | |
|----------------------------|--|---|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.7 (9.6691) years Time since type 2 diabetes diagnosed: 11.3 (6.356) years | Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 51.60% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-3 inhibitors: Not stated/unclear Sulfonylureas: 3.9% | change, Weight change Follow up: 6.4 months | |
| Gullaksen 2023 SEMPA | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 13.5 (67.5463) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 40 Empagliflozin (n=20) Placebo (n=20) Concomitant therapy: None reported Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 90% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 25% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Weight change Follow up: 7.3 months | Study location: Denmark Sources of funding: Novo Nordisk Foundation, Central Denmark Region Research Fund and Danish Medical Associations Research Foundation |
| Guo 2020 | Model 5: People with type 2 | Strategy: Adding N = 96 | All-cause mortality, | Study location: China |
| | , po 2 | | ortanty, | J |

| | | Intervention and | | |
|-------------|--|--|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 52.5667 (6.5977) years Time since type 2 diabetes diagnosed: Not stated/unclear | Insulin glargine (n=32) Liraglutide (n=32) Placebo (n=32) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 6 months | Sources of funding: Natural Science Foundation of Fujian Province and 900 Hospital of the Joint Logistics Team Internal Hospital Project |
| Gurkan 2014 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 52.65 (7.1263) | Strategy: Adding N = 34 Exenatide (n=17) Insulin glargine (n=17) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | HbA1c change, Weight change, BMI change Follow up: 6 months | Study location: Turkey Sources of funding: None |

| | | latementing and | | |
|--------------------|---|--|--|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| · | years Time since type 2 diabetes diagnosed: 7.235 (3.7931) years | | | |
| Guzman 2017 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.45 (8.5079) years Time since type 2 diabetes diagnosed: 10.55 (6.3755) years | Strategy: Adding N = 109 Sitagliptin (n=41) Placebo (n=68) Concomitant therapy: Metformin + sulfonylurea Antihyperglycaemic treatment received: No additional information available. | Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: Eli Lilly and Company |
| Handelsman 2019 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear | Strategy: Adding N = 461 Dapagliflozin + saxagliptin (n=232) Sitagliptin (n=229) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Acute kidney injury, Cardiac arrhythmia, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Hungary, Mexico, Poland, Romania, South Africa and the USA Sources of funding: Bristol- Myers Squibb and AstraZeneca |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|-----------------------------|--|--|---|--|
| | T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.85 (9.2543) years Time since type 2 diabetes diagnosed: 8.05 (5.4574) years | | | |
| Hanefeld 2004 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 60 (8.4089) years Time since type 2 diabetes diagnosed: 7.05 (5.6) years | Strategy: Adding N = 639 Pioglitazone 15-45 mg daily (n=319) Metformin 850-2550 mg daily (n=320) Concomitant therapy: Sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 31% | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 12 months | Study location: International (Canada, Belgium, Denmark, Estonia, Finland, Hungary, Italy, Lithuania, Netherlands, Slovak Republic, Sweden, UK) Sources of funding: Takeda Europe R&D Centre and Eli Lilly and Company, USA |
| Hanefeld 2011 PIOCOMB | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic | Strategy: Adding N = 121 Metformin (n=42) Pioglitazone (n=40) Pioglitazone + metformin (n=39) Concomitant therapy: Insulin glargine Antihyperglycaemic | Hypoglycaemia episodes, HbA1c change Follow up: 6 months | Study location: Germany Sources of funding: NR |

| | | Intervention and | | |
|----------|--|--|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Mean age (SD): 63 (7.4347) years Time since type 2 diabetes diagnosed: 11.0333 (6.1358) years | treatment received: No additional information available. | | |
| Hao 2022 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 51.85 (11.251) years Time since type 2 diabetes diagnosed: 6.3 (5.652) years | Strategy: Adding N = 360 Liraglutide 1.2mg/d (n=180) Dapagliflozin 10mg (n=180) Concomitant therapy: Metformin +/-SU Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Non-fatal myocardial infarction, Unstable angina, Hospitalisation for heart failure, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: China Sources of funding: Supported by the Exceptional Young Talents Fostering Foundation 2021 of the Tianjin Fourth Central Hospital |

| | | Intervention and | | |
|----------------------------------|--|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| Haring 2013 EMPA-REG METSU | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.1 (9.2327) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 669 Empagliflozin 25 mg (n=218) Empagliflozin 10 mg (n=226) Placebo (n=225) Concomitant therapy: Metformin + sulfonylurea Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multicenter Sources of funding: Boehringer Ingelheim and Eli Lilly and Company |
| Haring 2014 EMPA-REG MET | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 638 Empagliflozin 10 mg (n=217) Empagliflozin 25 mg (n=214) Placebo (n=207) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 17.5 months | Study location: Multicenter Sources of funding: Boehringer Ingelheim and Eli Lilly |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|-----------------------------|---|---|---|--|
| | Mean age (SD): 55.7 (9.9379) years Time since type 2 diabetes diagnosed: Not stated/unclear | | | |
| Harreiter 2021 EXENDA | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 60.15 (8.0081) years Time since type 2 diabetes diagnosed: 6.55 (4.9741) years | Strategy: Adding N = 30 Exenatide (n=16) Placebo (n=14) Concomitant therapy: Metformin and dapagliflozin Antihyperglycaemic treatment received: No additional information available. | Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 5.5 months | Study location: Austria Sources of funding: AstraZeneca |
| Hartemann- Heurtier 2009 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney | Strategy: Adding N = 28 Pioglitazone (n=14) Insulin (n=14) Concomitant therapy: Metformin + sulfonylurea Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: France Sources of funding: Public funds from Assistance Publique des Hopitaux de Paris. |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|--------------|---|---|--|---|
| Ciacy | disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 60 (10) years Time since type 2 diabetes diagnosed: 12 (5.3033) years | Companion | Cutcomes | |
| Hattori 2018 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.75 (11.0809) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 102 Empagliflozin (n=51) Placebo (n=51) Concomitant therapy: Medical treatment other than SGLT2 inhibitors Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 14.7% Biguanides: 23.5% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 20.6% | HbA1c change, BMI change Follow up: 12 months | Study location: Japan Sources of funding: None |
| Heine 2005 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and | Strategy: Adding N = 549 Exenatide (n=282) Insulin glargine (n=267) Concomitant therapy: Metformin + sulfonylurea Antihyperglycaemic | Health-related quality of life, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: Eli Lilly and Company, Inc., and Amylin Pharmaceuticals, Inc. |

| | | Intorvontion and | | |
|-------------------|--|---|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.9 (9.1471) years Time since type 2 diabetes diagnosed: 9.55 (5.856) years | treatment received: No additional information available. | | |
| Heise 2022 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 61.7333 (6.802) years Time since type 2 diabetes diagnosed: 11.3067 (6.1568) years | Strategy: Adding N = 117 Tirzepatide (n=45) Semaglutide (n=44) Placebo (n=28) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6.5 months | Study location: 2 centres in Germany Sources of funding: Eli Lilly |
| Henriksen 2011 | Model 5: People with type 2 diabetes at higher risk of | Strategy: Adding N = 211 Pioglitazone (n=102) | All-cause mortality, Cardiovascular mortality, Non- | Study location: Denmark, Sweden, Finland, |

| | | Intervention and | | |
|--|--|---|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 60.5 (8.1964) years Time since type 2 diabetes diagnosed: 13.2 (7.3485) years | Placebo (n=109) Concomitant therapy: Insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 20.5% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 1.5% | fatal myocardial infarction, HbA1c change, Weight change Follow up: 6 months | Sources of funding: Den Danske Forskningsfond [Authors were also employees of and owned stocks in Nordic Bioscience] |
| Hermansen 2007 - Stratum 1 Sitagliptin 035 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.8 (10.2501) years | Strategy: Adding N = 212 Stratum 1 - Sitagliptin (n=106) Stratum 1 - Placebo (n=106) Concomitant therapy: Glimepiride Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Methods state that the study is multinational, but no further information provided Sources of funding: Merck & Co. Inc |

| | | Intervention and | | |
|--|---|--|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | Time since type 2 diabetes diagnosed: 7.6 (5.7987) years | | | |
| Hermansen 2007 - Stratum 2 Sitagliptin 035 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.15 (8.8495) years Time since type 2 diabetes diagnosed: 9.95 (6.2669) years | Strategy: Adding N = 229 Stratum 2 - Sitagliptin (n=116) Stratum 2 - Placebo (n=113) Concomitant therapy: Glimepiride + metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Methods state that the study is multinational, but no further information provided Sources of funding: Merck & Co. Inc |
| Hiramatsu 2018 | Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher cardiovascular risk: Not | Strategy: Switching N = 139 Liraglutide 0.9 mg daily (n=45) Sitagliptin 50 mg daily (n=49) Linagliptin 5 mg daily (n=45) Concomitant therapy: None, insulin, alphaglucosidase inhibitor, or glinide Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 11.2% Biguanides: Not stated/unclear | All-cause mortality, Non- fatal myocardial infarction, Hospitalisation for heart failure, Development of end stage kidney disease, Cardiac arrhythmia, HbA1c change, BMI change Follow up: 48 months | Study location: Konan City, Japan Sources of funding: None reported |

| | | Intervention and | | |
|--------------------------------|--|--|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | Mean age (SD): 69.8 (7.4292) years Time since type 2 diabetes diagnosed: 8.7667 (6.3418) years | DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 9.2% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Hollander 2009 CV181-013 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.0333 (10.1333) years Time since type 2 diabetes diagnosed: 5.2 (5.2079) years | Strategy: Adding N = 565 Saxagliptin 2.5 mg (n=195) Saxagliptin 5 mg (n=186) Placebo (n=184) Concomitant therapy: Thiazolidinedione Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 17.5 months | Study location: Multicentre study in the US Sources of funding: Bristol- Myers Squibb and AstraZeneca |
| Hollander 2018 VERTIS SU | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not | Strategy: Adding N = 1326 Ertugliflozin 15 mg (n=441) Ertugliflozin 5 mg (n=448) Glimepiride (n=437) Concomitant therapy: Metformin and sitagliptin Antihyperglycaemic treatment received: | All-cause mortality, Cardiovascular mortality, Acute kidney injury, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change | Study location: Multicenter Sources of funding: Merck Sharp & Dohme Corp (subsidiary of Merck & Co.) and Pfizer Inc. |

| | | Intervention and | | |
|-----------------------|--|---|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.2 (9.6062) years Time since type 2 diabetes diagnosed: 7.4667 (5.6672) | No additional information available. | Follow up: 12 months | |
| Holman 2017 EXSCEL | Model 1: People with type 2 diabetes and heart failure Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Mixed population T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease | Strategy: Adding N = 14752 Exenatide (n=7356) Placebo (n=7396) Concomitant therapy: Up to three oral glucose-lowering agents or insulin plus two oral glucose-lowering agents Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 76.6% DPP-4 inhibitors: 15.00% GLP-1 receptor agonists: 0% Insulin: 13.8% SGLT-2 inhibitors: 1.00% Sulfonylureas: 36.60% | All-cause mortality, Cardiovascular mortality, 3-point MACE, Unstable angina, Hospitalisation for heart failure, Development of end stage kidney disease, Death from renal causes, Cardiac arrhythmia, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 38.4 months | Study location: Multicenter Sources of funding: Amylin Pharmaceuticals |

| | | 1.4 | | |
|------------------------|--|--|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | Includes results for a subgroup for people with heart failure and atherosclerotic cardiovascular disease. Mean age (SD): Not stated/unclear Time since type 2 diabetes diagnosed: Not stated/unclear | | | |
| Home 2015 HARMONY 5 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.7 (9.4576) years Time since type 2 diabetes diagnosed: 9.25 (6.1) years | Strategy: Adding N = 404 Pioglitazone (n=288) Placebo (n=116) Concomitant therapy: Metformin + sulfonylurea Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, 3-point MACE, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Multicenter Sources of funding: sponsored by GlaxoSmithKline. |
| Hong 2012 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not | Strategy: Adding N = 140 Sitagliptin (100 mg) (n=70) Insulin (n=70) Concomitant therapy: Insulin | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Korea Sources of funding: National Rearsch Foundation grant funded by the Korean government and |

| | | Intervention and | | |
|-----------|---|---|---|--|
| Study | stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.2 (13.6655) years Time since type 2 diabetes diagnosed: Not stated/unclear | Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 37% Biguanides: 43.6% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 24.2% | Outcomes | from a grant from the Seoul National University Bindang Hospital |
| Hong 2023 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.5 (9.5216) years Time since type 2 diabetes diagnosed: 12.4667 (7.0271) years | Strategy: Adding N = 78 Dapagliflozin 10 mg daily (n=26) Sitagliptin 100 mg daily (n=26) Lobeglitazone 0.5 mg daily (n=26) Concomitant therapy: Metformin + a sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 100% | Hospitalisation for heart failure, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, BMI change Follow up: 24 months | Study location: Seongnam, Gyeonggi, South Korea Sources of funding: Supported by grants from the Korean Diabetes Association (S.L., 2015F-7) and Seoul National University Bundang Hospital (14- 2015-0014) |

| | | Intervention and | | |
|--------------------------|---|---|---|--|
| Study | Population | comparison | Outcomes | Comments |
| Husain 2019 PIONEER 6 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Mixed population T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Mean age (SD): 66 (7) years Time since type 2 diabetes diagnosed: 14.9 (8.5) years | Strategy: Adding N = 3183 Semaglutide (n=1591) Placebo (n=1592) Concomitant therapy: Standard of care Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 2.5% Biguanides: 77.40% DPP-4 inhibitors: 0.00% GLP-1 receptor agonists: 0.00% Insulin: 60.6% SGLT-2 inhibitors: 9.6% Sulfonylureas: 32.20% | All-cause mortality, Cardiovascular mortality, 3-point MACE, 5-point MACE, Non-fatal myocardial infarction, Non-fatal stroke, Unstable angina, Hospitalisation for heart failure, Acute kidney injury, Severe hypoglycaemic episodes Follow up: 15.9 months | Study location: Multicenter Sources of funding: Novo Nordisk |
| lacobellis 2017 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular | Strategy: Adding N = 95 Liraglutide (n=54) Control (n=41) Concomitant therapy: None (all people received metformin before entering the trial) Antihyperglycaemic treatment received: No additional information available. | HbA1c change, BMI change Follow up: 6 months | Study location: US - Report describes that screening occurred at the University of Miami Sources of funding: Novo Nordisk |

| | | Intervention and | | |
|--------------------|---|--|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | risk: Not stated/unclear Mean age (SD): 51 (10) years Time since type 2 diabetes diagnosed: 3.65 (3.4434) years | | | |
| lacobellis 2020 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 51.5 (10.0499) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 100 Dapagliflozin (n=50) Placebo (n=50) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | HbA1c change, Weight change, BMI change Follow up: 5.5 months | Study location: US - report states that participants were screened at the University of Miami Sources of funding: AstraZeneca |
| lijima 2023 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear | Strategy: Switching N = 32 Semaglutide (n=16) Dulaglutide (n=16) Concomitant therapy: No additional information. Antihyperglycaemic treatment received: No additional | Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Japan Sources of funding: Not stated/unclear |

| | | 1 | | |
|--------------------|---|--|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 62.1 (11.4) years Time since type 2 diabetes diagnosed (SD): 12.9 (10.7) years | information available. | | |
| Ikonomidis 2020 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Mean age (SD): 57.5 (9.5131) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 160 Insulin (n=40) Liraglutide (n=40) Empagliflozin (n=40) Liraglutide + Empagliflozin (n=40) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | HbA1c change, Weight change, BMI change Follow up: 12 months | Study location: Study reports that recruitments was conducted at Attikon Hospital in Athens, Greece Sources of funding: Report states there were no sources of funding |
| Inagaki 2012 | Model 5: People with type 2 diabetes at higher risk of | Strategy: Adding N = 427 2 mg Exenatide QW | All-cause mortality, Cardiovascular mortality, Non- | Study location: Japan Sources of |
| | cardiovascular | (n=215) | fatal stroke, | funding: Eli Lilly |

| | | Intervention and | | |
|--------------|--|--|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.755 (10.8035) years Time since type 2 diabetes diagnosed: 9.035 (6.0253) years | Insulin glargine (n=212) Concomitant therapy: Biguanide ± thiazolidinedione ± sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 67% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | and Company and Amylin Pharmaceuticals Inc. Three authors are employees of Eli Lilly, two others have have received funding or honoraria from multiple pharmaceutical companies |
| Inagaki 2013 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 60.5 (10.8269) years Time since type 2 diabetes | Strategy: Adding N = 352 Linagliptin 5mg (n=228) Metformin (n=124) Concomitant therapy: Monotherapy with a sulfonylurea or a glucosidase inhibitor Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 41.1% Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SUlfonylureas: 59.00% | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 12 months | Study location: 43 centres in Japan Sources of funding: Medical writing funded by Boehringer Ingelheim. Five of the authors are employed by Boehringer Ingelheim. Authors declare multiple funding from numerous pharmaceutical companies |

| | | Intervention and | | |
|--------------------|--|--|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | diagnosed: Not stated/unclear | | | |
| Jabbour 2014 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.9 (10.3003) years Time since type 2 diabetes diagnosed: 5.67 (5.1424) years | Strategy: Adding N = 451 Dapagliflozin 10 mg (n=225) Placebo (n=226) Concomitant therapy: Sitagliptin ± metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: 49.40% GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 11 months | Study location: Conducted in Argentina, Germany, Mexico, Poland, UK and the US Sources of funding: Funded by AstraZeneca and Bristol-Myers Squibb. A number of authors are stockholders and/or employees of AstraZeneca. Primary author belongs to speakers' bureaus for Eli Lilly and Company and Amlyn. |
| Ji 2016B VISION | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not | Strategy: Adding N = 3084 Vildagliptin (n=2573) Metformin (n=511) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: 127 medical centres in China Sources of funding: Study funded by Novartis Pharmaceuticals. Two authors are also employees of Novartis Pharmaceuticals. |

| | | 1.4 | | |
|------------------------------|---|--|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | Mean age (SD): 56.35 (10.6334) years Time since type 2 diabetes diagnosed: 4.2 (4.2167) years | | | |
| Ji 2019 VERTIS Asia | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.4333 (9.1013) years Time since type 2 diabetes diagnosed: 6.9667 (5.0666) years | Strategy: Adding N = 506 Ertugliflozin 5 mg (n=170) Ertugliflozin 15 mg (n=169) Placebo (n=167) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 4.6% Biguanides: 100% DPP-4 inhibitors: 2.6% GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 19.4% | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicentre from China, Hong Kong, Republic of Korea, Philippines and Taiwan Sources of funding: Merck Sharp & Dohme Corp., a subsidiary of Merck & Co., Inc., USA, in collaboration with Pfizer Inc., USA. Several authors are employees of Merck Sharp and Dohme and Pfizer |
| Ji 2021A SUSTAIN China | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not | Strategy: Adding N = 868 Semaglutide 0.5 mg (n=288) Semaglutide 1.0 mg (n=290) Sitagliptin (100 mg) (n=290) Concomitant therapy: Metformin Antihyperglycaemic treatment received: | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, HbA1c change, Weight change, BMI change Follow up: 8.1 months | Study location: 65 sites in Brazil, China, Hong Kong, Taiwan, Republic of Korea, South Africa and Ukraine Sources of funding: Trial was funded by Novo Nordisk A/S Denmark. Multiple authors |

| | | Intervention and | | |
|-------------------------|---|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 53.0333 (10.8072) years Time since type 2 diabetes diagnosed: 6.3667 (5.1702) years | Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 99.8% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | declare employment and funding from Novo Nordisk |
| Ji 2023 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and risease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 60.2333 (8.2885) years Time since type 2 diabetes diagnosed: 14.6433 (7.2422) years | Strategy: Adding N = 219 Empagliflozin 10 mg (n=73) Empagliflozin 25 mg (n=73) Placebo (n=73) Concomitant therapy: Insulin Insulin ± up to 2 additional OADs Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 29.7% Biguanides: 68% DPP-4 inhibitors: 5.9% GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 8.7% | Acute kidney injury, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: 24 centres in China Sources of funding: Funded by Boehringer Ingelheim. Two of the authors are also employees of Boehringer Ingelheim. |
| Joubert 2021 EXEPUMP | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease | Strategy: Adding N = 46 Exenatide (n=28) Placebo (n=18) | Health-related quality of life, All- cause mortality, Cardiovascular mortality, Hospitalisation | Study location: Not clear, likely to be France Sources of |

| | | Intervention and | | |
|------------------|--|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.5 (7.4024) years Time since type 2 diabetes diagnosed: 6.45 (5.8754) years | Concomitant therapy: Insulin Antihyperglycaemic treatment received: No additional information available. | for heart failure, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 6 months | funding: AstraZeneca |
| Kadowaki 2011 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.0667 (9.9449) years Time since type 2 diabetes | Strategy: Adding N = 181 Exenatide 5 ug (n=72) Exenatide 10 ug (n=73) Placebo (n=36) Concomitant therapy: SU monotherapy, combination therapy with SU and BG, or SU and TZD Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: | All-cause mortality, Cardiovascular mortality, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Japan Sources of funding: Amylin Pharmaceuticals and Eli Lilly and Company |

| | | Intervention and | | |
|------------------|---|---|---|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | diagnosed:12.0 (6.5653) years | Not stated/unclear Sulfonylureas: 8.4% | | |
| Kadowaki 2017 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.2 (9.2005) years Time since type 2 diabetes diagnosed: 7.42 (6.1522) years | Strategy: Adding N = 138 Canagliflozin (n=70) Placebo (n=68) Concomitant therapy: Teneligliptin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Diabetic ketoacidosis, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Mulitcentre Sources of funding: Mitsubishi Tanabe Pharma Corporation. Numerous authors declare funding and honoraria from multiple pharmaceutical companies |
| Kaku 2009A | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular | Strategy: Adding N = 169 Pioglitazone + Metformin (n=83) Placebo + Metformin (n=86) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, HbA1c change Follow up: 6.5 months | Study location: Japan Sources of funding: Takeda Pharmaceutical Co., Ltd |

| | | Intervention and | | |
|------------|---|---|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | risk: Not stated/unclear Mean age (SD): 52.5 (8.0589) years Time since type 2 diabetes diagnosed: 5.05 (4.4098) years | | | |
| Kaku 2010 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.6667 (10.347) years Time since type 2 diabetes diagnosed: 10.3333 (6.9814) years | Strategy: Adding N = 264 Liraglutide 0.6 mg (n=88) Liraglutide 0.9 mg (n=88) Placebo (n=88) Concomitant therapy: Sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 69.7% | All-cause mortality, Cardiovascular mortality, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Japan Sources of funding: Novo Nordisk Pharmaceuticals Ltd |
| Kaku 2019A | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic | Strategy: Adding N = 819 Insulin degludec/liraglutide once daily (n=275) Degludec once daily (n=271) Liraglutide once daily (n=273) Concomitant therapy: OADs | Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Japan Sources of funding: Novo Nordisk funded medical writing and editorial support. Two Novo Nordisk employees also provided review |

| | | Intervention and | | |
|------------------|--|--|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | aligned with Japanese clinical practice guidelines: α-glucosidase inhibitors; thiazolidinediones; sodium-glucose co- transporter-2 inhibitors; glinides; metformin; or sulfonylureas Antihyperglycaemic | | and input to the manuscript. |
| | Mean age (SD): 57.1667 (10.0682) years Time since type 2 diabetes diagnosed: Not stated/unclear | treatment received: Alpha-glucosidase inhibitors: 40.7% Biguanides: 46.7% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: 61% Sulfonylureas: 42.3% | | |
| Kanazawa 2010 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 66.5 (10) years Time since type 2 diabetes | Strategy: Adding N = 45 Pioglitazone (n=22) Metformin (n=23) Concomitant therapy: Insulin, Sulfonylurea, Alphaglucosidase inhibitor Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 11.20% Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 55.5% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 24.5% | HbA1c change, Weight change, BMI change Follow up: 12 months | Study location: Shimane University Hospital, Japan Sources of funding: Supported by the Alumni Association of Shimane University School of Medicine and from the Ministry of Science, Education and Culture of Japan |

| | | 1.4 | | |
|-----------------------------|---|---|---|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| Ž | diagnosed: 13 (10.073) years | | | |
| Kaneto 2020 LixiLan JP-L | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 66.5 (10) years Time since type 2 | Strategy: Adding N = 512 IGlarLixi (n=255) Insulin glargine (n=257) Concomitant therapy: No additional information. Antihyperglycaemic treatment received: No additional information. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Japan Sources of funding: Sanofi |
| | diabetes diagnosed: 13 (10.073) years | | | |
| Kang 2021 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: People without chronic kidney disease | Strategy: Adding N = 159 Exenatide 10/20 mcg daily (n=79) Insulin glargine (n=80) Concomitant therapy: Metformin or sulfonylurea or both Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 29.6% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not | HbA1c change, Weight change, BMI change Follow up: 5.5 months | Study location: Chongqing, China Sources of funding: Supported by Project 2014YLC20 of the Xinqiao Hospital, and Project ctstc2015shmszx 120014 and ctstc2015jcsf100 03 of the Chongqing Science and Technology Commission. |

| | | Intervention and | | |
|-----------------------------|---|---|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 48.5 (9.0644) years Time since type 2 diabetes diagnosed: Not stated/unclear | stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 13.8% | | |
| Kawamori 2018 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.9 (10.2122) years Time since type 2 diabetes diagnosed: 8.85 (6.8491) years | Strategy: Adding N = 275 Empagliflozin (n=182) Placebo (n=93) Concomitant therapy: Linagliptin Antihyperglycaemic treatment received: No additional information available. | Cardiovascular mortality, Non-fatal stroke, Hospitalisation for heart failure, Acute kidney injury, Diabetic ketoacidosis, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 12 months | Study location: 40 sites in Japan Sources of funding: Funded by Boehringer Ingelheim and Eli Lilly and Company. Boehringer Ingelheim International GmbH and Nippon Boehringer Ingelheim Co. Ltd were involved in the study design, data collection, data analysis and preparation of the manuscript. A number of authors are employees of Boehringer Ingelheim and others disclose receiving multiple honoraria and funding grants from numerous pharmaceutical companies |
| Kellerer 2022 SUSTAIN 11 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and | Strategy: Adding N = 1748 Semaglutide (n=874) Insulin aspart (n=874) Concomitant therapy: Metformin. Antihyperglycaemic treatment received: | Health-related quality of life, All-cause mortality, Unstable angina, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change | Study location: Multicenter Sources of funding: Novo Nordisk A/S |

| | | Intomorphism and | | |
|---------------------------|---|---|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 61.2 (9.5) years Time since type 2 diabetes diagnosed: 13.4 (6.7) years | No additional information. | Follow up: 12 months | |
| Kendall 2005 Exendin-4 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.3333 (9.6773) years Time since type 2 diabetes diagnosed: 8.9333 (6.1689) years | Strategy: Adding N = 733 Exenatide 20 mcg daily (n=241) Exenatide 10 mcg daily (n=245) Placebo (n=247) Concomitant therapy: Metformin + sulfonylurea Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6.9 months | Study location: USA (91 sites) Sources of funding: Supported by Amylin Pharmaceuticals, CA, USA and Eli Lilly, IN, USA. |
| Kesavadev 2017 SWIM | Model 5: People with type 2 diabetes at higher risk of cardiovascular | Strategy: Adding N = 440 Glimepiride 1-3 mg daily (n=221) | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight | Study location: Kerala, India Sources of funding: Funded |

| | | Intervention and | | |
|-------------|--|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | disease | Sitagliptin 100 mg daily (n=219) | change, BMI change | by grant from Merck & Co., Inc. |
| | T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 50.6 (7.2349) years Time since type 2 diabetes diagnosed: 15.315 (7.265) years | Concomitant therapy: Metformin + sulfonylurea Antihyperglycaemic treatment received: No additional information available. | Follow up: 12 months | |
| Khaloo 2019 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 250 Pioglitazone 30 mg daily (n=125) Sitagliptin 100 mg daily (n=125) Concomitant therapy: Metformin + Gliclazide Antihyperglycaemic treatment received: No additional information available. | HbA1c change, Weight change, BMI change Follow up: 12 months | Study location: Vali-Asr Hospital, Tehran, Iran Sources of funding: Reports that study did not 'receive any specific grant from funding agencies in the public, commercial, or not-for-profit sectors' |

| | | lutementing and | | |
|-----------|---|--|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | Mean age (SD): 61.75 (8.1502) years Time since type 2 diabetes diagnosed: 12.8 (6.5593) years | | | |
| Khan 2022 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 50.85 (8.6026) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 120 Empagliflozin 10/20 mg daily (n=60) Vildagliptin 50/100 mg daily (n=60) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | HbA1c change, Weight change Follow up: 5.5 months | Study location: Karachi, Pakistan Sources of funding: Sponsored by Primary Care Diabetes Association, Pakistan |
| Kim 2018 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular | Strategy: Switching N = 168 Glimepiride + metformin sustained release (n=86) Glimepiride + metformin immediate release (n=82) Concomitant therapy: None Antihyperglycaemic | Hypoglycaemia episodes, At night hypoglycaemic episodes Follow up: 5.5 months | Study location: 11 centres in the Republic of Korea Sources of funding: HANDOK Pharm aceuticals |

| | | Intervention and | | |
|-----------------------|---|---|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | disease: Mixed population T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.8 (9.6119) years Time since type 2 diabetes diagnosed: 10.4 (6.8606) years | treatment received: No additional information available. | | |
| Kim 2020 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.6 (9.757) years Time since type 2 diabetes diagnosed: 10.15 (7.5045) years | Strategy: Adding N = 135 Pioglitazone 15 mg daily (n=69) Glimepiride 2 mg daily (n=66) Concomitant therapy: Metformin + Alogliptin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: 100% GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | Falls requiring hospitalisation, Hypoglycaemia episodes, HbA1c change Follow up: 6 months | Study location: Korea Sources of funding: This study was funded by Takeda Pharmaceuticals Korea Co. |
| Kimura 2023 COMING | Model 3: People with type 2 diabetes and chronic kidney disease | Strategy: Adding N = 120 Dulaglutide (n=59) Semaglutide (n=61) | Health-related quality of life, Hospitalisation for heart failure, Diabetic ketoacidosis, | Study location: Japan Sources of funding: Supported by |
| | T2DM and heart failure: Not | Concomitant therapy: Various | Progression of liver disease, | Research Project Grants from the |

| | | Intervention and | | |
|-------------------|--|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 62.7 (10.7586) years Time since type 2 diabetes diagnosed: 13.9 (7.4176) years | Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 3.8% Biguanides: 84.1% DPP-4 inhibitors: 75.60% GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: 64.4% Sulfonylureas: 22.4% | Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Kawasaki Medical School (R03B-058 and R04B-009). |
| Kinoshita 2020 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.5667 years Time since type 2 diabetes diagnosed: 7.2333 years | Strategy: Adding N = 110 Pioglitazone 7.5-15 mg daily (n=36) Glimepiride 0.5-1 mg daily (n=34) Dapagliflozin 5 mg daily (n=40) Concomitant therapy: Background anti-diabetic drugs Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 12.3% Biguanides: Not stated/unclear DPP-4 inhibitors: 66.2% GLP-1 receptor agonists: 1% Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 6.5 months | Study location: Japan (7 hospitals) Sources of funding: Supported in part by Research Project Grant 29G-002, Kawasaki Medical School, Japan. |
| Kohan 2014 | Model 3: People with type 2 | Strategy: Adding N = 352 | All-cause mortality, | Study location: 111 sites in |
| | | | • | |

| | | 1.4 | | |
|------------------------------|--|--|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 67 (8.4654) years Time since type 2 diabetes diagnosed: 16.9333 (9.5345) years | Dapagliflozin 5mg (n=83) Dapagliflozin 10mg (n=85) Placebo (n=184) Concomitant therapy: Antidiabetic drugs including insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 65.1% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 25% | Cardiovascular mortality, Acute kidney injury, Persistent signs of worsening kidney disease, Development of end stage kidney disease, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 24 months | United States, Argentina, Canada, India, Mexico, Peru, Italy, Australia, France, Spain, Denmark, Puerto Rico, and Singapore. Sources of funding: Bristol- Myers Squibb and AstraZeneca- supported study |
| Komorizono 2020 J-LINK | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 50 Linagliptin (n=25) Metformin (n=25) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, HbA1c change, Weight change Follow up: 12 months | Study location: 10 medical institutions in Kagoshima, Japan Sources of funding: Boehringer Ingelheim and Eli Lilly Company |

| | | Intervention and | | |
|-------------------|--|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | Mean age (SD): 52.5 (10.5043) years Time since type 2 diabetes diagnosed: Not stated/unclear | | | |
| Kooy 2009 HOME | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 61.5 (10.5093) years Time since type 2 diabetes diagnosed: 13 (8.5173) years | Strategy: Adding N = 390 Metformin (n=196) Placebo (n=194) Concomitant therapy: Insulin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hospitalisation for heart failure, HbA1c change, Weight change, BMI change Follow up: 52 months | Study location: 3 sites in the Netherlands Sources of funding: Supported by grants from Altana; Lifescan; E. Merck/Sante'; Merck, Sharpe, & Dohme; and Novo Nordisk |
| Kothny 2013 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People | Strategy: Adding N = 449 Vildagliptin 50 mg daily (n=228) Placebo (n=221) Concomitant therapy: Insulin ± metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 5.5 months | Study location: Multicentre trial conducted in Europe, Asia, Australia and Central America Sources of funding: Novartis Pharmaceuticals corporation for which two authors are also employees. Several authors declared |

| | | Intervention and | | |
|--------------------------------|--|--|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.2 (9.9989) years Time since type 2 diabetes diagnosed: 13.05 (7.4091) years | Biguanides: 61.5% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 60.6% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | honoraria for multiple pharmaceutical companies |
| Kothny 2015 | Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 66.8 (9.1593) years Time since type 2 diabetes diagnosed: 19.25 (10.2266) years | Strategy: Adding N = 148 Vildagliptin (n=83) Sitagliptin (n=65) Concomitant therapy: Pre-existing treatment Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 61.7% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 10.8% | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, HbA1c change Follow up: 5.5 months | Study location: 6 centres in Brazil and 81 centres in the US Sources of funding: Novartis Pharma |
| Kovacs 2014 EMPA-REG PIO | Model 5: People with type 2 diabetes at higher risk of cardiovascular | Strategy: Adding N = 498 Empagliflozin 10 mg (n=165) | All-cause mortality, Hypoglycaemia episodes, HbA1c change, Weight | Study location: Multicenter Sources of funding: |
| | disease | Empagliflozin 25 mg | change | Boehringer |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|------------------------|--|---|--|---|
| | T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.5 (9.7838) years Time since type 2 diabetes diagnosed: Not stated/unclear | (n=168) Placebo (n=165) Concomitant therapy: Pioglitazone ± Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Follow up: 17.6 months | Ingelheim and Eli Lilly. A number of authors are employees of Boehringer Ingelheim |
| Koyama 2014 PioRAGE | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 64.9 years | Strategy: Adding N = 63 Pioglitazone (n=31) Glimepiride (n=32) Concomitant therapy: Sulfonylurea or glinide or no treatment Antihyperglycaemic treatment received: No additional information available. | Non-fatal myocardial infarction, Non- fatal stroke, HbA1c change, Weight change Follow up: 5.5 months | Study location: Japan Sources of funding: Ministry of Education, Culture, Sports, Science and Technology, Japan. |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|-------------------------------|--|---|---|--|
| - Ciacy | Time since type 2 diabetes diagnosed: Not stated/unclear | Companion | | |
| Langenfeld 2005 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 62.5 (7.5307) years Time since type 2 diabetes diagnosed: 7.15 (7.2536) years | Strategy: Adding N = 179 Pioglitazone 45 mg daily (n=92) Glimepiride 1-6 mg daily (n=87) Concomitant therapy: Background anti-diabetic drugs Antihyperglycaemic treatment received: No additional information available. | Hospitalisation for heart failure, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, BMI change Follow up: 5.5 months | Study location: Germany Sources of funding: Unrestricted grant from Takeda Pharma GmbH, Germany |
| Lavalle- Gonzalez 2013A | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular disease T2DM and chronic kidney | Strategy: Adding N = 1284 Canagliflozin 300mg (n=367) Canagliflozin 100mg (n=368) Sitagliptin (n=366) Placebo (n=183) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: Janssen Research & Development, LLC |

| | | Intomos Com | | |
|-------------------------------|---|---|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.4 (9.5) years Time since type 2 diabetes diagnosed (SD): 6.9 (5.3) years | | | |
| Lavalle- Gonzalez 2013B | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular disease T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.4 (9.5) years Time since type 2 diabetes diagnosed (SD): 6.9 (5.3) years | Strategy: Adding N = 1284 Canagliflozin 300mg (n=367) Canagliflozin 100mg (n=368) Sitagliptin (n=366) Placebo (n=183) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemia episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Multicenter Sources of funding: Janssen Research & Development, LLC |
| Ledesma 2019 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease | Strategy: Adding N = 302 Linagliptin 20 mg daily (n=151) Placebo (n=151) Concomitant therapy: Basal | All-cause mortality, Cardiovascular mortality, 4-point MACE, Non-fatal myocardial infarction, Non- fatal stroke, Hospitalisation | Study location: Multicenter Sources of funding: Supported by Boehringer Ingelheim and Eli Lilly & Co. and |

| | | Intervention and | | |
|-----------|---|---|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 72.4 (5.3558) years Time since type 2 diabetes diagnosed: Not stated/unclear | insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 28.00% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | for heart failure, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 5.5 months | the Diabetes Alliance. |
| Lee 2013B | Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases: People with atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 61.1 (9.145) years Time since type 2 diabetes | Strategy: Adding N = 121 Pioglitazone 15 mg (n=60) Placebo (n=61) Concomitant therapy: pre-existing treatment: insulin , metformin , glimepride , sulfonylurea , α-glycosidase inhibitor Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 2.5% Biguanides: 34.80% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 10.00% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 61.20% | All-cause mortality, Non-fatal myocardial infarction, HbA1c change Follow up: 12 months | Study location: South Korea Sources of funding: This study was supported by a grant from the Korean Heath Technology R&D Project, Ministry of Health and Welfare, Republic of Korea (A070001). |

| | | Intervention and | | |
|-----------------------------|--|---|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | diagnosed: 5.765 (6.7493) years | | | |
| Lee 2022 DISTINCTIO N | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.75 (9.0529) years Time since type 2 diabetes diagnosed: 18.2 (9.0455) years | Strategy: Adding N = 60 Dapagliflozin 10 mg daily (n=30) Sitagliptin 100 mg daily (n=30) Concomitant therapy: Insulin +/-metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Diabetic ketoacidosis, HbA1c change, Weight change Follow up: 5.5 months | Study location: Hong Kong, P.R. of China Sources of funding: Supported in part by funding from AstraZeneca, and from Endowment Fund awarded to Dr K.CB. Tan. |
| Leiter 2014 | Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear | Strategy: Adding N = 962 Dapagliflozin 10 mg (n=480) Placebo (n=482) Concomitant therapy: Pre-existing treatment excluding rosiglitazone Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Hospitalisation for heart failure, Acute kidney injury, Persistent signs of worsening kidney disease, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Canada, Australia, Chile, Argentina, and five European countries (not specified) Sources of funding: Funded by Astra Zeneca and Bristol-Myers Squibb |

| | | lutama vers | | |
|----------|--|--|---|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 63.75 (7.3055) years Time since type 2 diabetes diagnosed: 13.25 (8.3008) years | Insulin: 94.5% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Li 2014A | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 47.1 (10.6557) years Time since type 2 diabetes diagnosed: 5.5333 (2.5818) years | Strategy: Adding N = 203 Liraglutide (n=68) Saxagliptin (n=68) Vildagliptin (n=67) Concomitant therapy: Metformin ± sulfonylurea ± alphaglucosidase inhibitor or thiazolidinedione Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 12.3% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 9.1% | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 5.5 months | Study location: Metabolic Disease Hospital of Tianjin Medical University, China Sources of funding: NR |
| Li 2014B | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart | Strategy: Adding N = 208 Saxagliptin (n=71) Vildagliptin (n=69) Sitagliptin (n=68) Concomitant therapy: Metformin + oral hypoglycaemic agent | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, HbA1c change Follow up: 5.5 months | Study location: Tianjin. China Sources of funding: Supported by the National Nature Science Foundation of China and grants from Tianjin |

| | | Intervention and | | |
|----------|--|--|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 46.6333 (10.2368) years Time since type 2 diabetes diagnosed: Not stated/unclear | Antihyperglycaemic treatment received: No additional information available. | | Health Bureau Technology Fund |
| Li 2014C | Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: People with chronic kidney disease: People with chronic kidney disease: T2DM and chronic kidney disease: T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.55 (12.3482) | Strategy: Adding N = 56 Glimepiride (n=29) Insulin (higher doses) (n=27) Concomitant therapy: Continuation of insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 36.4% Biguanides: 49.3% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Tianjin, China Sources of funding: National Nature Science Foundation of China, Tianjin Health Bureau Technology, Science and Technology Development Foundation of Tianjin Advanced College |

| | | Intervention and | | |
|---------------------------------|---|--|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | years Time since type 2 diabetes diagnosed: 15.5 (5.946) years | | | |
| Li 2017 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.4 (8.3961) year years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 33 Vildagliptin 50 mg twice daily (n=17) Placebo twice daily (n=16) Concomitant therapy: Insulin +/-Metformin Antihyperglycaemic treatment received: No additional information available. | Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: China Sources of funding: Science and Technology Support Program of Jiangsu Province (CN) (no. BL2014010) and by the China Postdoctoral Science Foundation (no. 2015M581829). |
| Lind 2015 MDI Liraglutide | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and | Strategy: Adding N = 124 Liraglutide 0.6 mg - 1.8 mg daily (n=64) Placebo (n=60) Concomitant therapy: Insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 71.00% DPP-4 inhibitors: Not stated/unclear | Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 5.5 months | Study location: Sweden Sources of funding: NovoNordisk provided financial support and study drugs but did not play a role in the design and execution of the trial. |

| | | Intervention and | | |
|---------------------------|--|---|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 40 (64.5653) years Time since type 2 diabetes diagnosed: 17.15 (7.8458) years | GLP-1 receptor agonists: Not stated/unclear Insulin: 100% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Lingvay 2016 DUAL V | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.75 (9.5528) years Time since type 2 diabetes diagnosed: 11.485 (7.0271) years | Strategy: Adding N = 557 Insulin Degludec/Liraglutide 50 U/1.8 mg once daily (n=278) Insulin glargine once daily (n=279) Concomitant therapy: Insulin glargine + metformin Antihyperglycaemic treatment received: No additional information available. | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Non-fatal stroke, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: NovoNordisk |
| Lingvay 2019 SUSTAIN 8 | Model 5: People with type 2 diabetes at higher risk of | Strategy: Adding N = 788 | All-cause mortality, Cardiovascular | Study location: Multicenter Sources of |
| | cardiovascular | Semag | mortality, Acute kidney injury, | Journes of |

| Study | Population | Intervention and | Outcomes | Commonts |
|----------|--|---|---|--|
| Study | T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Mixed population T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.6 (10.9018) years Time since type 2 diabetes diagnosed: 7.35 (5.6555) years | lutide 1.0 mg once weekly (n=394) Canagliflozin 300 mg once daily (n=394) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 394% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | Outcomes Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | funding: Novo Nordisk |
| Liu 2013 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.1 (8.6052) | Strategy: Adding N = 120 Pioglitazone 30 mg daily (n=60) Sitagliptin 100 mg daily (n=60) Concomitant therapy: Metformin or sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 91% | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Taiwan Sources of funding: The study was supported by the Mackay Memorial Hospital. The sponsor of the study was not directly involved in study design. |

| | | Intervention and | | |
|--------------|---|---|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | years Time since type 2 diabetes diagnosed: 7.8 (4.1049) years | | | |
| Liu 2021 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.55 (10.3005) years Time since type 2 diabetes diagnosed: 11.85 (6.1502) years | Strategy: Adding N = 106 Linagliptin (n=53) Empagliflozin (n=53) Concomitant therapy: Insulin +/-OAD Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 100% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Single centre Sources of funding: NR |
| Liutkus 2010 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher | Strategy: Adding N = 165 Exenatide 10 mcg twice daily (n=111) Placebo (n=54) Concomitant therapy: Metformin and thiazolidinedione Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: This study was sponsored by Amylin Pharmaceuticals, Inc. and Eli Lilly and Company. |

| | | Intervention and | | |
|--------------------------|--|--|--|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | cardiovascular risk: Not stated/unclear Mean age (SD): 54.5 (8.3383) years Time since type 2 diabetes diagnosed: 6.35 (4.3341) years | GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Ludvik 2018 AWARD-10 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.2733 (9.3205) years Time since type 2 diabetes diagnosed: 9.3767 (6.1526) years | Strategy: Adding N = 424 dulaglutide 1.5 mg (n=142) dulaglutide 0.75 mg (n=142) placebo (n=140) Concomitant therapy: SGLT2 inhibitor +/- metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 95.3% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Unstable angina, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Austria, Czechia, Germany, Hungary, Israel, Mexico, Puerto Rico, Spain, United States Sources of funding: Eli Lilly and Company |
| Ludvik 2021 SURPASS-3 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic | Strategy: Adding N = 1444 Tirzepatide (n=1079) Insulin degludec (n=365) Concomitant therapy: Metformin alone or in combination with an SGLT2 inhibitor | All-cause mortality, Cardiovascular mortality, 4-point MACE, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change | Study location: Multinational - Argentina, Austria, Greece, Hungary, Italy, Poland, Puerto Rico, Romania, South Korea, Spain, Taiwan, Ukraine, USA Sources of funding: |

| | | Intervention and | | |
|--|---|--|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.45 (10.0253) years Time since type 2 diabetes diagnosed: 8.3 (6.2256) years | Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 68% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | Follow up: 12 months | Conducted by employees and shareholders of Eli Lilly and Company |
| Lukashevich 2011 moderate renal impairment | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 68.7 (8.1764) years Time since type 2 diabetes diagnosed: 15.1 (9.505) years | Strategy: Adding N = 294 Moderate RI: Vildagliptin (n=165) Moderate RI: Placebo (n=129) Concomitant therapy: Untreated or treated with sulfonylurea, a glucosidase inhibitor, thiazolidinedione, insulin, meglitinide, or combination Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 55.20% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes Follow up: 52 months | Study location: 108 centres worldwide Sources of funding: Four of the five authors are employees of Novartis. The remaining authors declares honoraria and funding from multiple pharmaceutical companies Lukashevic 2011 moderate renal impairment and severe renal impairment are the same study, just reporting different populations. |
| Lukashevich 2011 severe | Model 5: People with type 2 | Strategy: Adding N = 221 | All-cause mortality, | Study location: 108 centres |
| renal impairment | diabetes at higher risk of | Severe RI: | Hypoglycaemia episodes, Severe | worldwide |

| | | Intervention and | | |
|---------------------|---|---|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 64.3 (9.9331) years Time since type 2 diabetes diagnosed: 18.15 (9.052) years | Vildagliptin (n=124) Severe RI: Placebo (n=97) Concomitant therapy: Untreated or treated with sulfonylurea, a glucosidase inhibitor, thiazolidinedione, insulin, meglitinide, or combination Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 69.1% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | hypoglycaemic episodes Follow up: 52 months | Sources of funding: Four of the five authors are employees of Novartis. The remaining authors declares honoraria and funding from multiple pharmaceutical companies Lukashevic 2011 moderate renal impairment and severe renal impairment are the same study, just reporting different populations. |
| Lukashevich 2014 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Mixed population T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.15 (10.6623) | Strategy: Adding N = 318 Vildagliptin 50 mg (n=158) Placebo (n=160) Concomitant therapy: Metformin + glimepiride Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes Follow up: 5.5 months | Study location: Australia, Germa ny,Hungary, India , Italy, Mexico, P hilippines, Roma nia Sources of funding: Novartis Pharmaceuticals Corporation |

| | | Intervention and | | |
|--|---|---|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | years Time since type 2 diabetes diagnosed: 7.3 (6.1499) years | | | |
| Lundby- Christensen 2016 CIMT | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 60.65 (8.9022) years Time since type 2 diabetes diagnosed: 12.85 (6.3518) years | Strategy: Adding N = 412 Metformin (n=206) Placebo (n=206) Concomitant therapy: Insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 28.5% | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 18 months | Study location: Eight hospitals in the greater Copenhagen region Sources of funding: Novo Nordisk A/S. Numerous authors declare multiple funding and honoraria from numerous pharmaceutical companies |
| Macauley 2015 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and rated/unclear T2DM and higher | Strategy: Adding N = 44 Vildagliptin 50 mg twice daily (n=22) Placebo twice daily (n=22) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 6 months | Study location: UK Sources of funding: Novartis Pharma AG |

| | | Intervention and | | |
|---|--|---|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | cardiovascular risk: Not stated/unclear Mean age (SD): 62.05 (1.2349) years Time since type 2 diabetes diagnosed: 5.7 (0.7) years | stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Mahaffey 2018 CANVAS/CA NVAS-R | Model 1: People with type 2 diabetes and heart failure Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Mixed population T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Includes results for a subgroup for people with or without heart failure and with atherosclerotic cardiovascular disease. | Strategy: Adding N = 10142 Canagliflozin (n=5795) Placebo (n=4347) Concomitant therapy: Monotherapy or combination therapy of any approved agent Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, 3-point MACE, Non-fatal myocardial infarction, Non-fatal stroke, Hospitalisation for heart failure, Persistent signs of worsening kidney disease, Development of end stage kidney disease, Cardiac arrhythmia, HbA1c change, Weight change Follow up: 43 months | Study location: Multicenter Sources of funding: Supported by Janssen Research & Development, LLC. Medical writing support was funded by Janssen Global Services, LLC. Canagliflozin has been developed by Janssen Research & Development, LLC, in collaboration with Mitsubishi Tanabe Pharma Corp. |

| | | Intomontion of | | |
|-----------------------|---|---|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | Mean age (SD): 63.35 (8.2573) years Time since type 2 diabetes diagnosed: 13.6 (7.743) years | | | |
| Marre 2009 LEAD-1 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.925 (9.6989) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 809 Liraglutide 0.6 mg daily (n=233) Liraglutide 1.2 mg daily (n=228) Liraglutide 1.8 mg daily (n=234) Placebo daily (n=114) Concomitant therapy: Oral drugs Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 202.20% | All-cause mortality, Cardiovascular mortality, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multicenter Sources of funding: Novo Nordisk |
| Marso 2016A LEADER | Model 1: People with type 2 diabetes and heart failure Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic | Strategy: Adding N = 9340 Liraglutide (n=4668) Placebo (n=4672) Concomitant therapy: One or more drugs, insulin, or combination Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 2.8% Biguanides: 76.40% DPP-4 inhibitors: 0.1% GLP-1 receptor | All-cause mortality, Cardiovascular mortality, 3-point MACE, Non-fatal myocardial infarction, Non-fatal stroke, Unstable angina, Hospitalisation for heart failure, Acute kidney injury, Persistent signs of worsening kidney disease, Development of end stage kidney disease, Death | Study location: Multicenter Sources of funding: Novo Nordisk and the National Institutes of Health |

| | | Intervention and | | |
|--------------------------|--|--|---|--|
| Study | Population | comparison | Outcomes | Comments |
| Mans = 2040D | cardiovascular disease: Mixed population T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Includes results for a subgroup for people with or without heart failure. Mean age (SD): 64.3 (7.2) years Time since type 2 diabetes diagnosed: 12.85 (8.0502) years | agonists: 0.00% Insulin: 44.60% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 50.7% | from renal causes, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 42 months | Charles In the Control of the Contro |
| Marso 2016B SUSTAIN 6 | Model 1: People with type 2 diabetes and heart failure Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Mixed population T2DM and chronic kidney disease: Mixed population T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease | Strategy: Adding N = 6594 Semaglutide 0.5 mg (n=826) Semaglutide 1.0 mg (n=822) Placebo 0.5 mg (n=824) Placebo 1.0 mg (n=825) Semaglutide 0.5mg and 1.0mg combined (n=1648) Placebo 0.5mg and 1.0mg combined (n=1649) Concomitant therapy: No more than two oral hypoglycemic drugs ± basal or premixed insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 1.3% Biguanides: 73.2% DPP-4 inhibitors: 0.1% GLP-1 receptor | Health-related quality of life, All-cause mortality, Cardiovascular mortality, 3-point MACE, 5-point MACE, Non-fatal myocardial infarction, Non-fatal stroke, Unstable angina, Hospitalisation for heart failure, Acute kidney injury, Persistent signs of worsening kidney disease, Cardiac arrhythmia, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 25.2 months | Study location: Multicenter Sources of funding: Novo Nordisk |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|------------------------------------|--|---|---|--|
| Siduy | Includes results for a subgroup for people with or without heart failure. Mean age (SD): 64.625 (5.2152) years Time since type 2 diabetes diagnosed: 13.9 (5.7153) years | agonists: Not stated/unclear Insulin: 58% SGLT-2 inhibitors: 0.1% Sulfonylureas: 42.8% | Outcomes | Comments |
| Mathieu 2014 BEGIN: VICTOZA ADD-ON | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular disease T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 61.0 (9.1) years Time since type 2 diabetes diagnosed (SD): 12.4 (6.5) years | Strategy: Adding N = 177 Insulin degludec/Liraglutide (n=88) Insulin degludec/Insulin aspart (n=89) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information. | All-cause mortality, Cardiovascular mortality, Non-fatal stroke, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: Novo Nordisk A/S |
| Mathieu 2015A | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease | Strategy: Adding N = 660 Sitagliptin (n=330) Placebo (n=330) Concomitant therapy: Metformin | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change | Study location: Multicenter Sources of funding: Merck & Co |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|------------------|---|--|--|---|
| | T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular disease T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.8 (9.3) years Time since type 2 diabetes diagnosed (SD): 13.5 (6.2) years | Antihyperglycaemic treatment received: No additional information. | Follow up: 12 months | |
| Mathieu 2015B | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.1 (9.1) years Time since type 2 diabetes | Strategy: Adding N = 320 Dapagliflozin (n=160) Placebo (n=160) Concomitant therapy: Saxagliptin + Metformin Antihyperglycaemic treatment received: No additional information. | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 6 months | Study location: Multicenter Sources of funding: Bristol- Myers Squibb and AstraZeneca. |

| | | Intervention and | | |
|-------------------|--|---|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | diagnosed (SD): 7.6 (6.2) years | | | |
| Matthaei 2015A | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.6 (9.5461) years Time since type 2 diabetes diagnosed: 7.75 (6.4109) years | Strategy: Adding N = 315 Saxagliptin 5 mg daily (n=153) Placebo daily (n=162) Concomitant therapy: Metformin + dapagliflozin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiac arrhythmia, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 12 months | Study location: USA Puerto Rico, Canada, R omania, Russia, Poland, Czech Republic, Mexico . Hungary Sources of funding: Bristol- Myers Squibb and Astra Zeneca |
| Matthaei 2015B | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 218 Dapagliflozin 10 mg daily (n=109) Placebo (n=109) Concomitant therapy: Metformin + sulfonylurea Antihyperglycaemic treatment received: No additional information available. | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: North America (Canada) and Europe (Czech Republic, Germany, Poland, Slovak Republic, and Spain) Sources of funding: Bristol- Myers Squibb and AstraZeneca |

| | | Intervention and | | |
|------------------|--|---|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | Mean age (SD): 61 (9.4533) years Time since type 2 diabetes diagnosed: Not stated/unclear | | | |
| Matthews 2005 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.5 (9.1015) years Time since type 2 diabetes diagnosed: 5.65 (5.1) years | Strategy: Adding N = 632 Pioglitazone 15-45 mg daily (n=319) Gliclazide 80-320 mg daily (n=313) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Non-fatal myocardial infarction, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 12 months | Study location: International (75 centres in Australia, Bulgaria, Czech Republic, France, Germany, Greece, Latvia, Poland, Romania, Turkey) Sources of funding: Takeda Europe R&D Centre and Eli Lilly and Company, USA |
| Matthews 2010 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without | Strategy: Adding N = 3118 Glimepiride 2-6 mg (n=1556) Vildagliptin 50 mg (n=1562) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 1559% | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 24 months | Study location: The study was conducted in 402 sites. Sources of funding: Novartis Pharmaceutical Corporation |

| | | Intervention and | | |
|----------------------------|--|---|--|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Mixed population T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.5 (9.1301) years Time since type 2 diabetes diagnosed: 5.7 (5.1012) years | DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Mattoo 2005 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.85 (6.6792) years Time since type 2 diabetes diagnosed: 162.15 (77.3725) months | Strategy: Adding N = 289 Pioglitazone 30 mg daily (n=142) Placebo (n=147) Concomitant therapy: Insulin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Not available Sources of funding: Sponsored by Eli Lilly and Company, Indianapolis, Indiana, and Takeda Europe R&D Centre, London, United Kingdom. |
| Mazzone 2006 CHICAGO | Model 5: People with type 2 diabetes at higher risk of cardiovascular | Strategy: Adding N = 462 Pioglitazone 15 - 45 mg daily (n=232) | All-cause mortality, Cardiovascular mortality, 3-point MACE, Non-fatal | Study location: US, Chicago at 28 clinical sites. Sources of |

| | | Intervention and | | |
|-------------------|---|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.6 (8.1002) years Time since type 2 diabetes diagnosed: 7.75 (7.2128) years | Glimepiride 1-4 mg daily (n=230) Concomitant therapy: Metformin ± sulfonylurea ± insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 70.5% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 29% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 76% | myocardial infarction, Non-fatal stroke, Unstable angina, Hospitalisation for heart failure, Hypoglycaemia episodes, Weight change Follow up: 18 months | funding: Takeda Pharmaceuticals North America Inc, Lincolnshire, Ill, sponsored and funded this study and provided the study drugs. |
| McCluskey 2004 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.5 (8.5493) years Time since type 2 | Strategy: Adding N = 40 Glimepiride 2-8 mg daily (n=25) Placebo daily (n=15) Concomitant therapy: Rosiglitazone Antihyperglycaemic treatment received: No additional information available. | Follow up: 7 months | Study location: US (17 sites) Sources of funding: No additional information. |

| | | Intervention and | | |
|----------------------------|---|--|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | diabetes diagnosed: 5.9 (7.403) years | | | |
| McGill 2013 | Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 64.45 (10.2854) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 133 Linagliptin (n=68) Placebo (n=65) Concomitant therapy: Insulin, sulfonylurea, glinides, pioglitazone, and a-glucosidase inhibitors Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 64.1% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 12% | All-cause mortality, Non-fatal myocardial infarction, Non-fatal stroke, Acute kidney injury, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Multicenter Sources of funding: Boehringer Ingelheim |
| McMurray 2018 VIVIDD | Model 1: People with type 2 diabetes and heart failure T2DM and heart failure: People with heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 254 Vildagliptin (n=128) Placebo (n=126) Concomitant therapy: Drug therapy, if any Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 2% Biguanides: 34.6% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Nonfatal stroke, Hospitalisation for heart failure, Cardiac arrhythmia, Hypoglycaemia episodes, HbA1c change Follow up: 12 months | Study location: Czechia, Denmark, Estonia, Germany, Greece, Guatemala, India, Italy, Latvia, Lithuania, Poland, Romania, Russian Federation, Singapore, Slovakia [Taken from Clinicaltials.gov] Sources of funding: Novartis |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|----------------------------|---|--|--|--|
| | Mean age (SD): 63.15 (9.3818) years Time since type 2 diabetes diagnosed: 9.3 (7.9526) years | Sulfonylureas: 50.00% | | |
| Meneghini 2010 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 52.35 (10.1997) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 247 Pioglitazone 15 mg - 45 mg daily (n=126) Insulin glargine titrated (n=121) Concomitant therapy: Metformin or sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 26% | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 11 months | Study location: US Sources of funding: One of the authors is employed by Sanofi Aventis group and editorial support was also provided by the Sanofi Aventis U.S. group which suggests that they provided at least some funding towards the study. |
| Meneilly 2017 GetGoal-O | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not | Strategy: Adding N = 350 Lixisenatide (n=176) Placebo (n=174) Concomitant therapy: Permitted therapies were metformin, sulfonylurea (except glibenclamide >10 mg and gliclazide >160 mg), meglitininde (excet | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multicenter Sources of funding: Sanofi. Numerous authors declare funding and honoraria from multiple pharmaceutical companies |

| | | Intervention and | | |
|------------------------|---|--|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 74.2 (3.9019) years Time since type 2 diabetes diagnosed: 14.1 (7.6042) years | repaglinide >6 mg), pioglitazone, and basal insulin) Antihyperglycaemic treatment received: No additional information available. | | |
| Miras 2019 GRAVITAS | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56 (8.9876) years Time since type 2 diabetes diagnosed: 18 (7.3485) years | Strategy: Adding N = 80 Liraglutide (n=53) Placebo (n=27) Concomitant therapy: Oral glucose lowering agents and or insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 25% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Health-related quality of life, All-cause mortality, Persistent signs of worsening kidney disease, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: London, UK Sources of funding: JP Moulton Charitable Foundation. Liraglutide and Placebo pens provided by Novo Nordisk. Multiple authors declare funding and honoraria from numerous pharmaceutical companies |
| Moeinzadeh 2021 | Model 3: People with type 2 diabetes and chronic kidney disease | Strategy: Adding N = 136 Linagliptin (n=68) Placebo (n=68) | HbA1c change, Weight change Follow up: 6 months | Study location: Iran Sources of funding: NR |
| | T2DM and heart failure: Not stated/unclear | Concomitant therapy: Current glucose-lowering | | |

| | | Intervention and | | |
|-----------|--|---|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 60.11 (12.986) years Time since type 2 diabetes diagnosed:Not stated/unclear | Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 19.80% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Moon 2014 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 53.1 (8.3435) years Time since type 2 diabetes diagnosed: 87.3 (66.0836) months | Strategy: Adding N = 75 Glimepiride 1-8 mg daily (n=36) Insulin glargine daily (n=39) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 11 months | Study location: Korea Sources of funding: NR |

| | | Intonvoction and | | |
|-------------------------------|--|---|--|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| Morikawa 2011 APRIME | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 62.45 (1.6593) years Time since type 2 diabetes diagnosed: 10.55 (1.1573) years | Strategy: Adding N = 63 Pioglitazone (n=32) Metformin (n=31) Concomitant therapy: Insulin or oral hypoglycemic agents other than thiazolidinediones/metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 19% Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 62.00% | All-cause mortality, Unstable angina, Persistent signs of worsening kidney disease, HbA1c change Follow up: 12 months | Study location: Japan Sources of funding: Takeda Pharmaceutical Company, Japan |
| Mosenzon 2019 PIONEER 5 | Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease T2DM and chronic kidney disease T2DM and higher | Strategy: Adding N = 324 Semaglutide (n=163) Placebo (n=161) Concomitant therapy: Metformin, a sulfonylurea, or both; or basal insulin with or without metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 74.60% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Hospitalisation for heart failure, Acute kidney injury, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 6 months | Study location: 88 sites in Denmark, Finland, Israel, Poland, Russia, Sweden, the UK, and the USA Sources of funding: Novo Nordisk |

| | | Intomiontion and | | |
|------------|--|--|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | cardiovascular risk: Not stated/unclear Mean age (SD): 70.5 (8) years Time since type 2 diabetes diagnosed: 14 (8.0262) years | stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 40.40% | | |
| Moses 2014 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57 (10.589) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 257 Saxagliptin 5 mg daily (n=129) Placebo daily (n=128) Concomitant therapy: Metformin + sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 100% | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multicenter Sources of funding: Bristol-Myers Squibb and Astra Zeneca |
| Moses 2017 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People | Strategy: Adding N = 427 Sitagliptin 100 mg daily (n=213) Placebo once daily (n=214) Concomitant therapy: Metformin + glimepiride/gliclazide Antihyperglycaemic treatment received: No additional | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: USA Sources of funding: Merck & Co., Inc. (Kenilworth, NJ, USA). |

| | | Intervention and | | |
|-------------------------|--|--|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): | information available. | | |
| | 54.9 (9.9052) years Time since type 2 diabetes diagnosed: 7.75 (5.3501) years | | | |
| Muller- Wieland 2018 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.4 (8.5909) years Time since type 2 diabetes | Strategy: Adding N = 939 Dapagliflozin (n=314) Dapagliflozin + Saxagliptin (n=312) Glimepiride (n=313) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Hospitalisation for heart failure, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: 194 centres in Germany, the Czech Republic, Hungary, Poland and Slovakia Sources of funding: AstraZeneca; Numerous authors declare honoraria and funding from multiple pharmaceutical companies. |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|-------------------|--|--|---|---|
| Nahra 2021 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.4 (9.6498) years Time since type 2 diabetes diagnosed: 7.6 (5.5722) years | Strategy: Adding N = 222 Liraglutide (n=110) Placebo (n=112) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, HbA1c change, Weight change Follow up: 12.5 months | Study location: Multicenter Sources of funding: AstraZeneca. A number of authors are also employees of AstraZeneca |
| Nakaguchi 2020 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 61 Liraglutide (n=30) Empagliflozin (n=31) Concomitant therapy: Insulin +/-OAD Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 22.80% Biguanides: 37.60% DPP-4 inhibitors: 42.7% GLP-1 receptor agonists: Not stated/unclear Insulin: 100% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 1.60% | Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 5.5 months | Study location: Yokahama Japan Sources of funding: Self- procurement with no subsidy |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|-----------------------------------|---|---|--|---|
| Nauck 2007A | Mean age (SD): 66.75 (9.2576) years Time since type 2 diabetes diagnosed: 18.9 (10.0022) years Model 5: People with type 2 | Strategy: Adding N = 501 | All-cause mortality, At night | Study location: Multicenter |
| | diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.5 (9) years Time since type 2 diabetes diagnosed: 9.9 (6.2507) years | Exenatide (n=253) Insulin (n=248) Concomitant therapy: Metformin + sulfonylurea Antihyperglycaemic treatment received: No additional information available. | hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Sources of funding: Industry initiated study. A number of authors are employees of Eli Lilly and Amlyn Pharmaceuticals or declare funding and / or honoraria from Eli Lilly and Amlyn Pharmaceuticals |
| Nauck 2007B Sitagliptin 024 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear | Strategy: Adding N = 1172 Sitagliptin (n=588) Glipizide (n=584) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 24 months | Study location: Multinational study Sources of funding: Merck & Co. |

| | | later a continue de | | |
|-----------------------|--|--|---|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.7 (9.5524) years Time since type 2 diabetes diagnosed: 6.35 (5.7618) years | | | |
| Nauck 2009A | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55 (11) years Time since type 2 diabetes diagnosed: 6 (4.6271) years | Strategy: Adding N = 527 Alogliptin 12.5 mg (n=213) Alogliptin 25 mg (n=210) Placebo (n=104) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: Takeda Global Research and Development Center, Inc. A number of authors are employees of Takeda Global Research and Development Center, Inc. The primary author declares honoraria from numerous pharmaceutical companies |
| Nauck 2009B LEAD-2 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic | Strategy: Adding N = 1091 Liraglutide 0.6 mg (n=242) Liraglutide 1.2 mg (n=241) Liraglutide 1.8 mg (n=242) Glimepiride (n=244) Placebo (n=122) Concomitant | Death from renal causes, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 24 months | Study location: Multicenter Sources of funding: A number of authors were supported by Novo Nordisk. Numerous authors declare |

| Study Population cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular T2DM and higher cardiovascular T2DM and higher cardiovascular Comments Comments Comments Comments Comments Comments Funding and honoraria Antihyperglycaemic treatment received: alientalication honoraria Displayed Comments Funding and honoraria Displayed Comments Funding and honoraria Comments Funding and honoraria Comments | |
|--|-------------|
| disease: Not +/- OADs honoraria stated/unclear T2DM and Antihyperglycaemic chronic kidney treatment received: disease: Not Alpha-glucosidase stated/unclear inhibitors: Not T2DM and higher stated/unclear cardiovascular Biguanides: 31.2% risk: Not DPP-4 inhibitors: | |
| GLP-1 receptor agonists: Not Mean age (SD): stated/unclear 56.75 (8.9876) Insulin: Not years stated/unclear Time since type 2 SGLT-2 inhibitors: diabetes Not stated/unclear diagnosed: 7.5 Sulfonylureas: 3.8% (4.7111) years | |
| Nauck 2011 Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart disease: People without atherosclerotic cardiovascular diseases People without gisease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and stated/unclear T2DM and stated/unclear T2DM and stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and stated/unclear T2DM and stated/unclear T2DM and stated/unclear T2DM and bigher cardiovascular risk: Not stated/unclear T2DM and stated/unclear Sulfonylureas: Not Sulfonylureas | oy a and |
| Nauck 2014 Model 5: People Strategy: Adding All-cause Study location Dulaglutide v with type 2 N = 1098 mortality, Severe US, Canada | |

| | | Intervention and | | |
|--------------------|---|---|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| Placebo AWARD-5 | diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.25 (9.846) year years Time since type 2 diabetes diagnosed: 7 (5.2959) years | Dulaglutide 1.5 mg weekly (n=304) Dulaglutide 0.75 mg weekly (n=302) Sitagliptin 100 mg daily (n=315) Placebo daily (n=177) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | hypoglycaemic episodes, HbA1c change Follow up: 24 months | France, Germany, India, Korea, Mexico, Poland, Puerto Rico, Romania, Russian, Spain and Taiwan. Sources of funding: Eli Lilly and company |
| Nauck 2016B | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.2 (10.3044) | Strategy: Adding N = 404 Liraglutide (n=202) Lixisenatide (n=202) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | Cardiac arrhythmia, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: Sponsored by Novo Nordisk A/S |

| | | Intervention and | | |
|------------------------------|---|--|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | years Time since type 2 diabetes diagnosed: 6.4 (5.1522) years | | | |
| Nesti 2022 EMPA- HEART | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 61.7 (9.8624) years Time since type 2 diabetes diagnosed: 9.45 (7.9421) years | Strategy: Adding N = 56 Empagliflozin (n=27) Sitagliptin (n=29) Concomitant therapy: Metformin and or basal insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 91% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 28.5% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | HbA1c change, Weight change Follow up: 6 months | Study location: Single centre in Pisa, Italy. Sources of funding: Supported at 49% by an unrestricted grant from Boehringer Ingelheim |
| Ning 2016 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear | Strategy: Adding N = 293 Vildagliptin (n=146) Placebo (n=147) Concomitant therapy: Metformin ± insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 71.00% DPP-4 inhibitors: | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 5.5 months | Study location: 22 centres in China, Thailand, Philippines, and Singapore Sources of funding: Novartis Pharma AG (Basel, Switzerland). Fiv e authors were also employed by Novartis and may be eligible for |

| | | Intonvention | | |
|--------------------------|--|---|---|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.1 (9.3542) years Time since type 2 diabetes diagnosed: 11.3 (7.0054) years | Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | Novartis stock and stock options |
| Nissen 2008 PERISCOPE | Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.85 (9.252) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 543 Glimepiride (n=273) Pioglitazone (n=270) Concomitant therapy: 1-2 oral drugs excluding thiazolidinedione Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 64.40% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 20.6% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, 3-point MACE, 5-point MACE, Non-fatal myocardial infarction, Non-fatal stroke, Unstable angina, Hospitalisation for heart failure, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 18 months | Study location: Multicenter trial. Sources of funding: Financially supported by Takeda Pharmaceuticals North America Inc. |
| Nogueira 2014 | Model 5: People with type 2 diabetes at higher risk of cardiovascular | Strategy: Adding N = 35 Sitagliptin (n=18) Insulin NPH (n=17) | HbA1c change, Weight change, BMI change | Study location: Unclear- appears to be Brazil Sources of |
| | disease | | | funding: grants |

| | | Intervention and | | |
|------------------|--|---|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.75 (6.7977) years Time since type 2 diabetes diagnosed: 10.9 (6.6785) | Concomitant therapy: Metformin; Glyburide Antihyperglycaemic treatment received: No additional information available. | Follow up: 5.5 months | from Fundação de Amparo à Pesquisa do Estado de São Paulo. (FAPESP) |
| Nowicki 2011A | Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 66.5 (8.7092) years Time since type 2 diabetes diagnosed: 16.65 (8.0156) years | Strategy: Adding N = 170 Saxagliptin (n=85) Placebo (n=85) Concomitant therapy: Excluded metformin therapy, and previous or current DPP-4 inhibitor or GLP1 receptor agonist Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 75.3% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 25.3% | All-cause mortality, Development of end stage kidney disease, Hypoglycaemia episodes, HbA1c change Follow up: 12 months | Study location: Multi-centre - Belarus, Croatia, Czech Republic, Estonia, Germany, Hungary, Latvia, Lithuania, Poland, Romania, Russia, Ukraine, USA Sources of funding: Bristol- Myers Squibb and AstraZeneca |

| | | lutement to the | | |
|------------------|--|---|---|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| Oh 2021 ELITE | Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease | Strategy: Adding N = 97 Empagliflozin 10 mg (n=48) Sitagliptin 100 mg (n=49) | HbA1c change, Weight change Follow up: 6 months | Study location: South Korea Sources of funding: No information available. |
| | T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: People without chronic kidney disease: T2DM and chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 64.95 (8.4296) years Time since type 2 diabetes diagnosed: 74.2 (69.9855) months | Concomitant therapy: Metformin, sulfonylurea, pioglitazone Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 67% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 33% | | |
| Ohira 2014A | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and ronic kidney disease: Not stated/unclear T2DM and higher | Strategy: Adding N = 60 Pioglitazone (n=30) Glimepiride (n=30) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | HbA1c change, Weight change, BMI change Follow up: 6 months | Study location: Not available Sources of funding: NR |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|-------------|---|---|---|---|
| | cardiovascular risk: Not stated/unclear Mean age (SD): 62.965 (10.3284) years Time since type 2 diabetes diagnosed: Not stated/unclear | | | |
| Ohira 2014B | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 60.37 (11.6992) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 70 Sitagliptin + metformin (n=35) Metformin (n=35) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | HbA1c change, Weight change, BMI change Follow up: 6 months | Study location: Japan Sources of funding: NR |
| Owens 2011 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular | Strategy: Adding N = 1058 Linagliptin (n=793) Placebo (n=265) Concomitant therapy: Metformin + sulfonylurea Antihyperglycaemic treatment received: No additional | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multicenter Sources of funding: Boehringer Ingelheim |

| | | Intoniontian | | |
|--------------------------------|--|---|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.95 (9.8504) years Time since type 2 diabetes diagnosed: Not stated/unclear | information available. | | |
| Pan 2012B | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.1333 (9.7696) years Time since type 2 diabetes diagnosed: 5.03 (4.6019) years | Strategy: Adding N = 438 Vildagliptin 50 mg qd (n=148) Vildagliptin 50 mg bid (n=146) Placebo (n=144) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Nonfatal stroke, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 5.5 months | Study location: Multicentre trial in China Sources of funding: Novartis Beijing |
| Pan 2014 GetGoal-M- Asia | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease | Strategy: Adding N = 391 Lixisenatide 20 mcg daily (n=196) Placebo (n=195) | All-cause mortality, Cardiovascular mortality, Non- fatal myocardial infarction, Non- | Study location: Multicenter Sources of funding: Funded |

| | | Intervention and | | |
|-------------------------|--|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.8 (10.4002) years Time since type 2 diabetes diagnosed: 6.65 (4.7008) years | Concomitant therapy: Metformin ± sulfonylurea Antihyperglycaemic treatment received: No additional information available. | fatal stroke, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | by Sanofi, France. |
| Papathanassi ou 2009 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 28 Pioglitazone 30 mg daily (n=14) Glimepiride 4 mg daily (n=14) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | HbA1c change, Weight change, BMI change Follow up: 6 months | Study location: Ioannina, Greece Sources of funding: Funded in part by Michaelidion Cardiac Center, University of Ioannina, Ioannina, Greece |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|-----------|---|--|---|---|
| | Mean age (SD): 63.2 (7.2502) years Time since type 2 diabetes diagnosed: 5.3 (5.254) years | , | | |
| Park 2011 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 62.7 (8.1994) years Time since type 2 diabetes diagnosed: 5 (5.1499) years | Strategy: Adding N = 67 Pioglitazone 15 mg daily (n=34) Metformin 1000 mg daily (n=33) Concomitant therapy: Glimepiride or other sulfonylurea Antihyperglycaemic treatment received: No additional information available. | HbA1c change, BMI change Follow up: 5.5 months | Study location: Seoul, South Korea Sources of funding: Supported by Faculty research grant of Yonsei University College of Medicine for 2007 and Yonsei University College of Medicine, Internal Medicine Research Grant 2007. |
| Park 2014 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney | Strategy: Adding N = 99 Metformin (n=33) Glimepiride (n=34) Metformin + Glimepiride (n=32) Concomitant therapy: Insulin glargine Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multicentre trial in Korea Sources of funding: Sanofi- Korea |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|---------------------|--|--|--|--|
| | disease: Not stated/unclear T2DM and higher cardiovascular risk: No information Mean age (SD): 56.6333 (10.2084) years Time since type 2 diabetes diagnosed: 12 (6.6126) years | | | |
| Park 2023 BEYOND | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.15 (9.1512) years Time since type 2 diabetes diagnosed: 6.25 (4.8503) years | Strategy: Adding N = 124 Dapagliflozin (n=62) Glimepiride (n=62) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, HbA1c change, Weight change, BMI change Follow up: 12 months | Study location: 14 centres in Korea Sources of funding: AstraZeneca |
| Pasquel 2021 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear | Strategy: Adding N = 287 Liraglutide (n=140) Insulin glargine (n=147) Concomitant therapy: Existing antidiabetic treatment | All-cause mortality, Acute kidney injury, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change | Study location: United States Sources of funding: Novo Nordisk |

| | | Intervention and | | |
|------------------------------|---|--|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56 (10.4056) years Time since type 2 diabetes diagnosed: 9.65 (8.4909) years | Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 15% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Follow up: 6 months | |
| Perkovic 2019 CREDENCE | Model 1: People with type 2 diabetes and heart failure Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: People without heart failure: T2DM and atherosclerotic cardiovascular disease: Mixed population T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Includes results for a subgroup for people with or without heart failure. Mean age (SD): 63.05 (9.2) years Time since type 2 | Strategy: Adding N = 4401 Canagliflozin (n=2202) Placebo (n=2199) Concomitant therapy: angiotensin-converting enzyme inhibitor or angiotensin II receptor blocker Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, 3-point MACE, 5-point MACE, Hospitalisation for heart failure, Acute kidney injury, Persistent signs of worsening kidney disease, Development of end stage kidney disease, Death from renal causes, Diabetic ketoacidosis, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 31.44 months | Study location: Multicenter Sources of funding: Janssen Research & Development, LLC |

| | | Intervention and | | |
|------------------------------|---|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | diabetes diagnosed: 15.75 (8.6502) years | | | |
| Perkovic 2024 FLOW | Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 66.7 (9.0) years Time since type 2 diabetes diagnosed: Majority at 15 years and greater | Strategy: Adding N = 3533 Semaglutide (subcutaneous) (n=1767) Placebo (n=1766) Concomitant therapy: Renin- angiotensin system inhibitor (antihypertensive). Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 51.9% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 61.3% SGLT-2 inhibitors: 15.6% Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, 3-point MACE, Non-fatal myocardial infarction, Non-fatal stroke, Unstable angina, Acute kidney injury, Persistent signs of worsening kidney disease, Development of end stage kidney disease, Death from renal causes, Cardiac arrhythmia, Diabetic ketoacidosis, Severe hypoglycaemic episodes, HbA1c change, Weight Change Follow up: 40.8 months | Study location: Multicenter Sources of funding: Novo Nordisk. |
| Pei 2021 DUAL II China | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 453 IDegLira (n=302) Insulin degludec (n=151) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information | All-cause mortality, Cardiovascular mortality, 3-point MACE, Non-fatal myocardial infarction, Non-fatal stroke, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: China and Hong Kong Sources of funding: Novo Nordisk |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|-----------------------------|--|--|--|---|
| | Mean age (SD): 54.8 (9.9) years Time since type 2 diabetes diagnosed: 11.5 (6.0) years | | | |
| Petrica 2011 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.85 (7.1415) years Time since type 2 diabetes diagnosed: 10.085 (4.4597) years | Strategy: Adding N = 78 Pioglitazone (n=39) Glimepiride (n=39) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | HbA1c change, BMI change Follow up: 12 months | Study location: Department of Diabetes and Metabolic Diseases, Romania Sources of funding: NR |
| Pfützner 2005 PIONEER | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not | Strategy: Adding N = 173 Pioglitazone (n=89) Glimepiride (n=84) Concomitant therapy: Any oral antidiabetic excluding thiazolidinedione treatment for the pioglitazone arm and metformin for the glimepiride arm | Hospitalisation for heart failure, Severe hypoglycaemic episodes, HbA1c change, BMI change Follow up: 6 months | Study location: Clinical Department of the Institute for Clinical Research and Development, Mainz, Germany Sources of funding: Takeda Pharma, Germany. A number of authors declare funding and |

| | | Intervention and | | |
|-----------------------------|--|--|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 62.6 (7.9304) years Time since type 2 diabetes diagnosed: 7.15 (7.2543) years | Antihyperglycaemic treatment received: No additional information available. | | honoraria from Takeda Pharma. |
| Pfützner 2011B PIOfix | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 62.6 (7.9304) years Time since type 2 diabetes diagnosed: 7.15 (7.2543) years | Strategy: Adding N = 288 Pioglitazone + Metformin (n=146) Glimepiride + Metformin (n=142) Concomitant therapy: No additional information. Antihyperglycaemic treatment received: No additional information available. | Hospitalisation for heart failure, Acute kidney injury, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Germany Sources of funding: Takeda Pharma. |
| Pfeffer 2015 ELIXA | Model 1: People with type 2 diabetes and heart failure Model 2: People with type 2 diabetes and | Strategy: Adding N = 6068 Lixisenatide (n=3034) Placebo (n=3034) | All-cause mortality, Cardiovascular mortality, 4-point MACE, 5-point MACE, Non-fatal myocardial | Study location: Multicenter Sources of funding: Funded by Sanofi |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|--------------------------------------|---|---|---|--|
| | atherosclerotic cardiovascular disease T2DM and heart failure: Mixed population T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: People without chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Includes results for a subgroup for people with or without heart failure. Mean age (SD): 60.25 (9.6501) years Time since type 2 diabetes diagnosed: 9.3 (8.2502) years | Concomitant therapy: Antidiabetic medications with the exception of other incretin therapies Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 66.3% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 39.1% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 33.00% | infarction, Non-fatal stroke, Unstable angina, Hospitalisation for heart failure, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 25 months | |
| Philis- Tsimikas 2013 BEGIN | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and | Strategy: Adding N = 458 Sitagliptin 100 mg daily (n=229) Insulin degludec 100 U/mL daily (n=229) Concomitant therapy: Background oral antidiabetic drugs Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not | All-cause mortality, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: Funded by Novo Nordisk, A/S, Denmark. |

| | | Intervention and | | |
|--|---|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.65 (10.8167) years Time since type 2 diabetes diagnosed: 7.75 (6.0519) years | stated/unclear Biguanides: 25.00% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Philis- Tsimikas 2019 DUAL IX | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.65 (10.3005) years Time since type 2 diabetes diagnosed: 9.55 (6.2502) years | Strategy: Adding N = 420 Insulin degludec/liraglutide daily titrated (n=210) Insulin glargine U100 daily titrated (n=210) Concomitant therapy: SGLT2 inhibitor ± other oral antidiabetic drugs Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, 3-point MACE, Non-fatal myocardial infarction, Acute kidney injury, Persistent signs of worsening kidney disease, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: Sponsored by Novo Nordisk, A/D, Denmark. |
| Phrommintiku I 2019 | Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease T2DM and heart failure: Mixed | Strategy: Adding Dapagliflozin 10 mg (n=25) Vildagliptin 50 - 100 mg (n=24) Concomitant therapy: Metformin and or sulfonylurea and or | All-cause mortality, Cardiovascular mortality, Non- fatal myocardial infarction, Non- fatal stroke, Hospitalisation for heart failure, Hypoglycaemia | Study location: Thailand Sources of funding: Thailand research fund |

| | | Intervention and | | |
|--------------------------|---|--|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | population T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Mean age (SD): 63.22 (7.91) years Time since type 2 diabetes diagnosed: Not stated/unclear | Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 89.80% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SUIfonylureas: 75.4% | episodes, HbA1c change, Weight change, BMI change Follow up: 6 months | |
| Pieber 2019 PIONEER 7 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.4 (9.9012) years Time since type 2 diabetes | Strategy: Adding N = 504 Semaglutide 3-14 mg daily (n=253) Sitagliptin 100 mg daily (n=251) Concomitant therapy: One or two glucose-lowering drugs (metformin, sulfonylureas, sodium glucose cotransporter-2 [SGLT2] inhibitors, or thiazolidinediones) Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 37.5% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Hospitalisation for heart failure, Acute kidney injury, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 12 months | Study location: Multicenter Sources of funding: Funded by Novo Nordisk A/S, Denmark. |

| | | Intonuantian and | | |
|--------------------------|--|--|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | diagnosed: 8.8 (6.2504) years | stated/unclear SGLT-2 inhibitors: 0.7% Sulfonylureas: 1.5% | | |
| Pinget 2013 GetGoal-P | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.65 (9.5) years Time since type 2 diabetes diagnosed: 8.1 (5.4672) | Strategy: Adding N = 484 Lixisenatide 80 mcg daily (n=323) Placebo (n=161) Concomitant therapy: Pioglitazone ± metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 81% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multicenter Sources of funding: Sanofi |
| Pollock 2019 DELIGHT | Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 448 Dapagliflozin + Saxagliptin (n=155) Dapagliflozin (n=145) Placebo (n=148) Concomitant therapy: Stable glucose-lowering therapy Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 60.3% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor | All-cause mortality, Persistent signs of worsening kidney disease, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multi-centre, multi-national study conducted at 116 research centres in Australia, Canada, Japan, South Korea, Mexico, South Africa, Spain, Taiwan and the USA. Sources of funding: Astra Zeneca |

| | | Intervention and | | |
|---|--|--|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | Mean age (SD): 64.4667 (8.7802) years Time since type 2 diabetes diagnosed: Not stated/unclear | agonists: Not stated/unclear Insulin: 71.3% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 32.7% | | |
| Pozzilli 2017 AWARD-9 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 60.4 (9.8046) years Time since type 2 diabetes diagnosed: 13.15 (7.6007) years | Strategy: Adding N = 300 Dulaglutide 1.5 mg weekly (n=150) Placebo (n=150) Concomitant therapy: Insulin glargine ± metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 88.3% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Nonfatal stroke, Unstable angina, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6.5 months | Study location: International (Czech Republic, Hungary, Italy, Puerto Rico, UK, USA Sources of funding: Sponsored by Eli Lilly and Co., Indianapolis, IN, USA. |
| Pratley 2009A Alogliptin Study 009 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and | Strategy: Adding N = 493 Alogliptin 12.5 mg (n=197) Alogliptin 25 mg (n=199) Placebo (n=97) Concomitant therapy: Thiazolidinedione ± metformin and or sulfonylurea Antihyperglycaemic treatment received: | All-cause mortality, Non- fatal myocardial infarction, Hospitalisation for heart failure, Hypoglycaemia episodes, Weight change Follow up: 6 months | Study location: 125 sites in the regions of United States, Western Europe, Australia and New Zealand, Latin America, plus Hungary, India and South Africa Sources of funding: Financial support provided by Takeda Global Research and Development |

| | | Intervention and | | |
|---|--|--|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: No information Mean age (SD): 55.3667 (10.0118) years Time since type 2 diabetes diagnosed: 7.6333 (5.7552) years | Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 56.4% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 20.7% | | Center, Inc,. USA. |
| Pratley 2009B Alogliptin Study 007 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.7 (11.138) years Time since type 2 diabetes diagnosed: 7.7 (5.9103) years | Strategy: Adding N = 500 Alogliptin 12.5 mg (n=203) Alogliptin 25 mg (n=198) Placebo (n=99) Concomitant therapy: Sulfonylurea Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Argentina, Australia, Brazil, Chile, Dominican Republic, Guatemala, India, Mexico, Netherlands, New Zealand, Peru, Poland, South Africa, United Kingdom, United States Sources of funding: Takeda |
| Pratley 2010 1860-LIRA- DPP-4 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease | Strategy: Adding N = 665 Liraglutide 1.2 mg (n=225) Liraglutide 1.8 mg (n=221) Sitagliptin (n=219) | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Weight | Study location: Multicenter Sources of funding: Funded by Novo Nordisk, Denmark. Numerous authors declare |
| | 12DW and heart | | change, Weight | authors declare |

| | | Intervention and | | |
|---|---|---|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.3 (9.24) years Time since type 2 diabetes diagnosed: 6.2333 (5.1132) years | Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | change Follow up: 12 months | funding and honoraria from numerous pharmaceutical companies |
| Pratley 2018A VERTIS FACTORIAL | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.1 (9.1115) years Time since type 2 diabetes diagnosed: 6.9 (5.36) years | Strategy: Adding N = 1233 Ertugliflozin 5 mg (n=250) Ertugliflozin 15 mg (n=248) Sitagliptin 100 mg (n=247) Ertugliflozin 5 mg + Sitagliptin 100 mg (n=243) Ertugliflozin 15 mg + Sitagliptin 100 mg (n=245) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Multicenter Sources of funding: Merck Sharp & Dohme Corp., a subsidiary of Merck & Co. Inc and Pfizer Inc. |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|-------------------------------|--|--|---|---|
| Pratley 2018B SUSTAIN 7 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.5 (10.6267) years Time since type 2 diabetes diagnosed: 7.4 (5.6771) years | Strategy: Adding N = 1201 Semaglutide 0.5 mg (n=301) Dulaglutide 0.75 mg (n=300) Semaglutide 1.0 mg (n=300) Dulaglutide 1.5 mg (n=300) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 10.5 months | Study location: Multicenter Sources of funding: Novo Nordisk. Numero us authors declare honoraria and funding from multiple pharmaceutical companies |
| Pratley 2019 PIONEER 4 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 711 Semaglutide (n=285) Liraglutide (n=284) Placebo (n=142) Concomitant therapy: Metfomin ± SGLT2 inhibitor Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Nonfatal stroke, Unstable angina, Acute kidney injury, Hypoglycaemia episodes, HbA1c change, Weight change, BMI change Follow up: 12 months | Study location: Multicenter Sources of funding: Novo Nordisk |

| | | 1.4 | | |
|---------------------------|--|---|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | Mean age (SD): 56.3333 (10) years Time since type 2 diabetes diagnosed: 7.6333 (5.5032) years | | | |
| Punthakee 2012 TIDE | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Mean age (SD): 66.35 (6.7167) years Time since type 2 diabetes diagnosed: 8.6 | Strategy: Adding N = 933 Pioglitazone (n=392) Placebo (n=541) Concomitant therapy: Two or fewer glucose-lowering drugs Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 81% DPP-4 inhibitors: 2.4% GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 47% | All-cause mortality, Cardiovascular mortality, 3-point MACE, Non-fatal myocardial infarction, Non-fatal stroke, Hospitalisation for heart failure, Persistent signs of worsening kidney disease, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.4 months | Study location: Multicenter Sources of funding: GlaxoSmithKline |
| Raman 2022 | (6.6) years Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not | Strategy: Adding N = 107 Empagliflozin + insulin (n=52) Linagliptin + insulin (n=55) Concomitant therapy: Insulin Antihyperglycaemic treatment received: No additional | Hypoglycaemia episodes, HbA1c change Follow up: 12 months | Study location: Eastern India Sources of funding: Unclear. Statement that the authors received no financial support for the research, authorship, and /or publication of this article. |

| Study Population Stated/unclear T2DM and chronic kidney disease: People with type 2 diabetes diagnosed: 13.415 (5.5487) years Raz 2008 | | | 1.4 | | |
|--|----------|--|---|--|--|
| stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease: People with chronic kidney disease: T2DM and higher cardiovascular risk: Not stated/unclear S13.415 (5.5487) years Raz 2008 Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear SUIfonylureas: Not Stated/uncl | Study | Population | | Outcomes | Comments |
| with type 2 diabetes at higher risk of cardiovascular disease N = 190 N = 190 N = 190 Multinational trial Cardiovascular mortality, Cardiovascular mortality, Hypoglycaemia episodes, HbA1c change Follow up: 6.9 months Follow | | stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 62.45 (7.4943) years Time since type 2 diabetes diagnosed: 13.415 (5.5487) | information | | |
| Retnakaran 2010 BEST Model 5: People with type 2 diabetes at higher risk of cardiovascular disease Model 5: People with type 2 strategy: Switching N = 21 Strategy: Switching Hypoglycaemia episodes Canada Study location: Canada Follow up: 11.2 Sources of funding: Samuel Lunenfeld | Raz 2008 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.85 (9.5) years Time since type 2 diabetes diagnosed: 7.85 | N = 190 Sitagliptin (n=96) Placebo (n=94) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 52.1% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not | mortality, Cardiovascular mortality, Hypoglycaemia episodes, HbA1c change | Multinational trial Sources of funding: Merck & |
| disease Lunenfeld | 2010 | Model 5: People with type 2 diabetes at higher risk of | N = 21 Sitagliptin (n=10) | episodes | Canada |
| | | | | - | Lunenfeld |

| | | Intervention and | | |
|--|---|--|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): Not stated/unclear Time since type 2 diabetes diagnosed: Not stated/unclear | therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 42.80% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 4.60% | | Institute, Mount Sinai Hospital |
| Ridderstrale 2014 EMPA-REG H2H-SU | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.95 (10.3506) year years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 1545 Glimepiride 1 - 4 mg once daily (n=780) Empagliflozin 25 mg once daily (n=765) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 772.5% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 48 months | Study location: Multicenter Sources of funding: Boehringer Ingelheim - involved in the study design, data gathering and analysis. Eli- Lilly co- sponsored the trial but was not involved in the study design, and data gathering analysis. |

| | | Intervention and | | |
|---|--|---|---|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| Riddle 1998 Glimepiride Combination Group 1998 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58 (8) years Time since type 2 diabetes diagnosed: 7 (4) years | Strategy: Adding N = 145 Glimepiride 16 mg daily (n=72) Placebo (n=73) Concomitant therapy: NPH Insulin 70%/Regular insulin 30% Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Nonfatal stroke, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Not reported but probably USA (multisite) Sources of funding: Funded by Hoechst Marion Roussel Pharmaceuticals. |
| Riddle 2013A GetGoal-L | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 496 Lixisenatide (n=329) Placebo (n=167) Concomitant therapy: Insulin ± metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 21% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 50% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multicenter Sources of funding: Sanofi |

| | | Into manting and | | |
|-----------------------------------|---|---|--|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | 57 (10) years Time since type 2 diabetes diagnosed: 12.45 (6.7729) years | | | |
| Riddle 2013B GetGoal- Duo-1 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56 (10) years Time since type 2 diabetes diagnosed: 9.15 (5.9008) years | Strategy: Adding N = 446 Lixisenatide (n=223) Placebo (n=223) Concomitant therapy: Insulin + Metformin ± thiazolidinedione + insulin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multicenter Sources of funding: Sanofi |
| Roberts 2005 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular | Strategy: Adding N = 170 Glimepiride 2-8 mg daily (n=85) Placebo (n=85) Concomitant therapy: Metformin and a thiazolidinedione Antihyperglycaemic treatment received: No additional information available. | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 6 months | Study location: USA (multisite) Sources of funding: Supported by Aventis Pharmaceuticals, Bridgewater, NJ, USA; Innovus Research Inc., Medford, MA, USA performed health-related quality of life analysis. |

| | | Intervention and | | |
|----------------------------|--|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | risk: Not stated/unclear Mean age (SD): 56.45 (9.9005) years Time since type 2 diabetes diagnosed: 8.3 (5.9266) years | | | |
| Rodbard 2016 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.45 (9.7063) years Time since type 2 diabetes diagnosed: 9.95 (5.6543) years | Strategy: Adding N = 213 Canagliflozin 100 mg/300 mg titrated (n=107) Placebo (n=106) Concomitant therapy: Metformin + sitagliptin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: Supported by Janssen Research and Development, LLC |
| Rodbard 2017 DUAL IV | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart | Strategy: Adding N = 435 Insulin degludec/liraglutide titrated (n=289) Placebo (n=146) Concomitant therapy: Sulfonylurea | All-cause mortality, Non- fatal myocardial infarction, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c | Study location: Multicenter Sources of funding: Sponsored by Novo Nordisk. |

| | | Intervention and | | |
|------------------------------|---|--|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.7 (10.0179) years Time since type 2 diabetes diagnosed: Not stated/unclear | Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 11% | change, Weight change Follow up: 6 months | |
| Rodbard 2018 SUSTAIN 5 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): Not stated/unclear Time since type 2 diabetes | Strategy: Adding N = 397 Semaglutide 1.0 mg weekly (n=132) Semaglutide 0.5 mg weekly (n=133) Concomitant therapy: Basal insulin with or without metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 86.7% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 53.8% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 0.3% | All-cause mortality, Cardiovascular mortality, Nonfatal stroke, Hospitalisation for heart failure, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: International (90 sites in Germany, Japan, Serbia, Slovakia, and USA) Sources of funding: Funded by Novo Nordisk |

| | | Intervention and | | |
|------------------------------|--|--|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | diagnosed: Not stated/unclear | | | |
| Rodbard 2019 PIONEER 2 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.5 (10) years Time since type 2 diabetes diagnosed: 7.45 | Strategy: Adding N = 822 Empagliflozin 25 mg daily (n=410) Semaglutide 14 mg (n=412) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Hospitalisation for heart failure, Acute kidney injury, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 12 months | Study location: Multicenter Sources of funding: Novo Nordisk A/S, Denmark. |
| Roden 2005 | (6.0558) years Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 353 Pioglitazone (n=316) Metformin (n=597) Concomitant therapy: Sulfonylurea Antihyperglycaemic treatment received: No additional information available. | Weight change Follow up: 12 months | Study location: Europe and Canada. Sources of funding: Eli Lily and Company and Takeda Europe R&D. |

| | | Intervention and | | |
|--|--|--|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | Mean age (SD): 60 (8) years Time since type 2 diabetes diagnosed: 7 (5.6) years | | | |
| Rosenstock 2006 Sitagliptin 019 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.25 (10.7587) years Time since type 2 diabetes diagnosed: 6.1 (5.5533) years | Strategy: Adding N = 353 Sitagliptin (n=175) Placebo (n=178) Concomitant therapy: Pioglitazone Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multinational study Sources of funding: Merck & Co., Inc. |
| Rosenstock 2009B | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic | Strategy: Adding N = 390 Alogliptin 12.5 mg (n=131) Alogliptin 25 mg (n=129) Placebo (n=130) Concomitant therapy: Insulin ± metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: Unclear, appears that the study could have been funded by Takeda Pharmaceuticals |

| | | Intervention and | | |
|---|---|--|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.4333 (10.2042) years Time since type 2 diabetes diagnosed: 12.5667 (6.8808) years | Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 41.3% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Rosenstock 2012 Study MB102030 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 53.5 (10.9064) years Time since type 2 diabetes diagnosed: 5.4867 (5.6489) years | Strategy: Adding N = 420 Dapagliflozin 5 mg (n=141) Dapagliflozin 10 mg (n=140) Placebo (n=139) Concomitant therapy: Pioglitazone Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 11 months | Study location: 105 sites in Argentina, Canada, India, Mexico, Peru, Philippines, Taiwan, and United States Sources of funding: Bristol- Myers Squibb and AstraZeneca. Numerous authors declare funding and honoraria from multiple pharmaceutical companies |
| Rosenstock 2013 GetGoal-X | Model 5: People with type 2 diabetes at higher risk of | Strategy: Adding N = 420 Dapagliflozin 5 mg | All-cause mortality, Hypoglycaemia episodes, Severe | Study location: Multicenter |

| | | Intervention and | | |
|----------------------------------|---|---|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.5 (10.0) years Time since type 2 diabetes diagnosed: Not stated/unclear | (n=141) Dapagliflozin 10 mg (n=140) Placebo (n=139) Concomitant therapy: Pioglitazone Antihyperglycaemic treatment received: No additional information | hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Sources of funding: Sanofi |
| Rosenstock 2014A GetGoal-S | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 859 Lixisenatide (n=573) Placebo (n=286) Concomitant therapy: Sulfonylurea ± metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 84.5% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 100% | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multicenter Sources of funding: Sanofi |

| | | Intervention and | | |
|--|---|---|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | 57.4 (9.9008) years Time since type 2 diabetes diagnosed: 9.45 (6.0672) years | | | |
| Rosenstock 2014B EMPA-REG MDI | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.6667 (9.4198) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 563 Empagliflozin 10 mg (n=186) Empagliflozin 25 mg (n=189) Placebo (n=188) Concomitant therapy: Insulin ± metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 29% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Multicenter Sources of funding: Boehringer Ingelheim and Eli Lilly. A number of authors are employees of Boehringer Ingelheim and others declare funding and honoraria from multiple pharmaceutical companies |
| Rosenstock 2015A | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic | Strategy: Adding N = 534 Dapagliflozin + Saxagliptin (n=179) Saxagliptin + Placebo (n=176) Dapagliflozin + Placebo (n=179) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional | All-cause mortality, Cardiovascular mortality, Persistent signs of worsening kidney disease, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Canada, Mexico, Poland, Puerto Rico, Republic of Korea, Romania, South Africa; United States Sources of funding: Bristol- Myers Squibb and AstraZeneca |

| | | 1 | | |
|--|---|--|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54 (10) years Time since type 2 diabetes diagnosed: 7.5667 (5.3033) years | information available. | | |
| Rosenstock 2015B EMPA-REG BASAL | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.8667 (9.8921) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 494 Empagliflozin 25 mg (n=155) Empagliflozin 10 mg (n=169) Placebo (n=170) Concomitant therapy: Insulin ± metformin ± sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 10% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 18 months | Study location: Multicenter Sources of funding: Boehringer Ingelheim and Eli Lilly and Company |
| Rosenstock 2016A | Model 5: People with type 2 diabetes at higher risk of | Strategy: Adding N = 894 Lixisenatide (n=298) | All-cause mortality, Cardiovascular mortality, | Study location: Multicenter |

| | | latement's a | | |
|----------------------------------|---|---|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| GetGoal- Duo-2 | cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.8 (8.9101) years Time since type 2 diabetes diagnosed: 12.2 (6.6693) years | Insulin glulisine QD (n=298) Insulin glulisine TID (n=298) Concomitant therapy: Metformin + insulin glargine Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 87.3% DPP-4 inhibitors: 12.1% GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 46.1% | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Sources of funding: Sanofi |
| Rosenstock 2016B LixiLan-O | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.4 (9.3053) years Time since type 2 | Strategy: Adding N = 1170 Lixisenatide/Insulin glargine (n=469) Lixisenatide (n=234) Insulin glargine (n=467) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: 2.4% GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: 0.3% Sulfonylureas: 53.7% | All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Nonfatal stroke, Unstable angina, Hospitalisation for heart failure, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 7 months | Study location: Multicenter Sources of funding: Sanofi |

| | | Intonoution and | | |
|------------------------------------|--|---|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| į | diabetes diagnosed: 8.8333 (5.7077) years | · | | |
| Rosenstock 2016C LixiLan PoC | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD):56.75 (9.45) years Time since type 2 diabetes diagnosed: 6.7 (4.8) years | Strategy: Adding Lixisentatide + Insulin glargine (n=161) Insulin glargine (n=162) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multicenter Sources of funding: Sanofi |
| Rosenstock 2018A FREEDOM-1 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not | Strategy: Adding N = 441 Exenatide 40 mcg/day (n=147) Exenatide 60 mcg/day (n=151) Placebo (n=143) Concomitant therapy: Diet and exercise alone or with metformin, sulfonylureas or pioglitazone monotherapy or in combination Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 9.1 months | Study location: 126 clinical sites in the U.S. Sources of funding: Intarcia Therapeutics |

| | | Intervention and | | |
|-----------------------------------|---|--|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | Mean age (SD): 54.9667 (9.6836) years Time since type 2 diabetes diagnosed: 8.8667 (6.3867) years | Biguanides: 41.3% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 3.5% | | |
| Rosenstock 2018B VERTIS MET | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.6667 (8.7471) years Time since type 2 diabetes diagnosed: 8 (5.9789) years | Strategy: Adding N = 621 Empagliflozin 5 mg (n=207) Empagliflozin 15 mg (n=205) Placebo (n=209) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 99.8% DPP-4 inhibitors: 3.4% GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, HbA1c change Follow up: 6 months | Study location: North America, South America, Europe, Asia, South Africa, Australia, New Zealand Sources of funding: Study funded by Pfizer.Numerous authors declare funding and honoraria from multiple pharmaceutical companies |
| Rosenstock 2019A CARMELINA | Model 1: People with type 2 diabetes and heart failure Model 3: People with type 2 diabetes and chronic kidney disease Model 5: People with type 2 diabetes at higher risk of cardiovascular | Strategy: Adding N = 6991 Linagliptin (n=3499) Placebo (n=3492) Concomitant therapy: Additional medications except DPP-4 inhibitors, GLP 1 receptor agonists or SGLT-2 inhibitors Antihyperglycaemic | All-cause mortality, Cardiovascular mortality, 3-point MACE, 4-point MACE, Non-fatal myocardial infarction, Non-fatal stroke, Unstable angina, Hospitalisation for heart failure, Acute kidney injury, Persistent signs of | Study location: Multicenter Sources of funding: Study was sponsored by Boehringer Ingelheim and Eli Lilly. |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|---------------------------------|--|--|--|--|
| | T2DM and heart failure: Mixed population T2DM and atherosclerotic cardiovascular disease: Mixed population T2DM and chronic kidney disease: Mixed population T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Includes results for a subgroup for people with or without heart failure and with or without chronic kidney disease. Mean age (SD): 65.85 (9.1) years Time since type 2 diabetes diagnosed: 14.75 (9.4513) years | treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 54.60% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 58% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 32.1% | worsening kidney disease, Development of end stage kidney disease, Death from renal causes, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 26.4 months | |
| Rosenstock 2019B CAROLINA | Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Mixed population | Strategy: Adding N = 6033 Linagliptin (n=3023) Glimepiride (n=3010) Concomitant therapy: Additional medications including adjustment of background therapy, or addition of pioglitazone, metformin, alpha glucosidase inhibitor or basal insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 3.20% Biguanides: 83.5% | All-cause mortality, Cardiovascular mortality, 3-point MACE, 4-point MACE, Non-fatal myocardial infarction, Non-fatal stroke, Unstable angina, Hospitalisation for heart failure, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 75.6 months | Study location: Multicenter Sources of funding: Boehringer Institute and Eli Lilly and Company. |

| | | Intervention and | | |
|----------------------------------|--|---|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Includes results for a subgroup for people with atherosclerotic cardiovascular disease. Mean age (SD): 64.05 (9.5) years Time since type 2 diabetes diagnosed: Not | DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 28.5% | | |
| | stated/unclear | | | |
| Rosenstock 2019C PIONEER 3 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.75 (10) years Time since type 2 diabetes diagnosed: 8.55 (6.0012) years | Strategy: Adding N = 1864 Semaglutide 3 mg/d (n=466) Semaglutide 7 mg/d (n=466) Semaglutide 14 mg/d (n=465) Sitagliptin 100 mg/d (n=467) Concomitant therapy: Metformin with or without sulfonylurea Antihyperglycaemic treatment received: No additional information available. | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Non-fatal myocardial infarction, Non-fatal stroke, Unstable angina, Hospitalisation for heart failure, Acute kidney injury, Death from renal causes, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 18 months | Study location: Multicenter Sources of funding: Novo Nordisk |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|---------------------------------|--|--|---|--|
| Rosenstock 2019D | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.7 (10.5073) years Time since type 2 diabetes diagnosed: 7.6333 (6.1369) years | Strategy: Adding N = 883 Dapagliflozin + Saxagliptin (n=293) Dapagliflozin (n=294) Saxagliptin (n=296) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: 119 centres in Canada, the Czech Republic, Germany, Mexico, Russia, and the USA Sources of funding: AstraZeneca |
| Rosenstock 2023 SURPASS-6 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and rated/unclear T2DM and higher | Strategy: Adding N = 1425 Tirzepatide (n=717) insulin lispro (n=708) Concomitant therapy: No additional information Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 84.5% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor | All-cause mortality, Acute kidney injury, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Globally - 135 centers in Argentina, Belgium, Brazil, Czech Republic, Germany, Greece, Hungary, Italy, Mexico, Romania, Russia, Slovakia, Spain, Turkey, and the US Sources of funding: Eli Lilly and Company |

| | | Intervention and | | |
|--|---|---|---|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | cardiovascular risk: Not stated/unclear Mean age (SD): 58.8 (9.7504) years Time since type 2 diabetes diagnosed: 13.8 (7.3001) years | agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Roussel 2019 CompoSIT-I | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.35 (9.6001) years Time since type 2 diabetes diagnosed: 10.75 (6.85) years | Strategy: Adding N = 743 Sitagliptin (n=373) Placebo (n=370) Concomitant therapy: Insulin, Metformin with or without DPP-4 inhibitor and/or sulfonylurea Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 7 months | Study location: Multicenter Sources of funding: Merck Sharp & Dohme Corp., a subsidiary of Merck & Co., Inc. The lead author declares support from multiple pharmaceutical companies and numerous authors are employees of Merck Sharp & Dohme Corp |
| Russell- Jones 2009 LEAD-5 met+SU | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular | Strategy: Adding N = 581 Liraglutide 1.8 mg daily (n=232) Placebo (n=115) Insulin glargine (n=234) Concomitant therapy: Metformin + glimepiride Antihyperglycaemic | Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: Funded by Novo Nordisk A/S |

| Study Population disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear risk: Not stated/unclear risk of cardiovascular failure: Not stated/unclear T2DM and heart failure: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and bigher cardiovascular risk: Not stated/unclear T2DM and bigher cardiovascular risk: Not stated/unclear SQLT-2 inhibitors: Not stated/unclear Insulin: Not stated/unclear Cardiovascular risk of Cardiovascular risk: Not stated/unclear Cardiovascu | | | Intervention and | | |
|--|-------|--|---|---|---|
| stated/unclear T2DM and chronic kidney disease. Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear risk of cardiovascular disease Not stated/unclear T2DM and heart failure: Not stated/unclear T2DM and chronic kidney disease. Not stated/unclear S2D diabetes diagnosed: Not S2D diabetes diagnosed: Not S2D diabetes diagnosed: Not S2D diabetes diagnosed: N | Study | Population | Intervention and comparison | Outcomes | Comments |
| ma 2011 with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear S2 years Time since type 2 diabetes diagnosed: Not stated/unclear Model 5: People with type 2 diabetes at higher risk of cardiovascular disease Savvidou Model 5: People with type 2 diabetes at higher risk of cardiovascular disease Medical Center of Diabetes Meditus in politors in sulin: (n=55) Follow up: 6 months Weight change, BMI change Follow up: 12 months Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Sulfonylureas: Not stated/unclear Sources of Follow up: 12 months Pharmaceuticas and Eli-Lilly supported the research through grants. Follow up: 12 months Pharmaceuticate Pharmaceuticate Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear Sources of Follow up: 12 months Pharmaceuticas and Eli-Lilly supported the research through grants. | | stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.5333 (9.9342) years Time since type 2 diabetes diagnosed: 9.4333 (6.1268) | No additional information | | |
| with type 2 diabetes at higher risk of Exenatide (n=55) cardiovascular disease N = 110 Weight change, BMI change BMI change Diabetes Mellitus in Follow up: 6 "Papageorgiou" University | | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 52 years Time since type 2 diabetes diagnosed: Not | N = 21 Pioglitazone 30 - 45 mg daily + Exenatide 10 µg twice daily (n=11) Pioglitazone 30 mg - 45 mg daily (n=10) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 81% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not | Weight change, BMI change Follow up: 12 | Sources of funding: Amylin Pharmaceuticals and Eli-Lilly supported the research through |
| | | Model 5: People with type 2 diabetes at higher risk of cardiovascular | N = 110 Exenatide (n=55) | Weight change, BMI change Follow up: 6 | Medical Center of Diabetes Mellitus in "Papageorgiou" |

| | | Intervention and | | |
|--|--|--|---|--|
| Study | Population | comparison | Outcomes | Comments |
| Schernthaner | T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 62.95 (7.1502) years Time since type 2 diabetes diagnosed: Not stated/unclear | therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause | Thessaloniki, Greece Sources of funding: None |
| Schernthaner 2013 CANTATA- D2 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.65 (9.4512) years Time since type 2 diabetes | Strategy: Adding N = 756 Canagliflozin (n=378) Sitagliptin (n=378) Concomitant therapy: Metformin + sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 30% | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 12 months | Study location: Multicenter Sources of funding: Janssen Global Services, LLC. |

| | | Intervention and | | |
|---|--|--|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | diagnosed: 9.55 (6.2008) years | | | |
| Schernthaner 2015A GENERATIO N | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 72.6 (5.552) years Time since type 2 diabetes diagnosed: 7.6 (6.2032) years | Strategy: Adding N = 720 Saxaglitpin (n=360) Glimepiride (n=360) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Weight change Follow up: 12 months | Study location: Multicenter Sources of funding: AstraZeneca and Bristol-Myers Squibb |
| Scirica 2013 SAVOR-TIMI 53 | Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Mixed population T2DM and | Strategy: Adding N = 16492 Saxagliptin (n=8280) Placebo (n=8212) Concomitant therapy: Antihyperglycemic therapy except DPP-4 inhibitor or GLP-1 receptor agonist Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 69.60% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 41.4% | All-cause mortality, Cardiovascular mortality, 3-point MACE, Non-fatal myocardial infarction, Non-fatal stroke, Unstable angina, Hospitalisation for heart failure, Persistent signs of worsening kidney disease, Development of end stage kidney disease, Death from renal causes, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI | Study location: Multicenter Sources of funding: AstraZeneca and Bristol-Myers Squibb. |

| | | Intervention and | | |
|--------------------------|---|--|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Includes results for a subgroup for people with atherosclerotic cardiovascular disease. Mean age (SD): 65.05 (8.5499) years Time since type 2 diabetes diagnosed: Not stated/unclear | SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 40.20% | change Follow up: 25.2 months | |
| Scott 2018 CompoSIT-R | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 67.15 (8.5501) years Time since type 2 diabetes diagnosed: 10.6 (7.2028) years | Strategy: Adding N = 614 Sitagliptin (n=307) Dapagliflozin (n=307) Concomitant therapy: Metformin with or without a sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 71.3% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 5.5 months | Study location: Multicenter Sources of funding: Merck Sharp & Dohme Corp., a subsidiary of Merck & Co., Inc., Kenilworth, NJ, USA. |

| | | Intervention and | | |
|---------------------------------|---|--|--|---|
| Study | Population | comparison | Outcomes | Comments |
| Seino 2012 GetGoal-L Asia | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.35 (10.1496) years Time since type 2 diabetes diagnosed: 13.9 (7.7) years | Strategy: Adding N = 311 Lixisenatide (n=154) Placebo (n=157) Concomitant therapy: Insulin ± sulfonylurea Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Nonfatal stroke, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multicenter Sources of funding: Sanofi |
| Seino 2016 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 257 Liraglutide (n=127) Placebo (n=130) Concomitant therapy: Insulin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 8.3 months | Study location: 23 sites in Japan Sources of funding: Novo Nordisk |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|------------------|---|--|---|---|
| · | Mean age (SD): 60.55 (11.1528) years Time since type 2 diabetes diagnosed: 14.505 (8.7445) years | | | |
| Seino 2021 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.5 (10.5946) years Time since type 2 diabetes diagnosed: 9.15 (4.9672) years | Strategy: Adding N = 141 Sitagliptin (n=70) Placebo (n=71) Concomitant therapy: Ipragliflozin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Japan Sources of funding: MSD K.K., a subsidiary of Merck & Co |
| Shankar 2017A | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic | Strategy: Adding N = 467 Sitagliptin (n=234) Placebo (n=233) Concomitant therapy: Insulin with or without metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: China Sources of funding: Merck & Co |

| | | Intervention and | | |
|------------------------------|---|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | cardiovascular disease T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.7 (8.8) years Time since type 2 diabetes diagnosed: 11.2 (5.4) years | | | |
| Sivalingam 2023 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 69.95 (8.0327) years Time since type 2 diabetes diagnosed: 18.5 (10.5119) years | Strategy: Adding N = 60 Semaglutide (n=30) Placebo (n=30) Concomitant therapy: Empagliflozin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 0% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 50% | All-cause mortality, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Copenhagen, Denmark Sources of funding: Novo Nordisk |
| Skrivanek 2014 AWARD-5 | Model 5: People with type 2 diabetes at higher risk of | Strategy: Adding N = 135 Dulaglutide 0.25 mg | HbA1c change, Weight change Follow up: 12 | Study location: US Sources of |
| | cardiovascular disease | once weekly (n=24) Dulaglutide 0.5 mg | months | funding: Eli Lily and company |

| | | Intervention and | | |
|-------------------------|---|--|--|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.75 (7.1388) years Time since type 2 diabetes diagnosed: 7 (3.3578) years | once weekly (n=25) Dulaglutide 0.75 mg once weekly (n=21) Dulaglutide 1.0 mg once weekly (n=10) Dulaglutide 1.5 mg once weekly (n=25) Dulaglutide 2.0 mg once weekly (n=30) Dulaglutide 3.0 mg once weekly (n=15) Sitagliptin 100 mg daily (n=42) Placebo daily/weekly (n=38) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Softeland 2017 NA | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not | Strategy: Adding N = 333 Empagliflozin 10 mg once daily (n=112) Empagliflozin 25 mg once daily (n=111) Placebo (n=110) Concomitant therapy: Meformin + linagliptin, Placebo Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multicenter Sources of funding: Funded by Boehringer Ingelheim and Eli Lilly and Company Diabetes Alliance |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|--------------|--|--|--|--|
| Olday | stated/unclear Mean age (SD): 55.2 (9.7338) years Time since type 2 diabetes diagnosed: Not stated/unclear | Companio | Cateomic | |
| Sone 2019 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.6667 (10.079) years Time since type 2 diabetes diagnosed: 13.8 (8.0144) years | Strategy: Adding N = 269 Empagliflozin 10 mg daily (n=89) Empagliflozin 25 mg daily (n=90) Placebo (n=90) Concomitant therapy: Insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 72.4% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Unstable angina, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Japan (51 sites) Sources of funding: Supported by Nippon Boehringer Ingelheim Co. Ltd. |
| Sridhar 2013 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not | Strategy: Adding N = 50 Pioglitazone 30 mg daily (n=25) Placebo (n=25) Concomitant therapy: Glimepiride + metformin Antihyperglycaemic treatment received: No additional | Hypoglycaemia episodes, HbA1c change, Weight change, BMI change Follow up: 6 months | Study location: India Sources of funding: Drug and placebo tablets provided by Sun Pharmaceutical Industries Ltd, Mumbai, India. |

| | | 1.4 | | |
|-------------------------|--|--|--|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 45.95 (6.5376) years Time since type 2 diabetes diagnosed: 2.55 (1.9105) years | information available. | | |
| Strain 2013 INTERVAL | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 74.75 (4.1527) years Time since type 2 diabetes diagnosed: 11.4 (7.4169) years | Strategy: Adding N = 278 Vildagliptin (n=139) Placebo (n=139) Concomitant therapy: Oral drugs Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Progression of liver disease, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 5.5 months | Study location: Multicenter (Belgium, Bulgaria, Germany, Finland, Slovakia, Spain and the UK). Sources of funding: Funded by Novartis Pharma AG. |
| Strojek 2011 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease | Strategy: Adding N = 592 Dapagliflozin 2.5 mg daily (n=154) Dapagliflozin 5 mg daily (n=142) Dapagliflozin 10 mg | All-cause mortality, Cardiovascular mortality, Non- fatal stroke, Persistent signs of worsening kidney disease, | Study location: Multicenter Sources of funding: Astra Zenaca and Bristol-Myers Squib funded |

| | | Intervention and | | |
|---------|---|---|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Mixed population T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.825 (9.5999) years Time since type 2 diabetes diagnosed: 7.425 (5.7299) years | daily (n=151) Placebo (n=145) Concomitant therapy: Glimepiride Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | medical writing and editorial assistance. |
| Su 2014 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 48.14 (13.0642) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 600 Vildagliptin 100 mg daily (n=300) Placebo daily (n=300) Concomitant therapy: Metformin + alpha glucosidase inhibitor Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: China Sources of funding: No additional information. |

| | | lutement's re- | | |
|-----------------------------|---|--|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| Takahashi 2023 | Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): Not stated/unclear Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Switching N = 110 Liraglutide 0.9-1.8 mg daily (n=20) Semaglutide 0.25-1.0 mg weekly A (n=20) Dulaglutide 0.75 mg weekly (n=35) Semaglutide 0.25-1.0 mg weekly B (n=35) Concomitant therapy: NA Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 15.7% Biguanides: 87.60% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: 43.20% Insulin: 51.40% SGLT-2 inhibitors: 77.8% Sulfonylureas: 23.20% | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 5.5 months | Study location: Japan (8 hospital sites) Sources of funding: Research did not receive any specific grant from funding agencies in the public, commercial, or not-for-profit sectors. |
| Takihata 2013 COMPASS | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 130 Pioglitazone 15 mg daily (n=65) Sitagliptin 50 mg daily (n=65) Concomitant therapy: Metformin ± sulfonylurea Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Japan Sources of funding: Grants- in-Aid for Scientific Research (B) 21390282 and (B) 24390235 from the Ministry of Education, Culture, Sports, Science and Technology (MEXT) of Japan, a Grant for the Strategic Japanese-Danish Cooperative Program on Molecular Diabetology from the Japan Science and Technology Agency, a Grant- in-Aid from the |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|------------------|--|--|---|---|
| | 60.5 (8.5586) years Time since type 2 diabetes diagnosed: Not stated/unclear | | | Uehara Memorial Foundation to one author, and a Grant-in-Aid from the Joint Research Association for Japanese Diabetes to another author. |
| Tan 2004 GLAD | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.4 (8.6797) years Time since type 2 diabetes diagnosed: 6.65 (6.7529) years | Strategy: Switching N = 244 Pioglitazone (n=121) Glimepiride (n=123) Concomitant therapy: NA Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 20.9% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 54.60% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Hypoglycaemia episodes, HbA1c change Follow up: 12 months | Study location: 16 centres in Mexico Sources of funding: This work relates to Eli Lilly and Company protocol H6E- MC-GLAD. The main author is an employee of Eli Lilly and Company |
| Tanaka 2017 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear | Strategy: Switching N = 132 Alogliptin (n=64) Vildagliptin (n=68) Concomitant therapy: Existing treatment (except for sitagliptin which was switched for study drug) Antihyperglycaemic treatment received: Alpha-glucosidase | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Japan Sources of funding: None declared |

| | | Intervention and | | |
|-----------------------|--|--|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 66.45 (10.0465) years Time since type 2 diabetes diagnosed: 11.25 (9.221) years | inhibitors: 15% Biguanides: 55.2% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 28.20% | | |
| Tanaka 2019 EMBLEM | Model 5: People with type 2 diabetes at higher risk of | Strategy: Adding Empagliflozin (n=58) Placebo (n=59) | HbA1c change, Weight change, BMI change | Study location: Japan. Sources of |
| | cardiovascular disease | Concomitant therapy: Standard therapy | Follow up: 5.5 months | funding: Funded by Boehringer Ingelheim and Eli Lilly and |
| | T2DM and heart failure: Mixed population T2DM and atherosclerotic cardiovascular disease: Mixed population T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 64.9 (10.4) years | Antihyperglycaemic treatment received: No additional information available. | | Company. |
| | Time since type 2 diabetes diagnosed: 13.2 (10.9) years | | | |
| Taskinen 2011 | Model 5: People with type 2 diabetes at higher | Strategy: Adding N = 700 | Hypoglycaemia episodes, Severe hypoglycaemic | Study location: Multicenter |
| | risk of cardiovascular disease | Linagliptin 5mg daily (n=523) Placebo (n=177) | episodes, HbA1c change, Weight change | Sources of funding: Funded by Boehringer Ingelheim. |
| | T2DM and heart failure: People | Concomitant therapy: Metformin with or without one | Follow up: 5.5 months | |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|---------------------------------------|--|---|---|---|
| Citaly | without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.55 (10.3076) years Time since type 2 diabetes diagnosed: Not stated/unclear | other blood-glucose lowering drug Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 68.5% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Cutcomics | |
| Terauchi 2020 LixiLan JP- 02 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.7 (10.7) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 226 Linagliptin 5 mg daily (n=106) Placebo (n=120) Concomitant therapy: Background blood-glucose lowering drug monotherapy Antihyperglycaemic treatment received: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Japan Sources of funding: Sanofi |
| Thrasher 2014 | Model 5: People with type 2 | Strategy: Adding N = 226 | All-cause mortality, | Study location: USA |

| | | Interpreting | | |
|-------------------|--|--|--|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 53.9 (9.9942) years Time since type 2 diabetes diagnosed: Not stated/unclear | Linagliptin 5 mg daily (n=106) Placebo (n=120) Concomitant therapy: Background blood-glucose lowering drug monotherapy Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 78.00% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 20.40% | Cardiovascular mortality, Non-fatal myocardial infarction, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Sources of funding: Funded by Boehringer Ingelheim Inc |
| Tinahones 2017 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.7 (10.7) years | Strategy: Adding N = 226 Linagliptin 5 mg + Empagliflozin 10 mg (n=126) (study 1) Linagliptin 5 mg + Empagliflozin 25 mg (n=114) (study 2) Placebo + Empagliflozin 10mg (n=130) (study 1) Placebo + Empagliflozin 25 mg (n=112) (study 2) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multicenter Sources of funding: Boehringer Ingelheim and Eli Lilly & Co. |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|------------------------|---|--|---|---|
| Ciacy | Time since type 2 diabetes diagnosed: Not stated/unclear | Companio | Culcomec | |
| Tripathy 2013 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.5 (2.5289) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 29 Pioglitazone 15 mg daily (n=15) Placebo (n=14) Concomitant therapy: Metformin with or without a sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 60.6% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | HbA1c change, Weight change, BMI change Follow up: 6 months | Study location: Texas, USA Sources of funding: Funded by Takeda. |
| Tuttle 2018 AWARD-7 | Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher | Strategy: Adding N = 577 Dulaglutide 1.5 mg weekly (n=193) Dulaglutide 0.75 mg weekly (n=190) Insulin glargine (n=194) Concomitant therapy: Insulin lispro +/- hypoglycaemic agent Antihyperglycaemic treatment received: No additional | All-cause mortality, Cardiovascular mortality, Persistent signs of worsening kidney disease, Development of end stage kidney disease, Death from renal causes, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight | Study location: Multicenter Sources of funding: Eli Lilly and Co. |

| | | Intervention and | | |
|-------------------|--|---|--|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | cardiovascular risk: Not stated/unclear Mean age (SD): 64.5667 (8.6012) years Time since type 2 diabetes diagnosed: 18.1 (8.7331) years | information available. | change Follow up: 12 months | |
| Umpierrez 2006 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 48.9 (10.4997) years Time since type 2 diabetes diagnosed: 5.4 (5.1247) years | Strategy: Adding N = 210 Pioglitazone 30-45 mg daily (n=109) Glimepiride 2-8 mg daily (n=101) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Hospitalisation for heart failure, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 6 months | Study location: USA (51 diabetes centres) Sources of funding: Sponsored by Sanofi-Aventis, Bridgewater, NJ, USA. |
| Vähätalo 2007 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular | Strategy: Adding Glipizide (n=15) Metformin (n=26) Concomitant therapy: Insulin Antihyperglycaemic treatment received: No additional information available. | HbA1c change, Weight change Follow up: 12 months | Study location: Finland. Sources of funding: No additional information. |

| | | 1.4 | | |
|---------------------------------|--|--|--|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): Not stated/unclear Time since type 2 diabetes diagnosed: Not stated/unclear | | | |
| van der Meer 2009 PIRAMID | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.6 (0.9513) years Time since type 2 diabetes diagnosed: 3.5 years | Strategy: Adding N = 78 Pioglitazone (n=39) Metformin (n=39) Concomitant therapy: Glimepiride Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 100% | Hospitalisation for heart failure, HbA1c change, Weight change Follow up: 5.5 months | Study location: The study was performed at two institutes in The Netherlands Sources of funding: Supported by Eli Lilly, the Netherlands, which has a partnership with Takeda, the manufacturer of pioglitazone. Metformin tablets and matching placebos were provided by Merck, the Netherlands. Multiple authors report receiving funding from numerous pharmaceutical companies |
| van Eyk 2019 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease | Strategy: Adding N = 47 Liraglutide (n=22) Placebo (n=25) Concomitant therapy: Concomitant | HbA1c change, Weight change, BMI change Follow up: 5.5 months | Study location: The Netherlands Sources of funding: The study was funded by Novo Nordisk (Bagsvaerd, Denmark) Roba |

| Study Population failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease: T2DM and higher cardiovascular risk: Not stated/unclear T2DM and the cardiovascular risk: Not stated/unclear disease: Not stated/unclear risk: Not stated/unclear disease: Not stated/unclear T2DM and heart failure: Not stated/unclear T2DM and and chronic kidney disease: Not stated/unclear T2DM and and chronic kidney disease: Not stated/unclear T2DM and bipher cardiovascular risk: Not stated/unclear T2D | | | Intervention and | | |
|--|-------|---|---|--|--|
| stated/unclear T2DM and atherosclerotic cardiovascular diseases: Not stated/unclear T2DM and chronic kidney diseases People without chronic kidney diseases T2DM and higher cardiovascular risk (10) years Van Gaal 2014 Vanderheide Vanderheide Vascular Imaging Group, Leiden University Medical Centre (Leiden, The Netherlands). Ilysselstein and the Cardio Vascular Imaging Group, Leiden University Medical Centre (Leiden, The Netherlands). International claim of the Netherlands of Study International claim of the Netherlands of Study International Centre (Leiden, The Netherlands). International Canada (Chile, Cardiovascular disease diagnosed: 18 (10) years Van Gaal 2014 Van Gaal 2014 Van Gaal 2014 Van Gaal 30 | Study | Population | | Outcomes | Comments |
| with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and bigher cardiovascular risk: Not stated/unclear T2DM and bigher cardiovascular risk: Not stated/unclear Wascaular risk: Not stated/unclear Vascaular risk: Not stated/unclear Vascaular ris | | stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55 (9.9833) years Time since type 2 diabetes diagnosed: 18 (10) years | metformin, sulfonylurea derivatives and insulin was optional Antihyperglycaemic treatment received: No additional information available. | | ljsselstein and the Cardio Vascular Imaging Group, Leiden University Medical Centre (Leiden, The Netherlands). |
| Vanderheide Model 5: People Strategy: Adding Health-related Study location: | | with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | N = 319 Lixisenatide 20 mcg daily (n=158) Sitagliptin 100 mg daily (n=161) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information | mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 | International (Australia, Brazil, Canada, Chile, Guatemala, Mexico, Peru, Poland, Romania, Russian Federation, Ukraine, USA) Sources of funding: Funded/supporte |
| | | | | | |

| | | lutom continue and | | |
|--|--|---|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.15 (7.3773) years Time since type 2 diabetes diagnosed: Not stated/unclear | Liraglutide 1.8 mg daily (n=35) Placebo (n=36) Concomitant therapy: Insulin Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 6 months | Sources of funding: Funded by Novo Nordisk |
| Verma 2019 EMPA- HEART CardioLink-6 | Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 97 Empagliflozin (n=49) Placebo (n=48) Concomitant therapy: Metformin +/- insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 94% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 25% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, 3-point MACE, Non-fatal myocardial infarction, Non-fatal stroke, Hospitalisation for heart failure, Persistent signs of worsening kidney disease, Cardiac arrhythmia, Diabetic ketoacidosis, Progression of liver disease, HbA1c change, Weight change, BMI change Follow up: 6 months | Study location: Canada. Sources of funding: Boehringer Ingelheim (Canada) Ltd. |

| | | Intervention and | | |
|---|---|--|---|--|
| Study | Population | comparison | Outcomes | Comments |
| | Mean age (SD): Not stated/unclear Time since type 2 diabetes diagnosed: Not stated/unclear | | | |
| Vianna 2018 BoneGLIC | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 61.9 (5.8758) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 42 Vildagliptin 100 mg (n=21) Gliclazide MR 120mg (n=21) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Nonfatal stroke, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: One centre, Brazil Sources of funding: investigator- initiated trial research funds from Novartis Pharmaceuticals |
| Vilsboll 2010 Sitagliptin Study 051 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney | Strategy: Adding N = 641 Sitagliptin 100 mg daily (n=322) Placebo (n=319) Concomitant therapy: Insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 72% DPP-4 inhibitors: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Unstable angina, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: Multicenter Sources of funding: Funded by Merck & Co., Inc. |

| | | Intervention and | | |
|-----------------------|--|--|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.75 (9.2001) years Time since type 2 diabetes diagnosed: 12.5 (6.5215) years | GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | |
| Vilsboll 2019 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.5 (9.5749) years Time since type 2 diabetes diagnosed: 9.45 (6.3525) years | Strategy: Adding N = 641 Dapagliflozin 10 mg daily + Saxagliptin 5 mg daily (n=322) Insulin (n=319) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 0.20% Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: 0.40% Insulin: 0.40% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Acute kidney injury, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Multicenter Sources of funding: Funded by AstraZeneca. |
| Wägner 2019 LIPER2 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear | Strategy: Adding N = 24 Liraglutide (n=12) Placebo (n=12) Concomitant therapy: Use of other glucose lowering agents | HbA1c change, Weight change, BMI change Follow up: 6 months | Study location: Single centre Sources of funding: Funding and Drug supplies; Novo Nordisk |

| | | Intervention and | | |
|------------|---|---|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 52.9 (11.9275) years Time since type 2 diabetes diagnosed: 8.71 (5.8588) years | Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 41.7% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 41.7% | | |
| Wada 2022 | Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 62.45 (10.8042) years Time since type 2 diabetes diagnosed: 15.96 (8.8116) years | Strategy: Adding N = 308 Canagliflozin (n=154) Placebo (n=154) Concomitant therapy: angiotensin-converting enzyme inhibitor or angiotensin II receptor blocker Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, 3-point MACE, 5-point MACE, Hospitalisation for heart failure, Diabetic ketoacidosis, Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 25 months | Study location: Japan Sources of funding: Mitsubishi Tanabe Pharma Corporation |
| Wang 2016B | Model 5: People with type 2 diabetes at higher risk of cardiovascular | Strategy: Adding N = 306 Linagliptin 5 mg daily (n=205) | All-cause mortality, Cardiovascular mortality, 5-point MACE, Non-fatal | Study location: International (19 centres in China, Philippines and Malaysia) |

| | | Intervention and | | |
|-----------|--|--|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.8 (10.086) years Time since type 2 diabetes diagnosed: Not stated/unclear | Placebo (n=101) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 4% Biguanides: 65.2% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 27.80% | myocardial infarction, Hospitalisation for heart failure, Development of end stage kidney disease, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Sources of funding: Funded by Boehringer Ingelheim |
| Wang 2017 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.15 (9.2033) years Time since type 2 | Strategy: Adding N = 380 Sitagliptin 100 mg daily (n=191) Placebo (n=189) Concomitant therapy: Acarbose Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 190% Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | Hypoglycaemia episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: China, Romania, Kore, Malaysia, India, Philippines Sources of funding: Merck & Co., Inc., Kenilworth, NJ, USA |

| | | Intonio d'ant | | |
|------------|--|--|---|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | diabetes diagnosed: 7.8 (5.313) years | | | |
| Wang 2019B | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.9667 (9.6045) years Time since type 2 diabetes diagnosed: 8.1333 (5.1387) years | Strategy: Adding N = 774 Dulaglutide 0.75 mg (n=257) Dulaglutide 1.5 mg (n=258) Insulin (n=259) Concomitant therapy: Metformin and /or a SU Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 40.7% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-1 inhibitors: Not stated/unclear Sulfonylureas: 11.7% | All-cause mortality, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: 45 sites in China, Russia, Mexico and South Korea Sources of funding: Elli Lilly and Company. A number of authors are employees of Elli Lilly and Company |
| Wang 2020A | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular | Strategy: Adding N = 95 Exenatide (n=49) Insulin (n=46) Concomitant therapy: existing oral antidiabetic therapy Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 46.00% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not | HbA1c change, Weight change Follow up: 5.5 months | Study location: Nantong University, Nantong, China Sources of funding: AstraZeneca and 3SBio Inc. |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|------------|---|--|--|---|
| • | risk: Not stated/unclear Mean age (SD): 58.235 (10.9911) years Time since type 2 diabetes diagnosed: Not stated/unclear | stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 36.8% | | |
| Wang 2020B | Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Mixed population T2DM and chronic kidney disease: People with chronic kidney disease: T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.05 (8.462) years Time since type 2 diabetes diagnosed: 11.15 (6.6121) years | Strategy: Adding N = 92 Exenatide 10 mcg twice daily (n=46) Insulin lispro thrice daily (n=46) Concomitant therapy: Insulin glargine Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 26.1% Biguanides: 38.00% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 75% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 15.2% | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: 4 hospitals in Guangzhou, China Sources of funding: AstraZeneca China and 3SBio Inc. funded study and provided drugs and examination items during follow up. |
| Wang 2020C | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear | Strategy: Adding N = 60 Sitagliptin (n=30) Liraglutide (n=30) Concomitant therapy: existing oral antidiabetic therapy Antihyperglycaemic treatment received: No additional information available. | HbA1c change Follow up: 6 months | Study location: Linyi Peoples Hospital, Linyi, China Sources of funding: NR |

| | | Intomiontion and | | |
|------------|--|--|--|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Mean age (SD): 66.65 (6.5276) years Time since type 2 diabetes diagnosed: 8.59 (2.857) years | | | |
| Wang 2022B | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney diseases T2DM and chronic kidney disease T2DM and ronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 720 Insulin degludec/liraglutide once daily (n=361) Insulin degludec once daily (n=179) Liraglutide 1.8 mg once daily (n=180) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Nonfatal stroke, Unstable angina, Hypoglycaemia episodes, At night hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: China Sources of funding: Novo Nordisk |

| | | Intervention and | | |
|---------------------------------|--|--|--|---|
| Study | Population | comparison | Outcomes | Comments |
| Wang 2023 AWARD- CHN3 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.05 (9.4016) years Time since type 2 diabetes diagnosed: 11.8 (6.4703) years | Strategy: Adding N = 291 Dulaglutide 5 mg once weekly (n=144) Placebo once weekly (n=147) Concomitant therapy: Basal insluin glargine + metformin +/or acarbose Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 13.5% Biguanides: 76% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Acute kidney injury, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6.4 months | Study location: China Sources of funding: Eli Lilly and Company |
| Watada 2019 DUAL II Japan | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 210 Insulin degludec/liraglutide titrated twice weekly (n=105) Insulin degludec titrated twice weekly (n=105) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, HbA1c change, Weight change Follow up: 6 months | Study location: Japan (Multicentre trial, 38 sites) Sources of funding: Funded by Novo Nordisk A/S |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|-----------------------|---|--|---|---|
| | Mean age (SD): 56.05 (10.202) years Time since type 2 diabetes diagnosed: 14.05 (7.6268) years | | | |
| Webb 2020 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Mean age (SD): 44.1 (6.4734) years Time since type 2 diabetes diagnosed: 4.45 (4.4503) years | Strategy: Adding N = 76 Liraglutide 0.6-1.8 mg weekly (n=38) Sitagliptin 100 mg daily (n=38) Concomitant therapy: Metformin and/or a sulfonylurea Antihyperglycaemic treatment received: No additional information available. | Non-fatal myocardial infarction, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 6 months | Study location: Diabetes Research Centre, University of Leicester, Leicester, UK Sources of funding: Funded by Novo Nordisk and supported by NIHR Leicester Biomedical Research Center, the NIHR CLAHRC-East Midlands, the NIHR Leicester Clinical Research Facility and The NIHR Leicester Clinical Trial Unit. |
| White 2013 EXAMINE | Model 1: People with type 2 diabetes and heart failure Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease T2DM and heart failure: Mixed population | Strategy: Adding N = 5380 Alogliptin (n=2701) Placebo (n=2679) Concomitant therapy: Anti- diabetic therapy other than DPP-4 inhibitor or GLP-1 receptor agonist Antihyperglycaemic treatment received: Alpha-glucosidase | All-cause mortality, Cardiovascular mortality, 3-point MACE, 4-point MACE, Non-fatal myocardial infarction, Nonfatal stroke, Unstable angina, Hospitalisation for heart failure, Development of end stage kidney disease, Hypoglycaemia | Study location: Multicenter Sources of funding: Takeda Development Center Americas |

| | | Intervention and | | |
|--------------------------|--|---|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Includes results for a subgroup for people with or without heart failure. Mean age (SD): Not stated/unclear Time since type 2 diabetes diagnosed: Not stated/unclear | inhibitors: Not stated/unclear Biguanides: 66.2% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 29.80% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 46.60% | episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 18 months | |
| Wilcox 2008 PROactive | Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of | Strategy: Adding N = 5238 Pioglitazone (n=2605) Placebo (n=2633) Concomitant therapy: Oral agents ± insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 10% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 0.20% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 19.5% | All-cause mortality, Cardiovascular mortality, 3-point MACE, Non-fatal myocardial infarction, Non-fatal stroke, Hospitalisation for heart failure, Cardiac arrhythmia, Hypoglycaemia episodes Follow up: 34.5 months | Study location: Multicenter Sources of funding: Takeda Europe R&D Centre Ltd, London, United Kingdom, and Eli Lilly and Company, Indianapolis, IN. |

| | | lutementing and | | |
|--------------------------------------|--|---|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| · | developing cardiovascular disease Mean age (SD): 61.75 (7.7012) years Time since type 2 diabetes diagnosed: Not stated/unclear | | | |
| Wilding 2012 Dapagliflozin 006 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Mixed population T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 59.3 (8.2289) years Time since type 2 diabetes diagnosed: 13.6 (7.269) years | Strategy: Adding N = 807 Dapagliflozin 10 mg daily (n=196) Dapagliflozin 5/10 mg daily (n=212) Dapagliflozin 2.5 mg daily (n=202) Placebo (n=197) Concomitant therapy: Insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 40.00% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Persistent signs of worsening kidney disease, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 24 months | Study location: Multicenter Sources of funding: Sponsored by Bristol-Myers Squibb and AstraZeneca. |
| Wilding 2013A CANTATA- MSU | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular | Strategy: Adding N = 469 Canagliflozin 100 mg (n=157) Canagliflozin 300 mg (n=156) Placebo (n=156) Concomitant therapy: Metformin + sulfonylurea Antihyperglycaemic | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change | Study location: Multicenter Sources of funding: Janssen Research & Development, LLC |

| | | Intervention and | | |
|-------------------------------------|--|--|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56.7667 (9.2827) years Time since type 2 diabetes diagnosed: 9.5667 (6.2795) years | treatment received: No additional information available. | | |
| Wilding 2013B | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57 (9.345) year years Time since type 2 diabetes diagnosed: 5.95 (4.8924) years | Strategy: Adding N = 182 Glipizide 5-20 mg (n=94) Placebo (n=88) Concomitant therapy: metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 100% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes Follow up: 6 months | Study location: 92 sites in: Germany, Hungary, Latvia, Lithuania, Poland, Romania, Sweden, UK, Chill e, Mexico, Peru Sources of funding: Astra Zeneca |
| Wiviott 2019 DECLARE- TIMI 58 | Model 1: People with type 2 diabetes and heart failure | Strategy: Adding N = 17160 Dapagliflozin (n=8582) Placebo (n=8578) | All-cause mortality, Cardiovascular mortality, 3-point MACE, Non-fatal myocardial | Study location: Multicenter Sources of |

| | | Intervention and | | |
|---------|---|--|---|--------------------------------|
| Study | Population | comparison | Outcomes | Comments |
| | Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease Model 3: People with type 2 diabetes and chronic kidney disease Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Mixed population T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Includes results for a subgroup for people with or without chronic kidney disease, heart failure and people with atherosclerotic cardiovascular disease. Mean age (SD): 63.95 (6.8) years Time since type 2 diabetes diagnosed: Not stated/unclear | Concomitant therapy: Use of other glucose lowering agents at discretion (other than SGLT-2 inhibitor, pioglitazone or rosiglitazone Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 82% DPP-4 inhibitors: 16.8% GLP-1 receptor agonists: 4.40% Insulin: 40.9% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 42.60% | infarction, Non-fatal stroke, Unstable angina, Hospitalisation for heart failure, Acute kidney injury, Persistent signs of worsening kidney disease, Development of end stage kidney disease, Death from renal causes, Cardiac arrhythmia, Diabetic ketoacidosis, Progression of liver disease, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 50.4 months | funding: Funded by AstraZeneca |
| Wu 2014 | Model 5: People with type 2 | Strategy: Adding N = 93 | HbA1c change, BMI change | Study location: China |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|---------------------------|--|--|---|---|
| | diabetes at higher risk of cardiovascular disease | Metformin 1500 mg daily (n=47) Pioglitazone 15 mg daily (n=46) | Follow up: 12 months | Sources of funding: No additional information. |
| | T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 60.2 (9.6496) years Time since type 2 diabetes diagnosed: Not stated/unclear | Concomitant therapy: Hypoglycaemic drugs (not specified) other than biguanides and thiazolidinediones Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 26.9% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 24.7% | | |
| Wysham 2014 AWARD-1 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 976 Dulaglutide 1.5 mg once weekly (n=279) Dulaglutide 0.75 mg once weekly (n=280) Exenatide 10 micrograms twice daily (n=276) Placebo twice daily (n=141) Concomitant therapy: Metformin + thiazolidinedione Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Severe hypoglycaemic episodes, HbA1c change Follow up: 12 months | Study location: USA Sources of funding: Eli Lilly and company |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|-----------|--|---|---|--|
| , | Mean age (SD): 55.5 (9.7235) year years Time since type 2 diabetes diagnosed: 9 (5.7308) years | , | | |
| Xiao 2015 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 53.855 (3.5036) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 120 Pioglitazone 15-45 mg daily (n=40) Glipizide 5-10 mg daily (n=40) Insulin (n=40) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | HbA1c change Follow up: 5.5 months | Study location: Anhui, China Sources of funding: Financially supported by the Natural Science Foundation of Anhui Province (09B117) |
| Xiao 2016 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney | Strategy: Adding N = 41 Glimepiride 4 mg daily (n=18) Sitagliptin 100 mg daily (n=23) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Non-fatal myocardial infarction, Non- fatal stroke, Hospitalisation for heart failure, HbA1c change, Weight change Follow up: 5.5 months | Study location: Qilu Hospital, Shandong University, Jinan, China Sources of funding: Supported by grants from special funds for scientific research projects of clinical medicine of the Chinese Medical Association (grant no. |

| | | lutam autian aud | | |
|---------|--|--|--|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 68.9 (6.3879) years Time since type 2 diabetes diagnosed: Not stated/unclear | | | 13060990484), the Medicine Health Care Science and Technology Development Project Program of Shandong Province (grant no. 2013WSC02036), Science Foundation of Qilu Hospital of Shandong University (grant no. 2015QLMS11) and Fundamental Research Funds of Shandong University (2601017561601 2). |
| Xu 2017 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 53.4 (9.8502) years Time since type 2 diabetes diagnosed: 5.65 (4.3011) years | Strategy: Adding N = 1103 Glimepiride (n=551) Gliclazide (n=552) Concomitant therapy: Metformin + sitagliptin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Nonfatal stroke, Unstable angina, Diabetic ketoacidosis, Falls requiring hospitalisation, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: 237 centres across 25 provinces in China Sources of funding: Merck & Co., Inc. |

| | | late was offered | | |
|-------------------------------|--|---|--|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| Yabe 2020 PIONEER 10 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.75 (10.1676) years Time since type 2 diabetes diagnosed: 9.425 (6.3286) years | Strategy: Adding N = 458 Semaglutide 3 mg weekly (n=131) Semaglutide 7 mg daily (n=132) Semaglutide 14 mg daily (n=130) Dulaglutide 0.75 mg weekly (n=65) Concomitant therapy: Glucose-lowering drugs Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 17% Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: 17% Sulfonylureas: 32% | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Acute kidney injury, Death from renal causes, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 12 months | Study location: Japan (36 clinics and hospitals) Sources of funding: Funded by Novo Nordisk, Denmark. |
| Yabe 2023 EMPA- ELDERLY | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 127 Empagliflozin (n=64) Placebo (n=63) Concomitant therapy: DPP-4 inhibitors, biguanides, sulfonylureas, thiazolidinediones, alpha-glucosidase inhibitors and/or meglitinides Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 6.3% Biguanides: 51.20% DPP-4 inhibitors: 67.7% GLP-1 receptor agonists: Not stated/unclear | Health-related quality of life, All-cause mortality, Persistent signs of worsening kidney disease, Diabetic ketoacidosis, Progression of liver disease, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 12 months | Study location: Japan. Sources of funding: Sponsored by Nippon Boehringer Ingelheim Co. Ltd and Eli Lilly K.K. |

| | | Into manting and | | |
|----------------------|---|---|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| · | Mean age (SD): 74.1 (5.0002) years Time since type 2 diabetes diagnosed: 12.1 (7.9081) years | Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 7.80% | | |
| Yale 2013 DIA3004 | Model 3: People with type 2 diabetes and chronic kidney disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Mixed population T2DM and chronic kidney disease: People with chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 68.5333 (8.2675) years Time since type 2 diabetes diagnosed: 16.3333 (8.5194) years | Strategy: Adding N = 269 Canagliflozin 100 (n=90) Canagliflozin 300 (n=89) Placebo (n=90) Concomitant therapy: None, monotherapy or combination therapy Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 1.5% DPP-4 inhibitors: 7.5% GLP-1 receptor agonists: Not stated/unclear Insulin: 74% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 31.2% | All-cause mortality, HbA1c change, Weight change Follow up: 12 months | Study location: Multicenter Sources of funding: Janssen Research & Development, LLC. |
| Yan 2019 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not | Strategy: Adding N = 75 Liraglutide 1.8 mg daily (n=24) Sitagliptin 100 mg daily (n=27) Insulin glargine (n=24) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, HbA1c change, Weight change, BMI change Follow up: 6 months | Study location: China (10 centres) Sources of funding: Supported by investigator- initiated trial research funds from Novo Nordisk, National Natural Science Foundation of China (81770821), Pearl River S&T Nova Program of Guangzhou |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|-----------|---|---|--|---|
| | stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 44.8 (8.8922) years Time since type 2 diabetes diagnosed: 4.4667 (3.9492) years | | | (201610010175) and Guangdong High-Level Talents Special Support Program (2016TQ03R590) |
| Yang 2011 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.1 (10.25) years Time since type 2 diabetes diagnosed: 5.1 (4.5242) years | Strategy: Adding N = 570 Saxagliptin 5 mg daily (n=283) Placebo (n=287) Concomitant therapy: Metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 79.20% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | Non-fatal stroke, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: International (40 sites in China, India, South Korea) Sources of funding: Funded by AstraZeneca LP and Bristol-Myers Squibb. |
| Yang 2012 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: People without heart failure | Strategy: Adding N = 395 Sitagliptin 100 mg daily (n=197) Placebo (n=198) Concomitant therapy: Metformin Antihyperglycaemic treatment received: | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change | Study location: China (17 sites) Sources of funding: Funded by Merck Sharp & Dohme Corp., subsidiary of Merck & Co, Inc. |

| | | Intervention and | | |
|-----------|--|---|--|---|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 54.6 (9.4095) years Time since type 2 diabetes diagnosed: 6.85 (4.5014) years | No additional information available. | Follow up: 5.5 months | |
| Yang 2015 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: People without atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.5 (9.5596) years Time since type 2 diabetes diagnosed: 6.9 (4.3635) years | Strategy: Adding N = 279 Vildagliptin 50 mg daily (n=143) Placebo (n=136) Concomitant therapy: Glimepiride Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Nonfatal stroke, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 5.5 months | Study location: China (multisite trial) Sources of funding: Novartis Pharmaceuticals |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|------------|--|--|--|---|
| Yang 2016 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 53.7333 (9.2712) years Time since type 2 diabetes diagnosed: 4.9333 (4.2834) years | Strategy: Adding N = 444 Dapagliflozin 10 mg daily (n=152) Dapagliflozin 5 mg daily (n=147) Placebo (n=145) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Development of end stage kidney disease, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: International (32 sites in China, India and South Korea) Sources of funding: Funded by Bristol-Myers Squibb, NJ, USA, and AstraZeneca, MD, USA |
| Yang 2018A | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear | Strategy: Adding N = 272 Dapagliflozin 10 mg daily (n=139) Placebo (n=133) Concomitant therapy: Insulin with or without oral antidiabetic drugs Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: 13.2% Biguanides: 45.2% DPP-4 inhibitors: 5.5% GLP-1 receptor agonists: Not stated/unclear Insulin: 39.7% SGLT-2 inhibitors: | All-cause mortality, Cardiovascular mortality, Nonfatal myocardial infarction, Development of end stage kidney disease, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: International (28 sites in China, Singapore and South Korea) Sources of funding: Funded by AstraZeneca |

| | | Into manting and | | |
|---------------------------|--|---|---|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | Mean age (SD): 57.55 (8.6481) years Time since type 2 diabetes diagnosed: 12.45 (6.96) years | Not stated/unclear Sulfonylureas: 11% | | |
| Yang 2018B GetGoal-L-C | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 55.1 (9.5084) years Time since type 2 diabetes diagnosed: 10.25 (6.1502) years | Strategy: Adding N = 448 Lixisenatide 20 mcg daily (n=224) Placebo (n=224) Concomitant therapy: Insulin with or without metformin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 88.6% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 82.80% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: International (51 centres in China, India, South Korea and Russian Federation) Sources of funding: Funded by Sanofi |
| Yang 2021 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney | Strategy: Adding N = 206 Linagliptin 5 mg daily (n=104) Placebo (n=102) Concomitant therapy: Insulin Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: China (25 sites) Sources of funding: Funded by Boehringer Ingelheim |

| | | Intervention and | | | | |
|---------------------------|---|---|---|--|--|--|
| Study | Population | comparison | Outcomes | Comments | | |
| | disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.6 (10.0597) years Time since type 2 diabetes diagnosed: Not stated/unclear | Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 30.2% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | | | | |
| Yang 2022 LixiLan-O-AP | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 56 (9.7) years Time since type 2 diabetes diagnosed: Not stated/unclear | Strategy: Adding N = 206 IGlarLixi (n=351) Insulin glargine (n=350) Lixisenatide (n=177) Concomitant therapy: All people had the opportunity to receive a second oral antihyperglycaemic drug Antihyperglycaemic treatment received: No additional information | All-cause mortality, Cardiovascular mortality, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 5.5 months | Study location: China Sources of funding: Sanofi | | |
| Yki-Järvinen 2013 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and | Strategy: Adding N = 1261 Linagliptin 5 mg daily (n=631) Placebo (n=630) Concomitant therapy: Insulin ± metformin and/or thiazolidinedione | All-cause mortality, Cardiovascular mortality, 5-point MACE, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change | Study location: Multicenter Sources of funding: Sponsored by Boehringer Ingelheim. | | |

| | | Intervention and | | |
|-----------------------------|---|---|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 60.05 (9.9501) years Time since type 2 diabetes diagnosed: Not stated/unclear | Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 75.60% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear Sulfonylureas: Not stated/unclear | Follow up: 12 months | |
| Yokoyama 2014 JDDM 33 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 61.3 (9.1609) years Time since type 2 diabetes diagnosed: 11.335 (6.9523) years | Strategy: Adding N = 99 Liraglutide 0.9 mg daily (n=50) Sitagliptin 50-100 mg daily (n=49) Concomitant therapy: Sulfonylurea Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: Not stated/unclear DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 46.40% | Health-related quality of life, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 5.5 months | Study location: Japan (21 primary care centres) Sources of funding: Supported by Japan Diabetes Foundation. |
| Yuan 2022 LixiLan-L-CN | Model 5: People with type 2 diabetes at higher | Strategy: Adding N = 426 | All-cause mortality, Cardiovascular | Study location: China (44 centres) |

| | | Intervention and | | |
|----------------------------------|---|---|---|---|
| Study | Population | comparison | Outcomes | Comments |
| | risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.45 (9.0064) years Time since type 2 diabetes diagnosed: 12.35 (6.1003) years | iGlarLixi (n=212) Insulin glargine (n=214) Concomitant therapy: Background oral antidiabetic treatment Antihyperglycaemic treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 78.20% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: 85% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: Not stated/unclear | mortality, Death from renal causes, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6.9 months | Sources of funding: Funded by Sanofi, Paris, France. |
| Zang 2016 LIRA-DPP-4 CHINA | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 51.55 (10.851) | Strategy: Adding N = 368 Liraglutide 1.8 mg daily (n=184) Sitagliptin 100 mg daily (n=184) Concomitant therapy: Metformin Antihyperglycaemic treatment received: No additional information available. | All-cause mortality, Cardiovascular mortality, Nonfatal stroke, Cardiac arrhythmia, Diabetic ketoacidosis, Hypoglycaemia episodes, At night hypoglycaemic episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 6 months | Study location: China (25 sites) Sources of funding: Funded by Novo Nordisk |

| | | Intervention and | | |
|-------------|---|--|---|---|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| | years Time since type 2 diabetes diagnosed: 5.25 (4.9254) years | | | |
| Zhang 2020B | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 58.44 (12.97) years Time since type 2 diabetes diagnosed: 7.38 (5.7114) years | Strategy: Adding N = 59 Exenatide 5-10 ι/₄g twice daily (n=27) Insulin (initially 0.2-0.4 IU/Kg then titrated) daily (n=32) Concomitant therapy: Any antihyperglycaemic other than sulfunylurea and nateglinide Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 12 months | Study location: China Sources of funding: Astra Zeneca and 3SBioInc. |
| Zhao 2017 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: People without chronic kidney disease | Strategy: Adding N = 100 Sitagliptin (n=50) Placebo (n=50) Concomitant therapy: Insulin, exenatide or metformin Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, HbA1c change, Weight change, BMI change Follow up: 9 months | Study location: China. Sources of funding: None declared. |

| | | 1 | | |
|------------------------------------|--|--|---|--|
| Study | Population | Intervention and comparison | Outcomes | Comments |
| · | T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 68.45 (7.9627) years Time since type 2 diabetes diagnosed: 5.65 (4.4045) years | | | |
| Zinman 2009 LEAD-4 Met+TZD | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 53.6667 (29.4694) years Time since type 2 diabetes diagnosed: 9 (6) years | Strategy: Adding N = 533 Liraglutide 1.2 mg (n=178) Liraglutide 1.8 mg (n=178) Placebo (n=177) Concomitant therapy: Metformin + TZD Antihyperglycaemic treatment received: No additional information available. | Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change Follow up: 6 months | Study location: Multi-centre (96 sites) study conducted in the USA and Canada. Sources of funding: Funding source not clearly stated. Statistical and writing assistance was provided by staff from Novo Nordisk. |
| Zinman 2015 EMPA-REG OUTCOME | Model 1: People with type 2 diabetes and heart failure Model 2: People with type 2 diabetes and atherosclerotic cardiovascular disease Model 3: People with type 2 | Strategy: Adding N = 7020 Empagliflozin (n=4687) Placebo (n=2333) Concomitant therapy: No or stable glucose lowering therapy Antihyperglycaemic | All-cause mortality, Cardiovascular mortality, 3-point MACE, 4-point MACE, Non-fatal myocardial infarction, Nonfatal stroke, Unstable angina, Hospitalisation for heart failure, Acute kidney | Study location: Multicenter Sources of funding: Supported by Boehringer Ingelheim and Eli Lilly |

| Study | Population | Intervention and comparison | Outcomes | Comments |
|------------------------------|--|--|---|--|
| | diabetes and chronic kidney disease T2DM and heart failure: People without heart failure T2DM and atherosclerotic cardiovascular disease: People with atherosclerotic cardiovascular diseases T2DM and chronic kidney disease: Mixed population T2DM and higher cardiovascular risk: People at higher risk of developing cardiovascular disease Includes results for a subgroup for people with or without heart failure and with or without chronic kidney disease. Mean age (SD): 63.15 (8.667) years Time since type 2 diabetes diagnosed: Not stated/unclear | treatment received: Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 74.00% DPP-4 inhibitors: 11.40% GLP-1 receptor agonists: 2.80% Insulin: 48.3% SGLT-2 inhibitors: Not stated/unclear Sulfonylureas: 42.80% | injury, Persistent signs of worsening kidney disease, Development of end stage kidney disease, Diabetic ketoacidosis, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change Follow up: 36 months | |
| Zinman 2019A SUSTAIN 9 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear | Strategy: Adding N = 302 Semaglutide 1.0 mg (n=151) Placebo (n=151) Concomitant therapy: SGLT-2 inhibitor as monotherapy or with a sulfonylurea or metformin Antihyperglycaemic treatment received: | Health-related quality of life, All-cause mortality, Cardiovascular mortality, Acute kidney injury, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 7 months | Study location: Multicenter Sources of funding: Novo Nordisk |

| | | Intervention and | | |
|------------------------------|---|---|--|--|
| Study | Population | comparison | Outcomes | Comments |
| | T2DM and chronic kidney disease: People without chronic kidney disease T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 57.05 (9.5189) years Time since type 2 diabetes diagnosed: 9.7 (6.1033) years | Alpha-glucosidase inhibitors: Not stated/unclear Biguanides: 71.5% DPP-4 inhibitors: Not stated/unclear GLP-1 receptor agonists: Not stated/unclear Insulin: Not stated/unclear SGLT-2 inhibitors: 99.60% Sulfonylureas: 12.9% | | |
| Zinman 2019B PIONEER 8 | Model 5: People with type 2 diabetes at higher risk of cardiovascular disease T2DM and heart failure: Not stated/unclear T2DM and atherosclerotic cardiovascular disease: Not stated/unclear T2DM and chronic kidney disease: Not stated/unclear T2DM and higher cardiovascular risk: Not stated/unclear Mean age (SD): 60.5 (9.7576) years Time since type 2 diabetes diagnosed: 15.05 (8.1036) years | Strategy: Adding N = 730 Semaglutide 3 mg (n=184) Semaglutide 7 mg (n=181) Semaglutide 14 mg (n=181) Placebo (n=184) Concomitant therapy: Insulin +/-metformin Antihyperglycaemic treatment received: No additional information available. | Health-related quality of life, All-cause mortality, Non-fatal myocardial infarction, Non-fatal stroke, Hospitalisation for heart failure, Acute kidney injury, Hypoglycaemia episodes, Severe hypoglycaemic episodes, HbA1c change, Weight change, BMI change Follow up: 12 months | Study location: Multicenter Sources of funding: PIONEER 8 was funded by Novo Nordisk A/S Denmark. |

1 See report 1.2 B, appendix D for full evidence tables.

1.1.6. Summary of the effectiveness evidence (network meta-analysis) – combined strategies

Table 4: Summary of effectiveness evidence (network meta-analysis) - people with type 2 diabetes and heart failure

| | , | | | | | 7 1 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 | | | |
|--|----------------|--------------|-------------|------------------|--------------|---|---------------|---------------------------|----------|
| Outcome | No. of studies | Study design | Sample size | Effect estimates | Risk of bias | Indirectness | Inconsistency | Imprecision | Quality |
| Cardiovascular mortality at 33 months | 6 | RCT | 8512 | See report F9 | Not serious | Not serious | Not serious | Very serious _a | Low |
| 3-point MACE at 36 months | 8 | RCT | 11956 | See report F9 | Not serious | Not serious | Not serious | Serious₅ | Moderate |
| Non-fatal myocardial infarction at 38 months | 3 | RCT | 4924 | See report F9 | Serious | Not serious | Not serious | Very serious _a | Very low |
| Non-fatal stroke at 32 months | 4 | RCT | 5178 | See report F9 | Serious | Not serious | Not serious | Very serious _a | Very Low |
| Hospitalisation for heart failure at 32 months | 10 | RCT | 15123 | See report F9 | Not serious | Not serious | Not serious | Very serious _a | Low |
| HbA1c change at 9 months | 2 | RCT | 55 | See report F9 | Seriousc | Not serious | Not serious | Very serious _a | Very low |

Footnotes:

1

2

9

10

- a) Downgraded by 2 increments as a significant proportion of inputs or measures produced by the network meta-analysis showed imprecision that had a substantial impact on the ability of the committee to draw conclusions from the results of the analysis.
- b) Downgraded by 1 increment as a significant proportion of inputs or measures produced by the network meta-analysis showed imprecision that had a moderate impact on the ability of the committee to draw conclusions from the results of the analysis.
- c) Downgraded by 1 increment as greater than 33.3% of the studies in the meta-analysis were at high or moderate risk of bias.

Table 5: Summary of effectiveness evidence (network meta-analysis) - people with type 2 diabetes and cardiovascular disease

| Outcome | No. of studies | Study design | Sample size | Effect estimates | Risk of bias | Indirectness | Inconsistency | Imprecision | Quality |
|---------------------------------------|----------------|-----------------|-------------|------------------|--------------|--------------|---------------|----------------------|---------|
| Cardiovascular mortality at 27 months | 12 | RCT | 62776 | See report F9 | Seriousa | No serious | No serious | Serious _b | Low |

| Outcome | No. of studies | Study design | Sample size | Effect estimates | Risk of bias | Indirectness | Inconsistency | Imprecision | Quality |
|--|----------------|-----------------|-------------|------------------|------------------------------|--------------|---------------|---------------------------|----------|
| 3-point MACE at 40 months | 12 | RCT | 83664 | See report F9 | Not serious | Not serious | Not serious | Serious _b | Moderate |
| Hospitalisation for heart failure at 29 months | 11 | RCT | 62683 | See report F9 | Not serious | Not serious | Not serious | Very serious | Low |
| Non-fatal myocardial infarction at 28 months | 11 | RCT | 55863 | See report F9 | Very serious _c | Not serious | Not serious | Serious _c | Very low |
| Non-fatal stroke at 31 months | 8 | RCT | 41022 | See report F9 | Not serious | Not serious | Not serious | Very serious _c | Low |
| Unstable angina at 24 months | 7 | RCT | 42392 | See report F9 | Very serious _d | Not serious | Not serious | Very serious _c | Very low |
| HbA1c change at 18 months | 18 | RCT | 34325 | See report F9 | Very serious _d | Not serious | Not serious | Serious _c | Very low |
| Weight change at 8 months | 5 | RCT | 1888 | See report F9 | Very serious _d | Not serious | Not serious | Not serious | Low |

1 Footnotes:

8

9

- a) Downgraded by 1 increment as greater than 33.3% of the studies in the meta-analysis were at high or moderate risk of bias.
- b) Downgraded by 1 increment as a significant proportion of inputs or measures produced by the network meta-analysis showed imprecision that had a moderate impact on the ability of the committee to draw conclusions from the results of the analysis.
- c) Downgraded by 2 increments as a significant proportion of inputs or measures produced by the network meta-analysis showed imprecision that had a substantial impact on the ability of the committee to draw conclusions from the results of the analysis.
- d) Downgraded by 2 increments as greater than 33.3% of the studies in the meta-analysis were at high risk of bias.

Table 6: Summary of effectiveness evidence (network meta-analysis) - people with type 2 diabetes and chronic kidney disease

| Outcome | No. of studies | Study design | Sample size | Effect estimates | Risk of bias | Indirectness | Inconsistency | Imprecision | Quality |
|---------------------------------------|----------------|-----------------|-------------|------------------|--------------|--------------|---------------|---------------------------|---------|
| Cardiovascular mortality at 22 months | 9 | RCT | 14523 | See report F9 | Not serious | Not serious | Not serious | Very serious _a | Low |

| Outcome | No. of studies | Study design | Sample size | Effect estimates | Risk of bias | Indirectness | Inconsistency | Imprecision | Quality |
|--|----------------|-----------------|-------------|------------------|------------------------------|--------------|---------------|---------------------------|----------|
| 3-point MACE at 35 months | 3 | RCT | 10593 | See report F9 | Serious _b | Not serious | Not serious | Serious _c | Low |
| Non-fatal myocardial infarction at 22 months | 3 | RCT | 632 | See report F9 | Very serious _d | Not serious | Not serious | Very serious _a | Very low |
| Non-fatal stroke at 8 months | 3 | RCT | 731 | See report F9 | Serious _b | Not serious | Not serious | Very serious _a | Very low |
| Hospitalisation for heart failure at 27 months | 9 | RCT | 18277 | See report F9 | Not serious | Not serious | Not serious | Very serious _a | Low |
| Development of end stage kidney disease at 22 months | 3 | RCT | 4823 | See report F9 | Serious₀ | Not serious | Not serious | Very serious _a | Very low |
| HbA1c change at 14 months | 24 | RCT | 10109 | See report F9 | Very serious _d | Not serious | Not serious | Serious _c | Very low |
| Weight change at 12 months | 6 | RCT | 1883 | See report F9 | Very serious _d | Not serious | Not serious | Very serious _a | Very low |

1 Footnotes:

- a) Downgraded by 2 increments as a significant proportion of inputs or measures produced by the network meta-analysis showed imprecision that had a substantial impact on the ability of the committee to draw conclusions from the results of the analysis.
- b) Downgraded by 1 increment as greater than 33.3% of the studies in the meta-analysis were at high or moderate risk of bias.
- c) Downgraded by 1 increment as a significant proportion of inputs or measures produced by the network meta-analysis showed imprecision that had a moderate impact on the ability of the committee to draw conclusions from the results of the analysis.
- d) Downgraded by 2 increments as greater than 33.3% of the studies in the meta-analysis were at high risk of bias.

9 Table 7: Summary of effectiveness evidence (network meta-analysis) - people with type 2 diabetes and high cardiovascular risk

| | No. of | Study | Sample | Effect | Risk of | | | | . |
|-----------------------------|---------|--------|--------|------------|-------------|--------------|---------------|-------------|----------|
| Outcome | studies | design | size | estimates | bias | Indirectness | Inconsistency | Imprecision | Quality |
| Cardiovascular mortality at | 84 | RCT | 164385 | See report | Not serious | Not serious | Not serious | Seriousa | Moderate |
| 15.9 months | | | | F10 | | | | | |

| Outcome | No. of studies | Study design | Sample size | Effect estimates | Risk of bias | Indirectness | Inconsistency | Imprecision | Quality |
|---|----------------|-----------------|-------------|------------------------|------------------|--------------|---------------|---------------------------|----------|
| 3-point MACE at 31.3 months | 17 | RCT | 106710 | See report F10 | Not serious | Not serious | Not serious | Serious _a | Moderate |
| 4-point MACE at 22.2 months | 7 | RCT | 20290 | See report F10 | Not serious | Not serious | Not serious | Very serious _b | Low |
| 5-point MACE at 18.5 months | 3 | RCT | 7741 | See report F10 | Not serious | Not serious | Not serious | Serious _a | Moderate |
| Non-fatal myocardial infarction at 15.1 months | 48 | RCT | 111619 | See report F10 | Not serious | Not serious | Not serious | Seriousa | Moderate |
| Non-fatal stroke at 15.6 months | 47 | RCT | 106740 | See report F10 | Not serious | Not serious | Not serious | Seriousa | Moderate |
| Unstable angina at 19.6 months | 29 | RCT | 105389 | See report F10 | Not serious | Not serious | Not serious | Serious _a | Moderate |
| Hospitalisation for heart failure (base case) at 22.8 months | 33 | RCT | 118970 | See report F10 | Not serious | Not serious | Not serious | Seriousa | Moderate |
| Hospitalisation for heart failure (sensitivity analysis) at 22.8 months | 33 | RCT | 120217 | See report F10 | Not serious | Not serious | Not serious | Very serious _b | Low |
| Development of end stage kidney disease at 34.7 months | 9 | RCT | 85882 | See report F10 | Not serious | Not serious | Not serious | Seriousa | Moderate |
| HbA1c change at 10 months | 335 | RCT | 264168 | See report F11-F12 | Very serious₅ | Not serious | Not serious | Not serious | Low |
| HbA1c change (regression analysis) at 10 months | 308 | RCT | 224786 | See report F11-F12 | Very serious₀ | Not serious | Not serious | Not serious | Low |
| Weight change at 9 months | 173 | RCT | 105529 | See report F11, F13 | Very serious₀ | Not serious | Not serious | Seriousa | Very low |

Note: Unless specified, the GRADE rating for the base case and sensitivity analyses are the same. The base case values are reported in the table.

Footnotes:

a) Downgraded by 1 increment as a significant proportion of inputs or measures produced by the network meta-analysis showed imprecision that had a moderate impact on the ability of the committee to draw conclusions from the results of the analysis.

- b) Downgraded by 2 increments as a significant proportion of inputs or measures produced by the network meta-analysis showed imprecision that had a substantial impact on the ability of the committee to draw conclusions from the results of the analysis.
- c) Downgraded by 2 increments as greater than 33.3% of the studies in the meta-analysis were at high risk of bias.

1 1.1.7. References

- 2 1. Abdul-Ghani M, Migahid O, Megahed A, Adams J, Triplitt C, DeFronzo RA et al.
 3 Combination Therapy With Exenatide Plus Pioglitazone Versus Basal/Bolus Insulin in
 4 Patients With Poorly Controlled Type 2 Diabetes on Sulfonylurea Plus Metformin: The
 5 Qatar Study. Diabetes Care. 2017; 40(3):325-331
- Abreu M, Tumyan A, Elhassan A, Peicher K, Papacostea O, Dimachkie P et al. A randomized trial comparing the efficacy and safety of treating patients with type 2 diabetes and highly elevated HbA1c levels with basal-bolus insulin or a glucagon-like peptide-1 receptor agonist plus basal-bolus insulin: the SIMPLE study. Diabetes Obes Metab. 2019; 21(9):2133-2141
- Adel SMH, Jorfi F, Mombeini H, Rashidi H, Fazeli S. Effect of a low dose of empagliflozin on short-term outcomes in type 2 diabetics with acute coronary syndrome after percutaneous coronary intervention. Saudi Medical Journal. 2022; 43(5):458-464
- Ahmann A, Rodbard HW, Rosenstock J, Lahtela JT, Loredo L, Tornoe K et al.
 Efficacy and safety of liraglutide versus placebo added to basal insulin analogues
 (with or without metformin) in patients with type 2 diabetes: A randomized, placebo-controlled trial. Diab Obes Metab. 2015; 17(11):1056-1064
- Ahmann AJ, Capehorn M, Charpentier G, Dotta F, Henkel E, Lingvay I et al. Efficacy
 and Safety of Once-Weekly Semaglutide Versus Exenatide ER in Subjects With Type
 Diabetes (SUSTAIN 3): A 56-Week, Open-Label, Randomized Clinical Trial.
 Diabetes Care. 2018; 41(2):258-266
- Ahr?n B, Gomis R, Standl E, Mills D, Schweizer A. Twelve- and 52-week efficacy of the dipeptidyl peptidase IV inhibitor LAF237 in metformin-treated patients with type 2 diabetes. Diabetes Care. 2004; 27(12):2874-2880
- Ahr?n B, Leguizamo Dimas A, Miossec P, Saubadu S, Aronson R. Efficacy and safety of lixisenatide once-daily morning or evening injections in type 2 diabetes inadequately controlled on metformin (GetGoal-M). Diabetes Care. 2013; 36(9):2543-2550
- 30 8. Ahr?n B, Masmiquel L, Kumar H, Sargin M, Karsbol JD, Jacobsen SH et al. Efficacy 31 and safety of once-weekly semaglutide versus once-daily sitagliptin as an add-on to 32 metformin, thiazolidinediones, or both, in patients with type 2 diabetes (SUSTAIN 2): 33 A 56-week, double-blind, phase 3a, randomised trial. Lancet Diabetes Endocrinol. 34 2017; 5(5):341-354
- 35 9. Ahren B, Johnson SL, Stewart M, Cirkel DT, Yang F, Perry C et al. HARMONY 3: 104-week randomized, double-blind, placebo- and active-controlled trial assessing the efficacy and safety of albiglutide compared with placebo, sitagliptin, and glimepiride in patients with type 2 diabetes taking metformin. Diabetes Care. 2014; 37(8):2141-2148
- 40 10. Ando Y, Shigiyama F, Hirose T, Kumashiro N. Simplification of complex insulin 41 regimens using canagliflozin or liraglutide in patients with well-controlled type 2 42 diabetes: A 24-week randomized controlled trial. Journal of Diabetes Investigation. 43 2021; 12(10):1816-1826
- 44 11. Araki E, Inagaki N, Tanizawa Y, Oura T, Takeuchi M, Imaoka T. Efficacy and safety 45 of once-weekly dulaglutide in combination with sulphonylurea and/or biguanide 46 compared with once-daily insulin glargine in Japanese patients with type 2 diabetes:

- a randomized, open-label, phase III, non-inferiority study. Diabetes Obes Metab. 2 2015; 17(10):994-1002
- Araki E, Tanizawa Y, Tanaka Y, Taniguchi A, Koiwai K, Kim G et al. Long-term treatment with empagliflozin as add-on to oral antidiabetes therapy in Japanese patients with type 2 diabetes mellitus. Diabetes Obes Metab. 2015; 17(7):665-674
- Arechavaleta R, Seck T, Chen Y, Krobot KJ, O'Neill EA, Duran L et al. Efficacy and safety of treatment with sitagliptin or glimepiride in patients with type 2 diabetes inadequately controlled on metformin monotherapy: a randomized, double-blind, non-inferiority trial. Diabetes Obes Metab. 2011; 13(2):160-168
- 14. Aroda VR, Bain SC, Cariou B, Piletic M, Rose L, Axelsen M et al. Efficacy and safety of once-weekly semaglutide versus once-daily insulin glargine as add-on to metformin (with or without sulfonylureas) in insulin-naive patients with type 2 diabetes (SUSTAIN 4): A randomised, open-label, parallel-group, multicentre, multinational, phase 3a trial. Lancet Diabetes Endocrinol. 2017; 5(5):355-366
- 15. Aroda VR, Gonzalez-Galvez G, Gron R, Halladin N, Haluzik M, Jermendy G et al.
 16. Durability of insulin degludec plus liraglutide versus insulin glargine U100 as initial
 17. injectable therapy in type 2 diabetes (DUAL VIII): a multicentre, open-label, phase 3b,
 18. randomised controlled trial. The Lancet Diabetes & Endocrinology. 2019; 7(8):59619. 605
- Aroda VR, Rosenstock J, Wysham C, Unger J, Bellido D, Gonzalez-Galvez G et al.
 Efficacy and Safety of LixiLan, a Titratable Fixed-Ratio Combination of Insulin
 Glargine Plus Lixisenatide in Type 2 Diabetes Inadequately Controlled on Basal
 Insulin and Metformin: The LixiLan-L Randomized Trial. Diabetes Care. 2016;
 39(11):1972-1980
- 25 17. Arturi F, Succurro E, Miceli S, Cloro C, Ruffo M, Maio R et al. Liraglutide improves 26 cardiac function in patients with type 2 diabetes and chronic heart failure. Endocrine. 27 2017; 57(3):464-473
- 28 18. Aschner P, Chan J, Owens DR, Picard S, Wang E, Dain MP et al. Insulin glargine 29 versus sitagliptin in insulin-naive patients with type 2 diabetes mellitus uncontrolled 30 on metformin (EASIE): a multicentre, randomised open-label trial. Lancet. 2012; 31 379(9833):2262-2269
- 32 19. Attaran F, Emami S, Sohrabi M, Malek M, Ajdarkosh H, Khoonsari M et al. Effect of
 33 Empagliflozin and Pioglitazone on left ventricular function in patients with type two
 34 diabetes and nonalcoholic fatty liver disease without established cardiovascular
 35 disease: a randomized single-blind clinical trial. BMC Gastroenterology. 2023;
 36 23(1):327
- 37 20. Avil?s-Santa L, Sinding J, Raskin P. Effects of metformin in patients with poorly controlled, insulin-treated type 2 diabetes mellitus. A randomized, double-blind, placebo-controlled trial. Annals of Internal Medicine. 1999; 131(3):182-188
- 40 21. Ba J, Han P, Yuan G, Mo Z, Pan C, Wu F et al. Randomized trial assessing the 41 safety and efficacy of sitagliptin in Chinese patients with type 2 diabetes mellitus 42 inadequately controlled on sulfonylurea alone or combined with metformin. J 43 Diabetes. 2017; 9:667-676
- Babar M, Hussain M, Ahmad M, Akhtar L. Comparison Of Efficacy And Safety Profile
 Of Empagliflozin As A Combination Therapy In Obese Type 2 Diabetic Patients.
 Journal of Ayub Medical College, Abbottabad: JAMC. 2021; 33(2):188-191

- Bae J, Huh JH, Lee M, Lee YH, Lee BW. Glycaemic control with add-on thiazolidinedione or a sodium-glucose co-transporter-2 inhibitor in patients with type 2 diabetes after the failure of an oral triple antidiabetic regimen: A 24-week, randomized controlled trial. Diabetes, Obesity & Metabolism. 2021; 23(2):609-618
- 5 24. Bailey CJ, Gross JL, Pieters A, Bastien A, List JF. Effect of dapagliflozin in patients with type 2 diabetes who have inadequate glycaemic control with metformin: a randomised, double-blind, placebo-controlled trial. Lancet. 2010; 375(9733):2223-2233
- 9 25. Bailey TS, Takacs R, Tinahones FJ, Rao PV, Tsoukas GM, Thomsen AB et al.
 10 Efficacy and safety of switching from sitagliptin to liraglutide in subjects with type 2
 11 diabetes (LIRA-SWITCH): a randomized, double-blind, double-dummy, active12 controlled 26-week trial. Diabetes Obes Metab. 2016; 18(12):1191-1198
- 26. Bajaj M, Gilman R, Patel S, Kempthorne-Rawson J, Lewis-D'Agostino D, Woerle HJ.
 Linagliptin improved glycaemic control without weight gain or hypoglycaemia in patients with Type 2 diabetes inadequately controlled by a combination of metformin and pioglitazone: A 24-week randomized, double-blind study. Diabetic Medicine.
 2014; 31(12):1505-1514
- 18 27. Barnett AH, Charbonnel B, Donovan M, Fleming D, Chen R. Effect of saxagliptin as add-on therapy in patients with poorly controlled type 2 diabetes on insulin alone or insulin combined with metformin. Current Medical Research and Opinion. 2012; 28(4):513-523
- 22 28. Barnett AH, Huisman H, Jones R, Eynatten M, Patel S, Woerle HJ. Linagliptin for patients aged 70 years or older with type 2 diabetes inadequately controlled with common antidiabetes treatments: a randomised, double-blind, placebo-controlled trial. Lancet. 2013; 382(9902):1413-1423
- 29. Barnett AH, Mithal A, Manassie J, Jones R, Rattunde H, Woerle HJ et al. Efficacy and safety of empagliflozin added to existing antidiabetes treatment in patients with type 2 diabetes and chronic kidney disease: A randomised, double-blind, placebo-controlled trial. Lancet Diabetes Endocrinol. 2014; 2(5):369-384
- 30. Bergenstal R, Lewin A, Bailey T, Chang D, Gylvin T, Roberts V. Efficacy and safety of biphasic insulin aspart 70/30 versus exenatide in subjects with type 2 diabetes failing to achieve glycemic control with metformin and a sulfonylurea. Current Medical Research and Opinion. 2009; 25(1):65-75
- 31. Bergenstal RM, Wysham C, Macconell L, Malloy J, Walsh B, Yan P et al. Efficacy and safety of exenatide once weekly versus sitagliptin or pioglitazone as an adjunct to metformin for treatment of type 2 diabetes (DURATION-2): a randomised trial.

 37. Lancet. 2010; 376(9739):431-439
- 38 32. Berndt-Zipfel C, Michelson G, Dworak M, Mitry M, Loffler A, Pfutzner A et al.
 39 Vildagliptin in addition to metformin improves retinal blood flow and erythrocyte
 40 deformability in patients with type 2 diabetes mellitus results from an exploratory
 41 study. Cardiovascular Diabetology. 2013; 12(1):59
- 42 33. Billings LK, Doshi A, Gouet D, Oviedo A, Rodbard HW, Tentolouris N et al. Efficacy 43 and Safety of IDegLira Versus Basal-Bolus Insulin Therapy in Patients With Type 2 44 Diabetes Uncontrolled on Metformin and Basal Insulin: The DUAL VII Randomized 45 Clinical Trial. Diabetes Care. 2018; 41(5):1009-1016
- 46 34. Bizino MB, Jazet IM, Westenberg JJM, Van Eyk HJ, Paiman EHM, Smit JWA et al.
 47 Effect of liraglutide on cardiac function in patients with type 2 diabetes mellitus:
 48 randomized placebo-controlled trial. Cardiovascular Diabetology. 2019; 18(1):55

- Blonde L, Belousova L, Fainberg U, Garcia-Hernandez PA, Jain SM, Kaltoft MS et al. Liraglutide as add-on to sodium-glucose co-transporter-2 inhibitors in patients with inadequately controlled type 2 diabetes: LIRA-ADD2SGLT2i, a 26-week, randomized, double-blind, placebo-controlled trial. Diab Obes Metab. 2020;
- 5 36. Blonde L, Jendle J, Gross J, Woo V, Jiang H, Fahrbach JL et al. Once-weekly dulaglutide versus bedtime insulin glargine, both in combination with prandial insulin lispro, in patients with type 2 diabetes (AWARD-4): a randomised, open-label, phase 3, non-inferiority study. Lancet. 2015; 385(9982):2057-2066
- 9 37. Bode B, Stenl?f K, Sullivan D, Fung A, Usiskin K. Efficacy and safety of canagliflozin 10 treatment in older subjects with type 2 diabetes mellitus: a randomized trial. Hosp 11 Pract (1995). 2013; 41(2):72-84
- 38. Bolinder J, Ljunggren, Kullberg J, Johansson L, Wilding J, Langkilde AM et al. Effects of dapagliflozin on body weight, total fat mass, and regional adipose tissue distribution in patients with type 2 diabetes mellitus with inadequate glycemic control on metformin. Journal of Clinical Endocrinology and Metabolism. 2012; 97(3):1020-1031
- 39. Bolli G, Dotta F, Rochotte E, Cohen SE. Efficacy and tolerability of vildagliptin vs.
 pioglitazone when added to metformin: a 24-week, randomized, double-blind study.
 Diabetes, Obesity & Metabolism. 2008; 10(1):82-90
- 40. Bolli GB, Munteanu M, Dotsenko S, Niemoeller E, Boka G, Wu Y et al. Efficacy and safety of lixisenatide once daily vs. placebo in people with Type 2 diabetes insufficiently controlled on metformin (GetGoal-F1). Diabetic Med. 2014; 31(2):176-184
- 24 41. Bosi E, Camisasca RP, Collober C, Rochotte E, Garber AJ. Effects of vildagliptin on glucose control over 24 weeks in patients with type 2 diabetes inadequately controlled with metformin. Diabetes Care. 2007; 30(4):890-895
- 27 42. Brown AJM, Gandy S, McCrimmon R, Houston JG, Struthers AD, Lang CC. A
 28 randomized controlled trial of dapagliflozin on left ventricular hypertrophy in people
 29 with type two diabetes: the DAPA-LVH trial. European Heart Journal. 2020;
 30 41(36):3421-3432
- 31 43. Bunck MC, Diamant M, Corn?r A, Eliasson B, Malloy JL, Shaginian RM et al. One-32 year treatment with exenatide improves beta-cell function, compared with insulin 33 glargine, in metformin-treated type 2 diabetic patients: a randomized, controlled trial. 34 Diabetes Care. 2009; 32(5):762-768
- 35 44. Buse JB, Bergenstal RM, Glass LC, Heilmann CR, Lewis MS, Kwan AY et al. Use of
 36 twice-daily exenatide in Basal insulin-treated patients with type 2 diabetes: a
 37 randomized, controlled trial. Annals of Internal Medicine. 2011; 154(2):103-112
- 38 45. Buse JB, Henry RR, Han J, Kim DD, Fineman MS, Baron AD. Effects of exenatide (Exendin-4) on glycemic control over 30 weeks in sulfonylurea-treated patients with type 2 diabetes. Diabetes Care. 2004; 27(11):2628-2635
- 41 46. Buse JB, Nauck M, Forst T, Sheu WHH, Shenouda SK, Heilmann CR et al. Exenatide once weekly versus liraglutide once daily in patients with type 2 diabetes (DURATION-6): a randomised, open-label study. Lancet (London, England). 2013; 381(9861):117-124
- 45 47. Buse JB, Rosenstock J, Sesti G, Schmidt WE, Montanya E, Brett JH et al. Liraglutide once a day versus exenatide twice a day for type 2 diabetes: a 26-week randomised,

- parallel-group, multinational, open-label trial (LEAD-6). Lancet (London, England). 2009; 374(9683):39-47
- 48. Buse JB, Vilsboll T, Thurman J, Blevins TC, Langbakke IH, Bottcher SG et al.
 Contribution of liraglutide in the fixed-ratio combination of insulin degludec and liraglutide (IDegLira). Diabetes Care. 2014; 37(11):2926-2933
- 6 49. Camerini-Davalos RA, Velasco CA, Reddi AS. Effect of insulin-glipizide combination 7 on skeletal muscle capillary basement membrane width in diabetic patients. Clinical 8 Therapeutics. 1994; 16(6):952-961
- 50. Cannon Christopher P, Pratley R, Dagogo-Jack S, Mancuso J, Huyck S,
 Masiukiewicz U et al. Cardiovascular Outcomes with Ertugliflozin in Type 2 Diabetes.
 The New England journal of medicine. 2020; 383(15):1425-1435
- 12 51. Capehorn MS, Catarig AM, Furberg JK, Janez A, Price HC, Tadayon S et al. Efficacy 13 and safety of once-weekly semaglutide 1.0mg vs once-daily liraglutide 1.2mg as add-14 on to 1-3 oral antidiabetic drugs in subjects with type 2 diabetes (SUSTAIN 10). 15 Diabetes and Metabolism. 2020; 46(2):100-109
- 52. Cefalu WT, Leiter LA, de Bruin TW, Gause-Nilsson I, Sugg J, Parikh SJ.
 Dapagliflozin's effects on glycemia and cardiovascular risk factors in high-risk patients with type 2 diabetes: A 24-week, multicenter, randomized, double-blind, placebo-controlled study with a 28-week extension. Diabetes Care. 2015; 38(7):1218-1227
- 53. Cefalu WT, Leiter LA, Yoon KH, Arias P, Niskanen L, Xie J et al. Efficacy and safety
 of canagliflozin versus glimepiride in patients with type 2 diabetes inadequately
 controlled with metformin (CANTATA-SU): 52 week results from a randomised,
 double-blind, phase 3 non-inferiority trial. Lancet. 2013; 382(9896):941-950
- Charbonnel B, Karasik A, Liu J, Wu M, Meininger G. Efficacy and safety of the
 dipeptidyl peptidase-4 inhibitor sitagliptin added to ongoing metformin therapy in
 patients with type 2 diabetes inadequately controlled with metformin alone. Diabetes
 Care. 2006; 29(12):2638-2643
- 28 55. Charbonnel B, Steinberg H, Eymard E, Xu L, Thakkar P, Prabhu V et al. Efficacy and safety over 26 weeks of an oral treatment strategy including sitagliptin compared with an injectable treatment strategy with liraglutide in patients with type 2 diabetes mellitus inadequately controlled on metformin: a randomised clinical trial.

 32 Diabetologia. 2013; 56(7):1503-1511
- 56. Charpentier G, Halimi S. Earlier triple therapy with pioglitazone in patients with type 2 diabetes. Diabetes Obes Metab. 2009; 11(9):844-854
- 57. Chen WJY, Diamant M, de Boer K, Harms HJ, Robbers LFHJ, van Rossum AC et al.
 Effects of exenatide on cardiac function, perfusion, and energetics in type 2 diabetic patients with cardiomyopathy: a randomized controlled trial against insulin glargine.
 Cardiovascular Diabetology. 2017; 16(1):67
- 58. Chen X, Wang J, Huang X, Tan Y, Deng S, Fu Y. Effects of vildagliptin versus
 saxagliptin on daily acute glucose fluctuations in Chinese patients with T2DM
 inadequately controlled with a combination of metformin and sulfonylurea. Current
 Medical Research and Opinion. 2016; 32(6):1131-1136
- Chen Y, Liu X, Li Q, Ma J, Lv X, Guo L et al. Saxagliptin add-on therapy in Chinese patients with type 2 diabetes inadequately controlled by insulin with or without metformin: results from the SUPER study, a randomized, double-blind, placebo-controlled trial. Diabetes Obes Metab. 2018; 20(4):1044-1049

- 1 60. Cho KY, Nakamura A, Omori K, Takase T, Miya A, Manda N et al. Effect of switching from pioglitazone to the sodium glucose co-transporter-2 inhibitor dapagliflozin on body weight and metabolism-related factors in patients with type 2 diabetes mellitus:
 4 An open-label, prospective, randomized, parallel-group comparison trial. Diabetes,
 5 Obesity & Metabolism. 2019; 21(3):710-714
- 61. Civera M, Merchante A, Salvador M, Sanz J, Mart?nez I. Safety and efficacy of repaglinide in combination with metformin and bedtime NPH insulin as an insulin treatment regimen in type 2 diabetes. Diabetes Research and Clinical Practice. 2008; 79(1):42-47
- 10 62. Cusi K, Bril F, Barb D, Polidori D, Sha S, Ghosh A et al. Effect of canagliflozin 11 treatment on hepatic triglyceride content and glucose metabolism in patients with type 12 2 diabetes. Diabetes Obes Metab. 2019; 21(4):812-821
- 13 63. D'Alessio D, Haring HU, Charbonnel B, de Pablos-Velasco P, Candelas C, Dain MP 14 et al. Comparison of insulin glargine and liraglutide added to oral agents in patients 15 with poorly controlled type 2 diabetes. Diabetes Obes Metab. 2015; 17(2):170-178
- da Silva GM, Nogueira KC, Fukui RT, Correia MRS, dos Santos RF, da Silva ME.
 Short and long term effects of a DPP-4 inhibitor versus bedtime NPH insulin as ADD-ON therapy in patients with type 2 diabetes. Curr Pharm Design. 2016; 22(44):6716-6721
- 20 65. Dagogo-Jack S, Liu J, Eldor R, Amorin G, Johnson J, Hille D et al. Efficacy and safety of the addition of ertugliflozin in patients with type 2 diabetes mellitus inadequately controlled with metformin and sitagliptin: the VERTIS SITA2 placebo-controlled randomized study. Diabetes Obes Metab. 2018; 20(3):530-540
- 24 66. Dahl D, Onishi Y, Norwood P, Huh R, Bray R, Patel H et al. Effect of Subcutaneous Tirzepatide vs Placebo Added to Titrated Insulin Glargine on Glycemic Control in Patients With Type 2 Diabetes: The SURPASS-5 Randomized Clinical Trial. JAMA. 2022; 327(6):534-545
- Davies M, F?rch L, Jeppesen OK, Pakseresht A, Pedersen SD, Perreault L et al. Semaglutide 2?4 mg once a week in adults with overweight or obesity, and type 2 diabetes (STEP 2): a randomised, double-blind, double-dummy, placebo-controlled, phase 3 trial. Lancet (London, England). 2021; 397(10278):971-984
- Davies M, Heller S, Sreenan S, Sapin H, Adetunji O, Tahbaz A et al. Once-weekly exenatide versus once- or twice-daily insulin detemir: randomized, open-label, clinical trial of efficacy and safety in patients with type 2 diabetes treated with metformin alone or in combination with sulfonylureas. Diabetes Care. 2013; 36(5):1368-1376
- 36 69. Davies M, Pieber TR, Hartoft-Nielsen ML, Hansen OKH, Jabbour S, Rosenstock J.
 37 Effect of Oral Semaglutide Compared With Placebo and Subcutaneous Semaglutide
 38 on Glycemic Control in Patients With Type 2 Diabetes: a Randomized Clinical Trial.
 39 JAMA. 2017; 318(15):1460-1470
- 40 70. Davies MJ, Bain SC, Atkin SL, Rossing P, Scott D, Shamkhalova MS et al. Efficacy 41 and safety of liraglutide versus placebo as add-on to glucose-lowering therapy in 42 patients with type 2 diabetes and moderate renal impairment (LIRA-RENAL): A 43 randomized clinical trial. Diabetes Care. 2016; 39(2):222-230
- 71. Davies MJ, Bergenstal R, Bode B, Kushner RF, Lewin A, Skjoth TV et al. Efficacy of liraglutide for weight loss among patients with type 2 diabetes: The SCALE diabetes randomized clinical trial. JAMA. 2015; 314(7):687-699

- Davies MJ, Donnelly R, Barnett AH, Jones S, Nicolay C, Kilcoyne A. Exenatide compared with long-acting insulin to achieve glycaemic control with minimal weight gain in patients with type 2 diabetes: results of the Helping Evaluate Exenatide in patients with diabetes compared with Long-Acting insulin (HEELA) study. Diabetes Obes Metab. 2009; 11(12):1153-1162
- 73. DeFronzo RA, Burant CF, Fleck P, Wilson C, Mekki Q, Pratley RE. Efficacy and tolerability of the DPP-4 inhibitor alogliptin combined with pioglitazone, in metformin-treated patients with type 2 diabetes. Journal of Clinical Endocrinology and Metabolism. 2012; 97(5):1615-1622
- 10 74. DeFronzo RA, Hissa MN, Garber AJ, Luiz Gross J, Yuyan Duan R, Ravichandran S 11 et al. The efficacy and safety of saxagliptin when added to metformin therapy in 12 patients with inadequately controlled type 2 diabetes with metformin alone. Diabetes 13 Care. 2009; 32(9):1649-1655
- 75. DeFronzo RA, Lewin A, Patel S, Liu D, Kaste R, Woerle HJ et al. Combination of
 empagliflozin and linagliptin as second-line therapy in subjects with type 2 diabetes
 inadequately controlled on metformin. Diabetes Care. 2015; 38(3):384-393
- 76. DeFronzo RA, Ratner RE, Han J, Kim DD, Fineman MS, Baron AD. Effects of exenatide (exendin-4) on glycemic control and weight over 30 weeks in metformin-treated patients with type 2. Diabetes Care. 2005; 28(5):1092-1100
- 77. Del Prato S, Camisasca R, Wilson C, Fleck P. Durability of the efficacy and safety of
 alogliptin compared with glipizide in type 2 diabetes mellitus: a 2-year study. Diabetes
 Obes Metab. 2014; 16(12):1239-1246
- 78. Del Prato S, Kahn SE, Pavo I, Weerakkody GJ, Yang Z, Doupis J et al. Tirzepatide versus insulin glargine in type 2 diabetes and increased cardiovascular risk (SURPASS-4): a randomised, open-label, parallel-group, multicentre, phase 3 trial. Lancet. 2021; 398(10313):1811-1824
- 27 79. DePaoli AM, Higgins LS, Henry RR, Mantzoros C, Dunn FL. Can a selective 28 PPARgamma modulator improve glycemic control in patients with type 2 diabetes 29 with fewer side effects compared with pioglitazone? Diabetes Care. 2014; 30 37(7):1918-1923
- 31 80. Derosa G, Bonaventura A, Bianchi L, Romano D, Fogari E, D'Angelo A et al.
 32 Comparison of vildagliptin and glimepiride: effects on glycaemic control, fat tolerance
 33 and inflammatory markers in people with type 2 diabetes. Diabetic medicine: a
 34 journal of the British Diabetic Association. 2014; 31(12):1515-1523
- 35 81. Derosa G, Bonaventura A, Bianchi L, Romano D, Fogari E, D'Angelo A et al.
 36 Vildagliptin compared to glimepiride on post-prandial lipemia and on insulin
 37 resistance in type 2 diabetic patients. Metabolism: Clinical and Experimental. 2014;
 38 63(7):957-967
- 39 82. Derosa G, Carbone A, D'Angelo A, Querci F, Fogari E, Cicero AF et al. A
 40 randomized, double-blind, placebo-controlled trial evaluating sitagliptin action on
 41 insulin resistance parameters and beta-cell function. Expert Opinion on
 42 Pharmacotherapy. 2012; 13(17):2433-2442
- 43 83. Derosa G, Franzetti IG, Querci F, Carbone A, Ciccarelli L, Piccinni MN et al.
 44 Exenatide plus metformin compared with metformin alone on beta-cell function in
 45 patients with Type 2 diabetes. Diabetic medicine : a journal of the British Diabetic
 46 Association. 2012; 29(12):1515-1523

- Derosa G, Maffioli P, Ferrari I, Mereu R, Ragonesi PD, Querci F et al. Effects of one year treatment of vildagliptin added to pioglitazone or glimepiride in poorly controlled type 2 diabetic patients. Hormone and metabolic research = Hormon- und Stoffwechselforschung = Hormones et metabolisme. 2010; 42(9):663-669
- 5 85. Derosa G, Maffioli P, Salvadeo SA, Ferrari I, Ragonesi PD, Querci F et al. Effects of sitagliptin or metformin added to pioglitazone monotherapy in poorly controlled type 2 diabetes mellitus patients. Metabolism: Clinical and Experimental. 2010; 59(6):887-895
- 9 86. Derosa G, Putignano P, Bossi AC, Bonaventura A, Querci F, Franzetti IG et al.
 10 Exenatide or glimepiride added to metformin on metabolic control and on insulin
 11 resistance in type 2 diabetic patients. European Journal of Pharmacology. 2011;
 12 666(13):251-256
- Berosa G, Ragonesi PD, Carbone A, Fogari E, Bianchi L, Bonaventura A et al.
 Vildagliptin added to metformin on beta-cell function after a euglycemic hyperinsulinemic and hyperglycemic clamp in type 2 diabetes patients. Diabetes Technology & Therapeutics. 2012; 14(6):475-484
- 17 88. Derosa G, Ragonesi PD, Carbone A, Fogari E, D'Angelo A, Cicero AFG et al.
 18 Vildagliptin action on some adipocytokine levels in type 2 diabetic patients: a 1219 month, placebo-controlled study. Expert Opinion on Pharmacotherapy. 2012;
 20 13(18):2581-2591
- 21 89. Derosa G, Ragonesi PD, Fogari E, Cicero AFG, Bianchi L, Bonaventura A et al.
 22 Sitagliptin added to previously taken antidiabetic agents on insulin resistance and
 23 lipid profile: A 2-year study evaluation. Fundamental and Clinical Pharmacology.
 24 2014; 28(2):221-229
- 90. Diamant M, Gaal L, Stranks S, Northrup J, Cao D, Taylor K et al. Once weekly
 exenatide compared with insulin glargine titrated to target in patients with type 2
 diabetes (DURATION-3): an open-label randomised trial. Lancet. 2010;
 375(9733):2234-2243
- 29 91. Diamant M, Nauck MA, Shaginian R, Malone JK, Cleall S, Reaney M et al. Glucagon-30 like peptide 1 receptor agonist or bolus insulin with optimized basal insulin in type 2 31 diabetes. Diabetes Care. 2014; 37(10):2763-2773
- 92. Dobs AS, Goldstein BJ, Aschner P, Horton ES, Umpierrez GE, Duran L et al. Efficacy
 and safety of sitagliptin added to ongoing metformin and rosiglitazone combination
 therapy in a randomized placebo-controlled 54-week trial in patients with type 2
 diabetes. J Diabetes. 2013; 5(1):68-79
- 93. Dorkhan M, Dencker M, Stagmo M, Groop L. Effect of pioglitazone versus insulin
 glargine on cardiac size, function, and measures of fluid retention in patients with type
 2 diabetes. Cardiovascular Diabetology. 2009; 8:15
- Jouek IF, Allen SE, Ewings P, Gale EA, Bingley PJ. Continuing metformin when
 starting insulin in patients with Type 2 diabetes: a double-blind randomized placebo controlled trial. Diabetic Med. 2005; 22(5):634-640
- 95. Dungan KM, Povedano ST, Forst T, Gonzalez JGG, Atisso C, Sealls W et al. Onceweekly dulaglutide versus once-daily liraglutide in metformin-treated patients with type 2 diabetes (AWARD-6): a randomised, open-label, phase 3, non-inferiority trial. Lancet (London, England). 2014; 384(9951):1349-1357
- 96. Dungan KM, Weitgasser R, Perez Manghi F, Pintilei E, Fahrbach JL, Jiang HH et al.
 A 24-week study to evaluate the efficacy and safety of once-weekly dulaglutide added

- on to glimepiride in type 2 diabetes (AWARD-8). Diabetes Obes Metab. 2016; 18(5):475-482
- Ferdinand KC, Izzo JL, Lee J, Meng L, George J, Salsali A et al. Antihyperglycemic
 and Blood Pressure Effects of Empagliflozin in Black Patients With Type 2 Diabetes
 Mellitus and Hypertension. Circulation. 2019; 139(18):2098-2109
- Fernandez M, Triplitt C, Wajcberg E, Sriwijilkamol AA, Musi N, Cusi K et al. Addition of pioglitazone and ramipril to intensive insulin therapy in type 2 diabetic patients improves vascular dysfunction by different mechanisms. Diabetes Care. 2008; 31(1):121-127
- 99. Ferrannini E, Fonseca V, Zinman B, Matthews D, Ahren B, Byiers S et al. Fifty-two-week efficacy and safety of vildagliptin vs. glimepiride in patients with type 2 diabetes mellitus inadequately controlled on metformin monotherapy. Diabetes, Obesity & Metabolism. 2009; 11(2):157-166
- 14 100. Filozof C, Gautier JF. A comparison of efficacy and safety of vildagliptin and gliclazide 15 in combination with metformin in patients with Type 2 diabetes inadequately 16 controlled with metformin alone: a 52-week, randomized study. Diabetic Med. 2010; 17 27(3):318-326
- 18 101. Filozof C, Schwartz S, Foley JE. Effect of vildagliptin as add-on therapy to a low-dose metformin. World journal of diabetes. 2010; 1(1):19-26
- 20 102. Fioretto P, Del Prato S, Buse JB, Goldenberg R, Giorgino F, Reyner D et al. Efficacy 21 and safety of dapagliflozin in patients with type 2 diabetes and moderate renal 22 impairment (chronic kidney disease stage 3A): the DERIVE Study. Diab Obes Metab. 23 2018; 20(11):2532-2540
- 103. Fonseca V, Schweizer A, Albrecht D, Baron MA, Chang I, Dejager S. Addition of vildagliptin to insulin improves glycaemic control in type 2 diabetes. Diabetologia.
 26 2007; 50(6):1148-1155
- 27 104. Fonseca V, Staels B, Morgan JD, Shentu Y, Golm GT, Johnson-Levonas AO et al.
 28 Efficacy and safety of sitagliptin added to ongoing metformin and pioglitazone
 29 combination therapy in a randomized, placebo-controlled, 26-week trial in patients
 30 with type 2 diabetes. Journal of Diabetes and Its Complications. 2013; 27(2):177-183
- 105. Forst T, Guthrie R, Goldenberg R, Yee J, Vijapurkar U, Meininger G et al. Efficacy and safety of canagliflozin over 52 weeks in patients with type 2 diabetes on background metformin and pioglitazone. Diabetes Obes Metab. 2014; 16(5):467-477
- Torst T, Hohberg C, Fuellert SD, L?bben G, Konrad T, L?big M et al.
 Pharmacological PPARgamma stimulation in contrast to beta cell stimulation results in an improvement in adiponectin and proinsulin intact levels and reduces intima media thickness in patients with type 2 diabetes. Hormone and Metabolic Research.
 37 2005; 37(8):521-527
- 107. Forst T, Koch C, Dworak M. Vildagliptin versus insulin in patients with type 2 diabetes mellitus inadequately controlled with sulfonylurea: Results from a randomized, 24 week study. Current Medical Research and Opinion. 2015; 31(6):1079-1084
- 42 108. Frias JP, Davies MJ, Rosenstock J, Perez Manghi FC, Fernandez Lando L, Bergman
 43 BK et al. Tirzepatide versus Semaglutide Once Weekly in Patients with Type 2
 44 Diabetes. New England Journal of Medicine. 2021; 385(6):503-515
- 45 109. Frias JP, Deenadayalan S, Erichsen L, Knop FK, Lingvay I, Macura S et al. Efficacy and safety of co-administered once-weekly cagrilintide 2.4 mg with once-weekly

- semaglutide 2.4 mg in type 2 diabetes: a multicentre, randomised, double-blind, active-controlled, phase 2 trial. Lancet (London, England). 2023; 402(10403):720-730
- 110. Frias JP, Gonzalez-Galvez G, Johnsson E, Maaske J, Testa MA, Simonson DC et al. Efficacy and safety of dual add-on therapy with dapagliflozin plus saxagliptin versus glimepiride in patients with poorly controlled type 2 diabetes on a stable dose of metformin: Results from a 52-week, randomized, active-controlled trial. Diabetes, Obesity & Metabolism. 2020; 22(7):1083-1093
- 8 111. Frias JP, Guja C, Hardy E, Ahmed A, Dong F, Ohman P et al. Exenatide once weekly plus dapagliflozin once daily versus exenatide or dapagliflozin alone in patients with type 2 diabetes inadequately controlled with metformin monotherapy (DURATION-8): a 28 week, multicentre, double-blind, phase 3, randomised controlled trial. The lancet Diabetes & endocrinology. 2016; 4(12):1004-1016
- 13 112. Frias JP, Nauck MA, Van J, Kutner ME, Cui X, Benson C et al. Efficacy and safety of LY3298176, a novel dual GIP and GLP-1 receptor agonist, in patients with type 2 diabetes: a randomised, placebo-controlled and active comparator-controlled phase 2 trial. Lancet. 2018; 392(10160):2180-2193
- 17 113. Fu Q, Zhou L, Fan Y, Liu F, Fan Y, Zhang X et al. Effect of SGLT-2 inhibitor, dapagliflozin, on left ventricular remodeling in patients with type 2 diabetes and HFrEF. BMC Cardiovascular Disorders. 2023; 23(1):544
- 114. Fujioka K, Pans M, Joyal S. Glycemic control in patients with type 2 diabetes mellitus
 switched from twice-daily immediate-release metformin to a once-daily extended-release formulation. Clinical Therapeutics. 2003; 25(2):515-529
- 115. G?ke B, Gallwitz B, Eriksson J, Hellqvist A, Gause-Nilsson I. Saxagliptin is non-inferior to glipizide in patients with type 2 diabetes mellitus inadequately controlled on metformin alone: a 52-week randomised controlled trial. International Journal of Clinical Practice. 2010; 64(12):1619-1631
- 116. Gadde KM, Vetter ML, Iqbal N, Hardy E, Ohman P. Efficacy and safety of
 autoinjected exenatide once-weekly suspension versus sitagliptin or placebo with
 metformin in patients with type 2 diabetes: The DURATION-NEO-2 randomized
 clinical study. Diabetes Obes Metab. 2017; 19(7):979-988
- 31 117. Galindo RJ, Moazzami B, Scioscia MF, Zambrano C, Albury BS, Saling J et al. A
 32 Randomized Controlled Trial Comparing the Efficacy and Safety of IDegLira Versus
 33 Basal-Bolus in Patients With Poorly Controlled Type 2 Diabetes and Very High
 34 HbA1c >=9-15%: DUAL HIGH Trial. Diabetes Care. 2023; 46(9):1640-1645
- Galle J, Kleophas W, Dellanna F, Schmid VHR, Forkel C, Dikta G et al. Comparison of the Effects of Pioglitazone versus Placebo when Given in Addition to Standard Insulin Treatment in Patients with Type 2 Diabetes Mellitus Requiring Hemodialysis: Results from the PIOren Study. Nephron extra. 2012; 2(1):104-114
- 39 119. Gallwitz B, B?hmer M, Segiet T, M?lle A, Milek K, Becker B et al. Exenatide twice
 40 daily versus premixed insulin aspart 70/30 in metformin-treated patients with type 2
 41 diabetes: a randomized 26-week study on glycemic control and hypoglycemia.
 42 Diabetes Care. 2011; 34(3):604-606
- 43 120. Gallwitz B, Guzman J, Dotta F, Guerci B, Sim R, Basson BR et al. Exenatide twice
 44 daily versus glimepiride for prevention of glycaemic deterioration in patients with type
 45 2 diabetes with metformin failure (EUREXA): an open-label, randomised controlled
 46 trial. Lancet. 2012; 379(9833):2270-2278

- 1 121. Gallwitz B, Rosenstock J, Rauch T, Bhattacharya S, Patel S, Eynatten M et al. 2-year efficacy and safety of linagliptin compared with glimepiride in patients with type 2 diabetes inadequately controlled on metformin: a randomised, double-blind, non-inferiority trial. Lancet. 2012; 380(9840):475-483
- 5 122. Gao L, Lee BW, Chawla M, Kim J, Huo L, Du L et al. Tirzepatide versus insulin glargine as second-line or third-line therapy in type 2 diabetes in the Asia-Pacific region: the SURPASS-AP-Combo trial. Nature Medicine. 2023; 29(6):1500-1510
- 8 123. Garber A, Henry R, Ratner R, Garcia-Hernandez PA, Rodriguez-Pattzi H, Olvera-9 Alvarez I et al. Liraglutide versus glimepiride monotherapy for type 2 diabetes (LEAD-10 3 Mono): a randomised, 52-week, phase III, double-blind, parallel-treatment trial. 11 Lancet. 2009; 373(9662):473-481
- 12 124. Garber AJ, Foley JE, Banerji MA, Ebeling P, Gudbj?rnsdottir S, Camisasca RP et al.
 13 Effects of vildagliptin on glucose control in patients with type 2 diabetes inadequately controlled with a sulphonylurea. Diabetes Obes Metab. 2008; 10(11):1047-1056
- 15 125. Garber AJ, Schweizer A, Baron MA, Rochotte E, Dejager S. Vildagliptin in combination with pioglitazone improves glycaemic control in patients with type 2 diabetes failing thiazolidinedione monotherapy: a randomized, placebo-controlled study. Diabetes Obes Metab. 2007; 9(2):166-174
- 126. Garvey WT, Birkenfeld AL, Dicker D, Mingrone G, Pedersen SD, Satylganova A et al.
 Efficacy and Safety of Liraglutide 3.0 mg in Individuals With Overweight or Obesity
 and Type 2 Diabetes Treated With Basal Insulin: The SCALE Insulin Randomized
 Controlled Trial. Diabetes Care. 2020; 43(5):1085-1093
- 127. Garvey WT, Frias JP, Jastreboff AM, le Roux CW, Sattar N, Aizenberg D et al.
 Tirzepatide once weekly for the treatment of obesity in people with type 2 diabetes
 (SURMOUNT-2): a double-blind, randomised, multicentre, placebo-controlled, phase
 3 trial. Lancet (London, England). 2023; 402(10402):613-626
- 27 128. Genovese S, Passaro A, Brunetti P, Comaschi M, Cucinotta D, Egan CG et al.
 28 Pioglitazone randomised Italian study on metabolic syndrome (PRISMA): Effect of
 29 pioglitazone withmetformin on HDL-C levels in type 2 diabetic patients. Journal of
 30 Endocrinological Investigation. 2013; 36(8):606-616
- 31 129. Gerstein Hertzel C, Colhoun Helen M, Dagenais Gilles R, Diaz R, Lakshmanan M,
 32 Pais P et al. Dulaglutide and cardiovascular outcomes in type 2 diabetes (REWIND):
 33 a double-blind, randomised placebo-controlled trial. Lancet (London, England). 2019;
 34 394(10193):121-130
- 35 130. Giorgino F, Benroubi M, Sun JH, Zimmermann AG, Pechtner V. Efficacy and safety of once-weekly dulaglutide versus insulin glargine in patients with type 2 diabetes on metformin and glimepiride (AWARD-2). Diabetes Care. 2015; 38(12):2241-2249
- 38 131. Giugliano D, Quatraro A, Consoli G, Minei A, Ceriello A, Rosa N et al. Metformin for obese, insulin-treated diabetic patients: improvement in glycaemic control and reduction of metabolic risk factors. European Journal of Clinical Pharmacology. 1993; 44(2):107-112
- 42 132. Gohari S, Reshadmanesh T, Khodabandehloo H, Karbalaee-Hasani A, Ahangar H,
 43 Arsang-Jang S et al. The effect of EMPAgliflozin on markers of inflammation in
 44 patients with concomitant type 2 diabetes mellitus and Coronary ARtery Disease: the
 45 EMPA-CARD randomized controlled trial. Diabetology & Metabolic Syndrome. 2022;
 46 14(1):170

- 1 133. Goodman M, Thurston H, Penman J. Efficacy and tolerability of vildagliptin in patients with type 2 diabetes inadequately controlled with metformin monotherapy. Hormone and Metabolic Research. 2009; 41(5):368-373
- 4 134. Gough SC, Bode B, Woo V, Rodbard HW, Linjawi S, Poulsen P et al. Efficacy and safety of a fixed-ratio combination of insulin degludec and liraglutide (IDegLira) compared with its components given alone: results of a phase 3, open-label, randomised, 26-week, treat-to-target trial in insulin-naive patients with type 2 diabetes. Lancet Diabetes Endocrinol. 2014; 2(11):885-893
- 9 135. Gram J, Henriksen JE, Grodum E, Juhl H, Hansen TB, Christiansen C et al.
 10 Pharmacological treatment of the pathogenetic defects in type 2 diabetes: the
 11 randomized multicenter South Danish Diabetes Study. Diabetes Care. 2011;
 12 34(1):27-33
- 136. Green Jennifer B, Bethel M A, Armstrong Paul W, Buse John B, Engel Samuel S,
 Garg J et al. Effect of Sitagliptin on Cardiovascular Outcomes in Type 2 Diabetes.
 The New England journal of medicine. 2015; 373(3):232-242
- 137. Grey A, Bolland M, Fenwick S, Horne A, Gamble G, Drury PL et al. The skeletal effects of pioglitazone in type 2 diabetes or impaired glucose tolerance: A randomized controlled trial. European Journal of Endocrinology of the European Federation of Endocrine Societies. 2014; 170(2):255-262
- 138. Groop PH, Cooper ME, Perkovic V, Hocher B, Kanasaki K, Haneda M et al.
 Linagliptin and its effects on hyperglycaemia and albuminuria in patients with type 2
 diabetes and renal dysfunction: the randomized MARLINA-T2D trial. Diabetes,
 Obesity and Metabolism. 2017; 19(11):1610-1619
- 24 139. Group GSR, Nathan DM, Lachin JM, Bebu I, Burch HB, Buse JB et al. Glycemia
 25 Reduction in Type 2 Diabetes Microvascular and Cardiovascular Outcomes. New
 26 England Journal of Medicine. 2022; 387(12):1075-1088
- 27 140. Grunberger G, Camp S, Johnson J, Huyck S, Terra SG, Mancuso JP et al.
 28 Ertugliflozin in patients with Stage 3 chronic kidney disease and type 2 diabetes
 29 mellitus: the VERTIS RENAL randomized study. Diabetes Therapy. 2018; 9(1):49-66
- 30 141. Gu T, Ma J, Zhang Q, Zhu L, Zhang H, Xu L et al. Comparative effect of saxagliptin 31 and glimepiride with a composite endpoint of adequate glycaemic control without 32 hypoglycaemia and without weight gain in patients uncontrolled with metformin 33 therapy: results from the SPECIFY study, a 48-week, multi-centre, randomized, 34 controlled trial. Diab Obes Metab. 2019; 21(4):939-948
- Guja C, Fr?as JP, Somogyi A, Jabbour S, Wang H, Hardy E et al. Effect of exenatide
 QW or placebo, both added to titrated insulin glargine, in uncontrolled type 2
 diabetes: the DURATION-7 randomized study. Diabetes, Obesity & Metabolism.
 2018; 20(7):1602-1614
- Gullaksen S, Vernstrom L, Sorensen SS, Ringgaard S, Laustsen C, Funck KL et al.
 Separate and combined effects of semaglutide and empagliflozin on kidney oxygenation and perfusion in people with type 2 diabetes: a randomised trial.
 Diabetologia. 2023; 66(5):813-825
- 43 144. Guo W, Tian W, Lin L, Xu X. Liraglutide or insulin glargine treatments improves 44 hepatic fat in obese patients with type 2 diabetes and nonalcoholic fatty liver disease 45 in twenty-six weeks: A randomized placebo-controlled trial. Diabetes Research and 46 Clinical Practice. 2020; 170:108487

- 1 145. Gurkan E, Tarkun I, Sahin T, Cetinarslan B, Canturk Z. Evaluation of exenatide versus insulin glargine for the impact on endothelial functions and cardiovascular risk markers. Diabetes Research and Clinical Practice. 2014; 106(3):567-575
- 4 146. Guzman CB, Zhang XM, Liu R, Regev A, Shankar S, Garhyan P et al. Treatment with LY2409021, a glucagon receptor antagonist, increases liver fat in patients with type 2 diabetes. Diabetes Obes Metab. 2017; 19(11):1521-1528
- 7 147. Handelsman Y, Mathieu C, Del Prato S, Johnsson E, Kurlyandskaya R, Iqbal N et al.
 8 Sustained 52-week efficacy and safety of triple therapy with dapagliflozin plus
 9 saxagliptin versus dual therapy with sitagliptin added to metformin in patients with
 10 uncontrolled type 2 diabetes. Diabetes, Obesity and Metabolism. 2019; 21(4):883-892
- 148. Hanefeld M, Brunetti P, Schernthaner GH, Matthews DR, Charbonnel BH. One-year
 glycemic control with a sulfonyurea plus pioglitazone versus a sulfonylurea plus
 metformin in patients with type 2 diabetes. Diabetes Care. 2004; 27(1):141-147
- 14 149. Hanefeld M, Pf?tzner A, Forst T, Kleine I, Fuchs W. Double-blind, randomized,
 multicentre, and active comparator controlled investigation of the effect of
 pioglitazone, metformin, and the combination of both on cardiovascular risk in
 patients with type 2 diabetes receiving stable basal insulin therapy: the PIOCOMB
 study. Cardiovascular Diabetology. 2011; 10:65
- 150. Hao Z, Huang X, Shao H, He F. Efficacy and Safety of Dapagliflozin versus
 Liraglutide in Patients with Overweight or Obesity and Type 2 Diabetes Mellitus: a
 Randomised Controlled Clinical Trial in Tianjin, China. Journal of Diabetes Research.
 2022; 2022:4126995
- 151. Haring HU, Merker L, Seewaldt-Becker E, Weimer M, Meinicke T, Broedl UC et al.
 Empaglif lozin as add-on to metformin in patients with type 2 diabetes: A 24-week,
 randomized, double-blind, placebo-controlled trial. Diabetes Care. 2014; 37(6):1650 1659
- 152. Haring HU, Merker L, Seewaldt-Becker E, Weimer M, Meinicke T, Woerle HJ et al.
 Empagliflozin as add-on to metformin plus sulfonylurea in patients with type 2
 diabetes: A 24-week, randomized, double-blind, placebo-controlled trial. Diabetes
 Care. 2013; 36(11):3396-3404
- Harreiter J, Just I, Leutner M, Bastian M, Brath H, Schelkshorn C et al. Combined
 exenatide and dapagliflozin has no additive effects on reduction of hepatocellular
 lipids despite better glycaemic control in patients with type 2 diabetes mellitus treated
 with metformin: EXENDA, a 24-week, prospective, randomized, placebo-controlled
 pilot trial. Diabetes, Obesity & Metabolism. 2021; 23(5):1129-1139
- Hartemann-Heurtier A, Halbron M, Golmard JL, Jacqueminet S, Bastard JP, Rouault
 C et al. Effects of bed-time insulin versus pioglitazone on abdominal fat accumulation,
 inflammation and gene expression in adipose tissue in patients with type 2 diabetes.
 Diabetes Research and Clinical Practice. 2009; 86(1):37-43
- 40 155. Hattori S. Anti-inflammatory effects of empagliflozin in patients with type 2 diabetes and insulin resistance. Diabetology & Metabolic Syndrome. 2018; 10(1):93
- Heine RJ, Gaal LF, Johns D, Mihm MJ, Widel MH, Brodows RG. Exenatide versus insulin glargine in patients with suboptimally controlled type 2 diabetes: a randomized trial. Annals of Internal Medicine. 2005; 143(8):559-569
- Heise T, Mari A, DeVries JH, Urva S, Li J, Pratt EJ et al. Effects of subcutaneous tirzepatide versus placebo or semaglutide on pancreatic islet function and insulin sensitivity in adults with type 2 diabetes: a multicentre, randomised, double-blind,

- parallel-arm, phase 1 clinical trial. The lancet Diabetes & endocrinology. 2022; 10(6):418-429
- Henriksen K, Byrjalsen I, Qvist P, Beck-Nielsen H, Hansen G, Riis BJ et al. Efficacy and safety of the PPARgamma partial agonist balaglitazone compared with pioglitazone and placebo: a phase III, randomized, parallel-group study in patients with type 2 diabetes on stable insulin therapy. Diabetes/Metabolism Research and Reviews. 2011; 27(4):392-401
- 8 159. Heo JH, Han KA, Hong JH, Seo H-A, Hong E-G, Yu JM et al. Pioglitazone as Add-on THERAPY in Patients with Type 2 Diabetes Mellitus Inadequately Controlled with Dapagliflozin and Metformin: Double-Blind, Randomized, Placebo-Controlled Trial. Diabetes & Metabolism Journal. 2024;
- 12 160. Hermansen K, Kipnes M, Luo E, Fanurik D, Khatami H, Stein P. Efficacy and safety
 13 of the dipeptidyl peptidase-4 inhibitor, sitagliptin, in patients with type 2 diabetes
 14 mellitus inadequately controlled on glimepiride alone or on glimepiride and metformin.
 15 Diabetes Obes Metab. 2007; 9(5):733-745
- 161. Hiramatsu T, Asano Y, Mabuchi M, Imai K, Iguchi D, Furuta S. Liraglutide relieves
 17 cardiac dilated function than DPP-4 inhibitors. European Journal of Clinical
 18 Investigation. 2018; 48(10):e13007
- 19 162. Hollander P, Li J, Allen E, Chen R. Saxagliptin added to a thiazolidinedione improves 20 glycemic control in patients with type 2 diabetes and inadequate control on 21 thiazolidinedione alone. Journal of Clinical Endocrinology and Metabolism. 2009; 22 94(12):4810-4819
- Hollander P, Liu J, Hill J, Johnson J, Jiang ZW, Golm G et al. Ertugliflozin Compared with Glimepiride in Patients with Type 2 Diabetes Mellitus Inadequately Controlled on Metformin: the VERTIS SU Randomized Study. Diabetes Therapy. 2018; 9(1):193-207
- Holman Rury R, Bethel M A, Mentz Robert J, Thompson Vivian P, Lokhnygina Y,
 Buse John B et al. Effects of Once-Weekly Exenatide on Cardiovascular Outcomes in
 Type 2 Diabetes. The New England journal of medicine. 2017; 377(13):1228-1239
- Home PD, Shamanna P, Stewart M, Yang F, Miller M, Perry C et al. Efficacy and tolerability of albiglutide versus placebo or pioglitazone over 1year in people with type 2 diabetes currently taking metformin and glimepiride: HARMONY 5. Diabetes Obes Metab. 2015; 17(2):179-187
- Hong ES, Khang AR, Yoon JW, Kang SM, Choi SH, Park KS et al. Comparison
 between sitagliptin as add-on therapy to insulin and insulin dose-increase therapy in
 uncontrolled Korean type 2 diabetes: CSI study. Diabetes, Obesity & Metabolism.
 2012; 14(9):795-802
- Hong JH, Moon JS, Seong K, Lim S. Comparison of therapeutic efficacy and safety of sitagliptin, dapagliflozin, or lobeglitazone adjunct therapy in patients with type 2 diabetes mellitus inadequately controlled on sulfonylurea and metformin: Third agent study. Diabetes Research and Clinical Practice. 2023; 203:110872
- Hooshmand Gharabagh L, Shargh A, Mohammad Hosseini Azar MR, Esmaeili A.
 Comparison between the effect of Empagliflozin and Pioglitazone added to metformin in patients with type 2 diabetes and nonalcoholic fatty liver disease. Clinics and research in hepatology and gastroenterology. 2024; 48(3):102279

- Husain M, Birkenfeld Andreas L, Donsmark M, Dungan K, Eliaschewitz Freddy G,
 Franco Denise R et al. Oral Semaglutide and Cardiovascular Outcomes in Patients
 with Type 2 Diabetes. The New England journal of medicine. 2019; 381(9):841-851
- 4 170. Iacobellis G, Gra-Menendez S. Effects of dapagliflozin on epicardial fat thickness in patients with type 2 diabetes and obesity. Obesity. 2020; 28(6):1068-1074
- 6 171. Iacobellis G, Mohseni M, Bianco SD, Banga PK. Liraglutide causes large and rapid epicardial fat reduction. Obesity. 2017; 25(2):311-316
- 8 172. Iijima T, Shibuya M, Ito Y, Terauchi Y. Effects of switching from liraglutide to 9 semaglutide or dulaglutide in patients with type 2 diabetes: A randomized controlled 10 trial. Journal of Diabetes Investigation. 2023; 14(6):774-781
- 173. Ikonomidis I, Pavlidis G, Thymis J, Birba D, Kalogeris A, Kousathana F et al. Effects
 of Glucagon-Like Peptide-1 Receptor Agonists, Sodium-Glucose Cotransporter-2
 Inhibitors, and Their Combination on Endothelial Glycocalyx, Arterial Function, and
 Myocardial Work Index in Patients With Type 2 Diabetes Mellitus After 12-Month
 Treatment. Journal of the American Heart Association. 2020; 9(9):e015716
- 174. Inagaki N, Atsumi Y, Oura T, Saito H, Imaoka T. Efficacy and safety profile of exenatide once weekly compared with insulin once daily in Japanese patients with type 2 diabetes treated with oral antidiabetes drug(s): results from a 26-week, randomized, open-label, parallel-group, multicenter, noninferiority study. Clinical Therapeutics. 2012; 34(9):1892-1908
- 175. Inagaki N, Watada H, Murai M, Kagimura T, Gong Y, Patel S et al. Linagliptin
 provides effective, well-tolerated add-on therapy to pre-existing oral antidiabetic
 therapy over 1year in Japanese patients with type 2 diabetes. Diabetes Obes Metab.
 2013; 15(9):833-843
- Jabbour SA, Hardy E, Sugg J, Parikh S. Dapagliflozin is effective as add-on therapy
 to sitagliptin with or withoutmetformin: A 24-Week, multicenter, randomized, double-blind, placebo-controlled study. Diabetes Care. 2014; 37(3):740-750
- 28 177. Ji L, Dong X, Li Y, Lim S, Liu M et al. Efficacy and safety of once-weekly 29 semaglutide versus once-daily sitagliptin as add-on to metformin in patients with type 30 2 diabetes in SUSTAIN China: A 30-week, double-blind, phase 3a, randomized trial. 31 Diabetes, Obesity & Metabolism. 2021; 23(2):404-414
- 32 178. Ji L, Liu Y, Miao H, Xie Y, Yang M, Wang W et al. Safety and efficacy of ertugliflozin 33 in Asian patients with type 2 diabetes mellitus inadequately controlled with metformin 34 monotherapy: VERTIS Asia. Diabetes Obes Metab. 2019; 21(6):1474-1482
- Ji L, Lu Y, Li Q, Fu L, Luo Y, Lei T et al. Efficacy and safety of empagliflozin in
 combination with insulin in Chinese patients with type 2 diabetes and insufficient
 glycaemic control: A Phase III, randomised, double-blind, placebo-controlled, parallel
 study. Diabetes, Obesity & Metabolism. 2023;
- 180. Ji LN, Pan CY, Lu JM, Li H, Zhu DL, Li Q et al. Efficacy and safety of combination therapy with vildagliptin and metformin versus metformin uptitration in Chinese patients with type 2 diabetes inadequately controlled with metformin monotherapy: a randomized, open-label, prospective study (VISION). Diabetes Obes Metab. 2016; 18(8):775-782
- Jiang J, Lin L, Chen P. Comparison of Dapaglifozin and Liraglutide in Patients with
 Poorly Controlled Type 2 Diabetes Mellitus: a 24-week, Open, Double-centered,
 Head to Head Trial. Endocrine, Metabolic & Immune Disorders Drug Targets. 2021;
 21(7):1366-1374

- 1 182. Joubert M, Opigez V, Pavlikova B, Peyro Saint Paul L, Jeandidier N, Briant AR et al. Efficacy and safety of exenatide as add-on therapy for patients with type 2 diabetes with an intensive insulin regimen: A randomized double-blind trial. Diabetes, Obesity & Metabolism. 2021; 23(2):374-381
- 5 183. Kadowaki T, Inagaki N, Kondo K, Nishimura K, Kaneko G, Maruyama N et al. Efficacy 6 and safety of canagliflozin as add-on therapy to teneligliptin in Japanese patients with 7 type 2 diabetes mellitus: Results of a 24-week, randomized, double-blind, placebo-8 controlled trial. Diabetes Obes Metab. 2017; 19(6):874-882
- 9 184. Kadowaki T, Namba M, Imaoka T, Yamamura A, Goto W, Boardman MK et al.
 10 Improved glycemic control and reduced bodyweight with exenatide: A double-blind,
 11 randomized, phase 3 study in Japanese patients with suboptimally controlled type 2
 12 diabetes over 24 weeks. J Diabetes Invest. 2011; 2(3):210-217
- 13 185. Kaku K. Efficacy and safety of therapy with metformin plus pioglitazone in the 14 treatment of patients with type 2 diabetes: a double-blind, placebo-controlled, clinical 15 trial. Current Medical Research and Opinion. 2009; 25(5):1111-1119
- 186. Kaku K, Araki E, Tanizawa Y, Ross Agner B, Nishida T, Ranthe M et al. Superior efficacy with a fixed-ratio combination of insulin degludec and liraglutide (IDegLira) compared with insulin degludec and liraglutide in insulin-naive Japanese patients with type 2 diabetes in a phase 3, open-label, randomized trial. Diab Obes Metab. 2019; 21(12):2674-2683
- 21 187. Kaku K, Rasmussen MF, Clauson P, Seino Y. Improved glycaemic control with 22 minimal hypoglycaemia and no weight change with the once-daily human glucagon-23 like peptide-1 analogue liraglutide as add-on to sulphonylurea in Japanese patients 24 with type 2 diabetes. Diabetes, Obesity & Metabolism. 2010; 12(4):341-347
- 188. Kanazawa I, Yamaguchi T, Yano S, Yamamoto M, Yamauchi M, Kurioka S et al.
 Baseline atherosclerosis parameter could assess the risk of bone loss during pioglitazone treatment in type 2 diabetes mellitus. Osteoporosis International. 2010; 21(12):2013-2018
- 29 189. Kaneto H, Takami A, Spranger R, Amano A, Watanabe D, Niemoeller E. Efficacy and safety of insulin glargine/lixisenatide fixed-ratio combination (iGlarLixi) in Japanese patients with type 2 diabetes mellitus inadequately controlled on basal insulin and oral antidiabetic drugs: The LixiLan JP-L randomized clinical trial. Diabetes, Obesity & Metabolism. 2020; 22suppl4:3-13
- 190. Kang C, Qiao Q, Tong Q, Bai Q, Huang C, Fan R et al. Effects of exenatide on urinary albumin in overweight/obese patients with T2DM: a randomized clinical trial.
 Scientific Reports. 2021; 11(1):20062
- 191. Kawamori R, Haneda M, Suzaki K, Cheng G, Shiki K, Miyamoto Y et al. Empagliflozin as add-on to linagliptin in a fixed-dose combination in Japanese patients with type 2 diabetes: glycaemic efficacy and safety profile in a 52-week, randomized, placebo-controlled trial. Diab Obes Metab. 2018; 20(9):2200-2209
- 41 192. Kellerer M, Kaltoft MS, Lawson J, Nielsen LL, Strojek K, Tabak et al. Effect of once-42 weekly semaglutide versus thrice-daily insulin aspart, both as add-on to metformin 43 and optimized insulin glargine treatment in participants with type 2 diabetes 44 (SUSTAIN 11): a randomized, open-label, multinational, phase 3b trial. Diabetes, 45 Obesity & Metabolism. 2022; 24(9):1788-1799
- 46 193. Kendall DM, Riddle MC, Rosenstock J, Zhuang D, Kim DD, Fineman MS et al. Effects of exenatide (exendin-4) on glycemic control over 30 weeks in patients with type 2

- diabetes treated with metformin and a sulfonylurea. Diabetes Care. 2005; 28(5):1083-1091
- 3 194. Kesavadev J, Pillai PBS, Shankar A, Krishnan G, Jothydev S. Sitagliptin 100 mg vs glimepiride 1-3 mg as an add-on to insulin and metformin in type 2 diabetes (SWIM). Endocrine connections. 2017; 6(8):748-757
- Khaloo P, Asadi Komeleh S, Alemi H, Mansournia MA, Mohammadi A, Yadegar A et al. Sitagliptin vs. pioglitazone as add-on treatments in patients with uncontrolled type 2 diabetes on the maximal dose of metformin plus sulfonylurea. Journal of Endocrinological Investigation. 2019; 42(7):851-857
- 10 196. Khan A, Khan IA, Abidi H, Ahmed M. Comparison of empagliflozin and vildagliptin for efficacy and safety in type 2 diabetes mellitus in the Pakistani population. Frontiers in Endocrinology. 2022; 13:926633
- 13 197. Kim J-D, Park C-Y, Cha B-Y, Ahn KJ, Kim IJ, Park KS et al. Comparison of
 14 Adherence to Glimepiride/Metformin Sustained Release Once-daily Versus
 15 Glimepiride/Metformin Immediate Release BID Fixed-combination Therapy Using the
 16 Medication Event Monitoring System in Patients With Type 2 Diabetes. Clinical
 17 Therapeutics. 2018; 40(5):752-761e752
- 18 198. Kim JM, Kim SS, Kim JH, Kim MK, Kim TN, Lee SH et al. Efficacy and safety of pioglitazone versus glimepiride after metformin and alogliptin combination therapy: A randomized, open-label, multicenter, parallel-controlled study. Diabetes Metabol J. 2020; 44(1):67-77
- 199. Kimura T, Katakura Y, Shimoda M, Kawasaki F, Yamabe M, Tatsumi F et al.
 Comparison of clinical efficacy and safety of weekly glucagon-like peptide-1 receptor agonists dulaglutide and semaglutide in Japanese patients with type 2 diabetes:
 Randomized, parallel-group, multicentre, open-label trial (COMING study). Diabetes,
 Obesity & Metabolism. 2023;
- 27 200. Kinoshita T, Shimoda M, Nakashima K, Fushimi Y, Hirata Y, Tanabe A et al.
 28 Comparison of the effects of three kinds of glucose-lowering drugs on non-alcoholic
 29 fatty liver disease in patients with type 2 diabetes: a randomized, open-label, three30 arm, active control study. J Diabetes Invest. 2020;
- 201. Kohan DE, Fioretto P, Tang W, List JF. Long-term study of patients with type 2
 diabetes and moderate renal impairment shows that dapagliflozin reduces weight and
 blood pressure but does not improve glycemic control. Kidney International. 2014;
 85(4):962-971
- Komorizono Y, Hosoyamada K, Imamura N, Kajiya S, Hashiguchi Y, Ueyama N et al.
 Metformin increase versus added linagliptin in nonalcoholic liver disease and type 2
 diabetes: An analysis of J-LINK study. Diabetes, Obesity & Metabolism. 2020;
- 38 203. Kooy A, Jager J, Lehert P, Bets D, Wulffel MG, Donker AJ et al. Long-term effects of metformin on metabolism and microvascular and macrovascular disease in patients with type 2 diabetes mellitus. Archives of Internal Medicine. 2009; 169(6):616-625
- 41 204. Kothny W, Foley J, Kozlovski P, Shao Q, Gallwitz B, Lukashevich V. Improved 42 glycaemic control with vildagliptin added to insulin, with or without metformin, in 43 patients with type 2 diabetes mellitus. Diabetes Obes Metab. 2013; 15(3):252-257
- 44 205. Kothny W, Lukashevich V, Foley JE, Rendell MS, Schweizer A. Comparison of 45 vildagliptin and sitagliptin in patients with type 2 diabetes and severe renal 46 impairment: a randomised clinical trial. Diabetologia. 2015; 58(9):2020-2026

- 1 206. Kovacs CS, Seshiah V, Swallow R, Jones R, Rattunde H, Woerle HJ et al.
 2 Empagliflozin improves glycaemic and weight control as add-on therapy to
 3 pioglitazone or pioglitazone plus metformin in patients with type 2 diabetes: A 244 week, randomized, placebo-controlled trial. Diabetes Obes Metab. 2014; 16(2):147158
- 6 207. Koyama H, Tanaka S, Monden M, Morioka T, Fukumoto S, Mori K et al. Comparison 7 of effects of pioglitazone and glimepiride on plasma soluble RAGE and RAGE 8 expression in peripheral mononuclear cells in type 2 diabetes: Randomized controlled 9 trial (PioRAGE). Atherosclerosis. 2014; 234(2):329-334
- 208. Langenfeld MR, Forst T, Hohberg C, Kann P, Lubben G, Konrad T. Pioglitazone decreases carotid intima-media thickness independently of glycaemic control in patients with type 2 diabetes mellitus. Results from a controlled randomized study. Circulation. 2005; 111:2525-2531
- Lavalle-Gonzalez FJ, Januszewicz A, Davidson J, Tong C, Qiu R, Canovatchel W et
 al. Efficacy and safety of canagliflozin compared with placebo and sitagliptin in
 patients with type 2 diabetes on background metformin monotherapy: A randomised
 trial. Diabetologia. 2013; 56(12):2582-2592
- 18 210. Ledesma G, Umpierrez GE, Morley JE, Lewis-D'Agostino D, Keller A, Meinicke T et 19 al. Efficacy and safety of linagliptin to improve glucose control in older people with 20 type 2 diabetes on stable insulin therapy: a randomized trial. Diabetes Obes Metab. 21 2019;
- 22 211. Lee CH, Wu MZ, Lui D-W, Chan D-H, Fong C-Y, Shiu S-M et al. Comparison of
 23 Serum Ketone Levels and Cardiometabolic Efficacy of Dapagliflozin versus Sitagliptin
 24 among Insulin-Treated Chinese Patients with Type 2 Diabetes Mellitus. Diabetes &
 25 Metabolism Journal. 2022;
- 26 212. Lee HW, Lee HC, Kim BW, Yang MJ, Park JS, Oh JH et al. Effects of low dose pioglitazone on restenosis and coronary atherosclerosis in diabetic patients undergoing drug eluting stent implantation. Yonsei Medical Journal. 2013; 54(6):1313-1320
- Leiter LA, Cefalu WT, De Bruin TWA, Gause-Nilsson I, Sugg J, Parikh SJ.
 Dapagliflozin added to usual care in individuals with type 2 diabetes mellitus with
 preexisting cardiovascular disease: A 24-week, multicenter, randomized, double-blind, placebo-controlled study with a 28-week extension. Journal of the American
 Geriatrics Society. 2014; 62(7):1252-1262
- Li C-J, Liu X-J, Bai L, Yu Q, Zhang Q-M, Yu P et al. Efficacy and safety of vildagliptin,
 Saxagliptin or Sitagliptin as add-on therapy in Chinese patients with type 2 diabetes
 inadequately controlled with dual combination of traditional oral hypoglycemic agents.
 Diabetology & Metabolic Syndrome. 2014; 6:69
- 215. Li C-J, Zhang J-Y, Yu D-M, Zhang Q-M. Adding glimepiride to current insulin therapy increases high-molecular weight adiponectin levels to improve glycemic control in poorly controlled type 2 diabetes. Diabetology & Metabolic Syndrome. 2014; 6(1):41
- 42 216. Li CJ, Yu Q, Yu P, Zhang QM, Ding M, Liu XJ et al. Efficacy and safety comparison of add-on therapy with liraglutide, saxagliptin and vildagliptin, all in combination with current conventional oral hypoglycemic agents therapy in poorly controlled Chinese type 2 diabetes. Experimental and Clinical Endocrinology and Diabetes. 2014; 122(8):469-476

- Li F, Shen Y, Sumn R, Zhang D, Jin X, Zhai X et al. Effects of vildagliptin add-on insulin therapy on nocturnal glycemic variations in uncontrolled type 2 diabetes.
 Diabetes Therapy. 2017; 8(5):1111-1122
- Lim S, Lee S-H, Min K-W, Lee CB, Kim SY, Yoo HJ et al. A multicentre, double-blind, placebo-controlled, randomized, parallel comparison, phase 3 trial to evaluate the efficacy and safety of pioglitazone add-on therapy in type 2 diabetic patients treated with metformin and dapagliflozin. Diabetes, Obesity & Metabolism. 2024;
- Lind M, Hirsch IB, Tuomilehto J, Dahlqvist S, Ahren B, Torffvit O et al. Liraglutide in people treated for type 2 diabetes with multiple daily insulin injections: randomised clinical trial (MDI Liraglutide trial). BMJ. 2015; 351:h5364
- Lingvay I, Catarig AM, Frias JP, Kumar H, Lausvig NL, le Roux CW et al. Efficacy and safety of once-weekly semaglutide versus daily canagliflozin as add-on to metformin in patients with type 2 diabetes (SUSTAIN 8): a double-blind, phase 3b, randomised controlled trial. Lancet Diabetes Endocrinol. 2019; 7(11):834-844
- Lingvay I, Perez Manghi F, Garcia-Hernandez P, Norwood P, Lehmann L, Tarp-Johansen MJ et al. Effect of Insulin Glargine Up-titration vs Insulin
 Degludec/Liraglutide on Glycated Hemoglobin Levels in Patients With Uncontrolled
 Type 2 Diabetes: The DUAL V Randomized Clinical Trial. JAMA. 2016; 315(9):898-907
- 22. Liu SC, Chien KL, Wang CH, Chen WC, Leung CH. Efficacy and safety of adding pioglitazone or sitagliptin to patients with type 2 diabetes insufficiently controlled with metformin and a sulfonylurea. Endocrine Practice. 2013; 19(6):980-988
- 23. Liu SC, Lee CC, Chuang SM, Sun FJ, Zeng YH. Comparison of efficacy and safety of 24 empagliflozin vs linagliptin added to premixed insulin in patients with uncontrolled 25 type 2 diabetes: A randomized, open-label study. Diabetes and Metabolism. 2021; 26 47(3):101184
- 27 224. Liutkus J, Rosas Guzman J, Norwood P, Pop L, Northrup J, Cao D et al. A placebo-28 controlled trial of exenatide twice-daily added to thiazolidinediones alone or in 29 combination with metformin. Diabetes Obes Metab. 2010; 12(12):1058-1065
- 225. Ludvik B, Fr?as JP, Tinahones FJ, Wainstein J, Jiang H, Robertson KE et al.
 Dulaglutide as add-on therapy to SGLT2 inhibitors in patients with inadequately
 controlled type 2 diabetes (AWARD-10): a 24-week, randomised, double-blind,
 placebo-controlled trial. Lancet Diabetes Endocrinol. 2018; 6(5):370-381
- 226. Ludvik B, Giorgino F, Jodar E, Frias JP, Fernandez Lando L, Brown K et al. Onceweekly tirzepatide versus once-daily insulin degludec as add-on to metformin with or without SGLT2 inhibitors in patients with type 2 diabetes (SURPASS-3): a randomised, open-label, parallel-group, phase 3 trial. Lancet. 2021; 398(10300):583-
- Lukashevich V, Del Prato S, Araga M, Kothny W. Efficacy and safety of vildagliptin in patients with type 2 diabetes mellitus inadequately controlled with dual combination of metformin and sulphonylurea. Diabetes, Obesity & Metabolism. 2014; 16(5):403-409
- Lukashevich V, Schweizer A, Shao Q, Groop PH, Kothny W. Safety and efficacy of vildagliptin versus placebo in patients with type 2 diabetes and moderate or severe renal impairment: a prospective 24-week randomized placebo-controlled trial.

 Diabetes Obes Metab. 2011; 13(10):947-954
- Lundby-Christensen L, Tarnow L, Boesgaard TW, Lund SS, Wiinberg N, Perrild H et al. Metformin versus placebo in combination with insulin analogues in patients with

- type 2 diabetes mellitus-the randomised, blinded Copenhagen Insulin and Metformin Therapy (CIMT) trial. BMJ Open. 2016; 6(2):e008376
- 3 230. Ma J, Liu M, Wang R, Du L, Ji L. Efficacy and safety of tirzepatide in people with type 2 diabetes by baseline body mass index: an exploratory subgroup analysis of SURPASS-AP-Combo. Diabetes, Obesity & Metabolism. 2024;
- 6 231. Macauley M, Hollingsworth KG, Smith FE, Thelwall PE, Al-Mrabeh A, Schweizer A et al. Effect of vildagliptin on hepatic steatosis. J Clin Endocrinol Metabol. 2015; 100(4):1578-1585
- 9 232. Mahaffey Kenneth W, Neal B, Perkovic V, de Zeeuw D, Fulcher G, Erondu N et al.
 10 Canagliflozin for Primary and Secondary Prevention of Cardiovascular Events:
 11 Results From the CANVAS Program (Canagliflozin Cardiovascular Assessment
 12 Study). Circulation. 2018; 137(4):323-334
- 233. Marre M, Shaw J, Br?ndle M, Bebakar WM, Kamaruddin NA, Strand J et al.
 Liraglutide, a once-daily human GLP-1 analogue, added to a sulphonylurea over 26 weeks produces greater improvements in glycaemic and weight control compared with adding rosiglitazone or placebo in subjects with Type 2 diabetes (LEAD-1 SU).
 Diabetic Medicine. 2009; 26(3):268-278
- 18 234. Marso SP, Poulter NR, Nissen SE, Nauck MA, Zinman B, Daniels GH et al. Design of 19 the liraglutide effect and action in diabetes: evaluation of cardiovascular outcome 20 results (LEADER) trial. American Heart Journal. 2013; 166(5):823-830e825
- 235. Marso Steven P, Bain Stephen C, Consoli A, Eliaschewitz Freddy G, Jodar E, Leiter Lawrence A et al. Semaglutide and Cardiovascular Outcomes in Patients with Type 2 Diabetes. The New England journal of medicine. 2016; 375(19):1834-1844
- 24 236. Marso Steven P, Daniels Gilbert H, Brown-Frandsen K, Kristensen P, Mann
 25 Johannes F E, Nauck Michael A et al. Liraglutide and Cardiovascular Outcomes in
 26 Type 2 Diabetes. The New England journal of medicine. 2016; 375(4):311-322
- 27 237. Marx N, Rosenstock J, Kahn SE, Zinman B, Kastelein JJ, Lachin JM et al. Design
 28 and baseline characteristics of the CARdiovascular Outcome Trial of LINAgliptin
 29 Versus Glimepiride in Type 2 Diabetes (CAROLINA R). Diabetes & Vascular Disease
 30 Research. 2015; 12(3):164-174
- Mathieu C, Ranetti AE, Li D, Ekholm E, Cook W, Hirshberg B et al. Randomized,
 Double-Blind, Phase 3 Trial of Triple Therapy With Dapagliflozin Add-on to
 Saxagliptin Plus Metformin in Type 2 Diabetes. Diabetes Care. 2015; 38(11):2009 2017
- 35 239. Mathieu C, Rodbard HW, Cariou B, Handelsman Y, Philis-Tsimikas A, Ocampo 36 Francisco AM et al. A comparison of adding liraglutide versus a single daily dose of 37 insulin aspart to insulin degludec in subjects with type 2 diabetes (BEGIN: VICTOZA 38 ADD-ON). Diabetes Obes Metab. 2014; 16(7):636-644
- 39 240. Mathieu C, Shankar RR, Lorber D, Umpierrez G, Wu F, Xu L et al. A randomized 40 clinical trial to evaluate the efficacy and safety of co-administration of sitagliptin with 41 intensively titrated insulin glargine. Diabetes Therapy. 2015; 6(2):127-142
- 42 241. Matthaei S, Bowering K, Rohwedder K, Grohl A, Parikh S, Study G. Dapagliflozin 43 improves glycemic control and reduces body weight as add-on therapy to metformin 44 plus sulfonylurea: a 24-week randomized, double-blind clinical trial. Diabetes Care. 45 2015; 38(3):365-372
- 46 242. Matthaei S, Catrinoiu D, Celinski A, Ekholm E, Cook W, Hirshberg B et al. Randomized, double-blind trial of triple therapy with saxagliptin add-on to

- dapagliflozin plus metformin in patients with type 2 diabetes. Diabetes Care. 2015; 38(11):2018-2024
- Matthews DR, Charbonnel BH, Hanefeld M, Brunetti P, Schernthaner G. Long-term therapy with addition of pioglitazone to metformin compared with the addition of gliclazide to metformin in patients with type 2 diabetes: a randomized, comparative study. Diabetes/Metabolism Research and Reviews. 2005; 21(2):167-174
- 7 244. Matthews DR, Dejager S, Ahren B, Fonseca V, Ferrannini E, Couturier A et al.
 8 Vildagliptin add-on to metformin produces similar efficacy and reduced
 9 hypoglycaemic risk compared with glimepiride, with no weight gain: results from a 210 year study. Diabetes Obes Metab. 2010; 12(9):780-789
- Mattoo V, Eckland D, Widel M, Duran S, Fajardo C, Strand J et al. Metabolic effects of pioglitazone in combination with insulin in patients with type 2 diabetes mellitus whose disease is not adequately controlled with insulin therapy: results of a sixmonth, randomized, double-blind, prospective, multicenter, parallel-group study.
 Clinical Therapeutics. 2005; 27(5):554-567
- Mazzone T, Meyer PM, Feinstein SB, Davidson MH, Kondos GT, D'Agostino RB et al.
 Effect of pioglitazone compared with glimepiride on carotid intima-media thickness in type 2 diabetes: a randomized trial. JAMA. 2006; 296(21):2572-2581
- McCluskey D, Touger MS, Melis R, Schleusener DS, McCluskey D. Results of a
 randomized, double-blind, placebo-controlled study administering glimepiride to
 patients with type 2 diabetes mellitus inadequately controlled with rosiglitazone
 monotherapy. Clinical Therapeutics. 2004; 26(11):1783-1790
- 248. McGill JB, Sloan L, Newman J, Patel S, Sauce C, Eynatten M et al. Long-term efficacy and safety of linagliptin in patients with type 2 diabetes and severe renal impairment: a 1-year, randomized, double-blind, placebo-controlled study. Diabetes Care. 2013; 36(2):237-244
- 27 249. McMurray JJV, Ponikowski P, Bolli GB, Lukashevich V, Kozlovski P, Kothny W et al. 28 Effects of vildagliptin on ventricular function in patients with type 2 diabetes mellitus 29 and heart failure: A randomized placebo-controlled trial. JACC: heart failure. 2018; 30 6(1):8-17
- 31 250. Meneghini LF, Traylor L, Schwartz SL. Improved glycemic control with insulin glargine 32 versus pioglitazone as add-on therapy to sulfonylurea or metformin in patients with 33 uncontrolled type 2 diabetes mellitus. Endocrine Practice. 2010; 16(4):588-599
- 34 251. Meneilly GS, Roy-Duval C, Alawi H, Dailey G, Bellido D, Trescoli C et al. Lixisenatide 35 therapy in older patients with type 2 diabetes inadequately controlled on their current 36 antidiabetic treatment: The GetGoal-O randomized trial. Diabetes Care. 2017; 37 40(4):485-493
- 38 252. Mentz RJ, Bethel MA, Gustavson S, Thompson VP, Pagidipati NJ, Buse JB et al.
 39 Baseline characteristics of patients enrolled in the Exenatide Study of Cardiovascular
 40 Event Lowering (EXSCEL). American Heart Journal. 2017; 187:1-9
- 41 253. Miras AD, Perez-Pevida B, Aldhwayan M, Kamocka A, McGlone ER, Al-Najim W et 42 al. Adjunctive liraglutide treatment in patients with persistent or recurrent type 2 43 diabetes after metabolic surgery (GRAVITAS): a randomised, double-blind, placebo-44 controlled trial. Lancet Diabetes Endocrinol. 2019; 7(7):549-559
- 45 254. Moeinzadeh F, Iraj B, Mortazavi M, Ramezani P. The Renoprotective Effect of 46 Linagliptin in Type 2 Diabetic Patients with Severely Increased Albuminuria. Iranian 47 Journal of Kidney Diseases. 2021; 15(5):344-350

- 1 255. Moon JS, Ha KS, Yoon JS, Lee HW, Lee HC, Won KC. The effect of glargine versus glimepiride on pancreatic beta-cell function in patients with type 2 diabetes uncontrolled on metformin monotherapy: Open-label, randomized, controlled study. Acta Diabetologica. 2014; 51(2):277-285
- 5 256. Morikawa A, Ishizeki K, Iwashima Y, Yokoyama H, Muto E, Oshima E et al.
 6 Pioglitazone reduces urinary albumin excretion in renin-angiotensin system inhibitor7 treated type 2 diabetic patients with hypertension and microalbuminuria: the APRIME study. Clinical and Experimental Nephrology. 2011; 15(6):848-853
- 9 257. Mosenzon O, Blicher TM, Rosenlund S, Eriksson JW, Heller S, Hels OH et al.
 10 Efficacy and safety of oral semaglutide in patients with type 2 diabetes and moderate
 11 renal impairment (PIONEER 5): a placebo-controlled, randomised, phase 3a trial.
 12 Lancet Diabetes Endocrinol. 2019; 7(7):515-527
- 13 258. Moses RG, Kalra S, Brook D, Sockler J, Monyak J, Visvanathan J et al. A 14 randomized controlled trial of the efficacy and safety of saxagliptin as add-on therapy 15 in patients with type 2 diabetes and inadequate glycaemic control on metformin plus a 16 sulphonylurea. Diabetes Obes Metab. 2014; 16(5):443-450
- 17 259. Moses RG, Round E, Shentu Y, Golm GT, O'Neill EA, Gantz I et al. A randomized clinical trial evaluating the safety and efficacy of sitagliptin added to the combination of sulfonylurea and metformin in patients with type 2 diabetes mellitus and inadequate glycemic control. J Diabetes. 2017; 8(5):701-711
- 21 260. Muller-Wieland D, Kellerer M, Cypryk K, Skripova D, Rohwedder K, Johnsson E et al. Efficacy and safety of dapagliflozin or dapagliflozin plus saxagliptin versus glimepiride as add-on to metformin in patients with type 2 diabetes. Diabetes Obes Metab. 2018; 20(11):2598-2607
- 25 261. Nahra R, Wang T, Gadde KM, Oscarsson J, Stumvoll M, Jermutus L et al. Effects of
 Cotadutide on Metabolic and Hepatic Parameters in Adults With Overweight or
 Obesity and Type 2 Diabetes: A 54-Week Randomized Phase 2b Study. Diabetes
 Care. 2021; 44(6):1433-1442
- 29 262. Nakaguchi H, Kondo Y, Kyohara M, Konishi H, Oiwa K, Terauchi Y. Effects of liraglutide and empagliflozin added to insulin therapy in patients with type 2 diabetes: a randomized controlled study ELLENA?IT study. J Diabetes Invest. 2020;
- 32 263. Nauck M, Frid A, Hermansen K, Shah NS, Tankova T, Mitha IH et al. Efficacy and safety comparison of liraglutide, glimepiride, and placebo, all in combination with metformin, in type 2 diabetes. Diabetes Care. 2009; 32(1):84-90
- 35 264. Nauck M, Rizzo M, Johnson A, Bosch-Traberg H, Madsen J, Cariou B. Once-Daily 36 Liraglutide Versus Lixisenatide as Add-on to Metformin in Type 2 Diabetes: A 26-37 Week Randomized Controlled Clinical Trial. Diabetes Care. 2016; 39(9):1501-1509
- 38 265. Nauck M, Weinstock RS, Umpierrez GE, Guerci B, Skrivanek Z, Milicevic Z. Efficacy 39 and safety of dulaglutide versus sitagliptin after 52 weeks in type 2 diabetes in a 40 randomized controlled trial (AWARD-5). Diabetes Care. 2014; 37(8):2149-2158
- 41 266. Nauck MA, Duran S, Kim D, Johns D, Northrup J, Festa A et al. A comparison of 42 twice-daily exenatide and biphasic insulin aspart in patients with type 2 diabetes who 43 were suboptimally controlled with sulfonylurea and metformin: a non-inferiority study. 44 Diabetologia. 2007; 50(2):259-267
- 45 267. Nauck MA, Ellis GC, Fleck PR, Wilson CA, Mekki Q. Efficacy and safety of adding the dipeptidyl peptidase-4 inhibitor alogliptin to metformin therapy in patients with type 2 diabetes inadequately controlled with metformin monotherapy: a multicentre,

- randomised, double-blind, placebo-controlled study. International Journal of Clinical Practice. 2009; 63(1):46-55
- 3 268. Nauck MA, Meininger G, Sheng D, Terranella L, Stein PP. Efficacy and safety of the dipeptidyl peptidase-4 inhibitor, sitagliptin, compared with the sulfonylurea, glipizide, in patients with type 2 diabetes inadequately controlled on metformin alone: a randomized, double-blind, non-inferiority trial. Diabetes Obes Metab. 2007; 9(2):194-205
- 8 269. Nauck MA, Prato S, Meier JJ, Dur?n-Garc?a S, Rohwedder K, Elze M et al.
 9 Dapagliflozin versus glipizide as add-on therapy in patients with type 2 diabetes who
 10 have inadequate glycemic control with metformin: a randomized, 52-week, double11 blind, active-controlled noninferiority trial. Diabetes Care. 2011; 34(9):2015-2022
- 12 270. Nesti L, Pugliese NR, Sciuto P, Trico D, Dardano A, Baldi S et al. Effect of 13 empagliflozin on left ventricular contractility and peak oxygen uptake in subjects with 14 type 2 diabetes without heart disease: results of the EMPA-HEART trial. 15 Cardiovascular Diabetology. 2022; 21(1)
- 16 271. Ning G, Li L, Ma J, Lv X, Yang M, Wang W et al. Vildagliptin as add-on therapy to insulin improves glycemic control without increasing risk of hypoglycemia in Asian, predominantly Chinese, patients with type 2 diabetes mellitus. J Diabetes. 2016; 8(3):345-353
- 272. Nissen SE, Nicholls SJ, Wolski K, Nesto R, Kupfer S, Perez A et al. Comparison of pioglitazone vs glimepiride on progression of coronary atherosclerosis in patients with type 2 diabetes: the PERISCOPE randomized controlled trial. JAMA. 2008; 299(13):1561-1573
- 24 273. Nogueira KC, Furtado M, Fukui RT, Correia MRS, Dos Santos RF, Andrade JL et al. 25 Left ventricular diastolic function in patients with type 2 diabetes treated with a 26 dipeptidyl peptidase-4 inhibitor- a pilot study. Diabetology & Metabolic Syndrome. 27 2014; 6(1)
- 28 274. Nowicki M, Rychlik I, Haller H, Warren M, Suchower L, Gause-Nilsson I et al. Long-29 term treatment with the dipeptidyl peptidase-4 inhibitor saxagliptin in patients with 30 type 2 diabetes mellitus and renal impairment: a randomised controlled 52-week 31 efficacy and safety study. International Journal of Clinical Practice. 2011; 32 65(12):1230-1239
- 33 275. Oh M, Choi JH, Kim SO, Lee PH, Ahn JM, Lee SW et al. Comparison of empagliflozin and sitagliptin therapy on myocardial perfusion reserve in diabetic patients with coronary artery disease. Nuclear Medicine Communications. 2021; 42(9):972-978
- Ohira M, Yamaguchi T, Saiki A, Ban N, Kawana H, Nagayama D et al. Metformin reduces circulating malondialdehyde-modified low-density lipoprotein in type 2 diabetes mellitus. Clinical and Investigative Medicine Medecine Clinique et Experimentale. 2014; 37(4):E243-251
- Ohira M, Yamaguchi T, Saiki A, Ban N, Kawana H, Nagumo A et al. Pioglitazone
 improves the cardio-ankle vascular index in patients with type 2 diabetes mellitus
 treated with metformin. Diabetes, Metabolic Syndrome and Obesity. 2014; 7:313-319
- 278. Omachi T, Ohara M, Fujikawa T, Kohata Y, Sugita H, Irie S et al. Comparison of Effects of Injectable Semaglutide and Dulaglutide on Oxidative Stress and Glucose Variability in Patients with Type 2 Diabetes Mellitus: A Prospective Preliminary Study. Diabetes therapy: research, treatment and education of diabetes and related disorders. 2024; 15(1):111-126

- Owens DR, Swallow R, Dugi KA, Woerle HJ. Efficacy and safety of linagliptin in persons with type 2 diabetes inadequately controlled by a combination of metformin and sulphonylurea: a 24-week randomized study. Diabetic Med. 2011; 28(11):1352-1361
- 5 280. Pan C, Xing X, Han P, Zheng S, Ma J, Liu J et al. Efficacy and tolerability of vildagliptin as add-on therapy to metformin in Chinese patients with type 2 diabetes mellitus. Diabetes Obes Metab. 2012; 14(8):737-744
- Pan CY, Han P, Liu X, Yan S, Feng P, Zhou Z et al. Lixisenatide treatment improves glycaemic control in Asian patients with type 2 diabetes mellitus inadequately controlled on metformin with or without sulfonylurea: a randomized, double-blind, placebo-controlled, 24-week trial (GetGoal-M-Asia). Diabetes/Metabolism Research and Reviews. 2014; 30(8):726-735
- 282. Papathanassiou K, Naka KK, Kazakos N, Kanioglou C, Makriyiannis D, Pappas K et al. Pioglitazone vs glimepiride: Differential effects on vascular endothelial function in patients with type 2 diabetes. Atherosclerosis. 2009; 205(1):221-226
- Park CY, Kang JG, Chon S, Noh J, Oh SJ, Lee CB et al. Comparison between the therapeutic effect of metformin, glimepiride and their combination as an add-on treatment to insulin glargine in uncontrolled patients with type 2 diabetes. PloS One. 2014; 9(3):e88779
- 284. Park HK, Kim K-A, Min K-W, Sohn T-S, Jeong IK, Ahn CW et al. Effects of dapagliflozin compared with glimepiride on body composition in Asian patients with type 2 diabetes inadequately controlled with metformin: The BEYOND study. Diabetes, Obesity & Metabolism. 2023;
- 24 285. Park JS, Cho MH, Nam JS, Yoo JS, Ahn CW, Cha BS et al. Effect of pioglitazone on serum concentrations of osteoprotegerin in patients with type 2 diabetes mellitus.
 26 European Journal of Endocrinology of the European Federation of Endocrine
 27 Societies. 2011; 164(1):69-74
- 28 286. Pasquel FJ, Urrutia MA, Cardona S, Coronado KWZ, Albury B, Perez-Guzman MC et al. Liraglutide hospital discharge trial: A randomized controlled trial comparing the safety and efficacy of liraglutide versus insulin glargine for the management of patients with type 2 diabetes after hospital discharge. Diabetes, Obesity & Metabolism. 2021; 23(6):1351-1360
- 287. Pei Y, Agner BR, Luo B, Dong X, Li D, Liu J et al. DUAL II China: Superior HbA1c reductions and weight loss with insulin degludec/liraglutide (IDegLira) versus insulin degludec in a randomized trial of Chinese people with type 2 diabetes inadequately controlled on basal insulin. Diabetes, Obesity & Metabolism. 2021; 23(12):2687-2696
- 288. Perkovic V, Jardine MJ, Neal B, Bompoint S, Heerspink HJL, Charytan DM et al.
 Canagliflozin and renal outcomes in type 2 diabetes and nephropathy. New England
 Journal of Medicine. 2019; 380(24):2295-2306
- 40 289. Perkovic V TKRRPMKWMJFEBGBFMMITB-TH, Committees FT, Investigators.
 41 Effects of Semaglutide on Chronic Kidney Disease in Patients with Type 2 Diabetes.
 42 New England Journal of Medicine. 2024; 391(2)
- 290. Petrica L, Vlad A, Petrica M, Jianu CD, Gluhovschi G, Gadalean F et al. Pioglitazone
 delays proximal tubule dysfunction and improves cerebral vessel endothelial
 dysfunction in normoalbuminuric people with type 2 diabetes mellitus. Diabetes
 Research and Clinical Practice. 2011; 94(1):22-32

- 1 291. Pf?tzner A, Marx N, L?bben G, Langenfeld M, Walcher D, Konrad T et al.
 2 Improvement of cardiovascular risk markers by pioglitazone is independent from
 3 glycemic control: results from the pioneer study. Journal of the American College of
 4 Cardiology. 2005; 45(12):1925-1931
- 5 292. Pf?tzner A, Sch?ndorf T, Tsch?pe D, Lobmann R, Merke J, M?ller J et al. PIOfix-6 study: effects of pioglitazone/metformin fixed combination in comparison with a 7 combination of metformin with glimepiride on diabetic dyslipidemia. Diabetes 8 Technology & Therapeutics. 2011; 13(6):637-643
- 9 293. Pfeffer Marc A, Claggett B, Diaz R, Dickstein K, Gerstein Hertzel C, Kober Lars V et al. Lixisenatide in Patients with Type 2 Diabetes and Acute Coronary Syndrome. The New England journal of medicine. 2015; 373(23):2247-2257
- 294. Philis-Tsimikas A, Billings LK, Busch R, Portillo CM, Sahay R, Halladin N et al.

 Superior efficacy of insulin degludec/liraglutide versus insulin glargine U100 as addon to sodium-glucose co-transporter-2 inhibitor therapy: a randomized clinical trial in
 people with uncontrolled type 2 diabetes. Diab Obes Metab. 2019; 21(6):1399-1408
- Philis-Tsimikas A, Del Prato S, Satman I, Bhargava A, Dharmalingam M, Skjoth TV et
 al. Effect of insulin degludec versus sitagliptin in patients with type 2 diabetes
 uncontrolled on oral antidiabetic agents. Diabetes Obes Metab. 2013; 15(8):760-766
- 296. Phrommintikul A, Wongcharoen W, Kumfu S, Jaiwongkam T, Gunaparn S,
 Chattipakorn S et al. Effects of dapagliflozin vs vildagliptin on cardiometabolic
 parameters in diabetic patients with coronary artery disease: a randomised study.
 British Journal of Clinical Pharmacology. 2019; 85(6):1337-1347
- 23 297. Pieber TR, Bode B, Mertens A, Cho YM, Christiansen E, Hertz CL et al. Efficacy and safety of oral semaglutide with flexible dose adjustment versus sitagliptin in type 2 diabetes (PIONEER 7): a multicentre, open-label, randomised, phase 3a trial. Lancet Diabetes Endocrinol. 2019; 7(7):528-539
- 27 298. Pinget M, Goldenberg R, Niemoeller E, Muehlen-Bartmer I, Guo H, Aronson R.
 28 Efficacy and safety of lixisenatide once daily versus placebo in type 2 diabetes
 29 insufficiently controlled on pioglitazone (GetGoal-P). Diabetes Obes Metab. 2013;
 30 15(11):1000-1007
- 299. Pollock C, Stefansson B, Reyner D, Rossing P, Sjostrom CD, Wheeler DC et al.
 Albuminuria-lowering effect of dapagliflozin alone and in combination with saxagliptin
 and effect of dapagliflozin and saxagliptin on glycaemic control in patients with type 2
 diabetes and chronic kidney disease (DELIGHT): a randomised, double-blind,
 placebo-controlled trial. Lancet Diabetes Endocrinol. 2019; 7(6):429-441
- 36 300. Pozzilli P, Norwood P, Jodar E, Davies MJ, Ivanyi T, Jiang H et al. Placebo 37 controlled, randomized trial of the addition of once-weekly glucagon-like peptide-1
 38 receptor agonist dulaglutide to titrated daily insulin glargine in patients with type 2
 39 diabetes (AWARD-9). Diabetes Obes Metab. 2017; 19(7):1024-1031
- 40 301. Pratley R, Amod A, Hoff ST, Kadowaki T, Lingvay I, Nauck M et al. Oral semaglutide 41 versus subcutaneous liraglutide and placebo in type 2 diabetes (PIONEER 4): a 42 randomised, double-blind, phase 3a trial. Lancet. 2019; 394(10192):39-50
- 43 302. Pratley RE, Aroda VR, Lingvay I, Ludemann J, Andreassen C, Navarria A et al.
 44 Semaglutide versus dulaglutide once weekly in patients with type 2 diabetes
 45 (SUSTAIN 7): a randomised, open-label, phase 3b trial. The lancet Diabetes &
 46 endocrinology. 2018; 6(4):275-286

- 1 303. Pratley RE, Eldor R, Raji A, Golm G, Huyck SB, Qiu Y et al. Ertugliflozin plus sitagliptin versus either individual agent over 52 weeks in patients with type 2 diabetes mellitus inadequately controlled with metformin: the VERTIS FACTORIAL randomized trial. Diabetes Obes Metab. 2018; 20(5):1111-1120
- 5 304. Pratley RE, Kipnes MS, Fleck PR, Wilson C, Mekki Q. Efficacy and safety of the dipeptidyl peptidase-4 inhibitor alogliptin in patients with type 2 diabetes inadequately controlled by glyburide monotherapy. Diabetes Obes Metab. 2009; 11(2):167-176
- 8 305. Pratley RE, Nauck M, Bailey T, Montanya E, Cuddihy R, Filetti S et al. Liraglutide 9 versus sitagliptin for patients with type 2 diabetes who did not have adequate 10 glycaemic control with metformin: a 26-week, randomised, parallel-group, open-label 11 trial. Lancet. 2010; 375(9724):1447-1456
- 12 306. Pratley RE, Reusch JE, Fleck PR, Wilson CA, Mekki Q. Efficacy and safety of the dipeptidyl peptidase-4 inhibitor alogliptin added to pioglitazone in patients with type 2 diabetes: a randomized, double-blind, placebo-controlled study. Current Medical Research and Opinion. 2009; 25(10):2361-2371
- 16 307. Punthakee Z, Bosch J, Dagenais G, Diaz R, Holman R, Probstfield J et al. Design, history and results of the Thiazolidinedione Intervention with vitamin D Evaluation (TIDE) randomised controlled trial. Diabetologia. 2012; 55(1):36-45
- 308. Raman RB, Kumar D, Roushan R. Comparative Study of Efficacy and Safety of
 Empagliflozin vs Linagliptin as Add on Therapy to Insulin in Patients of Type 2
 Diabetes Mellitus and Chronic Kidney Disease in Tertiary Care Centre of Eastern
 India. International Journal of Pharmaceutical Sciences Review and Research. 2022;
 77(2):139-145
- 24 309. Raz I, Chen Y, Wu M, Hussain S, Kaufman KD, Amatruda JM et al. Efficacy and safety of sitagliptin added to ongoing metformin therapy in patients with type 2 diabetes. Current Medical Research and Opinion. 2008; 24(2):537-550
- 27 310. Retnakaran R, Qi Y, Opsteen C, Vivero E, Zinman B. Initial short-term intensive 28 insulin therapy as a strategy for evaluating the preservation of beta-cell function with 29 oral antidiabetic medications: a pilot study with sitagliptin. Diabetes Obes Metab. 30 2010; 12(10):909-915
- 31 311. Ridderstrale M, Andersen KR, Zeller C, Kim G, Woerle HJ, Broedl UC. Comparison of empagliflozin and glimepiride as add-on to metformin in patients with type 2 diabetes:
 A 104-week randomised, active-controlled, double-blind, phase 3 trial. Lancet Diabetes Endocrinol. 2014; 2(9):691-700
- 35 312. Riddle MC, Aronson R, Home P, Marre M, Niemoeller E, Miossec P et al. Adding once-daily lixisenatide for type 2 diabetes inadequately controlled by established basal insulin: a 24-week, randomized, placebo-controlled comparison (GetGoal-L). Diabetes Care. 2013; 36(9):2489-2496
- 39 313. Riddle MC, Forst T, Aronson R, Sauque-Reyna L, Souhami E, Silvestre L et al.
 40 Adding once-daily lixisenatide for type 2 diabetes inadequately controlled with newly
 41 initiated and continuously titrated basal insulin glargine: a 24-week, randomized,
 42 placebo-controlled study (GetGoal-Duo 1). Diabetes Care. 2013; 36(9):2497-2503
- 43 314. Riddle MC, Schneider J. Beginning insulin treatment of obese patients with evening 70/30 insulin plus glimepiride versus insulin alone. Glimepiride Combination Group. Diabetes Care. 1998; 21(7):1052-1057
- 46 315. Roberts VL, Stewart J, Issa M, Lake B, Melis R. Triple therapy with glimepiride in patients with type 2 diabetes mellitus inadequately controlled by metformin and a

- thiazolidinedione: results of a 30-week, randomized, double-blind, placebo-controlled, parallel-group study. Clinical Therapeutics. 2005; 27(10):1535-1547
- 3 316. Rodbard Helena W, Rosenstock J, Canani Luis H, Deerochanawong C, Gumprecht J,
 4 Lindberg Soren O et al. Oral Semaglutide Versus Empagliflozin in Patients With Type
 5 2 Diabetes Uncontrolled on Metformin: The PIONEER 2 Trial. Diabetes Care. 2019;
 6 42(12):2272-2281
- 7 317. Rodbard HW, Bode BW, Harris SB, Rose L, Lehmann L, Jarlov H et al. Safety and efficacy of insulin degludec/liraglutide (IDegLira) added to sulphonylurea alone or to sulphonylurea and metformin in insulin-naive people with Type 2 diabetes: the DUAL IV trial. Diabetic medicine: a journal of the British Diabetic Association. 2017; 34(2):189-196
- 318. Rodbard HW, Lingvay I, Reed J, de la Rosa R, Rose L, Sugimoto D et al.
 Semaglutide Added to Basal Insulin in Type 2 Diabetes (SUSTAIN 5): a Randomized,
 Controlled Trial. J Clin Endocrinol Metabol. 2018; 103(6):2291-2301
- 15 319. Rodbard HW, Seufert J, Aggarwal N, Cao A, Fung A, Pfeifer M et al. Efficacy and safety of titrated canagliflozin in patients with type 2 diabetes mellitus inadequately controlled on metformin and sitagliptin. Diabetes Obes Metab. 2016; 18(8):812-819
- 18 320. Roden M, Laakso M, Johns D, Widel M, Urquhart R, Richardson C et al. Long-term effects of pioglitazone and metformin on insulin sensitivity in patients with Type 2 diabetes mellitus. Diabetic medicine : a journal of the British Diabetic Association. 2005; 22(8):1101-1106
- 22 321. Rosenstock J, Allison D, Birkenfeld AL, Blicher TM, Deenadayalan S, Jacobsen JB et 23 al. Effect of additional oral semaglutide vs sitagliptin on glycated hemoglobin in adults 24 with type 2 diabetes uncontrolled with metformin alone or with sulfonylurea: the 25 PIONEER 3 randomized clinical trial. JAMA. 2019; 321(15):1466-1480
- 26 322. Rosenstock J, Aronson R, Grunberger G, Hanefeld M, Piatti P, Serusclat P et al.
 27 Benefits of LixiLan, a Titratable Fixed-Ratio Combination of Insulin Glargine Plus
 28 Lixisenatide, Versus Insulin Glargine and Lixisenatide Monocomponents in Type 2
 29 Diabetes Inadequately Controlled on Oral Agents: The LixiLan-O Randomized Trial.
 30 Diabetes Care. 2016; 39(11):2026-2035
- 323. Rosenstock J, Brazg R, Andryuk PJ, Lu K, Stein P. Efficacy and safety of the dipeptidyl peptidase-4 inhibitor sitagliptin added to ongoing pioglitazone therapy in patients with type 2 diabetes: a 24-week, multicenter, randomized, double-blind, placebo-controlled, parallel-group study. Clinical Therapeutics. 2006; 28(10):1556-1568
- 36 324. Rosenstock J, Buse JB, Azeem R, Prabhakar P, Kjems L, Huang H et al. Efficacy and Safety of ITCA 650, a Novel Drug-Device GLP-1 Receptor Agonist, in Type 2
 38 Diabetes Uncontrolled With Oral Antidiabetes Drugs: the FREEDOM-1 Trial. Diabetes Care. 2018; 41(2):333-340
- 40 325. Rosenstock J, Diamant M, Aroda VR, Silvestre L, Souhami E, Zhou T et al. Efficacy 41 and Safety of LixiLan, a Titratable Fixed-Ratio Combination of Lixisenatide and 42 Insulin Glargine, Versus Insulin Glargine in Type 2 Diabetes Inadequately Controlled 43 on Metformin Monotherapy: The LixiLan Proof-of-Concept Randomized Trial. 44 Diabetes Care. 2016; 39(9):1579-1586
- 45 326. Rosenstock J, Frias J, Pall D, Charbonnel B, Pascu R, Saur D et al. Effect of 46 ertugliflozin on glucose control, body weight, blood pressure and bone density in type 47 2 diabetes mellitus inadequately controlled on metformin monotherapy (VERTIS 48 MET). Diabetes, Obesity & Metabolism. 2018; 20(3):520-529

- 1 327. Rosenstock J, Frias JP, Rodbard HW, Tofe S, Sears E, Huh R et al. Tirzepatide vs 2 Insulin Lispro Added to Basal Insulin in Type 2 Diabetes: The SURPASS-6 3 Randomized Clinical Trial. JAMA. 2023; 330(17):1631-1640
- 4 328. Rosenstock J, Guerci B, Hanefeld M, Gentile S, Aronson R, Tinahones FJ et al.
 5 Prandial options to advance basal insulin glargine therapy: Testing lixisenatide plus
 6 basal insulin versus insulin glulisine either as basal-plus or basal-bolus in type 2
 7 diabetes: The GetGoal Duo-2 Trial. Diabetes Care. 2016; 39(8):1318-1328
- 8 329. Rosenstock J, Hanefeld M, Shamanna P, Min KW, Boka G, Miossec P et al.
 9 Beneficial effects of once-daily lixisenatide on overall and postprandial glycemic
 10 levels without significant excess of hypoglycemia in Type 2 diabetes inadequately
 11 controlled on a sulfonylurea with or without metformin (GetGoal-S). Journal of
 12 Diabetes and Its Complications. 2014; 28(3):386-392
- 330. Rosenstock J, Hansen L, Zee P, Li Y, Cook W, Hirshberg B et al. Dual add-on therapy in type 2 diabetes poorly controlled with metformin monotherapy: a randomized double-blind trial of saxagliptin plus dapagliflozin addition versus single addition of saxagliptin or dapagliflozin to metformin. Diabetes Care. 2015; 38(3):376-383
- 18 331. Rosenstock J, Jelaska A, Frappin G, Salsali A, Kim G, Woerle HJ et al. Improved 19 glucose control with weight loss, lower insulin doses, and no increased hypoglycemia 20 with empagliflozin added to titrated multiple daily injections of insulin in obese 21 inadequately controlled type 2 diabetes. Diabetes Care. 2014; 37(7):1815-1823
- 22 332. Rosenstock J, Jelaska A, Zeller C, Kim G, Broedl UC, Woerle HJ et al. Impact of 23 empagliflozin added on to basal insulin in type 2 diabetes inadequately controlled on 24 basal insulin: a 78-week randomized, double-blind, placebo-controlled trial. Diabetes 25 Obes Metab. 2015; 17(10):936-948
- 333. Rosenstock J, Kahn S E, Johansen O E, Zinman B, Espeland M A, Woerle H J et al.
 Effect of Linagliptin vs Glimepiride on Major Adverse Cardiovascular Outcomes in
 Patients with Type 2 Diabetes: The CAROLINA Randomized Clinical Trial. JAMA Journal of the American Medical Association. 2019; 322(12):1155-1166
- 30 334. Rosenstock J, Perkovic V, Johansen Odd E, Cooper Mark E, Kahn Steven E, Marx N et al. Effect of Linagliptin vs Placebo on Major Cardiovascular Events in Adults With Type 2 Diabetes and High Cardiovascular and Renal Risk: The CARMELINA Randomized Clinical Trial. JAMA. 2019; 321(1):69-79
- 34 335. Rosenstock J, Perl S, Johnsson E, Garcia-Sanchez R, Jacob S. Triple therapy with low-dose dapagliflozin plus saxagliptin versus dual therapy with each monocomponent, all added to metformin, in uncontrolled type 2 diabetes. Diabetes Obes Metab. 2019; 21(9):2152-2162
- 38 336. Rosenstock J, Raccah D, Koranyi L, Maffei L, Boka G, Miossec P et al. Efficacy and safety of lixisenatide once daily versus exenatide twice daily in type 2 diabetes inadequately controlled on metformin: a 24-week, randomized, open-label, active-controlled study (GetGoal-X). Diabetes Care. 2013; 36(10):2945-2951
- 42 337. Rosenstock J, Rendell MS, Gross JL, Fleck PR, Wilson CA, Mekki Q. Alogliptin 43 added to insulin therapy in patients with type 2 diabetes reduces HbA(1C) without 44 causing weight gain or increased hypoglycaemia. Diabetes Obes Metab. 2009; 45 11(12):1145-1152
- 46 338. Rosenstock J, Vico M, Wei L, Salsali A, List JF. Effects of dapagliflozin, an SGLT2 inhibitor, on HbA(1c), body weight, and hypoglycemia risk in patients with type 2

- diabetes inadequately controlled on pioglitazone monotherapy. Diabetes Care. 2012; 35(7):1473-1478
- 339. Roussel R, Duran-Garcia S, Zhang Y, Shah S, Darmiento C, Shankar RR et al.
 4 Double-blind, randomized clinical trial comparing the efficacy and safety of continuing
 5 or discontinuing the dipeptidyl peptidase-4 inhibitor sitagliptin when initiating insulin
 6 glargine therapy in patients with type 2 diabetes: the CompoSIT-I Study. Diab Obes
 7 Metab. 2019; 21(4):781-790
- 8 340. Russell-Jones D, Vaag A, Schmitz O, Sethi BK, Lalic N, Antic S et al. Liraglutide vs insulin glargine and placebo in combination with metformin and sulfonylurea therapy in type 2 diabetes mellitus (LEAD-5 met+SU): a randomised controlled trial.

 11 Diabetologia. 2009; 52(10):2046-2055
- 341. Sathyanarayana P, Jogi M, Muthupillai R, Krishnamurthy R, Samson SL, Bajaj M.
 Effects of combined exenatide and pioglitazone therapy on hepatic fat content in type
 2 diabetes. Obesity (Silver Spring, Md). 2011; 19(12):2310-2315
- Savvidou S, Karatzidou K, Tsakiri K, Gagalis A, Hytiroglou P, Goulis J. Circulating adiponectin levels in type 2 diabetes mellitus patients with or without non-alcoholic fatty liver disease: Results of a small, open-label, randomized controlled intervention trial in a subgroup receiving short-term exenatide. Diabetes Research and Clinical Practice. 2016; 113:125-134
- 343. Schernthaner G, Duran-Garcia S, Hanefeld M, Langslet G, Niskanen L, Ostgren CJ et
 al. Efficacy and tolerability of saxagliptin compared with glimepiride in elderly patients
 with type 2 diabetes: A randomized, controlled study (GENERATION). Diabetes Obes
 Metab. 2015; 17(7):630-638
- 344. Schernthaner G, Gross JL, Rosenstock J, Guarisco M, Fu M, Yee J et al.
 Canagliflozin compared with sitagliptin for patients with type 2 diabetes who do not have adequate glycemic control with metformin plus sulfonylurea: A 52-week randomized trial. Diabetes Care. 2013; 36(9):2508-2515
- Scirica Benjamin M, Bhatt Deepak L, Braunwald E, Steg P G, Davidson J, Hirshberg
 B et al. Saxagliptin and cardiovascular outcomes in patients with type 2 diabetes
 mellitus. The New England journal of medicine. 2013; 369(14):1317-1326
- 31 346. Scott R, Morgan J, Zimmer Z, Lam RLH, O'Neill EA, Kaufman KD et al. A randomized clinical trial of the efficacy and safety of sitagliptin compared with dapagliflozin in patients with type 2 diabetes mellitus and mild renal insufficiency: the CompoSIT-R study. Diab Obes Metab. 2018; 20(12):2876-2884
- 35 347. Seino Y, Kaku K, Kadowaki T, Okamoto T, Sato A, Shirakawa M et al. A randomized, placebo-controlled trial to assess the efficacy and safety of sitagliptin in Japanese patients with type 2 diabetes and inadequate glycaemic control on ipragliflozin. Diabetes, Obesity & Metabolism. 2021; 23(6):1342-1350
- 39 348. Seino Y, Kaneko S, Fukuda S, Osonoi T, Shiraiwa T, Nishijima K et al. Combination 40 therapy with liraglutide and insulin in Japanese patients with type 2 diabetes: A 36-41 week, randomized, double-blind, parallel-group trial. J Diabetes Invest. 2016; 42 7(4):565-573
- 43 349. Seino Y, Min KW, Niemoeller E, Takami A. Randomized, double-blind, placebo-44 controlled trial of the once-daily GLP-1 receptor agonist lixisenatide in Asian patients 45 with type 2 diabetes insufficiently controlled on basal insulin with or without a 46 sulfonylurea (GetGoal-L-Asia). Diabetes Obes Metab. 2012; 14(10):910-917

- Shankar RR, Bao Y, Han P, Hu J, Ma J, Peng Y et al. Sitagliptin added to stable insulin therapy with or without metformin in Chinese patients with type 2 diabetes. J Diabetes Invest. 2017; 8(3):321-329
- 4 351. Sivalingam S, Wasehuus VS, Rotbain Curovic V, Blond MB, Hansen TW, Persson F et al. Albuminuria-lowering effect of adding semaglutide on top of empagliflozin in individuals with type 2 diabetes: A randomized and placebo-controlled study.

 7 Diabetes, Obesity & Metabolism. 2023;
- 8 352. Softeland E, Meier JJ, Vangen B, Toorawa R, Maldonado-Lutomirsky M, Broedl UC. 9 Empagliflozin as add-on therapy in patients with type 2 diabetes inadequately 10 controlled with linagliptin and metformin: A 24-week randomized, double-blind, 11 parallel-group trial. Diabetes Care. 2017; 40(2):201-209
- 353. Sone H, Kaneko T, Shiki K, Tachibana Y, Pfarr E, Lee J et al. Efficacy and safety of
 empagliflozin as add-on to insulin in Japanese patients with type 2 diabetes: a
 randomised, double-blind, placebo-controlled trial. Diabetes Obes Metab. 2019;
- Sridhar S, Walia R, Sachdeva N, Bhansali A. Effect of pioglitazone on testosterone in eugonadal men with type 2 diabetes mellitus: a randomized double-blind placebo-controlled study. Clin Endocrinol. 2013; 78(3):454-459
- 18 355. Strain WD, Lukashevich V, Kothny W, Hoellinger MJ, Pald?nius PM. Individualised 19 treatment targets for elderly patients with type 2 diabetes using vildagliptin add-on or 20 lone therapy (INTERVAL): a 24 week, randomised, double-blind, placebo-controlled 21 study. Lancet. 2013; 382(9890):409-416
- 356. Strojek K, Yoon KH, Hruba V, Elze M, Langkilde AM, Parikh S. Effect of dapagliflozin in patients with type 2 diabetes who have inadequate glycaemic control with glimepiride: a randomized, 24-week, double-blind, placebo-controlled trial. Diabetes
 Obes Metab. 2011; 13(10):928-938
- 357. Su Y, Su YL, Lv LF, Wang LM, Li QZ, Zhao ZG. A randomized controlled clinical trial
 of vildagliptin plus metformin combination therapy in patients with type II diabetes
 mellitus. Experimental and Therapeutic Medicine. 2014; 7(4):799-803
- 358. Takahashi Y, Nomoto H, Yokoyama H, Takano Y, Nagai S, Tsuzuki A et al.
 Improvement of glycaemic control and treatment satisfaction by switching from
 liraglutide or dulaglutide to subcutaneous semaglutide in patients with type 2
 diabetes: A multicentre, prospective, randomized, open-label, parallel-group
 comparison study (SWITCH-SEMA 1 study). Diabetes, Obesity & Metabolism. 2023;
 34
- 35 359. Takihata M, Nakamura A, Tajima K, Inazumi T, Komatsu Y, Tamura H et al.
 36 Comparative study of sitagliptin with pioglitazone in Japanese type 2 diabetic
 37 patients: the COMPASS randomized controlled trial. Diabetes Obes Metab. 2013;
 38 15(5):455-462
- 39 360. Tan M, Johns D, Gonz?lez G?lvez G, Ant?nez O, Fabi?n G, Flores-Lozano F et al.
 40 Effects of pioglitazone and glimepiride on glycemic control and insulin sensitivity in
 41 Mexican patients with type 2 diabetes mellitus: A multicenter, randomized, double42 blind, parallel-group trial. Clinical Therapeutics. 2004; 26(5):680-693
- 43 361. Tanaka A, Shimabukuro M, MacHii N, Teragawa H, Okada Y, Shima KR et al. Effect 44 of empagliflozin on endothelial function in patients with type 2 diabetes and 45 cardiovascular disease: results from the multicenter, randomized, placebo- controlled, 46 double-blind EMBLEM trial. Diabetes Care. 2019; 42(10):E159-E161

- Tanaka K, Okada Y, Mori H, Miyazaki M, Kuno F, Sonoda S et al. Comparative analysis of the effects of alogliptin and vildagliptin on glucose metabolism in type 2 diabetes mellitus. Endocrine Journal. 2017; 64(2):179-189
- Taskinen MR, Rosenstock J, Tamminen I, Kubiak R, Patel S, Dugi KA et al. Safety and efficacy of linagliptin as add-on therapy to metformin in patients with type 2 diabetes: a randomized, double-blind, placebo-controlled study. Diabetes Obes Metab. 2011; 13(1):65-74
- 8 364. Terauchi Y, Utsunomiya K, Yasui A, Seki T, Cheng G, Shiki K et al. Safety and
 9 Efficacy of Empagliflozin as Add-On Therapy to GLP-1 Receptor Agonist (Liraglutide)
 10 in Japanese Patients with Type 2 Diabetes Mellitus: A Randomised, Double-Blind,
 11 Parallel-Group Phase 4 Study. Diabetes therapy: research, treatment and education
 12 of diabetes and related disorders. 2019; 10(3):951-963
- 13 365. Thrasher J, Daniels K, Patel S, Whetteckey J, Woerle HJ. Efficacy and safety of linagliptin in black/African American patients with type 2 diabetes: A 6-month, randomized, double-blind, placebo-controlled study. Endocrine Pract. 2014; 20(5):412-420
- Tinahones FJ, Gallwitz B, Nordaby M, G?tz S, Maldonado-Lutomirsky M, Woerle HJ et al. Linagliptin as add-on to empagliflozin and metformin in patients with type 2 diabetes: two 24-week randomized, double-blind, double-dummy, parallel-group trials. Diabetes Obes Metab. 2017; 19(2):266-274
- 367. Tripathy D, Daniele G, Fiorentino TV, Perez-Cadena Z, Chavez-Velasquez A,
 Kamath S et al. Pioglitazone improves glucose metabolism and modulates skeletal
 muscle TIMP-3-TACE dyad in type 2 diabetes mellitus: A randomised, double-blind,
 placebo-controlled, mechanistic study. Diabetologia. 2013; 56(10):2153-2163
- 25 368. Tuttle KR, Lakshmanan MC, Rayner B, Busch RS, Zimmermann AG, Woodward DB et al. Dulaglutide versus insulin glargine in patients with type 2 diabetes and moderate-to-severe chronic kidney disease (AWARD-7): a multicentre, open-label, randomised trial. Lancet Diabetes Endocrinol. 2018; 6(8):605
- 369. Umpierrez G, Issa M, Vlajnic A. Glimepiride versus pioglitazone combination therapy
 in subjects with type 2 diabetes inadequately controlled on metformin monotherapy:
 results of a randomized clinical trial. Current Medical Research and Opinion. 2006;
 22(4):751-759
- 370. V?h?talo M, R?nnemaa T, Viikari J. Recognition of fasting or overall hyperglycaemia when starting insulin treatment in patients with type 2 diabetes in general practice.

 Scandinavian Journal of Primary Health Care. 2007; 25(3):147-153
- 36 371. van der Meer RW, Rijzewijk LJ, Jong HW, Lamb HJ, Lubberink M, Romijn JA et al.
 37 Pioglitazone improves cardiac function and alters myocardial substrate metabolism
 38 without affecting cardiac triglyceride accumulation and high-energy phosphate
 39 metabolism in patients with well-controlled type 2 diabetes mellitus. Circulation. 2009;
 40 119(15):2069-2077
- 41 372. Van Eyk HJ, Paiman EHM, Bizino MB, De Heer P, Geelhoed-Duijvestijn PH, Kharagjitsingh AV et al. A double-blind, placebo-controlled, randomised trial to assess the effect of liraglutide on ectopic fat accumulation in South Asian type 2 diabetes patients. Cardiovascular Diabetology. 2019; 18(1):87
- 45 373. Van Gaal L, Souhami E, Zhou T, Aronson R. Efficacy and safety of the glucagon-like 46 peptide-1 receptor agonist lixisenatide versus the dipeptidyl peptidase-4 inhibitor 47 sitagliptin in young (<50 years) obese patients with type 2 diabetes mellitus. J Clin 48 Transl Endocrinol. 2014; 1(2):31-37

- Vanderheiden A, Harrison L, Warshauer J, Li X, Adams-Huet B, Lingvay I. Effect of adding liraglutide vs placebo to a high-dose Insulin regimen in patients with type 2 diabetes a randomized clinical trial. JAMA Int Med. 2016; 176(7):939-947
- Verma S, Mazer CD, Yan AT, Mason T, Garg V, Teoh H et al. Effect of empagliflozin on left ventricular mass in patients with type 2 diabetes and coronary artery disease: the EMPA-HEART CardioLink-6 randomized clinical trial. Circulation. 2019;
- 7 376. Vianna AGD, Lacerda CS, Pechmann LM, Polesel MG, Marino EC, Faria-Neto JR. A randomized controlled trial to compare the effects of sulphonylurea gliclazide MR (modified release) and the DPP-4 inhibitor vildagliptin on glycemic variability and control measured by continuous glucose monitoring (CGM) in Brazilian women with type 2 diabetes. Diabetes Research and Clinical Practice. 2018; 139:357-365
- 377. Vilsb?ll T, Rosenstock J, Yki-J?rvinen H, Cefalu WT, Chen Y, Luo E et al. Efficacy
 and safety of sitagliptin when added to insulin therapy in patients with type 2
 diabetes. Diabetes Obes Metab. 2010; 12(2):167-177
- 378. Vilsboll T, Ekholm E, Johnsson E, Dronamraju N, Jabbour S, Lind M. Dapagliflozin
 Plus Saxagliptin Add-on Therapy Compared With Insulin in Patients With Type 2
 Diabetes Poorly Controlled by Metformin With or Without Sulfonylurea Therapy: A
 Randomized Clinical Trial. Diabetes Care. 2019; 42(8):1464-1472
- 379. W?gner AM, Miranda-Calderin G, Ugarte-Lopetegui MA, Marrero-Santiago H,
 Suarez-Castellano L, Lopez-Madrazo MJ et al. Effect of liraglutide on physical
 performance in type 2 diabetes: results of a randomized, double-blind, controlled trial
 (LIPER2). Diabetes and Metabolism. 2019; 45(3):268-275
- 23 380. Wada T, Mori-Anai K, Takahashi A, Matsui T, Inagaki M, Iida M et al. Effect of canagliflozin on the decline of estimated glomerular filtration rate in chronic kidney disease patients with type 2 diabetes mellitus: A multicenter, randomized, double-blind, placebo-controlled, parallel-group, phase III study in Japan. Journal of Diabetes Investigation. 2022; 13(12):1981-1989
- 381. Wagner AM, Miranda-Calderin G, Ugarte-Lopetegui MA, Marrero-Santiago H,
 Suarez-Castellano L, Alberiche-Ruano MDP et al. Effect of liraglutide on physical
 performance in type 2 diabetes (LIPER2): A randomised, double-blind, controlled trial.
 Contemporary clinical trials communications. 2016; 4:46-51
- 382. Wang Q, Wang D, Cheng A, Sun FY, Li Z. Comparison between the effects of sitagliptin and liraglutide on blood glucose and cognitive function of patients with both type 2 diabetes mellitus and post-stroke mild cognitive impairment. Int J Clin Experimental Med. 2020; 13(2):1219-1227
- 383. Wang W, Nevarez L, Filippova E, Song Ki H, Tao B, Gu L et al. Efficacy and safety of once-weekly dulaglutide versus insulin glargine in mainly Asian patients with type 2 diabetes mellitus on metformin and/or a sulphonylurea: a 52-week open-label, randomized phase III trial. Diabetes Obes Metab. 2019; 21(2):234-243
- 40 384. Wang W, Ning G, Ma J, Liu X, Zheng S, Wu F et al. A randomized clinical trial of the safety and efficacy of sitagliptin in patients with type 2 diabetes mellitus inadequately controlled by acarbose alone. Current Medical Research and Opinion. 2017; 33(4):693-699
- 44 385. Wang W, Yan X, Cheng Z, Zhang Q, Wang R, Deng Y et al. Efficacy and safety of 45 adding once-weekly dulaglutide to basal insulin for inadequately controlled type 2 46 diabetes in Chinese patients (AWARD-CHN3): A randomized, double-blind, placebo-47 controlled, phase III trial. Diabetes, Obesity & Metabolism. 2023;

- Wang W, Yang J, Yang G, Gong Y, Patel S, Zhang C et al. Efficacy and safety of linagliptin in Asian patients with type 2 diabetes mellitus inadequately controlled by metformin: A multinational 24-week, randomized clinical trial. J Diabetes. 2016; 8(2):229-237
- 5 387. Wang X, Zhang H, Zhang Q, Guan M, Sheng S, Mo W et al. Exenatide and Renal Outcomes in Patients with Type 2 Diabetes and Diabetic Kidney Disease. American Journal of Nephrology. 2020; 51(10):806-814
- 8 388. Wang X, Zhao X, Gu Y, Zhu X, Yin T, Tang Z et al. Effects of Exenatide and Humalog 9 Mix25 on Fat Distribution, Insulin Sensitivity, and beta-Cell Function in Normal BMI 10 Patients with Type 2 Diabetes and Visceral Adiposity. Journal of Diabetes Research. 11 2020; 2020:9783859
- 389. Watada H, Kaneko S, Komatsu M, Agner BR, Nishida T, Ranthe M et al. Superior
 HbA1c control with the fixed-ratio combination of insulin degludec and liraglutide
 (IDegLira) compared with a maximum dose of 50 units of insulin degludec in
 Japanese individuals with type 2 diabetes in a phase 3, double-blind, randomized
 trial. Diabetes, Obesity & Metabolism. 2019; 21(12):2694-2703
- 390. Webb DR, Htike ZZ, Swarbrick DJ, Brady EM, Gray LJ, Biglands J et al. A
 randomized, open-label, active comparator trial assessing the effects of 26 weeks of
 liraglutide or sitagliptin on cardiovascular function in young obese adults with type 2
 diabetes. Diabetes Obes Metab. 2020; 22:1187-1196
- 391. White William B, Cannon Christopher P, Heller Simon R, Nissen Steven E,
 Bergenstal Richard M, Bakris George L et al. Alogliptin after acute coronary
 syndrome in patients with type 2 diabetes. The New England journal of medicine.
 2013; 369(14):1327-1335
- 392. Wilcox R, Kupfer S, Erdmann E, Proactive S, investigators. Effects of pioglitazone on major adverse cardiovascular events in high-risk patients with type 2 diabetes: results from PROspective pioglitAzone Clinical Trial In macro Vascular Events (PROactive 10). American Heart Journal. 2008; 155(4):712-717
- 393. Wilding JP, Woo V, Soler NG, Pahor A, Sugg J, Rohwedder K et al. Long-term
 efficacy of dapagliflozin in patients with type 2 diabetes mellitus receiving high doses
 of insulin: a randomized trial. Annals of Internal Medicine. 2012; 156(6):405-415
- 394. Wilding JPH, Charpentier G, Hollander P, Gonzalez-Galvez G, Mathieu C,
 Wercruysse F et al. Efficacy and safety of canagliflozin in patients with type 2 diabetes
 mellitus inadequately controlled with metformin and sulphonylurea: A randomised
 trial. International Journal of Clinical Practice. 2013; 67(12):1267-1282
- 36 395. Wilding JPH, Leonsson-Zachrisson M, Wessman C, Johnsson E. Dose-ranging study 37 with the glucokinase activator AZD1656 in patients with type 2 diabetes mellitus on 38 metformin. Diabetes, Obesity & Metabolism. 2013; 15(8):750-759
- 39 396. Wiviott Stephen D, Raz I, Bonaca Marc P, Mosenzon O, Kato Eri T, Cahn A et al.
 40 Dapagliflozin and Cardiovascular Outcomes in Type 2 Diabetes. The New England
 41 journal of medicine. 2019; 380(4):347-357
- 42 397. Wu S, Li X, Zhang H. Effects of metformin on endothelial function in type 2 diabetes. 43 Experimental and Therapeutic Medicine. 2014; 7(5):1349-1353
- 44 398. Wysham C, Blevins T, Arakaki R, Colon G, Garcia P, Atisso C et al. Efficacy and 45 safety of dulaglutide added onto pioglitazone and metformin versus exenatide in type 46 2 diabetes in a randomized controlled trial (AWARD-1). Diabetes Care. 2014; 47 37(8):2159-2167

- Xiao CC, Ren A, Yang J, Ye SD, Xing XN, Li SM et al. Effects of pioglitazone and glipizide on platelet function in patients with type 2 diabetes. European Review for Medical and Pharmacological Sciences. 2015; 19(6):963-970
- 4 400. Xiao X, Cui X, Zhang J, Han Z, Xiao Y, Chen N et al. Effects of sitagliptin as initial therapy in newly diagnosed elderly type 2 diabetics: A randomized controlled study. Experimental and Therapeutic Medicine. 2016; 12(5):3002-3008
- 7 401. Xu W, Mu Y, Zhao J, Zhu D, Ji Q, Zhou Z et al. Efficacy and safety of metformin and sitagliptin based triple antihyperglycemic therapy (STRATEGY): a multicenter, randomized, controlled, non-inferiority clinical trial. Sci China Life Sci. 2017; 60(3):225-238
- Yabe D, Nakamura J, Kaneto H, Deenadayalan S, Navarria A, Gislum M et al. Safety and efficacy of oral semaglutide versus dulaglutide in Japanese patients with type 2 diabetes (PIONEER 10): an open-label, randomised, active-controlled, phase 3a trial.
 The lancet Diabetes & endocrinology. 2020; 8(5):392-406
- Yabe D, Shiki K, Homma G, Meinicke T, Ogura Y, Seino Y. Efficacy and safety of the sodium-glucose co-transporter-2 inhibitor empagliflozin in elderly Japanese adults
 (>=65 years) with type 2 diabetes: A randomized, double-blind, placebo-controlled,
 52-week clinical trial (EMPA-ELDERLY). Diabetes, Obesity & Metabolism. 2023;
- 404. Yale JF, Bakris G, Cariou B, Yue D, David-Neto E, Xi L et al. Efficacy and safety of canagliflozin in subjects with type 2 diabetes and chronic kidney disease. Diabetes Obes Metab. 2013; 15(5):463-473
- Yan J, Yao B, Kuang H, Yang X, Huang Q, Hong T et al. Liraglutide, Sitagliptin, and
 Insulin Glargine Added to Metformin: The Effect on Body Weight and Intrahepatic
 Lipid in Patients With Type 2 Diabetes Mellitus and Nonalcoholic Fatty Liver Disease.
 Hepatology. 2019; 69(6):2414-2426
- Yang HK, Min KW, Park SW, Chung CH, Park KS, Choi SH et al. A randomized,
 placebo-controlled, double-blind, phase 3 trial to evaluate the efficacy and safety of
 anagliptin in drug-na?ve patients with type 2 diabetes. Endocrine Journal. 2015;
 62(5):449-462
- Yang S, Zhao L, Mi Y, He W. Effects of sodium-glucose cotransporter-2 inhibitors and aldosterone antagonists, in addition to renin-angiotensin system antagonists, on major adverse kidney outcomes in patients with type 2 diabetes and chronic kidney disease: A systematic review and network meta-analysis. Diabetes, Obesity & Metabolism. 2022; 24(11):2159-2168
- 408. Yang W, Guan Y, Shentu Y, Li Z, Johnson-Levonas AO, Engel SS et al. The addition of sitagliptin to ongoing metformin therapy significantly improves glycemic control in Chinese patients with type 2 diabetes. J Diabetes. 2012; 4(3):227-237
- 38 409. Yang W, Han P, Min KW, Wang B, Mansfield T, T'Joen C et al. Efficacy and safety of dapagliflozin in Asian patients with type 2 diabetes after metformin failure: A randomized controlled trial. J Diabetes. 2016; 8(6):796-808
- 41 410. Yang W, Ma J, Li Y, Li Y, Zhou Z, Kim JH et al. Dapagliflozin as add-on therapy in 42 Asian patients with type 2 diabetes inadequately controlled on insulin with or without 43 oral antihyperglycemic drugs: A randomized controlled trial. J Diabetes. 2018; 44 10(7):589-599
- 45 411. Yang W, Min K, Zhou Z, Li L, Xu X, Zhu D et al. Efficacy and safety of lixisenatide in a predominantly Asian population with type 2 diabetes insufficiently controlled with

- basal insulin: the GetGoal-L-C randomized trial. Diab Obes Metab. 2018; 20(2):335-
- 412. Yang W, Pan CY, Tou C, Zhao J, Gause-Nilsson I. Efficacy and safety of saxagliptin added to metformin in Asian people with type 2 diabetes mellitus: a randomized controlled trial. Diabetes Research and Clinical Practice. 2011; 94(2):217-224
- 413. Yang W, Xu X, Lei T, Ma J, Li L, Shen J et al. Efficacy and safety of linagliptin as addon therapy to insulin in Chinese patients with type 2 diabetes mellitus: A randomized, double-blind, placebo-controlled trial. Diabetes, Obesity & Metabolism. 2021; 23(2):642-647
- 414. Yki-J?rvinen H, Rosenstock J, Dur?n-Garcia S, Pinnetti S, Bhattacharya S, Thiemann
 S et al. Effects of adding linagliptin to basal insulin regimen for inadequately
 controlled type 2 diabetes: a ?52-week randomized, double-blind study. Diabetes
 Care. 2013; 36(12):3875-3881
- Yokoyama H, Hirao K, Yamaguchi K, Oishi M, Lee G, Yagi N et al. Liraglutide Versus Sitagliptin in a 24-week, Multicenter, Open-label, Randomized, Parallel-group Study in Japanese Type 2 Diabetes Mellitus Patients Responding Inadequately to a Sulfonylurea and/or One or Two Other Oral Antidiabetic Drugs (JDDM 33). Japanese clinical medicine. 2014; 5:33-41
- 416. Yuan X, Guo X, Zhang J, Dong X, Lu Y, Pang W et al. Improved glycaemic control and weight benefit with iGlarLixi versus insulin glargine 100 U/mL in Chinese people with type 2 diabetes advancing their therapy from basal insulin plus oral antihyperglycaemic drugs: Results from the LixiLan-L-CN randomized controlled trial. Diabetes, Obesity and Metabolism. 2022; 24(11):2182-2191
- 24 417. Zang L, Liu Y, Geng J, Luo Y, Bian F, Lv X et al. Efficacy and safety of liraglutide
 25 versus sitagliptin, both in combination with metformin, in Chinese patients with type 2
 26 diabetes: a 26-week, open-label, randomized, active comparator clinical trial.
 27 Diabetes Obes Metab. 2016; 18(8):803-811
- 28 418. Zhang J, Xian TZ, Wu MX, Li C, Pan Q, Guo LX. Comparison of the effects of twice-29 daily exenatide and insulin on carotid intima-media thickness in type 2 diabetes 30 mellitus patients: a 52-week randomized, open-label, controlled trial. Cardiovascular 31 Diabetology. 2020; 19(1):48
- 32 419. Zhao L, Sun T, Wang L. Chitosan oligosaccharide improves the therapeutic efficacy
 33 of sitagliptin for the therapy of Chinese elderly patients with type 2 diabetes mellitus.
 34 Therapeutics and Clinical Risk Management. 2017; 13:739-750
- Zinman B, Aroda VR, Buse JB, Cariou B, Harris SB, Hoff ST et al. Efficacy, Safety,
 and Tolerability of Oral Semaglutide Versus Placebo Added to Insulin With or Without
 Metformin in Patients With Type 2 Diabetes: The PIONEER 8 Trial. Diabetes Care.
 2019; 42(12):2262-2271
- Zinman B, Bhosekar V, Busch R, Holst I, Ludvik B, Thielke D et al. Semaglutide once
 weekly as add-on to SGLT-2 inhibitor therapy in type 2 diabetes (SUSTAIN 9): a
 randomised, placebo-controlled trial. Lancet Diabetes Endocrinol. 2019; 7(5):356-367
- 42 Zinman B, Gerich J, Buse JB, Lewin A, Schwartz S, Raskin P et al. Efficacy and safety of the human glucagon-like peptide-1 analog liraglutide in combination with metformin and thiazolidinedione in patients with type 2 diabetes (LEAD-4 Met+TZD). Diabetes Care. 2009; 32(7):1224-1230

1

2

4

423. Zinman B, Wanner C, Lachin John M, Fitchett D, Bluhmki E, Hantel S et al. Empagliflozin, Cardiovascular Outcomes, and Mortality in Type 2 Diabetes. The New England journal of medicine. 2015; 373(22):2117-2128