

Thyroid disease: assessment and management

[I] Management of Thyrotoxicosis: Drugs vs Surgery vs Radioactive Iodine

NICE guideline

Intervention evidence review underpinning recommendations 1.6.1 to 1.6.24 in the guideline. See also evidence reviews J, K, L and D

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*This evidence review was developed by
the National Guideline Centre*

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1 Management of thyrotoxicosis: drugs vs surgery vs radioactive iodine

1.1 Review question: What is the clinical and cost effectiveness of using radioactive iodine vs antithyroid drugs vs surgery to treat thyrotoxicosis secondary to Graves' disease?

Review question: What is the clinical and cost effectiveness of using radioactive iodine vs surgery to treat thyrotoxicosis secondary to toxic nodular goitre?

1.2 Introduction

The three principal treatment modalities when managing the patient with thyrotoxicosis are medical therapy with antithyroid drug (ATD), radioactive iodine or surgery. There is uncertainty in terms of how these modalities are best used in relation to the type and dose of antithyroid drug, the dose of radioactive iodine and the nature of the surgical procedure (partial or total thyroidectomy). The aetiology of thyrotoxicosis (such as Graves' disease, toxic nodular goitre, toxic nodule or thyroiditis), the age of the patient, the other patient factors (such as pregnancy or planned pregnancy and small children at home) and the presence of complicating factors such as thyroid eye disease are additional considerations. Although many patients with Graves' thyrotoxicosis are managed with ATD (carbimazole or propylthiouracil) initially, a majority will relapse and become thyrotoxic again when the drug is stopped. Patients will then be faced with the prospect of long term ATD therapy or choosing radioactive iodine or surgery – both of which will usually result in hypothyroidism and a requirement for life-long thyroid hormone replacement.

Radioactive iodine has been used to treat thyrotoxicosis for many years. The attractions of this therapy include the fact that it is relatively cheap. Administration is straight-forward although guidelines that limit exposure to ionising radiation need to be followed when using radioactive agents and there is variation between centres in terms of when this modality is considered to be an appropriate therapeutic option. Some units are more proactive than others and consider this treatment more readily in the context of the younger patient and the individual with complicating factors such as thyroid eye disease. Establishing the circumstances and threshold for using this treatment is an important area because the therapeutic options for patients who fail to respond to anti-thyroid drug are limited.

1.3 PICO table

For full details see the review protocol in Appendix A:.

Table 1: PICO characteristics of review question

Population	People diagnosed with thyrotoxicosis
Interventions	Antithyroid drugs Radioactive iodine Surgery
Comparison	Any of the above compared with any other
Outcomes	Critical

	<ul style="list-style-type: none"> • Mortality • Quality of life <p>Important (general)</p> <ul style="list-style-type: none"> • Thyroid ophthalmopathy • Euthyroidism • Hypothyroidism • Relapse of hyperthyroidism • Cardiovascular morbidity • Arrhythmia • Osteoporosis • Cognitive impairment • Pain • Symptom scores • Patient/family/carer experience • Healthcare contacts <p>Important (surgical)</p> <ul style="list-style-type: none"> • Recurrent laryngeal nerve damage • Hypocalcaemia • Hypoparathyroidism • Bleeding • Infection <p>Important (pharmacological)</p> <ul style="list-style-type: none"> • Agranulocytosis • Liver failure • Minor drug related adverse effects • Teratogenesis <p>Important (radioactive iodine)</p> <ul style="list-style-type: none"> • Infertility • Malignancy • Thyrotoxic storm • Growth abnormalities • Hypocalcaemia • Hypoparathyroidism • Teratogenesis
Study design	<p>RCTs only, non-randomised studies only if key confounders (age, co-existing conditions, baseline thyroid hormones) taken into account</p> <p>Minimum duration 3 months</p>

1 1.4 Clinical evidence

2 1.4.1 Included studies

3 Six randomised controlled studies (in nine publications) were included in the review;^{1, 3, 11, 16,}
 4 28, 70, 124, 127, 128 these are summarised in Table 2 below. Evidence from these studies is
 5 summarised in the clinical evidence summary below (Table 3). One Cochrane review in this
 6 area was identified,⁸¹ the studies included in this review were checked against the protocol
 7 and included as appropriate.

8 Five studies compared antithyroid drugs vs radioactive iodine. One study compared
 9 antithyroid drugs vs radioactive iodine vs surgery.

1 Five studies were in the treatment naïve population (or previous treatment unspecified). One
2 study was in people who had previously used antithyroid drugs and relapsed.

3 No studies were found in children or older adults.

4 All six studies were either exclusively in people with Graves' disease or in a mixed population
5 in which the majority had Graves' disease. No studies were found exclusively in people with
6 toxic nodular goitre.

7 See also the study selection flow chart in Appendix C:, study evidence tables in Appendix D:,
8 forest plots in Appendix E: and GRADE tables in Appendix F:.

9 **1.4.2 Excluded studies**

10 See the excluded studies list in Appendix J:.

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1.4.3 Summary of clinical studies included in the evidence review

Table 2: Summary of studies included in the evidence review

Study	Intervention and comparison	Population	Outcomes	Comments
Azizi 2005 ¹¹	<p>Antithyroid drugs, n = 52 MMI, 10mg twice daily for first month, once daily for second month, 2.5-10mg daily thereafter, no discontinuation specified</p> <p>Radioactive iodine , n = 52 Calculated activity, no information on number of treatments</p>	<p>Adults (mean age 48, SD 6)</p> <p>Second line (relapsed 1 year after 18 months of antithyroid drug use)</p> <p>Graves' disease</p> <p>Percent with ophthalmopathy at baseline not specified</p> <p>Iran</p>	<p>Euthyroidism (at end of follow-up)</p> <p>Hypothyroidism (at end of follow-up)</p> <p>Hyperthyroidism (at end of follow-up)</p> <p>Agranulocytosis</p> <p>10 year follow-up</p>	<p>In all patients dosage of MMI/levothyroxine adjusted to maintain normal thyroid function</p> <p>No discontinuation period specified for ATDs</p>
Bartalena 1998 ¹⁶	<p>Antithyroid drugs, n = 148 Lowest dose that maintained euthyroidism, no discontinuation specified</p> <p>Radioactive iodine , n = 150 MMI given for 3 to 4 months prior to RAI, stopped 5 days before, dose of 120-150uCi per gram of thyroid tissue, if hypo/hyperthyroid after RAI corrected with levothyroxine or MMI as relevant, second dose of RAI at end of follow-up if persistent hyperthyroidism still</p>	<p>Adults (mean age 42, range 15-85)</p> <p>70% had received MMI prior to referral</p> <p>Graves' disease</p> <p>~50% with ophthalmopathy (mild) at baseline</p> <p>Italy</p>	<p>Ophthalmopathy (new/worsening)</p> <p>Euthyroidism (at end of follow-up)</p> <p>Hypothyroidism (at end of follow-up)</p> <p>Hyperthyroidism (at end of follow-up)</p> <p>1 year follow-up (no discontinuation period for ATDs)</p>	<p>Radioactive iodine arm given levothyroxine/MMI if required</p> <p>No discontinuation period for ATDs</p>
Chen 2009 ²⁸	Antithyroid drugs, n = 230	Adults (mean age 37, SD 14)	Mortality	No information on drug

Study	Intervention and comparison	Population	Outcomes	Comments
	<p>Either MMI or PTU, at least 18 months of treatment, dose based on severity of symptoms and TSH</p> <p>Radioactive iodine , n = 230 Calculated activity, no pre-treatment with ATDs, at 3 months could have 2nd treatment (10% participants) an at 6 months could have 3rd (2.5% participants)</p>	<p>Treatment naïve</p> <p>Mixed cause (75% Graves', 25% toxic nodular goitre)</p> <p>~25% with ophthalmopathy at baseline</p> <p>China</p>	<p>Ophthalmopathy (incidence)</p> <p>Euthyroidism (at end of follow-up)</p> <p>Hypothyroidism (at end of follow-up)</p> <p>Hyperthyroidism</p> <p>Agranulocytosis</p> <p>Severe liver damage</p> <p>Malignancy</p> <p>Thyroid storm</p> <p>9 year follow-up</p>	<p>supplementation of radioactive iodine arm</p> <p>Discontinuation period for ATDs</p>
Kansara 2017 ⁷⁰	<p>Antithyroid drugs, n = 30 CZL, 30mg initially and then tapered</p> <p>Radioactive iodine , n = 30 Single oral dose of 10mCi, no stated pre-treatment with ATDs</p>	<p>Adults (mean age 33, SD 4.2)</p> <p>Treatment naïve</p> <p>Mixed cause (85% Graves', 15% toxic nodular goitre)</p> <p>Percent with ophthalmopathy at baseline not specified</p> <p>India</p>	<p>Euthyroidism (at end of follow-up)</p> <p>Hypothyroidism (at end of follow-up)</p> <p>Hyperthyroidism (at end of follow-up)</p> <p>1 year follow-up</p>	No discontinuation period for ATDs
Torrington 1996 ^{1, 124, 127}	<p>Antithyroid drugs, n = 71 MMI, 18 months of treatment, block and replace</p> <p>Radioactive iodine , n = 39 Oral dose, calculated activity,</p>	<p>Adults (younger adults mean age 29, SD 4, older adults mean 45, SD 6)</p> <p>Previous treatment not specified</p>	<p>Ophthalmopathy (new/worsening)</p> <p>Hyperthyroidism</p> <p>Recurrent laryngeal nerve damage</p> <p>Hypoparathyroidism</p>	<p>Study stratified by age group, older adults (35-55) randomised to all 3 treatments, younger adults (20-34) only randomised to antithyroid drugs or surgery.</p> <p>Evidence combined across age</p>

Study	Intervention and comparison	Population	Outcomes	Comments
	(~50% of participants required more than one dose, given >10 weeks after first) Surgery, n = 64 Bilateral subtotal thyroidectomy with posterior capsule and 1g or less of each lobe left behind, thyroxine after surgery	Graves' disease ~13% with ophthalmopathy at baseline (non-severe) Sweden	Agranulocytosis Maximum 21 year follow-up	groups as per protocol, except where this would affect group age composition (i.e. not comparing older adults receiving radioactive iodine with mix of young and old adults receiving antithyroid drugs) Discontinuation of ATDs
Träisk 2009 ^{3, 128}	Antithyroid drugs, n = 150 MMI, 18 months of treatment, block and replace Radioactive iodine, n = 163 Oral outpatient dose, calculated activity, no information on number of doses, levothyroxine substitution as required, no prophylactic steroid use	Adults (mean 51, SD 8) Treatment naïve Graves' disease ~13% with ophthalmopathy at baseline (non-severe) Sweden	Ophthalmopathy (new/worsening) Hyperthyroidism (relapse) 4 years follow-up	Radioactive iodine arm given levothyroxine if required Discontinuation of ATDs

See Appendix D: for full evidence tables.

1.4.4 Quality assessment of clinical studies included in the evidence review

Table 3: Clinical evidence summary: Radioactive iodine vs antithyroid drugs, Graves' disease, first line treatment

Outcomes	No of Participants (studies) Follow up	Quality of the evidence (GRADE)	Relative effect (95% CI)	Anticipated absolute effects	
				Risk with ATD	Risk difference with RAI (95% CI)
Mortality	386 (1 study)	⊕⊕⊖⊖ LOW1,2	Not estimable	0 per 1000	not estimable ⁵

Outcomes	No of Participants (studies) Follow up	Quality of the evidence (GRADE)	Relative effect (95% CI)	Anticipated absolute effects	
				Risk with ATD	Risk difference with RAI (95% CI)
	9 years	due to risk of bias, imprecision			
Ophthalmopathy (new/worsening cases)	948 (4 studies) 1-9 years	⊕⊕⊕⊖ MODERATE ¹ due to risk of bias	RR 2.17 (1.64 to 2.88)	103 per 1000	121 more per 1000 (from 66 more to 194 more)
Euthyroidism (at end of follow-up)	741 (3 studies) 1-9 years	⊕⊖⊖⊖ VERY LOW ^{1,3} due to risk of bias, inconsistency	RR 0.78 (0.37 to 1.62)	759 per 1000	167 fewer per 1000 (from 478 fewer to 471 more)
Hypothyroidism (at end of follow-up)	741 (3 studies) 1-9 years	⊕⊕⊕⊖ MODERATE ¹ due to risk of bias	RR 5.89 (3.12 to 11.11)	34 per 1000	166 more per 1000 (from 72 more to 344 more)
Hyperthyroidism (persistence/recurrence)	1102 (5 studies) 1-9 years	⊕⊕⊖⊖ LOW ^{1,3} due to risk of bias, inconsistency	RR 0.25 (0.09 to 0.69)	241 per 1000	181 fewer per 1000 (from 75 fewer to 219 fewer)
Osteoporosis	70 (1 study) 14-21 years	⊕⊖⊖⊖ VERY LOW ^{1,4} due to risk of bias, imprecision	RR 1.27 (0.43 to 3.78)	139 per 1000	38 more per 1000 (from 79 fewer to 386 more)
Agranulocytosis	423 (1 study) 9 years	⊕⊕⊖⊖ LOW ^{1,2} due to risk of bias, imprecision	Peto OR 0.13 (0.03 to 0.6)	33 per 1000	29 fewer per 1000 (from 13 fewer to 32 fewer)
Severe liver damage	423 (1 study) 9 years	⊕⊕⊖⊖ LOW ^{1,4} due to risk of bias, imprecision	Peto OR 0.14 (0.02 to 0.79)	23 per 1000	20 fewer per 1000 (from 5 fewer to 23 fewer)

Outcomes	No of Participants (studies) Follow up	Quality of the evidence (GRADE)	Relative effect (95% CI)	Anticipated absolute effects	
				Risk with ATD	Risk difference with RAI (95% CI)
	(1 study) 9 years	LOW ^{1,2} due to risk of bias, imprecision	estimable	0 per 1000	not estimable ⁵
Thyroid storm	386 (1 study) 9 years	⊕⊕⊖⊖ LOW ^{1,2} due to risk of bias, imprecision	Not estimable	0 per 1000	not estimable ⁵

1 Downgraded by 1 increment if the majority of the evidence was at high risk of bias, and downgraded by 2 increments if the majority of the evidence was at very high risk of bias
2 Downgraded by 1 increment as zero events in at least one arm
3 Downgraded by 1 or 2 increments because the point estimate and or the confidence intervals varied widely across studies, unexplained by subgroup analysis
4 Downgraded by 1 increment if the confidence interval crossed one MID or by 2 increments if the confidence interval crossed both MIDs
5 Zero events in both arms

Table 4: Clinical evidence summary: Surgery vs antithyroid drugs, Graves' disease, first line treatment

Outcomes	No of Participants (studies) Follow up	Quality of the evidence (GRADE)	Relative effect (95% CI)	Anticipated absolute effects	
				Risk with ATD	Risk difference with SUR (95% CI)
Ophthalmopathy (new/worsening cases)	129 (1 study) 4 years	⊕⊕⊖⊖ LOW ¹ due to imprecision	RR 1.14 (0.47 to 2.78)	123 per 1000	17 more per 1000 (from 65 fewer to 219 more)
Osteoporosis	111 (1 study) 14-21 years	⊕⊖⊖⊖ VERY LOW ^{1,2} due to risk of bias, imprecision	RR 1.57 (0.55 to 4.51)	91 per 1000	52 more per 1000 (from 41 fewer to 319 more)
Hyperthyroidism (persistence/recurrence)	133 (1 study) 4 years	⊕⊕⊕⊕ HIGH	RR 0.16 (0.06 to 0.44)	382 per 1000	321 fewer per 1000 (from 214 fewer to 359 fewer)

1 Downgraded by 1 increment if the confidence interval crossed one MID or by 2 increments if the confidence interval crossed both MIDs
2 Downgraded by 1 increment if the majority of the evidence was at high risk of bias, and downgraded by 2 increments if the majority of the evidence was

Outcomes	No of Participants (studies) Follow up	Quality of the evidence (GRADE)	Relative effect (95% CI)	Anticipated absolute effects	
				Risk with ATD	Risk difference with SUR (95% CI)
at very high risk of bias					

Table 5: Clinical evidence summary: Radioactive iodine vs surgery, Graves' disease, first line treatment

Outcomes	No of Participants (studies) Follow up	Quality of the evidence (GRADE)	Relative effect (95% CI)	Anticipated absolute effects	
				Risk with SUR	Risk difference with RAI (95% CI)
Ophthalmopathy (new/worsening cases)	76 (1 study) 4 years	⊕⊕⊕⊖ MODERATE ¹ due to imprecision	RR 2.06 (0.87 to 4.84)	162 per 1000	172 more per 1000 (from 21 fewer to 622 more)
Osteoporosis	68 (1 study) 14-21 years	⊕⊖⊖⊖ VERY LOW ^{1,2} due to risk of bias, imprecision	RR 0.86 (0.32 to 2.29)	206 per 1000	29 fewer per 1000 (from 140 fewer to 266 more)
Hyperthyroidism (persistence/recurrence)	76 (1 study) 4 years	⊕⊕⊖⊖ LOW ¹ due to imprecision	RR 2.53 (0.73 to 8.82)	81 per 1000	124 more per 1000 (from 22 fewer to 633 more)

1 Downgraded by 1 increment if the confidence interval crossed one MID or by 2 increments if the confidence interval crossed both MIDs
2 Downgraded by 1 increment if the majority of the evidence was at high risk of bias, and downgraded by 2 increments if the majority of the evidence was at very high risk of bias

Table 6: Clinical evidence summary: Radioactive iodine vs antithyroid drugs, Graves' disease, second line treatment

Outcomes	No of Participants (studies) Follow up	Quality of the evidence (GRADE)	Relative effect (95% CI)	Anticipated absolute effects	
				Risk with ATD	Risk difference with RAI (95% CI)

Outcomes	No of Participants (studies) Follow up	Quality of the evidence (GRADE)	Relative effect (95% CI)	Anticipated absolute effects	
				Risk with ATD	Risk difference with RAI (95% CI)
(at end of follow-up)	(1 study) 10 years	LOW1 due to risk of bias	(0.28 to 0.62)	929 per 1000	539 fewer per 1000 (from 353 fewer to 669 fewer)
Hypothyroidism (at end of follow-up)	69 (1 study) 10 years	⊕⊕⊖⊖ LOW1 due to risk of bias	RR 17.07 (2.45 to 118.83)	36 per 1000	579 more per 1000 (from 52 more to 1000 more)
Hyperthyroidism (at end of follow-up)	69 (1 study) 10 years	⊕⊖⊖⊖ VERY LOW1 due to risk of bias, imprecision	Peto OR 0.09 (0 to 4.6)	36 per 1000	33 fewer per 1000 (from 36 fewer to 111 more)
Agranulocytosis	69 (1 study) 10 years	⊕⊖⊖⊖ VERY LOW1,2 due to risk of bias, imprecision	Not estimable		Not estimable ³

1 Downgraded by 1 increment if the majority of the evidence was at high risk of bias, and downgraded by 2 increments if the majority of the evidence was at very high risk of bias
2 Downgraded by 1 increment as at least one arm with zero events
3 Zero events in both arms

See Appendix F: for full GRADE tables.

1 **1.5 Economic evidence**

2 **1.5.1 Included studies**

3 **Management of thyrotoxicosis secondary to Graves' disease**

4 One health economic study was identified with the relevant comparison and has been
5 included in this review ³⁵. This is summarised in the health economic evidence profile below
6 (Table 7) and the health economic evidence tables in Appendix H:

7 **Management of thyrotoxicosis secondary to toxic nodular goitre**

8 No relevant health economic studies were identified.

9 **1.5.2 Excluded studies**

10 One health economic study that was relevant to this question was excluded due to
11 assessment of limited applicability.

12 See also the health economic study selection flow chart in Appendix G:.

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1.5.3 Summary of studies included in the economic evidence review

Table 7: Health economic evidence profile: Radioactive iodine vs anti-thyroid drugs vs surgery for Graves' disease

Study	Applicability	Limitations	Other comments	Incremental cost ^(a)	Incremental effects ^(a)	Cost effectiveness	Uncertainty
Donovan, 2016 ³⁵ (UK and Australia) (a)	Directly applicable ^(b)	Minor limitations ^(c)	<ul style="list-style-type: none"> • Cost utility analysis • Life time horizon • Patients received either; <ol style="list-style-type: none"> 1. Radioactive iodine (RAI) 2. Anti-thyroid drugs (ATD) 3. Total thyroidectomy (TT). 	Mean per patient: (2-1): £11,441 (3-1): £1,690 (3-2): saves £9,751	Mean per patient: (2-1): 0.44 QALYs (3-1): -0.8 QALYs (3-2): -1.24 QALYs	RAI dominated TT (less costly and more effective). ATD was not cost effective compared to RAI at the £20,000 threshold. (ICER for ATD vs RAI = £26,279 per QALY gained)	RAI was dominant over TT in all sensitivity analysis of all parameters assessed. ATD was a cost-effective alternative to RAI at the £30,000 threshold (ICER: £26,279 per QALY gained).

Abbreviations: ICER: incremental cost-effectiveness ratio; QALY: quality-adjusted life years

(a) The results presented are those of the UK analysis only.

(b) No downgrading for applicability.

(c) The estimates of relative treatment effects are not based on met-analysis of all the available evidence. Some costs have been based on the national tariff and maybe overestimated. The model has not been run probabilistically, to adequately assess parameter uncertainty.

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1 **1.5.4 Health economic modelling**

2 This area was not prioritised for new cost-effectiveness analysis.

3 **1.5.5 Resource costs**

4 Relevant unit costs are provided below to aid consideration of cost effectiveness for the
 5 management of thyrotoxicosis secondary to toxic nodular goitre.

6 **Table 8: UK costs of thyroid surgery and radioactive iodine**

Intervention	Unit cost	
Surgery (Thyroid Procedures with CC Score 0-4+)(a)	£3,689	
Radioactive iodine fixed dose(b)	£286.32	
Radioactive iodine calculated dose (c)	Procedures (pre and post therapy)	Unit costs
	Uptake measurement with probe ~15 mins Band 7	£10 (d)
	USS for volume calculation	£62
	Calculations, verification, report: ~ 3 hours Band 7	£75
	Total additional cost to fixed dose	£167 (e)

7 Source: NHS reference costs 2016-17, total HRG schedule ³⁴.

8 (a) Weighted average of all 3 combined thyroid procedures with CC scores 0-1, 2-3, 4+(KA09C, KA09D, KA09E)
 9 including excess bed days with an average length of stay of 1.6 days

10 (b) Cost of oral delivery of radiotherapy for thyroid ablation, cost code RN51Z

11 (c) Estimation obtained from committee specialists

12 (d) Ideally allow for 3 uptake measurements (adding another £20), practice varies

13 (e) Total cost = £453.32 Economic considerations: trade-off between net clinical effects and costs
 14

15 **1.6 Evidence statements**

16 **1.6.1 Clinical evidence statements**

17 **1.6.1.1 Radioactive iodine vs antithyroid drugs, Graves' disease, first line treatment**

18 No clinically important difference was identified for mortality (1 study, low quality),
 19 osteoporosis (1 study, very low quality), agranulocytosis (1 study, low quality), severe liver
 20 damage (1 study, low quality), malignancy (1 study, low quality), thyroid storm (1 study, low
 21 quality).

22 There was a clinically important benefit of radioactive iodine for hyperthyroidism (5 studies,
 23 low quality).

24 There was a clinically important harm of radioactive iodine for ophthalmopathy (4 studies,
 25 moderate quality), euthyroidism (3 studies, very low quality) and hypothyroidism (3 studies,
 26 moderate quality).

27 No evidence was identified for other outcomes.

1 **1.6.1.2 Surgery vs antithyroid drugs, Graves' disease, first line treatment**

2 No clinically important difference was identified for ophthalmopathy (1 study, low quality),
3 osteoporosis (1 study, very low quality).

4 There was a clinically important benefit of surgery for hyperthyroidism (1 study, high quality).

5 No evidence was identified for other outcomes.

6 **1.6.1.3 Radioactive iodine vs surgery, Graves' disease, first line treatment**

7 No clinically important difference was identified for osteoporosis (1 study, very low quality).

8 There was a clinically important harm of radioactive iodine for ophthalmopathy (1 study,
9 moderate quality) and hyperthyroidism (1 study, low quality).

10 No evidence was identified for other outcomes.

11 **1.6.1.4 Radioactive iodine vs antithyroid drugs, Graves' disease, second line treatment**

12 No clinically important difference was identified for hyperthyroidism (1 study, very low
13 quality), agranulocytosis (1 study, very low quality).

14 There was a clinically important harm of radioactive iodine for euthyroidism (1 study, low
15 quality) and hypothyroidism (1 study, low quality).

16 No evidence was identified for other outcomes.

17 **1.6.2 Health economic evidence statements**

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19 **Management of thyrotoxicosis secondary to Graves' disease**

20 One cost–utility analysis found that anti-thyroid drugs were not cost effective at a threshold of
21 £20,000 per QALY, compared to radioactive iodine for treating thyrotoxicosis secondary to
22 Graves' disease (ICER: £26,279 per QALY gained compared to radioactive iodine). It also
23 found that radioactive iodine was dominant (less costly and more effective) compared to total
24 thyroidectomy. This analysis was assessed as directly applicable with minor limitations.

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26 **Management of thyrotoxicosis secondary to toxic nodular goitre**

27 • No relevant economic evaluations were identified.

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2 Radioactive iodine safety

2.1 Review question: What are the long term adverse events of radioactive iodine treatment for thyrotoxicosis?

2.2 Introduction

Radioactive iodine has been used to treat thyrotoxicosis for many years. The attractions of this therapy include the fact that it is relatively cheap. Administration is straight-forward although guidelines that limit exposure to ionising radiation need to be followed when using radioactive agents and there is variation between centres in terms of when this modality is considered to be an appropriate therapeutic option. There are concerns about the potential long-term risk of developing cancer because of exposure to radiation and the impact of radiation on fertility. The purpose of his review is to establish the level of risk radiation on these outcomes.

2.3 PICO table

For full details see the review protocol in Appendix A:.

Table 9: PICO characteristics of review question

Population	People being treated with radioactive iodine for thyrotoxicosis
Intervention	Radioactive iodine
Comparisons	Antithyroid drug treatment of thyrotoxicosis Surgical treatment of thyrotoxicosis Healthy controls
Outcomes	Cancer <ul style="list-style-type: none">• Overall diagnoses• Diagnoses in organs that take up iodine (e.g. thyroid, small bowel)• Diagnoses in organs that do not take up iodine• Infertility
Study design	Only studies with follow-up >5 years and sample size >1000 (for adults) will be included Evidence will be considered according to the following hierarchy: <ul style="list-style-type: none">• Comparative studies with hyperthyroid controls and adequate adjustment for key confounders (age, smoking)• Comparative studies with hyperthyroid controls and without adequate adjustment for key confounders• Comparative studies with healthy controls and adequate adjustment for key confounders (age, smoking)• Comparative studies with healthy controls without adequate adjustment for key confounders

2.4 Clinical evidence

2.4.1 Included studies

Eight studies were included in the review,^{38, 40, 42, 44, 59, 62, 90, 115} these are summarised below. Evidence from these studies is summarised in the clinical evidence summary below (Table

1 3). All studies were non-randomised comparisons in adults. Three studies compared
2 radioactive iodine with thyroidectomy and the remaining five studies compared people
3 treated with radioactive iodine, with the general population.

4 Where there were studies assessing the same cohort these were handled so as to minimise
5 double counting within the same meta-analysis.

6 Death from cancer was considered a surrogate outcome for cancer diagnoses and was
7 extracted if diagnoses of cancer was not available for that cohort comparison. This outcome
8 was downgraded for indirectness.

9 **2.4.2 Excluded studies**

10 See the excluded studies list in Appendix J:.

11

12

2.4.3 Summary of clinical studies included in the evidence review

Table 10: Summary of studies included in the evidence review

Study	Intervention and comparison	Population	Outcomes	Comments
Franklyn 1999 ³⁸	Radioactive iodine, n = 7417 Fixed dose, mean 308Mbq (SD 232), 84.9% received only one dose Age, sex and year matched SIR	People with hyperthyroidism, treated in West Midlands with radioactive iodine (mean age at treatment 56.6) Cohort treated between 1950-1991 UK	Overall cancer incidence Site specific cancer incidence Mean follow-up 9.7 years,	
Franklyn 2005 ⁴⁰	Radioactive iodine, n = 2668 Fixed dose either 185 or 370MBq, 84.3% received one dose only Age, sex and year matched SMRs	People with hyperthyroidism, treated in West Midlands with radioactive iodine (median age at treatment start 62) Cohort treated between 1984-2002 UK	Cancer mortality Median follow-up 5.6 years	542 person overlap with Franklyn 1999, outcome different
Giesecke 2018 ⁴²	Radioactive iodine, n = 10250 Dose not specified Thyroidectomy, n = 742 Surgery not specified	People with hyperthyroidism (mean age of RAI group 64, mean of surgery group 47) Cohort treated between 1976-2013 Sweden	Cancer mortality Mean follow-up 16.3 years	Adjusted for potential confounders in regression of age at treatment, gender, year of treatment, aetiology, co-existing conditions
Goldman 1988 ⁴⁴	Radioactive iodine, n = 1762 Dose not specified	Women with hyperthyroidism (age not stated) treated with	Overall cancer incidence Site specific cancer	

Study	Intervention and comparison	Population	Outcomes	Comments
	Age, sex, race, year matched SIRs for Connecticut	RAI at Mass. Gen. Hospital Cohort treated between 1946 and 1964 USA	incidence Mean follow-up 17.2 years	
Hoffman 1982 ⁵⁹	Radioactive iodine, n = 1005 Mean dose 10.6mCi (~392MBq), mean number of doses 1.2 Thyroidectomy, n = 2141 Surgery not specified	White women with hyperthyroidism treated by Mayo clinic (mean age of RAI group at Tx 56.8, surgery 45.7) Cohort treated between 1946 and 1964 USA	Overall cancer incidence Site specific cancer incidence Mean follow-up 15 years for RAI group, 21 years for surgical group	Adjusted for age, year of treatment, duration of follow-up
Holm 1991 ⁶²	Radioactive iodine, n = 10207 Mean dose 506MBq Age, sex, region, year matched incidence for whole of Sweden	People with hyperthyroidism (mean age 57, range 13-74) Cohort treated between 1950 and 1975 Sweden	Overall cancer incidence Site specific cancer incidence Mean follow-up 15 years	
Metso 2007 ⁹⁰	Radioactive iodine, n = 2793 Mean dose 305MBq Age, sex matched control from Finnish population register	People with hyperthyroidism treated with RAI at Tampere hospital (median age 62 years) Cohort treated between 1965 and 2002 Finland	Overall cancer incidence Site specific cancer incidence Mean follow-up 9.8 years for patients and 10.0 years for controls	
Ryodi 2015 ¹¹⁵	Radioactive iodine, n = 1814	People with hyperthyroidism	All cancer diagnoses	Unspecified overlap with Metso

Study	Intervention and comparison	Population	Outcomes	Comments
	Dose not specified Thyroidectomy, n = 4334 Surgery not specified	(median age of radioactive iodine group 59, median age of thyroidectomy group 46) Cohort treated between 1986-2007 Finland	Median follow-up 10 years	2007, however comparison different Adjusted for aetiology, age, and gender

See Appendix D: for full evidence tables.

2.4.4 Quality assessment of clinical studies included in the evidence review

Table 11: Clinical evidence summary: radioactive iodine vs surgery

Outcomes	No of Participants (studies) Follow up	Quality of the evidence (GRADE)	Relative effect (95% CI)	Anticipated absolute effects	
				Risk with Surgery	Risk difference with Radioactive iodine (95% CI)
Total cancer diagnoses (RR)	3146 (1 study) 15 years	⊕⊕⊕⊕ VERY LOW ^{1,2} due to risk of bias, imprecision	RR 1.00 (0.7 to 1.43)	115 per 1000	0 fewer per 1000 (from 35 fewer to 49 more)
Total cancer diagnoses (HR)	6148 (1 study) 10 years	⊕⊕⊕⊕ VERY LOW ¹ due to risk of bias	HR 1.03 (0.86 to 1.23)	⁻³	Not estimable
Total cancer mortality	10992 (1 study) 16.3 years	⊕⊕⊕⊕ VERY LOW ^{1,2} due to risk of bias, imprecision	HR 0.96 (0.73 to 1.26)	⁻³	Not estimable
Lip, oral, pharynx cancer diagnoses	3146 (1 study) 15 years	⊕⊕⊕⊕ VERY LOW ^{1,2} due to risk of bias, imprecision	RR 1.3 (0.2 to 8.45)	4 per 1000	1 more per 1000 (from 3 fewer to 28 more)

Outcomes	No of Participants (studies) Follow up	Quality of the evidence (GRADE)	Relative effect (95% CI)	Anticipated absolute effects	
				Risk with Surgery	Risk difference with Radioactive iodine (95% CI)
	(1 study) 15 years	VERY LOW ^{1,2} due to risk of bias, imprecision	(0.6 to 2.02)	24 per 1000	2 more per 1000 (from 10 fewer to 24 more)
Respiratory cancer diagnoses	3146 (1 study) 15 years	⊕⊕⊕⊕ VERY LOW ^{1,2} due to risk of bias, imprecision	RR 1.3 (0.4 to 4.23)	7 per 1000	2 more per 1000 (from 4 fewer to 23 more)
Breast cancer diagnoses	3146 (1 study) 15 years	⊕⊕⊕⊕ VERY LOW ^{1,2} due to risk of bias, imprecision	RR 0.8 (0.5 to 1.28)	34 per 1000	7 fewer per 1000 (from 17 fewer to 10 more)
Genital cancer diagnoses	3146 (1 study) 15 years	⊕⊕⊕⊕ VERY LOW ^{1,2} due to risk of bias, imprecision	RR 1.1 (0.4 to 3.02)	21 per 1000	2 more per 1000 (from 13 fewer to 42 more)
Kidney and bladder cancer diagnoses	3146 (1 study) 15 years	⊕⊕⊕⊕ VERY LOW ^{1,2} due to risk of bias, imprecision	RR 3.4 (0.5 to 23.12)	2 per 1000	5 more per 1000 (from 1 fewer to 42 more)
Melanoma diagnoses	3146 (1 study) 15 years	⊕⊕⊕⊕ VERY LOW ^{1,2} due to risk of bias, imprecision	RR 0 (0 to 7.8)	0 per 1000	1 fewer per 1000 (from 1 fewer to 3 more)
CNS cancer diagnoses	3146 (1 study) 15 years	⊕⊕⊕⊕ VERY LOW ^{1,2} due to risk of bias, imprecision	RR 0.3 (0.05 to 1.9)	3 per 1000	2 fewer per 1000 (from 3 fewer to 3 more)
Thyroid cancer diagnoses	3146 (1 study) 15 years	⊕⊕⊕⊕ VERY LOW ^{1,2} due to risk of bias,	RR 9.1 (1.2 to 69.01)	0 per 1000	4 more per 1000 (from 0 more to 34 more)

Outcomes	No of Participants (studies) Follow up	Quality of the evidence (GRADE)	Relative effect (95% CI)	Anticipated absolute effects	
				Risk with Surgery	Risk difference with Radioactive iodine (95% CI)
Other solid tumour diagnoses	3146 (1 study) 15 years	⊕⊕⊕⊕ VERY LOW ^{1,2} due to risk of bias, imprecision	RR 0.3 (0.02 to 4.3)	3 per 1000	2 fewer per 1000 (from 3 fewer to 9 more)
Lymphatic cancer diagnoses	3146 (1 study) 15 years	⊕⊕⊕⊕ VERY LOW ^{1,2} due to risk of bias, imprecision	RR 0.3 (0.02 to 3.7)	3 per 1000	2 fewer per 1000 (from 3 fewer to 9 more)
Leukaemia diagnoses	3146 (1 study) 15 years	⊕⊕⊕⊕ VERY LOW ^{1,2} due to risk of bias, imprecision	RR 0.6 (0.16 to 2.2)	5 per 1000	2 fewer per 1000 (from 4 fewer to 6 more)

1 Default starting quality of low overall due to selection bias in non-randomised studies. Downgraded further for risk of bias if the majority of evidence was at additional risk of bias, either once if high risk of bias or twice if very high risk of bias
2 Downgraded by 1 increment if the confidence interval crossed one MID or by 2 increments if the confidence interval crossed both MIDs
3 No control group risk provided

Table 12: Clinical evidence summary: radioactive iodine treated population vs general population

Outcomes	No of Participants (studies) Follow up	Quality of the evidence (GRADE)	Relative effect (95% CI)	Anticipated absolute effects	
				Risk with General population	Risk difference with Radioactive iodine (95% CI)
Total cancer diagnoses	26485 (5 studies) 5-17 years	⊕⊕⊕⊕ VERY LOW ^{1,2} due to risk of bias, inconsistency	Rate ratio 0.99 (0.83 to 1.18)	74 per 1000	1 fewer per 1000 (from 13 fewer to 13 more)
Lip, oral, pharynx cancer diagnoses	23210 (3 studies) 5-15 years	⊕⊕⊕⊕ VERY LOW ^{1,3} due to risk of bias, imprecision	Rate ratio 0.92 (0.57 to	1 per 1000	0 fewer per 1000 (from 0 fewer to 0 more)

Outcomes	No of Participants (studies) Follow up	Quality of the evidence (GRADE)	Relative effect (95% CI)	Anticipated absolute effects	
				Risk with General population	Risk difference with Radioactive iodine (95% CI)
			1.49)		
Salivary gland cancer diagnoses	15793 (2 studies) 10-15 years	⊕⊖⊖⊖ VERY LOW ^{1,3} due to risk of bias, imprecision	Rate ratio 1.88 (0.33 to 10.62)	0 per 1000	0 more per 1000 (from 0 fewer to 1 more)
Digestive organs and peritoneum cancer diagnoses	23817 (4 studies) 5-17 years	⊕⊖⊖⊖ VERY LOW ^{1,2} due to risk of bias, inconsistency	Rate ratio 1.06 (0.87 to 1.30)	27 per 1000	2 more per 1000 (from 4 fewer to 8 more)
Bone, connective tissue and skin cancer diagnoses	13003 (2 studies) 5-10 years	⊕⊖⊖⊖ VERY LOW ^{1,2} due to risk of bias, imprecision	Rate ratio 0.88 (0.69 to 1.14)	13 per 1000	2 fewer per 1000 (from 4 fewer to 2 more)
Breast cancer diagnoses	23817 (4 studies) 5-17 years	⊕⊖⊖⊖ VERY LOW ¹ due to risk of bias	Rate ratio 1.09 (0.97 to 1.22)	17 per 1000	2 more per 1000 (from 1 fewer to 4 more)
Brain and other CNS cancer diagnoses	23817 (4 studies) 5-17 years	⊕⊖⊖⊖ VERY LOW ^{1,3} due to risk of bias, imprecision	Rate ratio 1.46 (1.03 to 2.06)	3 per 1000	1 more per 1000 (from 0 more to 3 more)
Respiratory cancer diagnoses	23210 (3 studies) 5-17 years	⊕⊖⊖⊖ VERY LOW ^{1,2,3} due to risk of bias, inconsistency, imprecision	Rate ratio 0.84 (0.52 to 1.35)	9 per 1000	1 fewer per 1000 (from 4 fewer to 3 more)
Genitourinary cancer diagnoses	23210 (3 studies) 5-17 years	⊕⊖⊖⊖ VERY LOW ^{1,2,3} due to risk of bias, inconsistency, imprecision	Rate ratio 0.95 (0.73 to 1.24)	16 per 1000	1 fewer per 1000 (from 4 fewer to 4 more)

Outcomes	No of Participants (studies) Follow up	Quality of the evidence (GRADE)	Relative effect (95% CI)	Anticipated absolute effects	
				Risk with General population	Risk difference with Radioactive iodine (95% CI)
	(3 studies) 5-17 years	VERY LOW ¹ due to risk of bias	2.17 (1.36 to 3.48)	1 per 1000	1 more per 1000 (from 0 more to 2 more)
Haematopoietic cancer diagnoses	23210 (3 studies) 5-17 years	⊕⊖⊖⊖ VERY LOW ^{1,2,3} due to risk of bias, inconsistency, imprecision	Rate ratio 0.81 (0.56 to 1.19)	5 per 1000	1 fewer per 1000 (from 2 fewer to 1 more)
Kidney cancer diagnoses	15793 (2 studies) 10-15 years	⊕⊖⊖⊖ VERY LOW ^{1,3} due to risk of bias, imprecision	Rate ratio 1.62 (1.18 to 2.24)	4 per 1000	2 more per 1000 (from 1 more to 5 more)
Parathyroid cancer diagnoses	10207 (1 study) 15 years	⊕⊖⊖⊖ VERY LOW ^{1,3} due to risk of bias, imprecision	Rate ratio 1.6 (0.9 to 2.84)	2 per 1000	1 more per 1000 (from 0 fewer to 4 more)
Prostate cancer diagnoses	5586 (1 study) 10 years	⊕⊖⊖⊖ VERY LOW ^{1,3} due to risk of bias, imprecision	Rate ratio 1.3 (0.69 to 2.45)	37 per 1000	11 more per 1000 (from 11 fewer to 54 more)

1 Default starting quality of low overall due to selection bias in non-randomised studies. Downgraded further for risk of bias if the majority of evidence was at additional risk of bias, either once if high risk of bias or twice if very high risk of bias
2 Downgraded by 1 or 2 increments because the point estimate and or the confidence intervals varied widely across studies, unexplained by subgroup analysis
3 Downgraded by 1 increment if the confidence interval crossed one MID or by 2 increments if the confidence interval crossed both MIDs

1
2 See Appendix F: for full GRADE tables.

1 **2.5 Economic evidence**

2 The committee agreed that health economic studies would not be relevant to this review
3 question, and so were not sought.

4 **2.6 Evidence statements**

5 **2.6.1 Clinical evidence statements**

6 **2.6.1.1 Radioactive iodine vs surgery**

7 No clinically important difference was identified for total cancer diagnoses (2 studies, very
8 low quality), total cancer mortality (1 study, very low quality), lip/oral/pharynx cancer
9 diagnoses (1 study, very low quality), digestive organ and peritoneum cancer diagnoses (1
10 study, very low quality), respiratory cancer diagnoses (1 study, very low quality), breast
11 cancer diagnoses (1 study, very low quality), genital cancer diagnoses (1 study, very low
12 quality), kidney and bladder cancer diagnoses (1 study, very low quality), melanoma
13 diagnoses (1 study, very low quality), CNS cancer diagnoses (1 study, very low quality),
14 thyroid cancer diagnoses (1 study, very low quality), other solid tumour diagnoses (1 study,
15 very low quality), lymphatic cancer diagnoses (1 study, very low quality), leukaemia
16 diagnoses (1 study, very low quality).

17 **2.6.1.2 Radioactive iodine vs general population**

18 No clinically important difference was identified for total cancer diagnoses (5 studies, very
19 low quality), lip/oral/pharynx cancer diagnoses (3 studies, very low quality), salivary gland
20 cancer diagnoses (2 studies, very low quality), digestive organ and peritoneum cancer
21 diagnoses (4 studies, very low quality), bone/connective tissue/skin cancer diagnoses (2
22 studies, very low quality), breast cancer diagnoses (4 studies, very low quality), brain and
23 other CNS cancer diagnoses (4 studies, very low quality), respiratory cancer diagnoses (3
24 studies, very low quality), genitourinary cancer diagnoses (3 studies, very low quality),
25 thyroid cancer diagnoses (1 study, very low quality), haematopoietic cancer diagnoses (3
26 studies, very low quality), kidney cancer diagnoses (2 studies, very low quality), parathyroid
27 cancer diagnoses (1 study, very low quality).

28 There was a clinically important harm of radioactive iodine for prostate cancer diagnoses (1
29 study, very low quality).

30 **2.6.2 Health economic evidence statements**

- 31 • No relevant economic evaluations were identified.

32

33 **2.7 The committee's discussion of the evidence**

34 **2.7.1 Interpreting the evidence**

35 **2.7.1.1 The outcomes that matter most**

36 **Drugs vs Surgery vs Radioactive Iodine**

37 The committee agreed that the critical outcomes for this review were mortality and quality of
38 life. Important outcomes for all interventions included thyroid ophthalmopathy, euthyroidism,

1 hypothyroidism, relapse of hyperthyroidism, cardiovascular morbidity, arrhythmia,
2 osteoporosis, cognitive impairment, pain, symptom scores, experience of care, healthcare
3 contacts. Important intervention specific outcomes were recurrent laryngeal nerve damage,
4 hypocalcaemia, hypoparathyroidism, bleeding, infection, agranulocytosis, liver failure, minor
5 drug related adverse effects, teratogenesis, infertility, malignancy, thyrotoxic storm, growth
6 abnormalities.

7 **Radioactive Iodine safety**

8 The committee considered cancer diagnoses and infertility to be critical outcomes for this
9 review. No evidence was found for infertility.

10 **2.7.1.2 The quality of the evidence**

11 **Drugs vs Surgery vs Radioactive Iodine**

12 The quality of the evidence in this review ranged from high to very low quality, with the
13 majority being moderate or low quality. Evidence was typically downgraded for imprecision
14 as studies were often small, some comparisons were downgraded for inconsistency that
15 could not be explained by any protocol subgroup analyses and some comparisons were
16 downgraded for risk of bias. The trials included in this review generally had long follow-up
17 periods, with some participants being followed for up to 21 years.

18 Thyroid status and ophthalmopathy at end of follow-up were the most commonly reported
19 outcome. There was no quality of life evidence identified. The majority of outcomes included
20 in the protocol by the committee were not reported.

21 The comparison between radioactive iodine and antithyroid drugs, used for Graves' disease
22 in first line treatment, included the most evidence. Comparisons involving surgery or second
23 line treatment were only supported by one study each.

24 No evidence was identified in children or older adults. No evidence was identified in studies
25 explicitly in toxic multinodular goitre and there was only one small study in people who had
26 failed first line treatment.

27 The committee noted that the studies included in this review were not designed to capture
28 the rare but well established adverse events of some treatment options (for example
29 agranulocytosis with antithyroid drugs).

30 The committee noted that the doses of radioactive iodine used in most studies were lower
31 than what would be used in the UK currently. Qualitatively, higher doses would be expected
32 to lead to more hypothyroidism and euthyroidism and less hyperthyroidism. Higher doses
33 could also lead to more adverse events, although these were not identified in this review.

34 **Radioactive Iodine safety**

35 The majority of the evidence was very low quality due to the non-randomised nature of the
36 included studies. Beyond the lack of randomisation, studies that compared radioactive iodine
37 with surgery were more informative as they reduced the confounding effect of the underlying
38 thyroid disease (as opposed to the studies that compared cancer diagnoses between a
39 radioactive iodine treated group and the general population). The majority of studies included
40 a population who had been treated many years ago, some cohorts including participants
41 treated as far back as 1946. The doses and strategies of radioactive iodine (for example
42 whether a fixed administered activity or calculated absorbed dose was used) was not always
43 provided but generally appeared to be a fixed approach and using lower doses (for example
44 ~300MBq) than those used in the UK currently (typically 400-600MBq).

1 The committee agreed that there were limitations to the evidence available but also noted
2 that the studies were large, had long follow-up times and it is unlikely that RCTs that are as
3 large and lengthy in follow-up will ever be conducted. Nevertheless, they agreed that a
4 registry of patients receiving RAI would further develop our understanding of the risks and
5 benefits associated with RAI therapy and decided to make a research recommendation.

6 All the included evidence was on adults, there was no evidence to consider in children.

7 **2.7.1.3 Benefits and harms**

8 **Drugs vs Surgery vs Radioactive Iodine**

9 The evidence shows that radioactive iodine has a clinically important harm compared with
10 antithyroid drugs for ophthalmopathy but a clinically important benefit in terms of reducing
11 persistence or recurrence of hyperthyroidism.

12 Compared with antithyroid drugs, radioactive iodine also appeared to lead to more people
13 ending up in a hypothyroid state as opposed to euthyroid. The committee discussed the
14 outcomes of hypothyroidism and euthyroidism. Euthyroidism is seen as a preferential goal of
15 treatment by some people with thyroid disease, and eliminates the need for concurrent
16 thyroid function replacement with levothyroxine or similar. However the committee was
17 aware of some evidence (not the focus of this review) that long term outcomes for people
18 who achieve hypothyroidism after radioactive iodine are better than for those who are
19 euthyroid. The committee noted that current guidance by other groups is to aim for
20 hypothyroidism when using radioactive iodine. Committee members in primary care noted
21 that from their experience, the people they treated for hyperthyroidism with antithyroid drugs
22 were generally more satisfied with their care than the people they treated for hypothyroidism
23 (secondary to radioactive iodine or surgery for hyperthyroidism).

24 There was less evidence available comparing surgery to either modality. In general surgery
25 appeared to have a clinically important benefit over radioactive iodine or antithyroid drugs in
26 terms of the likelihood of relapse or persistence of hyperthyroidism, however the smaller
27 trials made this difficult to interpret. The committee noted that although there was no
28 evidence in this review on hypothyroidism as a result of surgery, this is conceptually a likely
29 outcome. The one study in this review reporting on surgical outcomes assessed the efficacy
30 of subtotal thyroidectomy, as opposed to total thyroidectomy as in the economic evidence.
31 As discussed in the review of different types of surgery, these two options are likely to have
32 different benefits and harms.

33 Beyond the impact of ophthalmopathy and thyroid state, the review did not identify definitive
34 evidence on the harms of radioactive iodine, antithyroid drugs or surgery. The committee
35 agreed that each form of treatment is associated with some harm. Some of these harms are
36 more definitive than others. Surgery is associated with the general harms of surgery (for
37 example bleeding, infection) as well as specific harms related to surgery on the thyroid gland
38 (for example hypoparathyroidism and recurrent laryngeal nerve damage). Antithyroid drugs
39 have a combination of common minor adverse events (for example skin rash) and rare but
40 severe adverse events (for example agranulocytosis and liver failure) which are documented
41 in the summary of product characteristics. Radioactive iodine treatment has theoretical
42 harms beyond those identified in this review, in terms of secondary malignancies and effects
43 on fertility or teratogenesis. None of these harms were identified in the RCTs in this review
44 and the committee's view overall was that while these were important risks to discuss with
45 people considering treatment, there was not information available on their likelihood.

46 The committee noted, based on their experience, that there may be particular features of a
47 person's hyperthyroidism that may suggest one treatment option is preferable to others. If
48 there was any uncertainty around the potential for thyroid cancer or if there were significant
49 compressive symptoms from a large goitre, then surgery was typically considered the most

1 appropriate option. If there was a significant degree of pre-existing ophthalmopathy this may
2 promote treatment options other than radioactive iodine. If people's hyperthyroidism
3 generally appeared likely to respond well to antithyroid drug treatment, this may make them a
4 better candidate for non-definitive treatment with drugs as opposed to potentially causing
5 long term hypothyroidism with either radioactive iodine or surgery. The committee noted that
6 there was no evidence in this review to suggest which groups might respond particularly well
7 to antithyroid drugs. In their experience, people with very mild hyperthyroidism and in
8 particular T3 hyperthyroidism did tend to respond well to antithyroid drugs.

9 The committee discussed the extrapolation of evidence and experience from adults to
10 children. Concerns over potential adverse effects of definitive treatment with radioactive
11 iodine or surgery were generally greater for children than adults. Surgery may be technically
12 more demanding in children. The potential long term risks of radioactive iodine in terms of
13 secondary malignancy are more relevant in children, given their greater life expectancy after
14 treatment compared with older adults. However at the same time, children and their families
15 are often keen to explore definitive treatment options. From the committee's experience,
16 hyperthyroidism in children may be more aggressive than in adults and require lengthier
17 treatment with antithyroid drugs (up to 10 years in children as opposed to 12-18 months in
18 adults).

19 **Radioactive iodine safety**

20 Overall the evidence in this review did not show a clinically important harm of radioactive
21 iodine treatment compared with either surgery or a general population in terms of increased
22 risk of cancer diagnoses. There was no clinically important effect for overall cancer
23 diagnoses, the outcome with the greatest event rates in both arms. When considering site
24 specific cancer diagnoses, due to the much smaller event rates there was generally more
25 imprecision and lower quality evidence with relative effects more likely to appear to show an
26 effect but the absolute effects remained small, with all but one remaining below the threshold
27 of 10 per 1000 people treated. The committee agreed that the one outcome, for which this
28 threshold was breached, prostate cancer diagnoses in the radioactive iodine versus general
29 population comparison, was likely to reflect statistical uncertainty more than a true effect and
30 noted the very low quality of the evidence.

31 The committee agreed that there was insufficient evidence to determine in this review if
32 dosing strategy affected safety as the studies generally did not provide adequate information
33 on the radioactive iodine strategies used.

34 Balanced against this evidence of no important harm of radioactive iodine, the committee
35 noted the underlying biological principles that any exposure to radiation is likely to increase
36 cancer risk to some degree. However the evidence in this review suggests that the risk
37 associated with the radiation involved in treatment of thyroid disease is not clinically
38 impactful.

39 **2.7.2 Cost effectiveness and resource use**

40 Resource use implications were considered through the published cost-effectiveness
41 evidence included in the review.

42 This was a UK cost-utility analysis that compared three options for the management of
43 thyrotoxicosis secondary to Graves' disease: radioactive iodine (RAI); antithyroid drugs
44 (ATD); and surgery (total thyroidectomy).

45 The analysis found that RAI was the most cost effective option at a cost effectiveness
46 threshold of £20,000 per QALY gained. RAI had the lowest mean cost per patient over a
47 lifetime horizon (£5,425) and a mean 34.73 QALYs per patient. Total thyroidectomy had
48 higher costs (£7,115) and lower QALYs (33.93 QALYs) than RAI. ATDs had higher costs
49 (£16,866) than RAI but also higher QALYs (35.17 QALYs); however, it had an incremental

1 cost effectiveness ratio compared to RAI of £26,279 per QALY gained and so was not
2 considered cost effective.

3 This supported a strong recommendation to offer radioactive iodine as the first line treatment
4 options for the management of thyrotoxicosis secondary to Graves' disease. The committee
5 noted that the results of the economic evidence were in line with the clinical evidence.

6 The committee noted that the model accurately captures the following key adverse events
7 associated with each of the three interventions: hypothyroidism secondary to total
8 thyroidectomy, the excess risk of ophthalmopathy when using radioactive iodine and
9 increased risk of relapse when using antithyroid drugs. However, it was noted that some
10 potential adverse events have not been reflected in the model structure, e.g. malignancy,
11 thyroid eye disease. This was justified in the study, though, as the authors explained that the
12 evidence supporting causal association between the use of radioactive iodine and
13 malignancy is limited which was confirmed by the committee during their discussions.
14 However, as a result of this concern, the committee chose to restrict the recommendation to
15 people in whom there is no risk of malignancy, thyroid eye disease and compression. The
16 risk of infertility was another potential adverse event, not captured in the model, and relating
17 to the use of radioactive iodine in women of childbearing age. Therefore, the committee
18 agreed that it was important to discuss treatment options with people with Graves' disease to
19 minimise these risks.

20 The committee also made consider recommendations where it assumed certain subgroups
21 would benefit more from a course of antithyroid drugs or in some rare cases surgery rather
22 than radioactive iodine. For example, people with mild uncomplicated Graves' disease whom
23 are likely to achieve remission with a course of antithyroid drug therefore unlikely to be
24 rendered hypothyroid, reducing the need for long-term hormone replacement therapy which
25 in turn saves money and improves patients quality of life.

26 There was no economic or clinical evidence for the management of thyrotoxicosis in people
27 with toxic nodular goitre. Hence, the committee extrapolated the findings from the study and
28 made a weaker recommendation to consider radioactive iodine as first line treatment except
29 in instances where there are concerns around malignancy.

30 In children, the committee were uncertain about the long-term health risk associated with
31 radioactive iodine and surgery, and agreed to offer antithyroid drugs as first line treatment,
32 which is in line with current practice and unlikely to have a substantial cost impact. However,
33 the committee noted that definitive options should be discussed with a multi-disciplinary team
34 especially when they have relapsed hyperthyroidism after a course of antithyroid drugs or in
35 children with a single toxic nodule. The population of children with single toxic nodule is very
36 small hence unlikely to result in a cost impact. Overall, the recommendation for the use of
37 RAI as first line is a change to current practice, which is likely to have a substantial cost
38 saving as shown by the economic evidence and agreed by the committee. Furthermore, in
39 children due to the uncertainty around the potential risk and benefit around radioactive iodine
40 treatment, the cost-effectiveness was considered uncertain. Other factors the committee took
41 into account

42 The committee noted that none of the currently available treatment options addressed the
43 potential underlying causes of hyperthyroidism (for example the immunological basis for
44 Graves' disease). While immunomodulatory treatment options were not a focus of this review
45 and therefore specific research recommendations could not be made, the committee were
46 keen to see this area be developed in the future.

47 The committee noted that although pregnancy is outside the scope of this guideline,
48 radioactive iodine would not be considered appropriate for anyone considering pregnancy,
49 currently pregnant or breast-feeding.

1 The committee made recommendations on toxic multinodular goitre based on extrapolations
2 from the evidence on Graves' disease (noting that some studies in predominantly Graves'
3 disease populations did include a minority of people with toxic multinodular goitre) and on
4 their own experience. The committee's experience was that antithyroid drugs would not be
5 an appropriate option for this population.

6 The committee noted that by the point in the treatment pathway that people arrive at
7 radioactive iodine currently, they have typically been started on antithyroid drugs in primary
8 care. The review on the use of radioactive iodine considers this issue further.

9

References

1. Abraham-Nordling M, Törring O, Hamberger B, Lundell G, Tallstedt L, Calissendorff J et al. Graves' disease: a long-term quality-of-life follow up of patients randomized to treatment with antithyroid drugs, radioiodine, or surgery. *Thyroid*. 2005; 15(11):1279-1286
2. Abraham-Nordling M, Wallin G, Lundell G, Törring O. Thyroid hormone state and quality of life at long-term follow-up after randomized treatment of Graves' disease. *European Journal of Endocrinology*. 2007; 156(2):173-179
3. Abraham-Nordling M, Wallin G, Träisk F, Berg G, Calissendorff J, Hallengren B et al. Thyroid-associated ophthalmopathy; quality of life follow-up of patients randomized to treatment with antithyroid drugs or radioiodine. *European Journal of Endocrinology*. 2010; 163(4):651-657
4. Abraham P, Avenell A, McGeoch SC, Clark LF, Bevan JS. Antithyroid drug regimen for treating Graves' hyperthyroidism. *Cochrane Database of Systematic Reviews* 2010, Issue 1. Art. No.: CD003420. DOI: 10.1002/14651858.CD003420.pub4.
5. Allanic H, Fauchet R, Orgiazzi J, Madec AM, Genetet B, Lorcy Y et al. Antithyroid drugs and Graves' disease: a prospective randomized evaluation of the efficacy of treatment duration. *Journal of Clinical Endocrinology and Metabolism*. 1990; 70(3):675-679
6. Andrade VA, Gross JL, Maia AL. Effect of methimazole pretreatment on serum thyroid hormone levels after radioactive treatment in Graves' hyperthyroidism. *Journal of Clinical Endocrinology and Metabolism*. 1999; 84(11):4012-4016
7. Andrade VA, Gross JL, Maia AL. The effect of methimazole pretreatment on the efficacy of radioactive iodine therapy in Graves' hyperthyroidism: one-year follow-up of a prospective, randomized study. *Journal of Clinical Endocrinology and Metabolism*. 2001; 86(8):3488-3493
8. Andrade VA, Gross JL, Maia AL. Serum thyrotropin-receptor autoantibodies levels after I therapy in Graves' patients: effect of pretreatment with methimazole evaluated by a prospective, randomized study. *European Journal of Endocrinology*. 2004; 151(4):467-474
9. Angusti T, Codegone A, Pellerito R, Favero A. Thyroid cancer prevalence after radioiodine treatment of hyperthyroidism. *Journal of Nuclear Medicine*. 2000; 41(6):1006-9
10. Azizi F, Amouzegar A. Management of thyrotoxicosis in children and adolescents: 35 years' experience in 304 patients. *Journal of Pediatric Endocrinology and Metabolism*. 2018; 31(2):159-165
11. Azizi F, Ataie L, Hedayati M, Mehrabi Y, Sheikholeslami F. Effect of long-term continuous methimazole treatment of hyperthyroidism: comparison with radioiodine. *European Journal of Endocrinology*. 2005; 152(5):695-701
12. Azizi F, Yousefi V, Bahrainian A, Sheikholeslami F, Tohidi M, Mehrabi Y. Long-term continuous methimazole or radioiodine treatment for hyperthyroidism. *Archives of Iranian Medicine*. 2012; 15(8):477-484
13. Barczyński M, Konturek A, Hubalewska-Dydejczyk A, Gołkowski F, Nowak W. Randomized clinical trial of bilateral subtotal thyroidectomy versus total thyroidectomy

- 1 for Graves' disease with a 5-year follow-up. *British Journal of Surgery*. 2012;
2 99(4):515-522
- 3 14. Barczynski M, Konturek A, Hubalewska-Dydejczyk A, Golkowski F, Miklaszewska G,
4 Romanowska-Dixon B. Five-year follow up of a randomized clinical trial of bilateral
5 subtotal thyroidectomy versus total thyroidectomy for Graves' disease. *Langenbeck's*
6 *Archives of Surgery*. 2010; 395(4):471
- 7 15. Barczynski M, Konturek A, Hubalewska-Dydejczyk A, Golkowski F, Nowak W. Ten-
8 year follow-up of a randomized clinical trial of total thyroidectomy versus Dunhill
9 operation versus bilateral subtotal thyroidectomy for multinodular non-toxic goiter.
10 *World Journal of Surgery*. 2018; 42(2):384-392
- 11 16. Bartalena L, Marcocci C, Bogazzi F, Manetti L, Tanda ML, Dell'Unto E et al. Relation
12 between therapy for hyperthyroidism and the course of Graves' ophthalmopathy. *New*
13 *England Journal of Medicine*. 1998; 338(2):73-78
- 14 17. Benker G, Reinwein D, Kahaly G, Tegler L, Alexander WD, Fassbinder J et al. Is
15 there a methimazole dose effect on remission rate in Graves' disease? Results from a
16 long-term prospective study. The European Multicentre Trial Group of the Treatment
17 of Hyperthyroidism with Antithyroid Drugs. *Clinical Endocrinology*. 1998; 49(4):451-
18 457
- 19 18. Benker G, Vitti P, Kahaly G, Raue F, Tegler L, Hirche H et al. Response to
20 methimazole in Graves' disease. The European Multicenter Study Group. *Clinical*
21 *Endocrinology*. 1995; 43(3):257-263
- 22 19. Bonnema SJ, Bennedbaek FN, Gram J, Veje A, Marving J, Hegedus L. Resumption
23 of methimazole after 131I therapy of hyperthyroid diseases: effect on thyroid function
24 and volume evaluated by a randomized clinical trial. *European Journal of*
25 *Endocrinology*. 2003; 149(6):485-492
- 26 20. Bonnema SJ, Bennedbaek FN, Veje A, Marving J, Hegedüs L. Propylthiouracil before
27 131I therapy of hyperthyroid diseases: effect on cure rate evaluated by a randomized
28 clinical trial. *Journal of Clinical Endocrinology and Metabolism*. 2004; 89(9):4439-
29 4444
- 30 21. Bonnema SJ, Grupe P, Boel-Jørgensen H, Brix TH, Hegedüs L. A randomized trial
31 evaluating a block-replacement regimen during radioiodine therapy. *European*
32 *Journal of Clinical Investigation*. 2011; 41(7):693-702
- 33 22. Braga M, Walpert N, Burch HB, Solomon BL, Cooper DS. The effect of methimazole
34 on cure rates after radioiodine treatment for Graves' hyperthyroidism: a randomized
35 clinical trial. *Thyroid*. 2002; 12(2):135-139
- 36 23. Burch HB, Solomon BL, Cooper DS, Ferguson P, Walpert N, Howard R. The effect of
37 antithyroid drug pretreatment on acute changes in thyroid hormone levels after (131)I
38 ablation for Graves' disease. *Journal of Clinical Endocrinology and Metabolism*. 2001;
39 86(7):3016-3021
- 40 24. Buscemi S, Verga S, Cottone S, Andronico G, D'Orio L, Mannino V et al. Favorable
41 clinical heart and bone effects of anti-thyroid drug therapy in endogenous subclinical
42 hyperthyroidism. *Journal of Endocrinological Investigation*. 2007; 30(3):230-235
- 43 25. Canto AU, Dominguez PN, Jimeno CA, Obaldo JM, Ogbac RV. Comparison of Fixed
44 versus Calculated Activity of Radioiodine for the Treatment of Graves Disease in
45 Adults. *Endocrinol metab*. 2016; 31(1):168-173

- 1 26. Cevallos JL, Hagen GA, Maloof F, Chapman EM. Low-dosage ¹³¹I therapy of
2 thyrotoxicosis (diffuse goiters). A five-year follow-up study. *New England Journal of*
3 *Medicine*. 1974; 290(3):141-3
- 4 27. Chao M, Jiawei X, Guoming W, Jianbin L, Wanxia L, Driedger A et al. Radioiodine
5 treatment for pediatric hyperthyroid Grave's disease. *European Journal of Pediatrics*.
6 2009; 168(10):1165-1169
- 7 28. Chen DY, Jing J, Schneider PF, Chen TH. Comparison of the long-term efficacy of
8 low dose ¹³¹I versus antithyroid drugs in the treatment of hyperthyroidism. *Nuclear*
9 *Medicine Communications*. 2009; 30(2):160-168
- 10 29. Chen DY, Schneider PF, Zhang XS, He ZM, Jing J, Chen TH. Striving for
11 euthyroidism in radioiodine therapy of Graves' disease: a 12-year prospective,
12 randomized, open-label blinded end point study. *Thyroid*. 2011; 21(6):647-654
- 13 30. Chen DY, Schneider PF, Zhang XS, Luo XY, He ZM, Chen TH. Changes in graves'
14 ophthalmopathy after radioiodine and anti-thyroid drug treatment of graves' disease
15 from 2 prospective, randomized, open-label, blinded end point studies. *Experimental*
16 *and clinical endocrinology & diabetes*. 2014; 122(1):1-6
- 17 31. Chi SY, Hsei KC, Sheen-Chen SM, Chou FF. A prospective randomized comparison
18 of bilateral subtotal thyroidectomy versus unilateral total and contralateral subtotal
19 thyroidectomy for graves' disease. *World Journal of Surgery*. 2005; 29(2):160-163
- 20 32. Connell JM, Hilditch TE, Robertson J, Coghill G, Alexander WD. Radioprotective
21 action of carbimazole in radioiodine therapy for thyrotoxicosis--influence of the drug
22 on iodine kinetics. *European Journal of Nuclear Medicine*. 1987; 13(7):358-361
- 23 33. De Luca F, Valenzise M. Controversies in the pharmacological treatment of Graves'
24 disease in children. *Expert Review of Clinical Pharmacology*. 2018:1-9
- 25 34. Department of Health. NHS reference costs 2016-17. 2017. Available from:
26 <https://www.gov.uk/government/collections/nhs-reference-costs> Last accessed:
27 15/02/2019
- 28 35. Donovan PJ, McLeod DS, Little R, Gordon L. Cost-utility analysis comparing
29 radioactive iodine, anti-thyroid drugs and total thyroidectomy for primary treatment of
30 Graves' disease. *European Journal of Endocrinology*. 2016; 175(6):595-603
- 31 36. Edmonds CJ, Tellez M. Treatment of Graves' disease by carbimazole: high dose with
32 thyroxine compared to titration dose. *European Journal of Endocrinology*. 1994;
33 131(2):120-124
- 34 37. Esfahani AF, Kakhki VR, Fallahi B, Eftekhari M, Beiki D, Saghari M et al.
35 Comparative evaluation of two fixed doses of 185 and 370 MBq ¹³¹I, for the
36 treatment of Graves' disease resistant to antithyroid drugs. *Hellenic Journal of*
37 *Nuclear Medicine*. 2005; 8(3):158-161
- 38 38. Franklyn JA, Maisonneuve P, Sheppard M, Betteridge J, Boyle P. Cancer incidence
39 and mortality after radioiodine treatment for hyperthyroidism: a population-based
40 cohort study. *Lancet*. 1999; 353(9170):2111-5
- 41 39. Franklyn JA, Maisonneuve P, Sheppard MC, Betteridge J, Boyle P. Mortality after the
42 treatment of hyperthyroidism with radioactive iodine. *New England Journal of*
43 *Medicine*. 1998; 338(11):712-8
- 44 40. Franklyn JA, Sheppard MC, Maisonneuve P. Thyroid function and mortality in
45 patients treated for hyperthyroidism. *JAMA*. 2005; 294(1):71-80

- 1 41. García-Mayor RV, Páramo C, Luna Cano R, Pérez Mendez LF, Galofré JC, Andrade
2 A. Antithyroid drug and Graves' hyperthyroidism. Significance of treatment duration
3 and TRAb determination on lasting remission. *Journal of Endocrinological*
4 *Investigation*. 1992; 15(11):815-820
- 5 42. Giesecke P, Frykman V, Wallin G, Lonn S, Discacciati A, Topping O et al. All-cause
6 and cardiovascular mortality risk after surgery versus radioiodine treatment for
7 hyperthyroidism. *British Journal of Surgery*. 2018; 105(3):279-286
- 8 43. Glinoe D, Nayer P, Bex M. Effects of l-thyroxine administration, TSH-receptor
9 antibodies and smoking on the risk of recurrence in Graves' hyperthyroidism treated
10 with antithyroid drugs: a double-blind prospective randomized study. *European*
11 *Journal of Endocrinology*. 2001; 144(5):475-483
- 12 44. Goldman MB, Maloof F, Monson RR, Aschengrau A, Cooper DS, Ridgway EC.
13 Radioactive iodine therapy and breast cancer. A follow-up study of hyperthyroid
14 women. *American Journal of Epidemiology*. 1988; 127(5):969-80
- 15 45. Goni Iriarte MJ, Forga Llenas L, Iriarte Beroiz A, Anda Apinaniz E, Rodriguez
16 Erdozain R, Menendez Torre E. Recurrence of Graves' disease: the influence of
17 treatment schedule. *Medicina Clínica*. 1995; 104(1):11-14
- 18 46. Grebe SK, Feek CM, Ford HC, Fagerström JN, Cordwell DP, Delahunt JW et al. A
19 randomized trial of short-term treatment of Graves' disease with high-dose
20 carbimazole plus thyroxine versus low-dose carbimazole. *Clinical Endocrinology*.
21 1998; 48(5):585-592
- 22 47. Hall P, Berg G, Bjelkengren G, Boice JD, Jr., Ericsson UB, Hallquist A et al. Cancer
23 mortality after iodine-131 therapy for hyperthyroidism. *International Journal of Cancer*.
24 1992; 50(6):886-90
- 25 48. Hall P, Boice JD, Jr., Berg G, Bjelkengren G, Ericsson UB, Hallquist A et al.
26 Leukaemia incidence after iodine-131 exposure. *Lancet*. 1992; 340(8810):1-4
- 27 49. Hall P, Holm LE. Cancer in iodine-131 exposed patients. *Journal of Endocrinological*
28 *Investigation*. 1995; 18(2):147-9
- 29 50. Hall P, Holm LE. Late consequences of radioiodine for diagnosis and therapy in
30 Sweden. *Thyroid*. 1997; 7(2):205-8
- 31 51. Hall P, Lundell G, Holm LE. Mortality in patients treated for hyperthyroidism with
32 iodine-131. *Acta Endocrinologica*. 1993; 128(3):230-4
- 33 52. Hamide AKAHDSJS. Radioiodine therapy in patients with Graves' disease and the
34 effects of prior carbimazole therapy. *Indian Journal of Endocrinology and Metabolism*.
35 2014; 18(5):688-693
- 36 53. Hashizume K, Ichikawa K, Sakurai A, Suzuki S, Takeda T, Kobayashi M et al.
37 Administration of thyroxine in treated Graves' disease. Effects on the level of
38 antibodies to thyroid-stimulating hormone receptors and on the risk of recurrence of
39 hyperthyroidism. *New England Journal of Medicine*. 1991; 324(14):947-953
- 40 54. He CT, Hsieh AT, Pei D, Hung YJ, Wu LY, Yang TC et al. Comparison of single daily
41 dose of methimazole and propylthiouracil in the treatment of Graves' hyperthyroidism.
42 *Clinical Endocrinology*. 2004; 60(6):676-681
- 43 55. Hieu TT, Russell AW, Cuneo R, Clark J, Kron T, Hall P et al. Cancer risk after
44 medical exposure to radioactive iodine in benign thyroid diseases: A meta-analysis.
45 *Endocrine-Related Cancer*. 2012; 19(5):645-655

- 1 56. Hoermann R, Quadbeck B, Roggenbuck U, Szabolcs I, Pfeilschifter J, Meng W et al.
2 Relapse of Graves' disease after successful outcome of antithyroid drug therapy:
3 results of a prospective randomized study on the use of levothyroxine. *Thyroid*. 2002;
4 12(12):1119-1128
- 5 57. Hoffman DA, McConahey WM. Breast cancer following iodine-131 therapy for
6 hyperthyroidism. *Journal of the National Cancer Institute*. 1983; 70(1):63-7
- 7 58. Hoffman DA, McConahey WM, Diamond EL, Kurland LT. Mortality in women treated
8 for hyperthyroidism. *American Journal of Epidemiology*. 1982; 115(2):243-54
- 9 59. Hoffman DA, McConahey WM, Fraumeni JF, Jr., Kurland LT. Cancer incidence
10 following treatment of hyperthyroidism. *International Journal of Epidemiology*. 1982;
11 11(3):218-24
- 12 60. Holm LE. Thyroid cancer after exposure to radioactive 131I. *Acta Oncologica*. 2006;
13 45(8):1037-40
- 14 61. Holm LE, Dahlqvist I, Israelsson A, Lundell G. Malignant thyroid tumors after iodine-
15 ¹³¹ therapy: a retrospective cohort study. *New England Journal of Medicine*. 1980;
16 303(4):188-91
- 17 62. Holm LE, Hall P, Wiklund K, Lundell G, Berg G, Bjelkengren G et al. Cancer risk after
18 iodine-131 therapy for hyperthyroidism. *Journal of the National Cancer Institute*.
19 1991; 83(15):1072-7
- 20 63. Homsanit M, Sriussadaporn S, Vannasaeng S, Peerapatdit T, Nitiyanant W,
21 Vichayanrat A. Efficacy of single daily dosage of methimazole vs. propylthiouracil in
22 the induction of euthyroidism. *Clinical Endocrinology*. 2001; 54(3):385-390
- 23 64. Howarth D, Epstein M, Lan L, Tan P, Booker J. Determination of the optimal minimum
24 radioiodine dose in patients with Graves' disease: a clinical outcome study. *European*
25 *Journal of Nuclear Medicine*. 2001; 28(10):1489-1495
- 26 65. Jaiswal AK, Bal C, Damle NA, Ballal S, Goswami R, Hari S et al. Comparison of
27 clinical outcome after a fixed dose versus dosimetry-based radioiodine treatment of
28 Graves' disease: Results of a randomized controlled trial in Indian population. *Indian*
29 *Journal of Endocrinology and Metabolism*. 2014; 18(5):648-54
- 30 66. Järhult J, Rudberg C, Larsson E, Selvander H, Sjövall K, Winsa B et al. Graves'
31 disease with moderate-severe endocrine ophthalmopathy-long term results of a
32 prospective, randomized study of total or subtotal thyroid resection. *Thyroid*. 2005;
33 15(10):1157-1164
- 34 67. Jorde R, Ytre-Arne K, Størmer J, Sundsfjord J. Short-term treatment of Graves'
35 disease with methimazole in high versus low doses. *Journal of Internal Medicine*.
36 1995; 238(2):161-165
- 37 68. Journy NMY, Bernier MO, Doody MM, Alexander BH, Linet MS, Kitahara CM.
38 Hyperthyroidism, hypothyroidism, and cause-specific mortality in a large cohort of
39 women. *Thyroid*. 2017; 27(8):1001-1010
- 40 69. Kallner G, Vitols S, Ljunggren JG. Comparison of standardized initial doses of two
41 antithyroid drugs in the treatment of Graves' disease. *Journal of Internal Medicine*.
42 1996; 239(6):525-529
- 43 70. Kansara S, Kotwal N, Kumar K, Singh Y, Upreti V, Nachankar A. Effect of Antithyroid
44 Therapies on Bone and Body Composition: A Prospective, Randomized, Clinical
45 Study Comparing Antithyroid Drugs with Radioiodine Therapy. *Indian Journal of*
46 *Endocrinology and Metabolism*. 2017; 21(4):531-534

- 1 71. Kung AW, Yau CC, Cheng AC. The action of methimazole and L-thyroxine in
2 radioiodine therapy: a prospective study on the incidence of hypothyroidism. *Thyroid*.
3 1995; 5(1):7-12
- 4 72. Leclere J. Treatment of Basedow disease with synthetic antithyroid drugs. Evaluation
5 of the dose on the efficacy of the long term treatment. *Annales d'Endocrinologie*.
6 1994; 55(1):11-14
- 7 73. Leslie WD, Ward L, Salamon EA, Ludwig S, Rowe RC, Cowden EA. A randomized
8 comparison of radioiodine doses in graves' hyperthyroidism. *Journal of Clinical*
9 *Endocrinology and Metabolism*. 2003; 88(3):978-983
- 10 74. Leung AKC, Leung AAC. Evaluation and Management of Children with
11 Thyrotoxicosis. *Recent Patents on Endocrine, Metabolic & Immune Drug Discovery*.
12 2017; 11(1):22-31
- 13 75. Li HX, Xiang N, Hu WK, Jiao XL. Relation between therapy options for Graves'
14 disease and the course of Graves' ophthalmopathy: a systematic review and meta-
15 analysis. *Journal of Endocrinological Investigation*. 2016; 39(11):1225-1233
- 16 76. Liu Y, Liu B, Liu RL, Jiang H, Huang ZN, Huang Y. A new method of subtotal
17 thyroidectomy for Graves' disease leaving a unilateral remnant based on the upper
18 pole. *Medicine*. 2017; 96(6):e5919
- 19 77. Liu ZW, Masterson L, Fish B, Jani P, Chatterjee K. Thyroid surgery for Graves'
20 disease and Graves' ophthalmopathy. *Cochrane Database of Systematic Reviews*
21 2015, Issue 11. Art. No.: CD010576. DOI: 10.1002/14651858.CD010576.pub2.
- 22 78. Ljunggren JG, Törring O, Wallin G, Taube A, Tallstedt L, Hamberger B et al. Quality
23 of life aspects and costs in treatment of Graves' hyperthyroidism with antithyroid
24 drugs, surgery, or radioiodine: results from a prospective, randomized study. *Thyroid*.
25 1998; 8(8):653-659
- 26 79. Lucas A, Salinas I, Rius F, Pizarro E, Granada ML, Foz M et al. Medical therapy of
27 Graves' disease: does thyroxine prevent recurrence of hyperthyroidism? *Journal of*
28 *Clinical Endocrinology and Metabolism*. 1997; 82(8):2410-2413
- 29 80. Ma C, Kuang A, Xie J, Liu GJ. Radioiodine treatment for pediatric Graves' disease.
30 *Cochrane Database of Systematic Reviews* 2008, Issue 3. Art. No.: CD006294. DOI:
31 10.1002/14651858.CD006294.pub2.
- 32 81. Ma C, Xie J, Wang H, Li J, Chen S. Radioiodine therapy versus antithyroid
33 medications for Graves' disease. *Cochrane Database of Systematic Reviews* 2016,
34 Issue 2. Art. No.: CD010094. DOI:
35 <https://dx.doi.org/10.1002/14651858.CD010094.pub2>.
- 36 82. Marcocci C, Bartalena L, Bogazzi F, Panicucci M, Bruno-Bossio G, Lepri A et al.
37 Radioiodine treatment of Graves' hyperthyroidism and progression of
38 ophthalmopathy: protective effect of systemic corticosteroids. *Acta endocrinologica*,
39 supplement. 1989; 121(2):145-148
- 40 83. Mashio Y, Beniko M, Matsuda A, Koizumi S, Matsuya K, Mizumoto H et al. Treatment
41 of hyperthyroidism with a small single daily dose of methimazole: a prospective long-
42 term follow-up study. *Endocrine Journal*. 1997; 44(4):553-558
- 43 84. Mastorakos G, Doufas AG, Mantzos E, Mantzos J, Koutras DA. T4 but not T3
44 administration is associated with increased recurrence of Graves' disease after
45 successful medical therapy. *Journal of Endocrinological Investigation*. 2003;
46 26(10):979-984

- 1 85. Maugendre D, Gatel A, Campion L, Massart C, Guilhem I, Lorcy Y et al. Antithyroid
2 drugs and Graves' disease--prospective randomized assessment of long-term
3 treatment. *Clinical Endocrinology*. 1999; 50(1):127-132
- 4 86. McIver B, Rae P, Beckett G, Wilkinson E, Gold A, Toft A. Lack of effect of thyroxine in
5 patients with Graves' hyperthyroidism who are treated with an antithyroid drug. *New*
6 *England Journal of Medicine*. 1996; 334(4):220-224
- 7 87. McTiernan AM, Weiss NS, Daling JR. Incidence of thyroid cancer in women in
8 relation to previous exposure to radiation therapy and history of thyroid disease.
9 *Journal of the National Cancer Institute*. 1984; 73(3):575-81
- 10 88. Menconi F, Marinò M, Pinchera A, Rocchi R, Mazzi B, Nardi M et al. Effects of total
11 thyroid ablation versus near-total thyroidectomy alone on mild to moderate Graves'
12 orbitopathy treated with intravenous glucocorticoids. *Journal of Clinical Endocrinology*
13 *and Metabolism*. 2007; 92(5):1653-1658
- 14 89. Metso S, Auvinen A, Huhtala H, Salmi J, Oksala H, Jaatinen P. Erratum: Increased
15 cancer incidence after radioiodine treatment for hyperthyroidism. *Cancer*. 2007;
16 110(8):1875
- 17 90. Metso S, Auvinen A, Huhtala H, Salmi J, Oksala H, Jaatinen P. Increased cancer
18 incidence after radioiodine treatment for hyperthyroidism. *Cancer*. 2007;
19 109(10):1972-9
- 20 91. Metso S, Jaatinen P, Huhtala H, Auvinen A, Oksala H, Salmi J. Erratum: Increased
21 cardiovascular and cancer mortality after radioiodine treatment for hyperthyroidism.
22 *Journal of Clinical Endocrinology and Metabolism*. 2007; 92(10):4008
- 23 92. Metso S, Jaatinen P, Huhtala H, Auvinen A, Oksala H, Salmi J. Increased
24 cardiovascular and cancer mortality after radioiodine treatment for hyperthyroidism.
25 *Journal of Clinical Endocrinology and Metabolism*. 2007; 92(6):2190-6
- 26 93. Metso S, Jaatinen P, Huhtala H, Luukkaala T, Oksala H, Salmi J. Long-term follow-up
27 study of radioiodine treatment of hyperthyroidism. *Clinical Endocrinology*. 2004;
28 61(5):641-648
- 29 94. Miranda-Padua ML, Cunanan EC, Kho SA, Marcelo M, Torres JF, Monzon OP et al.
30 A randomized double-blind comparison of fixed versus calculated radioiodine dose in
31 the treatment of graves' hyperthyroidism. *Phillippine Journal of Internal Medicine*.
32 2014; 52(3)
- 33 95. Müller PE, Bein B, Robens E, Bein HS, Spelsberg F. Thyroid surgery according to
34 Enderlen-Hotz or Dunhill: a comparison of two surgical methods for the treatment of
35 Graves' disease. *International Surgery*. 2001; 86(2):112-116
- 36 96. Nakamura H, Noh JY, Itoh K, Fukata S, Miyauchi A, Hamada N. Comparison of
37 methimazole and propylthiouracil in patients with hyperthyroidism caused by Graves'
38 disease. *Journal of Clinical Endocrinology and Metabolism*. 2007; 92(6):2157-2162
- 39 97. National Institute for Health and Care Excellence. Developing NICE guidelines: the
40 manual [updated October 2018]. London. National Institute for Health and Care
41 Excellence, 2014. Available from:
42 <http://www.nice.org.uk/article/PMG20/chapter/1%20Introduction%20and%20overview>
- 43 98. Nedrebo BG, Holm PI, Uhrlving S, Sorheim JI, Skeie S, Eide GE et al. Predictors of
44 outcome and comparison of different drug regimens for the prevention of relapse in
45 patients with Graves' disease. *European Journal of Endocrinology*. 2002; 147(5):583-
46 589

- 1 99. Noh JY, Sato S, Suzuki M, Yasuda S, Matsumoto M, Kunii Y et al. Comparison of
2 efficacy and adverse effects between methimazole 15 mg+inorganic iodine 38
3 mg/day and methimazole 30 mg/day as initial therapy for graves' disease patients
4 with moderate to severe hyperthyroidism. *Thyroid*. 2015; 25(1):43-50
- 5 100. Orsini F, Traino AC, Grosso M, Guidoccio F, Boni G, Volterrani D et al.
6 Personalization of radioiodine treatment for Graves' disease: a prospective,
7 randomized study with a novel method for calculating the optimal ¹³¹I-iodide activity
8 based on target reduction of thyroid mass. *Quarterly Journal of Nuclear Medicine and*
9 *Molecular Imaging*. 2012; 56(6):496-502
- 10 101. Patel NN, Abraham P, Buscombe J, Vanderpump MP. The cost effectiveness of
11 treatment modalities for thyrotoxicosis in a U.K. center. *Thyroid*. 2006; 16(6):593-8
- 12 102. Peixoto MC, Buescu A, Goncalves MRB, Albernaz MDS, Coeli CM, Vaisman M.
13 Antithyroid drugs for the treatment of graves disease: a randomized clinical trial.
14 *Endocrinologist*. 2006; 16(6):344-348
- 15 103. Peters H, Fischer C, Bogner U, Reiners C, Schleusener H. Radioiodine therapy of
16 Graves' hyperthyroidism: standard vs. calculated ¹³¹I activity. Results from a
17 prospective, randomized, multicentre study. *European Journal of Clinical*
18 *Investigation*. 1995; 25(3):186-193
- 19 104. Peters H, Fischer C, Bogner U, Reiners C, Schleusener H. Reduction in thyroid
20 volume after radioiodine therapy of Graves' hyperthyroidism: results of a prospective,
21 randomized, multicentre study. *European Journal of Clinical Investigation*. 1996;
22 26(1):59-63
- 23 105. Peters H, Fischer C, Bogner U, Reiners C, Schleusener H. Treatment of Graves'
24 hyperthyroidism with radioiodine: results of a prospective randomized study. *Thyroid*.
25 1997; 7(2):247-251
- 26 106. Pfeilschifter J, Ziegler R. Suppression of serum thyrotropin with thyroxine in patients
27 with Graves' disease: effects on recurrence of hyperthyroidism and thyroid volume.
28 *European Journal of Endocrinology*. 1997; 136(1):81-86
- 29 107. Pirnat E, Zaletel K, Gaber??ek S, Hojker S. The outcome of ¹³¹I treatment in Graves'
30 patients pretreated or not with methimazole. *Hellenic Journal of Nuclear Medicine*.
31 2011; 14(1):25-29
- 32 108. Pusuwan P, Tuntawiroon M, Sritongkul N, Chaudakshetrin P, Nopmaneejumruslers
33 C, Komoltri C et al. A prospective randomized study of the efficacy and cost-
34 effectiveness of high and low dose regimens of I-131 treatment in hyperthyroidism.
35 Chotmaiht thangkaet [Journal of the Medical Association of Thailand]. 2011;
36 94(3):361-368
- 37 109. Raber W, Kmen E, Waldhäusl W, Vierhapper H. Medical therapy of Graves' disease:
38 effect on remission rates of methimazole alone and in combination with
39 triiodothyronine. *European Journal of Endocrinology*. 2000; 142(2):117-124
- 40 110. Reinwein D, Benker G, Lazarus JH, Alexander WD. A prospective randomized trial of
41 antithyroid drug dose in Graves' disease therapy. European Multicenter Study Group
42 on Antithyroid Drug Treatment. *Journal of Clinical Endocrinology and Metabolism*.
43 1993; 76(6):1516-1521
- 44 111. Rittmaster RS, Abbott EC, Douglas R, Givner ML, Lehmann L, Reddy S et al. Effect
45 of methimazole, with or without L-thyroxine, on remission rates in Graves' disease.
46 *Journal of Clinical Endocrinology and Metabolism*. 1998; 83(3):814-818

- 1 112. Rokni H, Sadeghi R, Moossavi Z, Treglia G, Zakavi SR. Efficacy of different protocols of
2 radioiodine therapy for treatment of toxic nodular goiter: systematic review and
3 meta-analysis of the literature. *International Journal of Endocrinology and*
4 *Metabolism*. 2014; 12(2):e14424
- 5 113. Romaldini JH, Bromberg N, Werner RS, Tanaka LM, Rodrigues HF, Werner MC et al.
6 Comparison of effects of high and low dosage regimens of antithyroid drugs in the
7 management of Graves' hyperthyroidism. *Journal of Clinical Endocrinology and*
8 *Metabolism*. 1983; 57(3):563-570
- 9 114. Ron E, Doody MM, Becker DV, Brill AB, Curtis RE, Goldman MB et al. Cancer
10 mortality following treatment for adult hyperthyroidism. *Journal of the American*
11 *Medical Association*. 1998; 280(4):347-355
- 12 115. Ryodi E, Metso S, Jaatinen P, Huhtala H, Saaristo R, Valimaki M et al. Cancer
13 Incidence and Mortality in Patients Treated Either With RAI or Thyroidectomy for
14 Hyperthyroidism. *Journal of Clinical Endocrinology and Metabolism*. 2015;
15 100(10):3710-7
- 16 116. Santos RB, Romaldini JH, Ward LS. Propylthiouracil reduces the effectiveness of
17 radioiodine treatment in hyperthyroid patients with Graves' disease. *Thyroid*. 2004;
18 14(7):525-530
- 19 117. Santos RB, Romaldini JH, Ward LS. A randomized controlled trial to evaluate the
20 effectiveness of 2 regimens of fixed iodine (¹³¹I) doses for Graves disease
21 treatment. *Clinical Nuclear Medicine*. 2012; 37(3):241-244
- 22 118. Sapienza MT, Coura-Filho GB, Willegaignon J, Watanabe T, Duarte PS, Buchpiguel
23 CA. Clinical and Dosimetric Variables Related to Outcome After Treatment of Graves'
24 Disease With 550 and 1110 MBq of ¹³¹I: results of a Prospective Randomized Trial.
25 *Clinical Nuclear Medicine*. 2015; 40(9):715-719
- 26 119. Schneider P, Biko J, Hänscheid H, Hilliger S, Koutsampelas C, Kranzfelder M et al.
27 The route of administration (oral vs intravenous) does not influence dose or outcome
28 in Graves' disease and unifocal autonomy. *European Journal of Nuclear Medicine*
29 *and Molecular Imaging*. 2005; 32(7):788-793
- 30 120. Singer RB. Long-term comparative cancer mortality after use of radio-iodine in the
31 treatment of hyperthyroidism, a fully reported multicenter study. *Journal of Insurance*
32 *Medicine (Seattle)*. 2001; 33(2):138-42
- 33 121. Singer RB. Long-term comparative mortality in hyperthyroid patients treated with
34 radio-iodine, a cohort study in England. *Journal of Insurance Medicine (Seattle)*.
35 2001; 33(2):133-7
- 36 122. Singhal T, Bansal S, Singhal A, McDonald S, Bal CS. Adjunctive antithyroid drugs in
37 radioiodine therapy for hyperthyroidism. *Cochrane Database of Systematic Reviews*
38 2014, Issue 7. Art. No.: CD005447. DOI: 10.1002/14651858.CD005447.pub2.
- 39 123. Taïeb D, Bournaud C, Eberle MC, Catargi B, Schwartz C, Cavarec MB et al. Quality of
40 life, clinical outcomes and safety of early prophylactic levothyroxine administration in
41 patients with Graves' hyperthyroidism undergoing radioiodine therapy: a randomized
42 controlled study. *European Journal of Endocrinology*. 2016; 174(4):491-502
- 43 124. Tallstedt L, Lundell G, Tørring O, Wallin G, Ljunggren JG, Blomgren H et al.
44 Occurrence of ophthalmopathy after treatment for Graves' hyperthyroidism. The
45 Thyroid Study Group. *New England Journal of Medicine*. 1992; 326(26):1733-1738
- 46 125. Thientunyakit T, Thongmak S, Premprapha T. Comparative evaluation of two different
47 dosage calculation protocols of iodine-131 in the treatment of hyperthyroidism.

- 1 Chotmaihet thangphaet [Journal of the Medical Association of Thailand]. 2010;
2 93(8):969-977
- 3 126. Tian R, Kuang A. A study comparing 131I versus 131I plus antithyroid drug in the
4 management of Graves' disease. Journal of West China University of medical
5 sciences. 2001; 32(3):449-451
- 6 127. Törring O, Tallstedt L, Wallin G, Lundell G, Ljunggren JG, Taube A et al. Graves'
7 hyperthyroidism: treatment with antithyroid drugs, surgery, or radioiodine--a
8 prospective, randomized study. Thyroid Study Group. Journal of Clinical
9 Endocrinology and Metabolism. 1996; 81(8):2986-2993
- 10 128. Träisk F, Tallstedt L, Abraham-Nordling M, Andersson T, Berg G, Calissendorff J et
11 al. Thyroid-associated ophthalmopathy after treatment for Graves' hyperthyroidism
12 with antithyroid drugs or iodine-131. Journal of Clinical Endocrinology and
13 Metabolism. 2009; 94(10):3700-3707
- 14 129. Unalp HR, Erbil Y, Akguner T, Kamer E, Derici H, Issever H. Does near total
15 thyroidectomy offer advantage over total thyroidectomy in terms of postoperative
16 hypocalcemia? International Journal of Surgery (London, England). 2009; 7(2):120-
17 125
- 18 130. Verburg FA, Luster M, Lassmann M, Reiners C. 131-I therapy in patients with benign
19 thyroid disease does not conclusively lead to a higher risk of subsequent
20 malignancies. Nuklearmedizin. 2011; 50(3):93-9; quiz N20
- 21 131. Walter MA, Christ-Crain M, Schindler C, Müller-Brand J, Müller B. Outcome of
22 radioiodine therapy without, on or 3 days off carbimazole: a prospective interventional
23 three-group comparison. European Journal of Nuclear Medicine and Molecular
24 Imaging. 2006; 33(6):730-737
- 25 132. Wang J, Qin L. Radioiodine therapy versus antithyroid drugs in Graves' disease: a
26 meta-analysis of randomized controlled trials. British Journal of Radiology. 2016; 89
- 27 133. Weetman AP, Pickerill AP, Watson P, Chatterjee VK, Edwards OM. Treatment of
28 Graves' disease with the block-replace regimen of antithyroid drugs: the effect of
29 treatment duration and immunogenetic susceptibility on relapse. QJM: An
30 International journal of Medicine. 1994; 87(6):337-341
- 31 134. Witte J, Goretzki PE, Dotzenrath C, Simon D, Felis P, Neubauer M et al. Surgery for
32 Graves' disease: total versus subtotal thyroidectomy-results of a prospective
33 randomized trial. World Journal of Surgery. 2000; 24(11):1303-1311
- 34 135. Yousefi V, Bahreynian A, Sheikholeslami F, Tohidi M, Mehrabi Y, Azizi F. Effect of
35 long-term continuous methimazole treatment of thyrotoxicosis: comparison with
36 radioiodine. Iranian journal of endocrinology and metabolism. 2011; 12(5):466-
37 475+555
- 38 136. Yuan J, Lu X, Yue Y. Comparison of curative effect of 131I and antithyroid drugs in
39 Graves' disease: a meta analysis. Minerva Endocrinologica. 2017; Epublication

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Appendices

Appendix A: Review protocols

Table 13: Review protocol: Management of Thyrotoxicosis: Drugs vs Surgery vs Radioactive iodine

ID	Field	Content
I	Review question	<p>What is the clinical and cost effectiveness of using radioactive iodine vs antithyroid drugs (ATD) vs surgery to treat thyrotoxicosis secondary to Graves' disease?</p> <p>What is the clinical and cost effectiveness of using radioactive iodine vs surgery to treat thyrotoxicosis secondary to toxic nodular goitre?</p> <p>When antithyroid drugs are used, what is the most clinically and cost-effective way of using these drugs to treat thyrotoxicosis (for example choice of drugs, different treatment regimens)?</p> <p>When radioactive iodine is used, what is the most clinically and cost-effective way of using this treatment to treat thyrotoxicosis (for example different dosing strategies)?</p> <p>When surgery is indicated, what is the most clinically and cost-effective way of using surgery to treat thyrotoxicosis (for example total vs subtotal thyroidectomy)?</p>
II	Type of review question	<p>Intervention</p> <p>A review of health economic evidence related to the same review question was conducted in parallel with this review. For details see the health economic review protocol for this NICE guideline.</p>
III	Objective of the review	Provide clinically and cost effective recommendations on how to manage thyrotoxicosis
IV	Eligibility criteria – population / disease / condition / issue / domain	People diagnosed with thyrotoxicosis (TSH below normal reference ranges, free T3/T4 above normal reference range)
V	Eligibility criteria – intervention(s) / exposure(s) / prognostic factor(s)	<ul style="list-style-type: none"> • Radioactive iodine <ul style="list-style-type: none"> ○ Fixed administered activity strategy vs calculated absorbed radiation dose strategy ○ Pre-/post- treatment with ATD vs no pre-/post- treatment • Antithyroid drugs <ul style="list-style-type: none"> ○ Carbimazole/methimazole vs propylthiouracil ○ Block and replace (including levothyroxine) vs titration regimen ○ Duration of treatment: 6-<12 months vs 12-18 months vs >18 months • Surgery <ul style="list-style-type: none"> ○ Total thyroidectomy vs subtotal thyroidectomy vs near total (Dunhill) thyroidectomy vs one sided only (hemithyroidectomy/lobectomy/isthmectomy)
VI	Eligibility criteria – comparator(s)	<ul style="list-style-type: none"> • Comparisons between modalities • Comparisons between submodalities

	/ control or reference (gold) standard	
VII	Outcomes and prioritisation	<p>Critical</p> <ul style="list-style-type: none"> • Mortality (dichotomous, ≥ 1 year) • Quality of life (continuous) <p>Important (general)</p> <ul style="list-style-type: none"> • Thyroid ophthalmopathy (dichotomous) • Euthyroidism (dichotomous) • Hypothyroidism (dichotomous) • Relapse of hyperthyroidism (dichotomous) • Cardiovascular morbidity (ischaemic heart disease, dichotomous) • Arrhythmia (dichotomous) • Osteoporosis (dichotomous) • Cognitive impairment (dichotomous) • Pain (continuous) • Symptom scores (continuous) • Patient/family/carer experience (continuous) • Healthcare contacts (rates/dichotomous) <p>Important (surgical)</p> <ul style="list-style-type: none"> • Recurrent laryngeal nerve damage (dichotomous) • Hypocalcaemia (dichotomous) • Hypoparathyroidism (dichotomous) • Bleeding (dichotomous) • Infection (dichotomous) <p>Important (pharmacological)</p> <ul style="list-style-type: none"> • Agranulocytosis (dichotomous) • Liver failure (dichotomous) • Minor drug related adverse effects (dichotomous) • Teratogenesis (dichotomous) <p>Important (radioactive iodine)</p> <ul style="list-style-type: none"> • Infertility (dichotomous) • Malignancy (dichotomous) • Thyrotoxic storm (dichotomous) • Growth abnormalities (dichotomous) • Hypocalcaemia (dichotomous) • Hypoparathyroidism (dichotomous) • Teratogenesis (dichotomous) <p>Minimum duration as for the minimum duration for inclusion of studies unless specified.</p>
VIII	Eligibility criteria – study design	<ul style="list-style-type: none"> • Minimum follow-up of 3 months • RCTs • Non-randomised cohort studies to be considered if adjusted for key confounders (age, co-existing conditions, baseline T4, size of goitre) and insufficient RCTs evidence found, on an intervention by intervention basis
IX	Other inclusion / exclusion criteria	<ul style="list-style-type: none"> • Excluding studies in pregnancy • Excluding studies aimed specifically at treating thyroid eye disease • Excluding studies in context of thyroid malignancy

X	Proposed sensitivity / subgroup analysis, or meta-regression	<p>Stratifications</p> <ul style="list-style-type: none"> • Age – young children (0-4), children and young people (4-18), adults (>18-65), older adults (>65) • For antithyroid drugs vs radioactive iodine vs surgery - Cause of thyrotoxicosis (Graves' disease, toxic nodular goitre, thyroiditis) • Treatment stage – naïve/general (non-naïve, downgraded for indirectness), second line (remain symptomatic despite previous treatment, as defined by studies) <p>Subgroup analyses</p> <ul style="list-style-type: none"> • Gender (male only vs female only) • Age subdivisions (4-12, 12-18, 18-50, 50-65, 65-85, >85) • Comparison not under investigation (for example for block and replace vs titration, if some studies use methimazole and others use propylthiouracil)
XI	Selection process – duplicate screening / selection / analysis	<ul style="list-style-type: none"> • A sample of at least 10% of the abstract lists were double-sifted by a senior research fellow and discrepancies rectified, with committee input where consensus could not be reached, for more information please see the separate Methods report for this guideline.
XII	Data management (software)	<p>Pairwise meta-analyses were performed using Cochrane Review Manager (RevMan5). GRADEpro was used to assess the quality of evidence for each outcome. Endnote was used for bibliography, citations, sifting and reference management</p>
XIII	Information sources – databases and dates	<ul style="list-style-type: none"> • Medline, Embase and the Cochrane Library
XIV	Identify if an update	Not an update
XV	Author contacts	https://www.nice.org.uk/guidance/indevelopment/gid-ng10074
XVI	Highlight if amendment to previous protocol	Not amendment
XVI I	Search strategy – for one database	For details please see Appendix B:.
XVI II	Data collection process – forms / duplicate	A standardised evidence table format will be used, and published as appendix D of the evidence report.
XIX	Data items – define all variables to be collected	For details please see evidence tables in Appendix D: (clinical evidence tables) or Appendix H: (health economic evidence tables).
XX	Methods for assessing bias at outcome / study level	<p>Standard study checklists were used to critically appraise individual studies. For details please see section 6.2 of Developing NICE guidelines: the manual</p> <p>The risk of bias across all available evidence was evaluated for each outcome using an adaptation of the 'Grading of Recommendations Assessment, Development and Evaluation (GRADE) toolbox' developed by the international GRADE working group</p>

		http://www.gradeworkinggroup.org/
XXI	Criteria for quantitative synthesis	For details please see section 6.4 of Developing NICE guidelines: the manual.
XXI I	Methods for quantitative analysis – combining studies and exploring (in)consistency	For details please see the separate Methods report for this guideline.
XXI II	Meta-bias assessment – publication bias, selective reporting bias	For details please see section 6.2 of Developing NICE guidelines: the manual.
XXI V	Confidence in cumulative evidence	For details please see sections 6.4 and 9.1 of Developing NICE guidelines: the manual.
XX V	Rationale / context – what is known	For details please see the introduction to the evidence review.
XX VI	Describe contributions of authors and guarantor	A multidisciplinary committee [to add link to history page of the guideline after publication] developed the evidence review. The committee was convened by the National Guideline Centre (NGC) and chaired by Sarah Fishburn in line with section 3 of Developing NICE guidelines: the manual. Staff from NGC undertook systematic literature searches, appraised the evidence, conducted meta-analysis and cost-effectiveness analysis where appropriate, and drafted the evidence review in collaboration with the committee. For details please see Developing NICE guidelines: the manual.
XX VII	Sources of funding / support	NGC is funded by NICE and hosted by the Royal College of Physicians.
XX VIII	Name of sponsor	NGC is funded by NICE and hosted by the Royal College of Physicians.
XXI X	Roles of sponsor	NICE funds NGC to develop guidelines for those working in the NHS, public health and social care in England.
XX X	PROSPERO registration number	Not registered

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Table 14: Review protocol: Radioactive iodine safety

ID	Field	Content
I	Review question	What are the long term adverse events of radioactive iodine treatment for thyrotoxicosis?
II	Type of review question	Intervention A review of health economic evidence related to the same review question was conducted in parallel with this review. For details see the health economic review protocol for this NICE guideline.
III	Objective of the review	To determine the long term adverse event profile of radioactive iodine treatment for thyrotoxicosis
IV	Eligibility criteria – population / disease / condition / issue / domain	<ul style="list-style-type: none"> • No population restrictions (see below for prioritising of evidence)
V	Eligibility criteria – intervention(s) / exposure(s) / prognostic factor(s)	<ul style="list-style-type: none"> • Radioactive iodine
VI	Eligibility criteria – comparator(s) / control or reference (gold) standard	<ul style="list-style-type: none"> • Antithyroid drug treatment • Surgical treatment • Healthy controls (see below for prioritising of evidence)
VII	Outcomes and prioritisation	<ul style="list-style-type: none"> • Cancer <ul style="list-style-type: none"> ○ Overall ○ Organs group specific • Infertility
VIII	Eligibility criteria – study design	<ul style="list-style-type: none"> • Evidence will be considered according to the following hierarchy: <ul style="list-style-type: none"> ○ Cohort studies with hyperthyroid controls and adequate adjustment for key confounders (age, smoking) ○ Cohort studies with hyperthyroid controls and without adequate adjustment for key confounders ○ Cohort studies with healthy controls and adequate adjustment for key confounders (age, smoking) ○ Cohort studies with healthy controls without adequate adjustment for key confounders
IX	Other inclusion exclusion criteria	<ul style="list-style-type: none"> • Only included if: <ul style="list-style-type: none"> ○ For adults sample size >1000 ○ Length of follow-up >5 years
X	Proposed sensitivity / subgroup analysis, or meta-regression	<p>Stratifications</p> <ul style="list-style-type: none"> • Age – infants (<4), children and young people (4-18), adults (>18-65), older adults (>65) <p>Subgroup analyses</p> <ul style="list-style-type: none"> • Dose of radioactive iodine – fixed administered activity 200-<400 MBq, fixed 400-800 MBq, calculated absorbed dose strategy
XI	Selection process –	<ul style="list-style-type: none"> • A sample of at least 10% of the abstract lists were double-sifted by a senior research fellow and discrepancies rectified, with committee input

	duplicate screening / selection / analysis	where consensus could not be reached, for more information please see the separate Methods report for this guideline.
XII	Data management (software)	<ul style="list-style-type: none"> • Pairwise meta-analyses were performed using Cochrane Review Manager (RevMan5). • GRADEpro was used to assess the quality of evidence for each outcome. • Endnote was used for bibliography, citations, sifting and reference management
XIII	Information sources – databases and dates	<ul style="list-style-type: none"> • Medline (OVID), Embase (OVID and the Cochrane Library (Wiley)
XIV	Identify if an update	Not an update
XV	Author contacts	https://www.nice.org.uk/guidance/indevelopment/gid-ng10074
XVI	Highlight if amendment to previous protocol	Not an amendment
XVI I	Search strategy – for one database	For details please see Appendix B:
XVI II	Data collection process – forms / duplicate	A standardised evidence table format will be used, and published as Appendix D: of the evidence report.
XIX	Data items – define all variables to be collected	For details please see evidence tables in Appendix D: (clinical evidence tables) or Appendix H: (health economic evidence tables).
XX	Methods for assessing bias at outcome / study level	<p>Standard study checklists were used to critically appraise individual studies. For details please see section 6.2 of Developing NICE guidelines: the manual</p> <p>The risk of bias across all available evidence was evaluated for each outcome using an adaptation of the 'Grading of Recommendations Assessment, Development and Evaluation (GRADE) toolbox' developed by the international GRADE working group http://www.gradeworkinggroup.org/</p>
XXI	Criteria for quantitative synthesis	For details please see section 6.4 of Developing NICE guidelines: the manual.
XXI I	Methods for quantitative analysis – combining studies and exploring (in)consistency	For details please see the separate Methods report for this guideline.
XXI II	Meta-bias assessment – publication bias, selective reporting bias	For details please see section 6.2 of Developing NICE guidelines: the manual.
XXI	Confidence in	For details please see sections 6.4 and 9.1 of Developing NICE

V	cumulative evidence	guidelines: the manual.
XX V	Rationale / context – what is known	For details please see the introduction to the evidence review.
XX VI	Describe contributions of authors and guarantor	A multidisciplinary committee [to add link to history page of the guideline after publication] developed the evidence review. The committee was convened by the National Guideline Centre (NGC) and chaired by Sarah Fishburn in line with section 3 of Developing NICE guidelines: the manual. Staff from NGC undertook systematic literature searches, appraised the evidence, conducted meta-analysis and cost-effectiveness analysis where appropriate, and drafted the evidence review in collaboration with the committee. For details please see Developing NICE guidelines: the manual.
XX VII	Sources of funding / support	NGC is funded by NICE and hosted by the Royal College of Physicians.
XX VIII	Name of sponsor	NGC is funded by NICE and hosted by the Royal College of Physicians.
XXI X	Roles of sponsor	NICE funds NGC to develop guidelines for those working in the NHS, public health and social care in England.
XX X	PROSPERO registration number	Not registered

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Table 15: Health economic review protocol

Review question	All questions – health economic evidence
Objectives	To identify health economic studies relevant to any of the review questions.
Search criteria	<ul style="list-style-type: none"> • Populations, interventions and comparators must be as specified in the clinical review protocol above. • Studies must be of a relevant health economic study design (cost–utility analysis, cost-effectiveness analysis, cost–benefit analysis, cost–consequences analysis, comparative cost analysis). • Studies must not be a letter, editorial or commentary, or a review of health economic evaluations. (Recent reviews will be ordered although not reviewed. The bibliographies will be checked for relevant studies, which will then be ordered.) • Unpublished reports will not be considered unless submitted as part of a call for evidence. • Studies must be in English.
Search strategy	A health economic study search will be undertaken using population-specific terms and a health economic study filter – see appendix B below.
Review strategy	<p>Studies not meeting any of the search criteria above will be excluded. Studies published before 2003, abstract-only studies and studies from non-OECD countries or the USA will also be excluded.</p> <p>Each remaining study will be assessed for applicability and methodological limitations using the NICE economic evaluation checklist which can be found in appendix H of Developing NICE guidelines: the manual (2014).⁹⁷</p> <p>Inclusion and exclusion criteria</p> <ul style="list-style-type: none"> • If a study is rated as both ‘Directly applicable’ and with ‘Minor limitations’ then it will be included in the guideline. A health economic evidence table will be completed and it will be included in the health economic evidence profile. • If a study is rated as either ‘Not applicable’ or with ‘Very serious limitations’ then it will usually be excluded from the guideline. If it is excluded then a health economic evidence table will not be completed and it will not be included in the health economic evidence profile. • If a study is rated as ‘Partially applicable’, with ‘Potentially serious limitations’ or both then there is discretion over whether it should be included. <p>Where there is discretion</p> <p>The health economist will make a decision based on the relative applicability and quality of the available evidence for that question, in discussion with the guideline committee if required. The ultimate aim is to include health economic studies that are helpful for decision-making in the context of the guideline and the current NHS setting. If several studies are considered of sufficiently high applicability and methodological quality that they could all be included, then the health economist, in discussion with the committee if required, may decide to include only the most applicable studies and to selectively exclude the remaining studies. All studies excluded on the basis of applicability or methodological limitations will be listed with explanation in the excluded health economic studies appendix below.</p> <p>The health economist will be guided by the following hierarchies.</p> <p><i>Setting:</i></p> <ul style="list-style-type: none"> • UK NHS (most applicable). • OECD countries with predominantly public health insurance systems (for example, France, Germany, Sweden). • OECD countries with predominantly private health insurance systems (for example, Switzerland).

- Studies set in non-OECD countries or in the USA will be excluded before being assessed for applicability and methodological limitations.

Health economic study type:

- Cost–utility analysis (most applicable).
- Other type of full economic evaluation (cost–benefit analysis, cost-effectiveness analysis, cost–consequences analysis).
- Comparative cost analysis.
- Non-comparative cost analyses including cost-of-illness studies will be excluded before being assessed for applicability and methodological limitations.

Year of analysis:

- The more recent the study, the more applicable it will be.
- Studies published in 2003 or later but that depend on unit costs and resource data entirely or predominantly from before 2003 will be rated as ‘Not applicable’.
- Studies published before 2003 will be excluded before being assessed for applicability and methodological limitations.

Quality and relevance of effectiveness data used in the health economic analysis:

- The more closely the clinical effectiveness data used in the health economic analysis match with the outcomes of the studies included in the clinical review the more useful the analysis will be for decision-making in the guideline.

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Appendix B: Literature search strategies

The literature searches for this review are detailed below and complied with the methodology outlined in Developing NICE guidelines: the manual 2014, updated 2018
<https://www.nice.org.uk/guidance/pmg20/resources/developing-nice-guidelines-the-manual-pdf-72286708700869>

For more detailed information, please see the Methodology Review. [\[Add cross reference\]](#)

B.1 Clinical search literature search strategy

Searches were constructed using a PICO framework where population (P) terms were combined with Intervention (I) and in some cases Comparison (C) terms. Outcomes (O) are rarely used in search strategies for interventions as these concepts may not be well described in title, abstract or indexes and therefore difficult to retrieve. Search filters were applied to the search where appropriate.

Table 16: Database date parameters and filters used

Database	Dates searched	Search filter used
Medline (OVID)	1946 – 07 January 2019	Exclusions Randomised controlled trials Systematic review studies Observational studies
Embase (OVID)	1974 – 07 January 2019	Exclusions Randomised controlled trials Systematic review studies Observational studies
The Cochrane Library (Wiley)	Cochrane Reviews to 2019 Issue 1 or 12 CENTRAL to 2019 Issue 1 or 12 DARE, and NHSEED to 2015 Issue 2 of 4 HTA to 2016 Issue 2 of 4	None

Medline (Ovid) search terms

1.	exp goiter/
2.	exp Hyperthyroidism/
3.	(hyperthyroid* or thyrotoxicosis).ti,ab.
4.	(toxic adj4 (node* or nodul* or multi?nodul* or goitre or goiter)).ti,ab.
5.	(graves' disease or plummer's disease).ti,ab.
6.	5
7.	letter/
8.	editorial/
9.	news/
10.	exp historical article/
11.	Anecdotes as Topic/
12.	comment/
13.	case report/
14.	(letter or comment*).ti.

15.	or/7-14
16.	randomized controlled trial/ or random*.ti,ab.
17.	15 not 16
18.	animals/ not humans/
19.	exp Animals, Laboratory/
20.	exp Animal Experimentation/
21.	exp Models, Animal/
22.	exp Rodentia/
23.	(rat or rats or mouse or mice).ti.
24.	or/17-23
25.	randomized controlled trial.pt.
26.	controlled clinical trial.pt.
27.	randomi#ed.ti,ab.
28.	placebo.ab.
29.	randomly.ti,ab.
30.	Clinical Trials as topic.sh.
31.	trial.ti.
32.	or/25-31
33.	Meta-Analysis/
34.	exp Meta-Analysis as Topic/
35.	(meta analy* or metanaly* or metaanaly* or meta regression).ti,ab.
36.	((systematic* or evidence*) adj3 (review* or overview*)).ti,ab.
37.	(reference list* or bibliograph* or hand search* or manual search* or relevant journals).ab.
38.	(search strategy or search criteria or systematic search or study selection or data extraction).ab.
39.	(search* adj4 literature).ab.
40.	(medline or pubmed or cochrane or embase or psychlit or psyclit or psychinfo or psycinfo or cinahl or science citation index or bids or cancerlit).ab.
41.	cochrane.jw.
42.	((multiple treatment* or indirect or mixed) adj2 comparison*).ti,ab.
43.	or/33-42
44.	Epidemiologic studies/
45.	Observational study/
46.	exp Cohort studies/
47.	(cohort adj (study or studies or analys* or data)).ti,ab.
48.	((follow up or observational or uncontrolled or non randomi#ed or epidemiologic*) adj (study or studies or data)).ti,ab.
49.	((longitudinal or retrospective or prospective or cross sectional) and (study or studies or review or analys* or cohort* or data)).ti,ab.
50.	Controlled Before-After Studies/
51.	Historically Controlled Study/
52.	Interrupted Time Series Analysis/
53.	(before adj2 after adj2 (study or studies or data)).ti,ab.
54.	or/4-53
55.	exp case control study/
56.	case control*.ti,ab.

57.	or/55-56
58.	54 or 57
59.	Cross-sectional studies/
60.	(cross sectional and (study or studies or review or analys* or cohort* or data)).ti,ab.
61.	or/59-60
62.	54 or 61
63.	54 or 57 or 61
64.	6 not 24
65.	limit 64 to English language
66.	65 and (32 or 43 or 64)

1

Embase (Ovid) search terms

1.	goiter/
2.	hyperthyroidism/ or graves disease/ or thyrotoxicosis/ or toxic goiter/
3.	(hyperthyroid* or thyrotoxicosis).ti,ab.
4.	(toxic adj4 (node* of nodul* or multi?nodul* or goitre or goiter)).ti,ab.
5.	(graves' disease or plummer's disease).ti,ab.
6.	or/1-5
7.	letter.pt. or letter/
8.	note.pt.
9.	editorial.pt.
10.	case report/ or case study/
11.	(letter or comment*).ti.
12.	or/7-11
13.	randomized controlled trial/ or random*.ti,ab.
14.	12 not 13
15.	animal/ not human/
16.	nonhuman/
17.	exp Animal Experiment/
18.	exp Experimental Animal/
19.	animal model/
20.	exp Rodent/
21.	(rat or rats or mouse or mice).ti.
22.	or/14-21
23.	6 not 22
24.	random*.ti,ab.
25.	factorial*.ti,ab.
26.	(crossover* or cross over*).ti,ab.
27.	((doubl* or singl*) adj blind*).ti,ab.
28.	(assign* or allocat* or volunteer* or placebo*).ti,ab.
29.	crossover procedure/
30.	single blind procedure/
31.	randomized controlled trial/
32.	double blind procedure/
33.	or/24-32
34.	systematic review/

35.	meta-analysis/
36.	(meta analy* or metanaly* or metaanaly* or meta regression).ti,ab.
37.	((systematic* or evidence*) adj3 (review* or overview*)).ti,ab.
38.	(reference list* or bibliograph* or hand search* or manual search* or relevant journals).ab.
39.	(search strategy or search criteria or systematic search or study selection or data extraction).ab.
40.	(search* adj4 literature).ab.
41.	(medline or pubmed or cochrane or embase or psychlit or psyclit or psychinfo or psycinfo or cinahl or science citation index or bids or cancerlit).ab.
42.	cochrane.jw.
43.	((multiple treatment* or indirect or mixed) adj2 comparison*).ti,ab.
44.	or/34-43
45.	Clinical study/
46.	Observational study/
47.	family study/
48.	longitudinal study/
49.	retrospective study/
50.	prospective study/
51.	cohort analysis/
52.	follow-up/
53.	cohort*.ti,ab.
54.	52 and 53
55.	(cohort adj (study or studies or analys* or data)).ti,ab.
56.	((follow up or observational or uncontrolled or non randomi#ed or epidemiologic*) adj (study or studies or data)).ti,ab.
57.	((longitudinal or retrospective or prospective or cross sectional) and (study or studies or review or analys* or cohort* or data)).ti,ab.
58.	(before adj2 after adj2 (study or studies or data)).ti,ab.
59.	or/45-51,54-58
60.	exp case control study/
61.	case control*.ti,ab.
62.	or/60-61
63.	59 or 62
64.	cross-sectional study/
65.	(cross sectional and (study or studies or review or analys* or cohort* or data)).ti,ab.
66.	or/64-65
67.	59 or 66
68.	59 or 62 or 66
69.	23 and (33 or 44 or 68)
70.	limit 69 to English language

1

Cochrane Library (Wiley) search terms

#1.	MeSH descriptor: [Goiter] explode all trees
#2.	MeSH descriptor: [Hyperthyroidism] explode all trees
#3.	(hyperthyroid* or thyrotoxicosis):ti,ab
#4.	(toxic near/4 (node* or nodul* or multinodul* or multi-nodul* or goitre or goiter)):ti,ab

#5.	MeSH descriptor: [Graves Disease] explode all trees
#6.	(grave* near/4 (thyrotoxicos* or hyperthyr*)):ti,ab
#7.	graves' disease:ti,ab
#8.	(or #1-#7)

1 B.2 Health Economics literature search strategy

2 Health economic evidence was identified by conducting a broad search relating to a thyroid
3 disease population in NHS Economic Evaluation Database (NHS EED – this ceased to be
4 updated after March 2015) and the Health Technology Assessment database (HTA) with no
5 date restrictions. NHS EED and HTA databases are hosted by the Centre for Research and
6 Dissemination (CRD). Additional searches were run on Medline and Embase for health
7 economics, economic modelling and quality of life studies.

8 **Table 17: Database date parameters and filters used**

Database	Dates searched	Search filter used
Medline	2014 – 07 January 2019	Exclusions Health economics studies Health economics modelling studies Quality of life studies
Embase	2014 – 07 January 2019	Exclusions Health economics studies Health economics modelling studies Quality of life studies
Centre for Research and Dissemination (CRD)	HTA - Inception – 07 January 2019 NHSEED - Inception to March 2015	None

9 **Medline (Ovid) search terms**

1.	exp thyroid diseases/
2.	hyperthyroid*.ti,ab.
3.	hypothyroid*.ti,ab.
4.	thyrotoxicosis.ti,ab.
5.	(thyroid adj3 (swell* or dysfunction* or enlarg* or nodule* or node* or disease* or condition* or disorder*)):ti,ab.
6.	or/1-5
7.	letter/
8.	editorial/
9.	news/
10.	exp historical article/
11.	Anecdotes as Topic/
12.	comment/
13.	case report/
14.	(letter or comment*).ti.
15.	or/7-14
16.	randomized controlled trial/ or random*.ti,ab.

17.	15 not 16
18.	animals/ not humans/
19.	exp Animals, Laboratory/
20.	exp Animal Experimentation/
21.	exp Models, Animal/
22.	exp Rodentia/
23.	(rat or rats or mouse or mice).ti.
24.	or/17-23
25.	6 not 24
26.	limit 25 to English language
27.	Economics/
28.	Value of life/
29.	exp "Costs and Cost Analysis"/
30.	exp Economics, Hospital/
31.	exp Economics, Medical/
32.	Economics, Nursing/
33.	Economics, Pharmaceutical/
34.	exp "Fees and Charges"/
35.	exp Budgets/
36.	budget*.ti,ab.
37.	cost*.ti.
38.	(economic* or pharmaco?economic*).ti.
39.	(price* or pricing*).ti,ab.
40.	(cost* adj2 (effective* or utilit* or benefit* or minimi* or unit* or estimat* or variable*)),ab.
41.	(financ* or fee or fees).ti,ab.
42.	(value adj2 (money or monetary)).ti,ab.
43.	or/27-42
44.	exp models, economic/
45.	*Models, Theoretical/
46.	*Models, Organizational/
47.	markov chains/
48.	monte carlo method/
49.	exp Decision Theory/
50.	(markov* or monte carlo).ti,ab.
51.	econom* model*.ti,ab.
52.	(decision* adj2 (tree* or analy* or model*)).ti,ab.
53.	or/44-52
54.	quality-adjusted life years/
55.	sickness impact profile/
56.	(quality adj2 (wellbeing or well being)).ti,ab.
57.	sickness impact profile.ti,ab.
58.	disability adjusted life.ti,ab.
59.	(qal* or qtime* or qwb* or daly*).ti,ab.
60.	(euroqol* or eq5d* or eq 5*).ti,ab.

61.	(qol* or hql* or hqol* or h qol* or hrqol* or hr qol*).ti,ab.
62.	(health utility* or utility score* or disutilit* or utility value*).ti,ab.
63.	(hui or hui1 or hui2 or hui3).ti,ab.
64.	(health* year* equivalent* or hye or hyes).ti,ab.
65.	discrete choice*.ti,ab.
66.	rosser.ti,ab.
67.	(willingness to pay or time tradeoff or time trade off or tto or standard gamble*).ti,ab.
68.	(sf36* or sf 36* or short form 36* or shortform 36* or shortform36*).ti,ab.
69.	(sf20 or sf 20 or short form 20 or shortform 20 or shortform20).ti,ab.
70.	(sf12* or sf 12* or short form 12* or shortform 12* or shortform12*).ti,ab.
71.	(sf8* or sf 8* or short form 8* or shortform 8* or shortform8*).ti,ab.
72.	(sf6* or sf 6* or short form 6* or shortform 6* or shortform6*).ti,ab.
73.	or/54-72
74.	26 and (43 or 53 or 73)

1

Embase (Ovid) search terms

1.	exp thyroid diseases/
2.	hyperthyroid*.ti,ab.
3.	hypothyroid*.ti,ab.
4.	thyrotoxicosis*.ti,ab.
5.	(thyroid adj3 (swell* or dysfunction* or enlarg* or nodule* or node* or disease* or condition* or disorder*)).ti,ab.
6.	or/1-5
7.	letter.pt. or letter/
8.	note.pt.
9.	editorial.pt.
10.	case report/ or case study/
11.	(letter or comment*).ti.
12.	or/7-11
13.	randomized controlled trial/ or random*.ti,ab.
14.	12 not 13
15.	animal/ not human/
16.	nonhuman/
17.	exp Animal Experiment/
18.	exp Experimental Animal/
19.	animal model/
20.	exp Rodent/
21.	(rat or rats or mouse or mice).ti.
22.	or/14-21
23.	6 not 22
24.	limit 23 to English language
25.	health economics/
26.	exp economic evaluation/

27.	exp health care cost/
28.	exp fee/
29.	budget/
30.	funding/
31.	budget*.ti,ab.
32.	cost*.ti.
33.	(economic* or pharmaco?economic*).ti.
34.	(price* or pricing*).ti,ab.
35.	(cost* adj2 (effective* or utilit* or benefit* or minimi* or unit* or estimat* or variable*)).ab.
36.	(financ* or fee or fees).ti,ab.
37.	(value adj2 (money or monetary)).ti,ab.
38.	or/25-37
39.	statistical model/
40.	exp economic aspect/
41.	39 and 40
42.	*theoretical model/
43.	*nonbiological model/
44.	stochastic model/
45.	decision theory/
46.	decision tree/
47.	monte carlo method/
48.	(markov* or monte carlo).ti,ab.
49.	econom* model*.ti,ab.
50.	(decision* adj2 (tree* or analy* or model*)).ti,ab.
51.	or/41-50
52.	quality adjusted life year/
53.	"quality of life index"/
54.	short form 12/ or short form 20/ or short form 36/ or short form 8/
55.	sickness impact profile/
56.	(quality adj2 (wellbeing or well being)).ti,ab.
57.	sickness impact profile.ti,ab.
58.	disability adjusted life.ti,ab.
59.	(qal* or qtime* or qwb* or daly*).ti,ab.
60.	(euroqol* or eq5d* or eq 5*).ti,ab.
61.	(qol* or hql* or hqol* or h qol* or hrqol* or hr qol*).ti,ab.
62.	(health utility* or utility score* or disutilit* or utility value*).ti,ab.
63.	(hui or hui1 or hui2 or hui3).ti,ab.
64.	(health* year* equivalent* or hye or hyes).ti,ab.
65.	discrete choice*.ti,ab.
66.	rosser.ti,ab.

67.	(willingness to pay or time tradeoff or time trade off or tto or standard gamble*).ti,ab.
68.	(sf36* or sf 36* or short form 36* or shortform 36* or shortform36*).ti,ab.
69.	(sf20 or sf 20 or short form 20 or shortform 20 or shortform20).ti,ab.
70.	(sf12* or sf 12* or short form 12* or shortform 12* or shortform12*).ti,ab.
71.	(sf8* or sf 8* or short form 8* or shortform 8* or shortform8*).ti,ab.
72.	(sf6* or sf 6* or short form 6* or shortform 6* or shortform6*).ti,ab.
73.	or/52-72
74.	24 and (38 or 51 or 73)

1

NHS EED and HTA (CRD) search terms

#1.	MeSH DESCRIPTOR Thyroid Diseases EXPLODE ALL TREES
#2.	hyperthyroid*
#3.	hypothyroid*
#4.	thyrotoxicosis*
#5.	(thyroid adj3 (swell* or dysfunction* or enlarg* or nodule* or node* or disease* or condition* or disorder*))
#6.	#1 OR #2 OR #3 OR #4 or #5

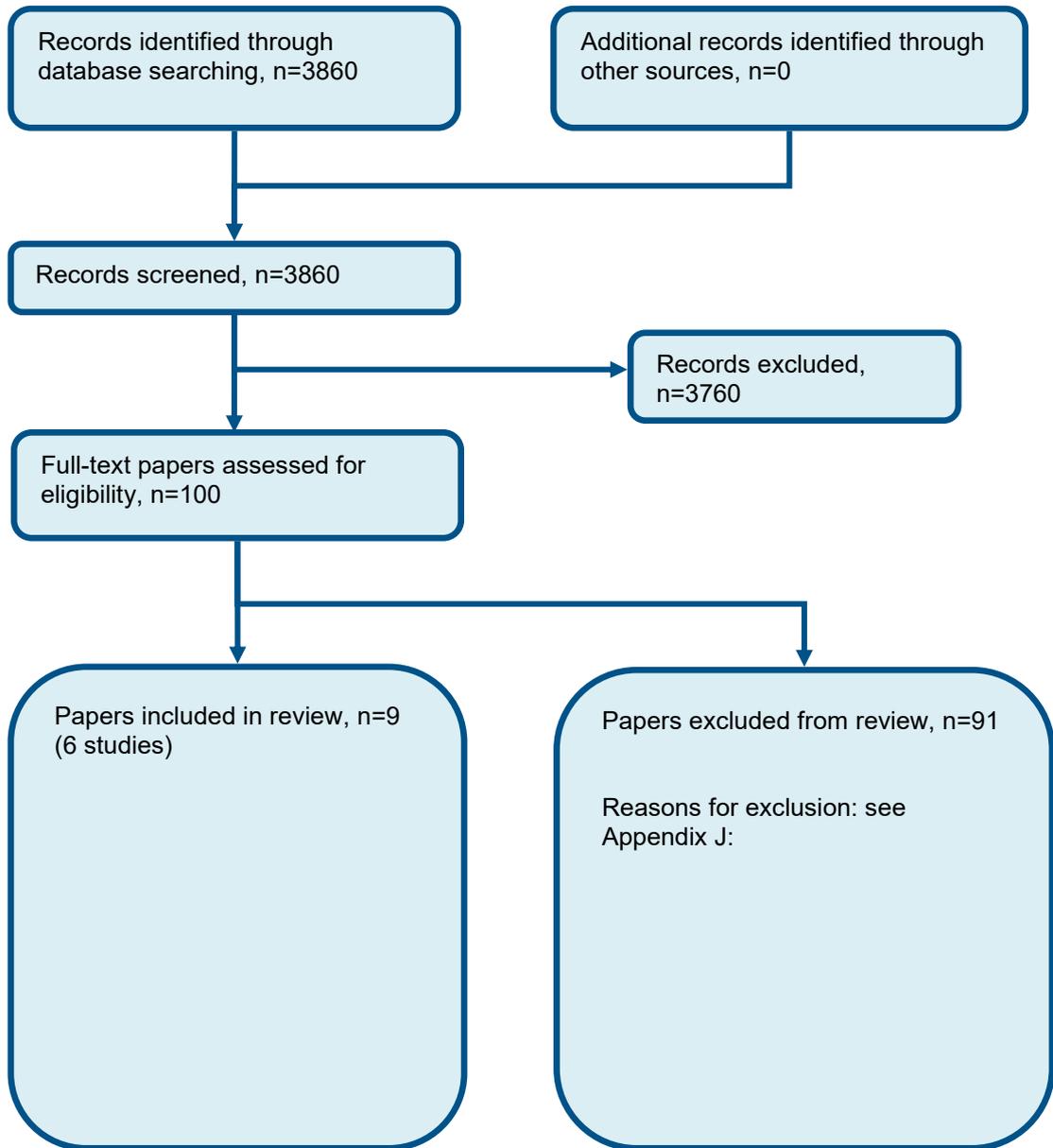
2

3

4

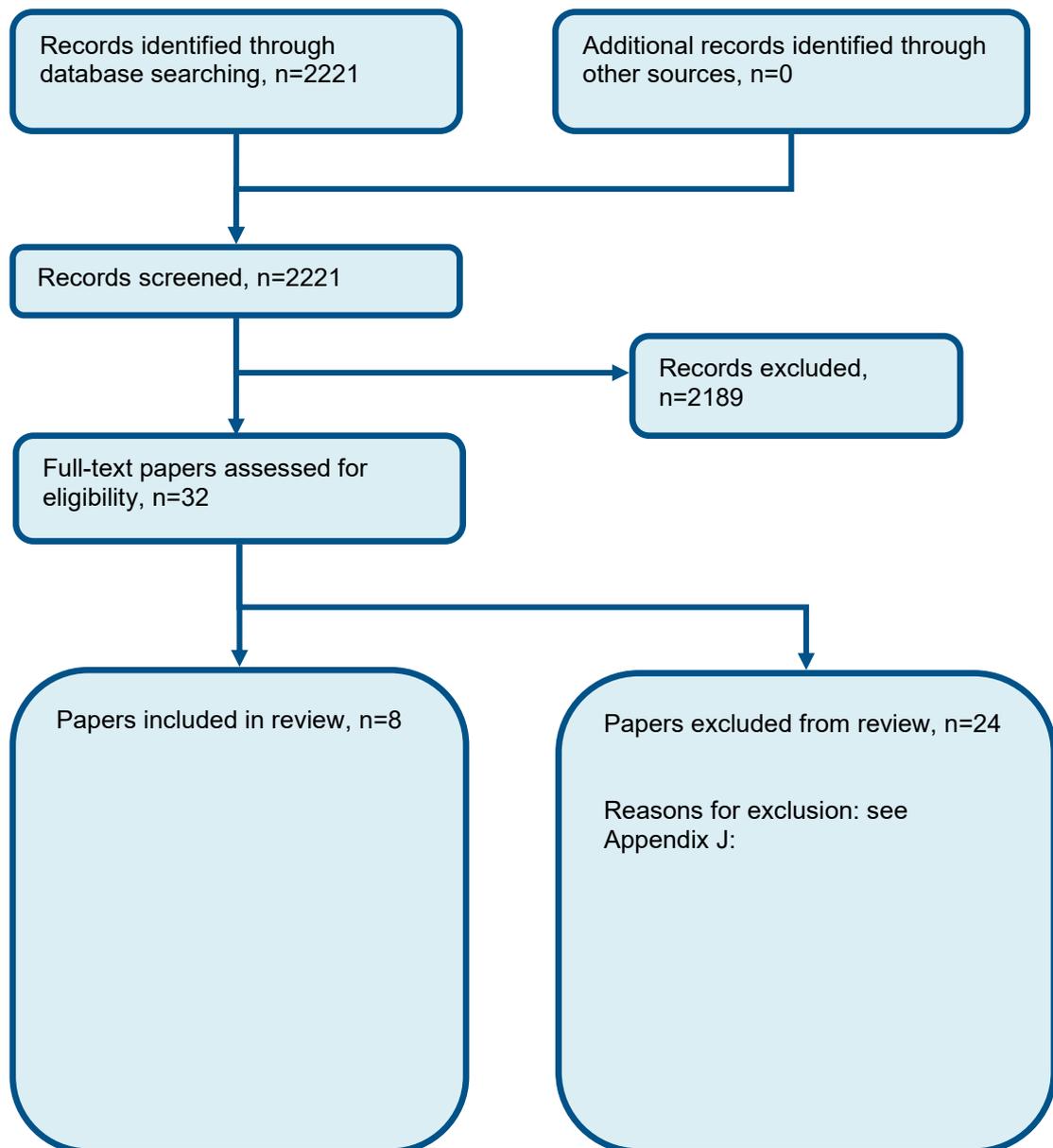
Appendix C: Clinical evidence selection

Figure 1: Flow chart of clinical study selection for the review of thyrotoxicosis (drugs vs surgery vs radioactive iodine)



1
2

Figure 2: Flow chart of clinical study selection for the review of radioactive iodine safety



Appendix D: Clinical evidence tables

D.1 Drugs vs Surgery vs Radioactive Iodine

Study	Azizi 2005 ¹¹
Study type	RCT (Patient randomised; Parallel)
Number of studies (number of participants)	1 (n=104)
Countries and setting	Conducted in Iran; Setting: Not specified
Line of therapy	2nd line
Duration of study	Intervention + follow up: 10 years
Method of assessment of guideline condition	Adequate method of assessment/diagnosis:
Stratum	Failed first line treatment
Subgroup analysis within study	Not applicable
Inclusion criteria	Older than 40, diffuse toxic goitre (Graves'), treated to euthyroidism with MMI for at least 18 months, relapse to hyperthyroidism within 1 year of discontinuation
Exclusion criteria	Did not accept randomisation
Recruitment/selection of patients	Not specified
Age, gender and ethnicity	Age - Mean (SD): 48 (6). Gender (M:F): 69/16. Ethnicity: Not stated
Further population details	1. Age: 18-50 2. Gender: Not stated / Unclear
Indirectness of population	No indirectness

Interventions	<p>(n=52) Intervention 1: Antithyroid drugs. MMI, 10mg twice daily for first month, 10mg daily during second month, maintenance of 2.5-10mg daily from third month on, no discontinuation specified. Duration 10 years. Concurrent medication/care: Usual care</p> <p>(n=52) Intervention 2: Radioactive iodine. Calculated activity based on thyroid weight and iodine uptake, mean dose delivered 7.9 mCi. Duration 10 years. Concurrent medication/care: Usual care. Indirectness: No indirectness</p>
Funding	Funding not stated
RESULTS (NUMBERS ANALYSED) AND RISK OF BIAS FOR COMPARISON: RADIOACTIVE IODINE versus ANTITHYROID DRUGS	
<p>Protocol outcome 1: Euthyroidism - Actual outcome for Failed first line treatment: Euthyroidism at end of follow-up at 10 years; Group 1: 16/41, Group 2: 26/34 Risk of bias: All domain - Very high, Selection - High, Blinding - Low, Incomplete outcome data - High, Outcome reporting - Low, Measurement - High, Crossover - Low; Indirectness of outcome: No indirectness ; Group 1 Number missing: 11, Reason: 10 lost to follow-up, 1 did not accept randomisation; Group 2 Number missing: 24, Reason: 6 lost to follow-up, 18 did not accept randomisation</p> <p>Protocol outcome 2: Hypothyroidism - Actual outcome for Failed first line treatment: Hypothyroidism at end of follow-up at 10 years; Group 1: 25/41, Group 2: 1/28 Risk of bias: All domain - Very high, Selection - High, Blinding - Low, Incomplete outcome data - High, Outcome reporting - Low, Measurement - High, Crossover - Low; Indirectness of outcome: No indirectness ; Group 1 Number missing: 11, Reason: 10 lost to follow-up, 1 did not accept randomisation; Group 2 Number missing: 24, Reason: 6 lost to follow-up, 18 did not accept randomisation</p> <p>Protocol outcome 3: Relapse of hyperthyroidism - Actual outcome for Failed first line treatment: Hyperthyroidism at end of follow-up at 10 years; Group 1: 0/41, Group 2: 1/28 Risk of bias: All domain - Very high, Selection - High, Blinding - Low, Incomplete outcome data - High, Outcome reporting - Low, Measurement - High, Crossover - Low; Indirectness of outcome: No indirectness ; Group 1 Number missing: 11, Reason: 10 lost to follow-up, 1 did not accept randomisation; Group 2 Number missing: 24, Reason: 6 lost to follow-up, 18 did not accept randomisation</p> <p>Protocol outcome 4: Agranulocytosis - Actual outcome for Failed first line treatment: Agranulocytosis at 10 years; Group 1: 0/41, Group 2: 0/28 Risk of bias: All domain - Very high, Selection - High, Blinding - Low, Incomplete outcome data - High, Outcome reporting - Low, Measurement - High, Crossover - Low; Indirectness of outcome: No indirectness ; Group 1 Number missing: 11, Reason: 10 lost to follow-up, 1 did not accept randomisation; Group 2 Number missing: 24, Reason: 6 lost to follow-up, 18 did not accept randomisation</p>	
Protocol outcomes not reported by the study	Quality of life ; Mortality ; Thyroid ophthalmopathy ; Ischaemic heart disease ; Heart failure ; Arrhythmia ; Osteoporosis ; Impaired cognitive function ; Growth ; Pain ; Svmotom scores ; Experience of care ; Healthcare

contacts ; Recurrent laryngeal nerve damage ; Hypocalcaemia ; Hypoparathyroidism ; Bleeding ; Infection ; Liver failure ; Minor drug related adverse events ; Teratogenesis ; Infertility ; Malignancy ; Thyrotoxic storm

Study	Bartalena 1998 ¹⁶
Study type	RCT (Patient randomised; Parallel)
Number of studies (number of participants)	1 (n=443)
Countries and setting	Conducted in Italy; Setting: Not specified
Line of therapy	1st line
Duration of study	Intervention + follow up: 2.5 years
Method of assessment of guideline condition	Adequate method of assessment/diagnosis
Stratum	Treatment naive/general population
Subgroup analysis within study	Not applicable
Inclusion criteria	Graves' disease, mild or no ophthalmopathy
Exclusion criteria	Severe ophthalmopathy, large goitres, CI to glucocorticoid treatment
Recruitment/selection of patients	Not specified
Age, gender and ethnicity	Age - Mean (range): 42 (15-85). Gender (M:F): 20:80. Ethnicity:
Further population details	1. Age: 18-50 2. Gender: Not applicable
Extra comments	~50% with ophthalmopathy
Indirectness of population	No indirectness
Interventions	(n=150) Intervention 1: Radioactive iodine. MMI was discontinued 5 days before administration of RAI, with dose of 120-150uCi per gram of thyroid tissue, if hypo or hyperthyroid after treatment - corrected with levothyroxine/MMI as appropriate. Duration 1 year. Concurrent medication/care: All given MMI for 3 to 4 months (70% had been given prior to trial achieving euthyroidism in roughly 1/3rd of the 70%). Indirectness: No indirectness (n=148) Intervention 2: Antithyroid drugs. Methimazole given at lowest dose that achieved euthyroidism, no discontinuation specified. Duration 1 year. Concurrent medication/care: Usual care. Indirectness: No indirectness
Funding	Funding not stated
RESULTS (NUMBERS ANALYSED) AND RISK OF BIAS FOR COMPARISON: RADIOACTIVE IODINE versus ANTITHYROID DRUGS	
Protocol outcome 1: Thvroid ophthalmopathv	

- Actual outcome for Treatment naive/general population: Development or worsening of thyroid ophthalmopathy at 1 year; Group 1: 23/150, Group 2: 4/148
 Risk of bias: All domain - Low, Selection - Low, Blinding - Low, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low;
 Indirectness of outcome: No indirectness ; Group 1 Number missing: ; Group 2 Number missing:

Protocol outcome 2: Euthyroidism

- Actual outcome for Treatment naive/general population: Euthyroidism at end of follow-up (including RAI patients requiring levothyroxine/MMI) at 1 year; Group 1: 128/150, Group 2: 145/148
 Risk of bias: All domain - High, Selection - Low, Blinding - High, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low;
 Indirectness of outcome: No indirectness ; Group 1 Number missing: ; Group 2 Number missing:

Protocol outcome 3: Hypothyroidism

- Actual outcome for Treatment naive/general population: Hypothyroidism at end of follow-up (including RAI patients requiring levothyroxine/MMI) at 1 year; Group 1: 20/150, Group 2: 2/148
 Risk of bias: All domain - High, Selection - Low, Blinding - High, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low;
 Indirectness of outcome: No indirectness ; Group 1 Number missing: ; Group 2 Number missing:
 - Actual outcome for Treatment naive/general population: Hyperthyroidism at end of follow-up (including RAI patients requiring levothyroxine/MMI) at 1 year; Group 1: 2/150, Group 2: 1/148
 Risk of bias: All domain - High, Selection - Low, Blinding - High, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low;
 Indirectness of outcome: No indirectness ; Group 1 Number missing: ; Group 2 Number missing:

Protocol outcomes not reported by the study

Quality of life ; Mortality ; Relapse of hyperthyroidism ; Ischaemic heart disease ; Heart failure ; Arrhythmia ; Osteoporosis ; Impaired cognitive function ; Growth ; Pain ; Symptom scores ; Experience of care ; Healthcare contacts ; Recurrent laryngeal nerve damage ; Hypocalcaemia ; Hypoparathyroidism ; Bleeding ; Infection ; Agranulocytosis ; Liver failure ; Minor drug related adverse events ; Teratogenesis ; Infertility ; Malignancy ; Thyrotoxic storm

Study	Chen 2009 ²⁸
Study type	RCT (Patient randomised; Parallel)
Number of studies (number of participants)	1 (n=460)
Countries and setting	Conducted in China; Setting: Not stated
Line of therapy	1st line
Duration of study	Intervention + follow up: 9 years
Method of assessment of guideline condition	Adequate method of assessment/diagnosis
Stratum	Overall
Subgroup analysis within study	Not applicable
Inclusion criteria	Newly diagnosed hyperthyroidism, no previous thyroid treatment, 24 hour uptake of ¹³¹ I >40%
Exclusion criteria	Severe liver or kidney damage, agranulocytosis, pregnancy or lactation, less than 8 years of age
Recruitment/selection of patients	Screened 2021, excluding 1519 with previous treatment, others refused or exclusion criteria
Age, gender and ethnicity	Age - Mean (SD): 37 (14). Gender (M:F): 33:67. Ethnicity:
Further population details	1. Age: 18-50 2. Gender: Systematic review: mixed
Extra comments	75% GD, 23% MNTG, 2% UNTG
Indirectness of population	No indirectness
Interventions	<p>(n=230) Intervention 1: Antithyroid drugs. Either MMI or PTU, for at least 18 months, initial dose based on severity of symptoms and titrated throughout to TSH. If recurrence after withdrawal, reinstated. . Duration 9 years . Concurrent medication/care: All also received propranolol as necessary. Advised to restrict iodine rich foods in diet. Examined 2-4 weekly in first year, 3-6 months thereafter if stable. Indirectness: No indirectness</p> <p>(n=230) Intervention 2: Radioactive iodine. No pre-treatment with ATD. Therapeutic activity from 1.85-4.44MBq per gram thyroid/lesion weight, calculated activity (based on weight, and 24hr iodine uptake). Maximum activity limited to 555MBq. At 3 months and 6 months determined if 2nd (10%) or 3rd (2.5%) treatment required. Duration 9 years. Concurrent medication/care: All also received propranolol as necessary. Advised to restrict iodine rich foods in diet. Examined 2-4 weekly in first year, 3-6 months thereafter if stable. Indirectness: No indirectness</p>
Funding	Academic or government funding

RESULTS (NUMBERS ANALYSED) AND RISK OF BIAS FOR COMPARISON: RADIOACTIVE IODINE versus MMI/PTU

Protocol outcome 1: Mortality

- Actual outcome for Treatment naive/general population: Mortality at 9 years; Group 1: 0/209, Group 2: 0/177

Risk of bias: All domain - High, Selection - Low, Blinding - Low, Incomplete outcome data - High, Outcome reporting - Low, Measurement - Low, Crossover - Low;
Indirectness of outcome: No indirectness ; Group 1 Number missing: 21, Reason: Loss to follow up; Group 2 Number missing: 53, Reason: 16 loss to follow-up, 37 excluded due to AEs

Protocol outcome 2: Thyroid ophthalmopathy

- Actual outcome for Treatment naive/general population: New cases of thyroid ophthalmopathy at 9 years; Group 1: 26/151, Group 2: 14/138

Risk of bias: All domain - High, Selection - Low, Blinding - Low, Incomplete outcome data - High, Outcome reporting - Low, Measurement - Low, Crossover - Low;
Indirectness of outcome: No indirectness ; Group 1 Number missing: 21, Reason: Loss to follow up; Group 2 Number missing: 53, Reason: 16 loss to follow-up, 37 excluded due to AEs

Protocol outcome 3: Euthyroidism

- Actual outcome for Treatment naive/general population: Normal T3+T4, no medication required at 9 years; Group 1: 146/209, Group 2: 73/177

Risk of bias: All domain - High, Selection - Low, Blinding - Low, Incomplete outcome data - High, Outcome reporting - Low, Measurement - Low, Crossover - Low;
Indirectness of outcome: No indirectness ; Group 1 Number missing: 21, Reason: Loss to follow up; Group 2 Number missing: 53, Reason: 16 loss to follow-up, 37 excluded due to AEs

Protocol outcome 4: Hypothyroidism

- Actual outcome for Treatment naive/general population: Clinical hypothyroidism, abnormal T3/T4 and TSH at 9 years; Group 1: 19/209, Group 2: 6/177

Risk of bias: All domain - High, Selection - Low, Blinding - Low, Incomplete outcome data - High, Outcome reporting - Low, Measurement - Low, Crossover - Low;
Indirectness of outcome: No indirectness ; Group 1 Number missing: 21, Reason: Loss to follow up; Group 2 Number missing: 53, Reason: 16 loss to follow-up, 37 excluded due to AEs

Protocol outcome 5: Relapse of hyperthyroidism

- Actual outcome for Treatment naive/general population: Relapse or persistence of hyperthyroidism, abnormal T3/T4 or TSH at 9 years; Group 1: 18/209, Group 2: 88/177

Risk of bias: All domain - High, Selection - Low, Blinding - Low, Incomplete outcome data - High, Outcome reporting - Low, Measurement - Low, Crossover - Low;
Indirectness of outcome: No indirectness ; Group 1 Number missing: 21, Reason: Loss to follow up; Group 2 Number missing: 53, Reason: 16 loss to follow-up, 37 excluded due to AEs

Protocol outcome 6: Agranulocytosis

- Actual outcome for Treatment naive/general population: Agranulocytosis at 9 years; Group 1: 0/209, Group 2: 7/214

Risk of bias: All domain - High. Selection - Low. Blinding - Low. Incomplete outcome data - High. Outcome reporting - Low. Measurement - Low. Crossover - Low:

Indirectness of outcome: No indirectness ; Group 1 Number missing: 21, Reason: Loss to follow up; Group 2 Number missing: 16, Reason: Loss to follow up

Protocol outcome 7: Liver failure

- Actual outcome for Treatment naive/general population: Severe liver damage at 9 years; Group 1: 0/209, Group 2: 5/214

Risk of bias: All domain - High, Selection - Low, Blinding - Low, Incomplete outcome data - High, Outcome reporting - Low, Measurement - Low, Crossover - Low;

Indirectness of outcome: No indirectness ; Group 1 Number missing: 21, Reason: Loss to follow up; Group 2 Number missing: 16, Reason: Loss to follow up

Protocol outcome 8: Malignancy

- Actual outcome for Treatment naive/general population: Malignancy at 9 years; Group 1: 0/209, Group 2: 0/177

Risk of bias: All domain - High, Selection - Low, Blinding - Low, Incomplete outcome data - High, Outcome reporting - Low, Measurement - Low, Crossover - Low;

Indirectness of outcome: No indirectness ; Group 1 Number missing: 21, Reason: Loss to follow up; Group 2 Number missing: 53, Reason: 16 loss to follow-up, 37 excluded due to AEs

- Actual outcome for Treatment naive/general population: Thyroid storm at 9 years; Group 1: 0/209, Group 2: 0/177

Risk of bias: All domain - High, Selection - Low, Blinding - Low, Incomplete outcome data - High, Outcome reporting - Low, Measurement - Low, Crossover - Low;

Indirectness of outcome: No indirectness ; Group 1 Number missing: 21, Reason: Loss to follow up; Group 2 Number missing: 53, Reason: 16 loss to follow-up, 37 excluded due to AEs

Protocol outcomes not reported by the study

Quality of life ; Ischaemic heart disease ; Heart failure ; Arrhythmia ; Osteoporosis ; Impaired cognitive function ; Growth ; Pain ; Symptom scores ; Experience of care ; Healthcare contacts ; Recurrent laryngeal nerve damage ; Hypocalcaemia ; Hypoparathyroidism ; Bleeding ; Infection ; Minor drug related adverse events ; Teratogenesis ; Infertility ; Thyrotoxic storm

Study	Kansara 2017 ⁷⁰
Study type	RCT (Patient randomised; Parallel)
Number of studies (number of participants)	1 (n=60)
Countries and setting	Conducted in India; Setting: Tertiary level referral centre
Line of therapy	1st line
Duration of study	Intervention + follow up: 1 year
Method of assessment of guideline condition	Adequate method of assessment/diagnosis
Stratum	Overall
Subgroup analysis within study	Not applicable
Inclusion criteria	20-50 years old, treatment naive
Exclusion criteria	History of thyroid disease, significant ophthalmopathy (clinical activity score >1), malignancy, previous exposure to RAI, known systemic disorders, long term use of corticosteroids or insulin
Age, gender and ethnicity	Age - Mean (SD): 33 (4.2). Gender (M:F): Not stated. Ethnicity:
Further population details	1. Age: 18-50 2. Gender: Not stated / Unclear
Indirectness of population	No indirectness
Interventions	(n=30) Intervention 1: Antithyroid drugs. Carbimazole, 30mg initially for 2 months, tapering as per clinical status. Duration 1 year. Concurrent medication/care: Usual care. Indirectness: No indirectness (n=30) Intervention 2: Radioactive iodine. Orally, single dose of 131I 10mCi, capsule form with water, . Duration 1 year. Concurrent medication/care: Usual care. Indirectness: No indirectness
Funding	Funding not stated

RESULTS (NUMBERS ANALYSED) AND RISK OF BIAS FOR COMPARISON: RADIOACTIVE IODINE versus CARBIMAZOLE

Protocol outcome 1: Euthyroidism

- Actual outcome for Treatment naive/general population: Biochemical and clinical euthyroidism at 1 year; Group 1: 4/28, Group 2: 22/29

Risk of bias: All domain - Low, Selection - Low, Blinding - Low, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low;

Indirectness of outcome: No indirectness ; Group 1 Number missing: 2, Reason: Lost to follow-up; Group 2 Number missing: 1, Reason: Lost to follow-up

- Actual outcome for Treatment naive/general population: Clinical hypothyroidism at 1 year: Group 1: 24/28. Group 2: 2/29

Risk of bias: All domain - Low, Selection - Low, Blinding - Low, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low;
 Indirectness of outcome: No indirectness ; Group 1 Number missing: 2, Reason: Lost to follow-up; Group 2 Number missing: 1, Reason: Lost to follow-up

Protocol outcome 2: Relapse of hyperthyroidism

- Actual outcome for Treatment naive/general population: Relapse/persistent hyperthyroidism at 1 year; Group 1: 0/28, Group 2: 0/29

Risk of bias: All domain - Low, Selection - Low, Blinding - Low, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low;
 Indirectness of outcome: No indirectness ; Group 1 Number missing: 2, Reason: Lost to follow-up; Group 2 Number missing: 1, Reason: Lost to follow-up

Protocol outcomes not reported by the study

Quality of life ; Mortality ; Thyroid ophthalmopathy ; Hypothyroidism ; Ischaemic heart disease ; Heart failure ;
 Arrhythmia ; Osteoporosis ; Impaired cognitive function ; Growth ; Pain ; Symptom scores ; Experience of care ;
 Healthcare contacts ; Recurrent laryngeal nerve damage ; Hypocalcaemia ; Hypoparathyroidism ; Bleeding ; Infection
 ; Agranulocytosis ; Liver failure ; Minor drug related adverse events ; Teratogenesis ; Infertility ; Malignancy ;
 Thyrotoxic storm

Study (subsidiary papers)	Törring 1996 ¹²⁷ (Abraham-nordling 2005 ¹ , Tallstedt 1992 ¹²⁴)
Study type	RCT (Patient randomised; Parallel)
Number of studies (number of participants)	1 (n=179)
Countries and setting	Conducted in Sweden; Setting: Not specified
Line of therapy	1st line
Duration of study	Intervention + follow up: Maximum 21 years
Method of assessment of guideline condition	Adequate method of assessment/diagnosis
Stratum	Treatment naive/general population
Subgroup analysis within study	Not applicable
Inclusion criteria	Graves' disease
Exclusion criteria	Previous thyroid disease
Recruitment/selection of patients	Not specified
Age, gender and ethnicity	Age - Mean (SD): Younger group = 29 (4), older group 45 (6). Gender (M:F): 16:84. Ethnicity:
Further population details	1. Age: Not applicable 2. Gender: Not applicable
Indirectness of population	No indirectness
Interventions	<p>(n=71) Intervention 1: Antithyroid drugs. 10mg MMI 4x daily for 18 months, thyroxine 0.1 to 0.3mg daily after 3-5 weeks to provide normal T3 and low TSH. Beta blockers given for initial weeks. Examined monthly for 2 months after initiation, then 3 monthly. After discontinuation examined twice in first year, once yearly. Duration Max 21 years follow-up. Concurrent medication/care: Usual care. Indirectness: No indirectness</p> <p>(n=67) Intervention 2: Surgery. Beta blockers before surgery for ~1 month, bilateral subtotal thyroidectomy, leaving posterior capsule and 1g or less of each lobe, thyroxine 0.1 to 0.3mg daily afterwards, seen after 5 weeks and then every 3 months during 1st year after surgery and once yearly thereafter. Duration Max 21 years follow-up. Concurrent medication/care: Usual care. Indirectness: No indirectness</p> <p>(n=41) Intervention 3: Radioactive iodine. First single oral dose of iodine 131I, dose based on size of thyroid, uptake and half-life aiming at 120Gy dose delivered. Beta blockers also given unless CI. 18 patients needed more than 1 one dose of RAI. Duration Max 21 years follow-up. Concurrent medication/care: Usual care. Indirectness: No indirectness</p>
Funding	Academic or government funding

RESULTS (NUMBERS ANALYSED) AND RISK OF BIAS FOR COMPARISON: SURGERY versus ANTITHYROID DRUGS

Protocol outcome 1: Thyroid ophthalmopathy

- Actual outcome for Treatment naive/general population: New or worsening ophthalmopathy at ~4 years follow-up; Group 1: 9/64, Group 2: 8/65

Risk of bias: All domain - Low, Selection - Low, Blinding - Low, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low;

Indirectness of outcome: No indirectness ; Group 1 Number missing: -, Reason: Not fully specified across publications; Group 2 Number missing: -, Reason: Not fully specified across publications

Protocol outcome 2: Relapse of hyperthyroidism

- Actual outcome for Treatment naive/general population: Relapse or persistence of hyperthyroidism at ~4 years follow-up; Group 1: 4/65, Group 2: 26/68

Risk of bias: All domain - Low, Selection - Low, Blinding - Low, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low;

Indirectness of outcome: No indirectness ; Group 1 Number missing: 2, Reason: Did not have surgery; Group 2 Number missing: 3, Reason: Did not comply/randomisation error

Protocol outcome 3: Osteoporosis

- Actual outcome for Treatment naive/general population: Osteoporosis (self-reported) at 14-21 years follow-up; Group 1: 8/56, Group 2: 5/55

Risk of bias: All domain - High, Selection - Low, Blinding - Low, Incomplete outcome data - High, Outcome reporting - Low, Measurement - High, Crossover - Low;

Indirectness of outcome: No indirectness ; Group 1 Number missing: -, Reason: Not fully specified across publications; Group 2 Number missing: -, Reason: Not fully specified across publications

RESULTS (NUMBERS ANALYSED) AND RISK OF BIAS FOR COMPARISON: RADIOACTIVE IODINE versus ANTITHYROID DRUGS

Protocol outcome 1: Thyroid ophthalmopathy

- Actual outcome for Treatment naive/general population: New or worsening ophthalmopathy at ~4 years follow-up; Group 1: 13/39, Group 2: 4/38

Risk of bias: All domain - Low, Selection - Low, Blinding - Low, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low;

Indirectness of outcome: No indirectness ; Group 1 Number missing: -, Reason: Not fully specified across publications; Group 2 Number missing: -, Reason: Not fully specified across publications

Protocol outcome 2: Relapse of hyperthyroidism

- Actual outcome for Treatment naive/general population: Relapse or persistence of hyperthyroidism at ~4 years follow-up; Group 1: 8/39, Group 2: 16/38

Risk of bias: All domain - Low, Selection - Low, Blinding - Low, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low;

Indirectness of outcome: No indirectness ; Group 1 Number missing: 2, Reason: Rejected assignment; Group 2 Number missing: 3, Reason: Did not comply/randomisation error

Protocol outcome 3: Osteoporosis

- Actual outcome for Treatment naive/general population: Osteoporosis (self-reported) at 14-21 years follow-up; Group 1: 6/34, Group 2: 5/36
 Risk of bias: All domain - High, Selection - Low, Blinding - Low, Incomplete outcome data - High, Outcome reporting - Low, Measurement - High, Crossover - Low;
 Indirectness of outcome: No indirectness ; Group 1 Number missing: -, Reason: Not fully specified across publications; Group 2 Number missing: -, Reason: Not fully specified across publications

RESULTS (NUMBERS ANALYSED) AND RISK OF BIAS FOR COMPARISON: RADIOACTIVE IODINE versus SURGERY

Protocol outcome 1: Thyroid ophthalmopathy

- Actual outcome for Treatment naive/general population: New or worsening ophthalmopathy at ~4 years follow-up; Group 1: 13/39, Group 2: 6/37
 Risk of bias: All domain - Low, Selection - Low, Blinding - Low, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low;
 Indirectness of outcome: No indirectness ; Group 1 Number missing: -, Reason: Not fully specified across publications; Group 2 Number missing: -, Reason: Not fully specified across publications

Protocol outcome 2: Relapse of hyperthyroidism

- Actual outcome for Treatment naive/general population: Relapse or persistence of hyperthyroidism at ~4 years follow-up; Group 1: 8/39, Group 2: 3/37
 Risk of bias: All domain - Low, Selection - Low, Blinding - Low, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low;
 Indirectness of outcome: No indirectness ; Group 1 Number missing: 2, Reason: Rejected assignment; Group 2 Number missing: 0

Protocol outcome 3: Osteoporosis

- Actual outcome for Treatment naive/general population: Osteoporosis (self-reported) at 14-21 years follow-up; Group 1: 6/34, Group 2: 7/34
 Risk of bias: All domain - High, Selection - Low, Blinding - Low, Incomplete outcome data - High, Outcome reporting - Low, Measurement - High, Crossover - Low;
 Indirectness of outcome: No indirectness ; Group 1 Number missing: -, Reason: Not fully specified across publications; Group 2 Number missing: -, Reason: Not fully specified across publications

Protocol outcomes not reported by the study

Quality of life ; Mortality ; Euthyroidism ; Hypothyroidism ; Ischaemic heart disease ; Heart failure ; Arrhythmia ; Impaired cognitive function ; Growth ; Pain ; Symptom scores ; Experience of care ; Healthcare contacts ; Recurrent laryngeal nerve damage ; Hypocalcaemia ; Hypoparathyroidism ; Bleeding ; Infection ; Agranulocytosis ; Liver failure ; Minor drug related adverse events ; Teratogenesis ; Infertility ; Malignancy ; Thyrotoxic storm

Study (subsidiary papers)	Träisk 2009 ¹²⁸ (Abraham-nordling 2010 ³)
Study type	RCT (Patient randomised; Parallel)
Number of studies (number of participants)	1 (n=313)
Countries and setting	Conducted in Sweden; Setting: Sweden, outpatients for RAI
Line of therapy	1st line
Duration of study	Intervention + follow up: 4 years
Method of assessment of guideline condition	Adequate method of assessment/diagnosis
Stratum	Treatment naive/general population
Subgroup analysis within study	Not applicable
Inclusion criteria	35-69 years old, symptomatic Graves' disease, activity of oral dose of radioactive iodine \leq 600MBq
Exclusion criteria	Previous treatment with thyroid drugs/surgery/radioactive iodine , severe ophthalmopathy, incipient toxic crisis, coronary heart disease, pregnancy, breast-feeding, pregnancy planned within 2 years
Recruitment/selection of patients	Not specified
Age, gender and ethnicity	Age - Mean (SD): 51 (8). Gender (M:F): 11:89. Ethnicity: Not stated
Further population details	1. Age: 50-65 2. Gender: Not applicable
Extra comments	Ophthalmopathy at baseline in 13%
Indirectness of population	No indirectness
Interventions	(n=163) Intervention 1: Radioactive iodine. Beta blocker pre-treatment, aim for one dose, calculated activity based on mass, estimated uptake and effective half-life. Duration 4 years. Concurrent medication/care: Usual care. Indirectness: No indirectness (n=150) Intervention 2: Antithyroid drugs. MMI given 15mg twice daily for 2 weeks, then 50ug of thyroxine added and increased to 100ug 2 weeks later. At 6 weeks adjusted to normalise T3/T4 and bring TSH to less than 0.4mIU/litre. Beta blockers used for symptomatic treatment. MMI replaced by PTU in people showing serious adverse reactions. Discontinued after 18 months, levothyroxine continued for 1 more month. Duration 4 years. Concurrent medication/care: Usual care. Indirectness: No indirectness
Funding	Academic or government funding

RESULTS (NUMBERS ANALYSED) AND RISK OF BIAS FOR COMPARISON: RADIOACTIVE IODINE versus ANTITHYROID DRUGS	
Protocol outcome 1: Thyroid ophthalmopathy	
- Actual outcome for Treatment naive/general population: Relapse of hyperthyroidism at 3 years; Group 1: 2/147, Group 2: 33/137	
Risk of bias: All domain - High, Selection - Low, Blinding - High, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low;	
Indirectness of outcome: No indirectness ; Group 1 Number missing: 16, Reason: Lost to follow-up; Group 2 Number missing: 10, Reason: Lost to follow-up	
Protocol outcomes not reported by the study	Quality of life ; Mortality ; Euthyroidism ; Hypothyroidism ; Relapse of hyperthyroidism ; Ischaemic heart disease ; Heart failure ; Arrhythmia ; Osteoporosis ; Impaired cognitive function ; Growth ; Pain ; Symptom scores ; Experience of care ; Healthcare contacts ; Recurrent laryngeal nerve damage ; Hypocalcaemia ; Hypoparathyroidism ; Bleeding ; Infection ; Agranulocytosis ; Liver failure ; Minor drug related adverse events ; Teratogenesis ; Infertility ; Malignancy ; Thyrotoxic storm

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D.2 Radioactive iodine safety

Study	Franklyn 1999 ³⁸
Study type	Non randomised study

Number of studies (number of participants)	1 (n=7417)
Countries and setting	Conducted in United Kingdom; Setting: Nil else stated
Line of therapy	1st line
Duration of study	--:
Method of assessment of guideline condition	Adequate method of assessment/diagnosis
Stratum	Fixed dose <400MBq
Subgroup analysis within study	Not applicable
Inclusion criteria	Treated with RAI in WM in UK between 1950 and 1991, did not die before 1971, found on register with ONS, not emigrated, registered with GP
Exclusion criteria	Nil else stated
Recruitment/selection of patients	Nil else stated
Age, gender and ethnicity	Age - Mean (SD): 56.6 (12.7). Gender (M:F): 17:83. Ethnicity: Not stated
Further population details	1. Age: 2. Gender:
Extra comments	Nil else stated
Indirectness of population	No indirectness
Interventions	(n=7417) Intervention 1: RAI - RAI alone. Mean dose 308MBq (SD 232). Duration Mean follow-up 9.7 years. Concurrent medication/care: Not specified . Indirectness: No indirectness (n=7417) Intervention 2: General population. Age, sex and period matched SIR from UK regional cancer registries. Duration Mean follow-up 9.7 years. Concurrent medication/care: Nil else stated. Indirectness: No indirectness
Funding	Other (Government + BUPA foundation)
RESULTS (NUMBERS ANALYSED) AND RISK OF BIAS FOR COMPARISON: RAI ALONE versus GENERAL POPULATION	
<p>Protocol outcome 1: Total cancer diagnoses - Actual outcome for Fixed dose <400MBq: All cancer diagnoses at Mean follow-up 9.7 years; RR; 0.83 (95%CI 0.77 to 0.9); Risk of bias: All domain - High, Selection - High, Blinding - Low, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low; Indirectness of outcome: No indirectness ; Group 1 Number missing: ; Group 2 Number missing:</p>	
Protocol outcomes not reported by the study	Quality of life ; Cancer diagnoses in iodine uptake glands ; Cancer diagnoses in non-iodine uptake glands ; Infertility

Study	Franklyn 2005 ⁴⁰
Study type	Non randomised study
Number of studies (number of participants)	1 (n=2668)
Countries and setting	Conducted in United Kingdom; Setting: Nil else
Line of therapy	1st line
Duration of study	Intervention + follow up: Median 5.6 years
Method of assessment of guideline condition	Adequate method of assessment/diagnosis
Stratum	Fixed dose <400MBq
Subgroup analysis within study	Not applicable
Inclusion criteria	>40, hyperthyroidism, treated in West Midlands with radioiodine between 1984 and 2002, records available
Exclusion criteria	Nil else
Recruitment/selection of patients	Not specified
Age, gender and ethnicity	Age - Median (range): 62 (40 to >80). Gender (M:F): 19:81. Ethnicity: Not stated
Further population details	1. Age: 2. Gender:
Indirectness of population	No indirectness
Interventions	(n=2668) Intervention 1: RAI - RAI alone. Fixed dose, either 185 or 370MBq, 84.3% received one dose only. Duration Median follow-up 5.6 years. Concurrent medication/care: Nil else stated. Indirectness: No indirectness (n=2668) Intervention 2: General population. From WHO databank, age, sex and year matched cohort. Duration Median follow-up 5.6 years. Concurrent medication/care: Nil else stated. Indirectness: No indirectness
Funding	Study funded by industry (Some funding from BUPA)

RESULTS (NUMBERS ANALYSED) AND RISK OF BIAS FOR COMPARISON: RAI ALONE versus GENERAL POPULATION

Protocol outcome 1: Total cancer diagnoses

- Actual outcome for Fixed dose <400MBq: Cancer mortality at 5.6 years; RR; 0.99 (95%CI 0.82 to 1.2);

Risk of bias: All domain - High, Selection - High, Blinding - Low, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low;

Indirectness of outcome: Serious indirectness ; Key confounders: Age, sex and year matched SMR; Group 1 Number missing: ; Group 2 Number missing:

Protocol outcomes not reported by the study | Quality of life ; Cancer diagnoses in iodine uptake glands ; Cancer diagnoses in non-iodine uptake glands ; Infertility

Study	Giesecke 2018 ⁴²
Study type	Non randomised study
Number of studies (number of participants)	1 (n=10992)
Countries and setting	Conducted in Sweden; Setting: Nil else
Line of therapy	1st line
Duration of study	Intervention + follow up: Mean 16.3 years
Method of assessment of guideline condition	Adequate method of assessment/diagnosis
Stratum	Cause not specified
Subgroup analysis within study	Not applicable
Inclusion criteria	RAI at Karolinska University Hospital or surgery in Stockholm between 1976 and 2013, older than 35, certain aetiology of hyperthyroidism, not treated with both RAI and surgery
Exclusion criteria	Nil else
Recruitment/selection of patients	Nil else
Age, gender and ethnicity	Age - Other: Mean for RAI 64, mean for surgery 47. Gender (M:F): 15:85. Ethnicity: Not stated
Further population details	1. Age: 2. Gender:
Extra comments	50% Graves disease in RAI arm, 63% in surgery arm
Indirectness of population	No indirectness
Interventions	(n=10250) Intervention 1: RAI - RAI alone. Dose not stated. Duration 16.3 years. Concurrent medication/care: Nil else stated. Indirectness: No indirectness (n=742) Intervention 2: ATD/SUR - SUR. No details provided. Duration 16.3 years. Concurrent medication/care: No details provided . Indirectness: No indirectness
Funding	Academic or government funding
RESULTS (NUMBERS ANALYSED) AND RISK OF BIAS FOR COMPARISON: RAI ALONE versus SUR	
Protocol outcome 1: Total cancer diagnoses	
- Actual outcome for Fixed dose <400MBq: Cancer mortality at 16.3 years follow-up: HR: 0.96 (95%CI 0.73 to 1.26):	

Risk of bias: All domain - Low, Selection - Low, Blinding - Low, Incomplete outcome data - High, Outcome reporting - Low, Measurement - Low, Crossover - Low;
Indirectness of outcome: Serious indirectness ; Group 1 Number missing: ; Group 2 Number missing:

Protocol outcomes not reported by the study	Quality of life ; Cancer diagnoses in iodine uptake glands ; Cancer diagnoses in non-iodine uptake glands ; Infertility
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Study	Goldman 1988 ⁴⁴
Study type	Non randomised study
Number of studies (number of participants)	1 (n=1762)
Countries and setting	Conducted in USA; Setting: None stated
Line of therapy	1st line
Duration of study	Intervention + follow up: 17.2 years
Method of assessment of guideline condition	Adequate method of assessment/diagnosis
Stratum	Cause not specified
Subgroup analysis within study	Not applicable
Inclusion criteria	Women with hyperthyroidism treated at MGH between 1946 and 1964 with I131
Exclusion criteria	None stated
Recruitment/selection of patients	None stated
Age, gender and ethnicity	Age - Other: Not stated. Gender (M:F): All women. Ethnicity: Not stated
Further population details	1. Age: 2. Gender:
Indirectness of population	No indirectness
Interventions	(n=1762) Intervention 1: RAI - RAI alone. None stated. Duration 17.2 years. Concurrent medication/care: None stated. Indirectness: No indirectness (n=1762) Intervention 2: General population. Age, sex, race, year matched incidence from state cancer register. Duration 17.2 years. Concurrent medication/care: None stated. Indirectness: No indirectness
Funding	Academic or government funding

RESULTS (NUMBERS ANALYSED) AND RISK OF BIAS FOR COMPARISON: RAI ALONE versus GENERAL POPULATION

Protocol outcome 1: Total cancer diagnoses

- Actual outcome for Cause not specified: Total cancer diagnoses, SIR at 17.2 years; RR; 0.8 (95%CI 0.6 to 1.1);

Risk of bias: All domain - Very high, Selection - High, Blinding - Low, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - High, Crossover - Low;

Indirectness of outcome: No indirectness ; Group 1 Number missing: ; Group 2 Number missing:

Protocol outcomes not reported by the study | Quality of life ; Cancer diagnoses in iodine uptake glands ; Cancer diagnoses in non-iodine uptake glands ; Infertility

Study	Hoffman 1982 ⁵⁹
Study type	Non randomised study
Number of studies (number of participants)	1 (n=3146)
Countries and setting	Conducted in USA; Setting: Nil else stated
Line of therapy	1st line
Duration of study	Intervention + follow up: 15 years mean for RAI, 21 years mean for surgery
Method of assessment of guideline condition	Adequate method of assessment/diagnosis
Stratum	Fixed dose <400MBq
Subgroup analysis within study	Not applicable
Inclusion criteria	White, female, treated for hyperthyroidism at Mayo clinic between 1946 and 1964, confirmed diagnosis of hyperthyroidism, no other isotope treatment, resident of USA
Exclusion criteria	Nil else stated
Recruitment/selection of patients	Nil else stated
Age, gender and ethnicity	Age - Other: Mean age at Tx 56.8 for RAI, 45.7 for surgery. Gender (M:F): Only women. Ethnicity: Only white patients
Further population details	1. Age: 2. Gender:
Extra comments	73% mild-moderate disease, ~50% Graves disease
Indirectness of population	No indirectness
Interventions	(n=1005) Intervention 1: RAI - RAI alone. Mean number of treatments 1.2, mean dose 10.6mCi (~392 MBq). Duration Mean 15 years follow-up. Concurrent medication/care: Nil else stated. Indirectness: No indirectness (n=2141) Intervention 2: ATD/SUR - SUR. Nil else stated. Duration Mean 21 years follow-up. Concurrent medication/care: Nil else stated. Indirectness: No indirectness
Funding	Funding not stated
RESULTS (NUMBERS ANALYSED) AND RISK OF BIAS FOR COMPARISON: RAI ALONE versus SUR	
Protocol outcome 1: Total cancer diagnoses - Actual outcome for Fixed dose <400MBq: Cancer incidence at all sites. adjusted for age. year of treatment and duration of follow-up at Mean follow-up 15 years for RAI.	

21 years for surgery; RR; 1.0 (95%CI 0.7 to 1.3);

Risk of bias: All domain - High, Selection - High, Blinding - Low, Incomplete outcome data - High, Outcome reporting - Low, Measurement - High, Crossover - Low;

Indirectness of outcome: No indirectness ; Group 1 Number missing: ; Group 2 Number missing:

Protocol outcomes not reported by the study

Quality of life ; Cancer diagnoses in iodine uptake glands ; Cancer diagnoses in non-iodine uptake glands ; Infertility

Study	Holm 1991 ⁶²
Study type	Non randomised study
Number of studies (number of participants)	1 (n=10207)
Countries and setting	Conducted in Sweden; Setting: Not stated
Line of therapy	1st line
Duration of study	Intervention + follow up: Mean 15 years
Method of assessment of guideline condition	Adequate method of assessment/diagnosis
Stratum	Fixed dose >400MBq
Subgroup analysis within study	Not applicable
Inclusion criteria	Under 75, treated for hyperthyroidism with RAI at one of 7 departments in Sweden, sufficient information on names and DoB
Exclusion criteria	Not stated
Recruitment/selection of patients	Not stated
Age, gender and ethnicity	Age - Mean (range): 57 (13-74). Gender (M:F): Not stated. Ethnicity: Not stated
Further population details	1. Age: 2. Gender:
Extra comments	51% with Graves' disease
Indirectness of population	No indirectness
Interventions	(n=10207) Intervention 1: RAI - RAI alone. Mean dose 506MBq, 59% received one treatment. Duration 15 years. Concurrent medication/care: Not stated. Indirectness: No indirectness (n=10207) Intervention 2: General population. Age, sex, region and year matched SIR based on Swedish Cancer register. Duration 15 years. Concurrent medication/care: Not stated. Indirectness: No indirectness
Funding	Academic or government funding
RESULTS (NUMBERS ANALYSED) AND RISK OF BIAS FOR COMPARISON: RAI ALONE versus GENERAL POPULATION	
Protocol outcome 1: Total cancer diagnoses - Actual outcome for Fixed dose >400MBq: Total cancer diagnoses. SIR at 15 year follow-up: RR: 1.10 (95%CI 1.02 to 1.17):	

Risk of bias: All domain - High, Selection - High, Blinding - Low, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low;
Indirectness of outcome: No indirectness ; Group 1 Number missing: ; Group 2 Number missing:

Protocol outcomes not reported by the study Quality of life ; Cancer diagnoses in iodine uptake glands ; Cancer diagnoses in non-iodine uptake glands ; Infertility

Study	Metso 2007 ⁹⁰
Study type	Non randomised study
Number of studies (number of participants)	1 (n=2793)
Countries and setting	Conducted in Finland; Setting: Nil else stated
Line of therapy	1st line
Duration of study	Intervention + follow up: 10 years
Method of assessment of guideline condition	Adequate method of assessment/diagnosis
Stratum	Fixed dose <400MBq
Subgroup analysis within study	Not applicable
Inclusion criteria	Treated with RAI for hyperthyroidism at Tampere hospital between 1965 and 2002
Exclusion criteria	Nil else stated
Recruitment/selection of patients	Nil else stated
Age, gender and ethnicity	Age - Median (range): 62 (50-75). Gender (M:F): 16:84. Ethnicity: Not stated
Further population details	1. Age: 2. Gender:
Indirectness of population	No indirectness
Interventions	(n=2793) Intervention 1: RAI - RAI alone. Mean dose 305MBq, 80.3% received a single dose. Duration 9.8 years follow-up. Concurrent medication/care: Nil else stated (n=2793) Intervention 2: General population. Age and sex matched control selected from Population register. Duration 9.8 years follow-up. Concurrent medication/care: Nil else stated
Funding	Academic or government funding

RESULTS (NUMBERS ANALYSED) AND RISK OF BIAS FOR COMPARISON: RAI ALONE versus GENERAL POPULATION

Protocol outcome 1: Total cancer diagnoses

- Actual outcome for Fixed dose <400MBq: Cancer, all diagnoses SIR at 10 years follow-up; RR; 1.25 (95%CI 1.08 to 1.46);

Risk of bias: All domain - High, Selection - High, Blinding - Low, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low;

Indirectness of outcome: No indirectness ; Group 1 Number missing: ; Group 2 Number missing:

Protocol outcomes not reported by the study | Quality of life ; Cancer diagnoses in iodine uptake glands ; Cancer diagnoses in non-iodine uptake glands ; Infertility

Study	Ryodi 2015 ¹¹⁵
Study type	Non randomised study
Number of studies (number of participants)	1 (n=6148)
Countries and setting	Conducted in Finland; Setting: Not stated
Line of therapy	1st line
Duration of study	Intervention + follow up: Median follow-up 10 years
Method of assessment of guideline condition	Adequate method of assessment/diagnosis
Stratum	Cause not specified
Subgroup analysis within study	Not applicable
Inclusion criteria	People treated with surgery for hyperthyroidism in Finland between 1986 and 2007, people treated with RAI for hyperthyroidism at Tampere University Hospital, reference population randomly chosen from national population register with 3 age and sex matched control subjects
Exclusion criteria	Not stated
Recruitment/selection of patients	Not stated
Age, gender and ethnicity	Age - Median (range): 46 for thyroidectomy, 59 for RAI. Gender (M:F): 16:84. Ethnicity:
Further population details	1. Age: 2. Gender:
Indirectness of population	No indirectness
Interventions	(n=1814) Intervention 1: RAI - RAI alone. No details provided. Duration Median follow-up 10 years. Concurrent medication/care: No details provided. Indirectness: No indirectness (n=4334) Intervention 2: ATD/SUR - SUR. No details provided. Duration Median follow-up 10 years. Concurrent medication/care: No details provided. Indirectness: No indirectness (n=18432) Intervention 3: General population. No details provided. Duration Median follow-up 10 years. Concurrent medication/care: No details provided. Indirectness: No indirectness
Funding	No funding
RESULTS (NUMBERS ANALYSED) AND RISK OF BIAS FOR COMPARISON: RAI ALONE versus SUR	

Protocol outcome 1: Total cancer diagnoses

- Actual outcome for Cause not specified: Total cancer diagnoses at Please enter a time period.; RR; 1.03 (95%CI 0.86 to 1.23);

Risk of bias: All domain - Very high, Selection - Very high, Blinding - Low, Incomplete outcome data - Low, Outcome reporting - Low, Measurement - Low, Crossover - Low; Indirectness of outcome: No indirectness, Comments: Obtained from Finnish cancer registry which captures 98% of cancer diagnoses, excluding benign, uncertain or borderline tumours; Key confounders: Adjusted for etiology, age, gender; Group 1 Number missing: ; Group 2 Number missing:

Protocol outcomes not reported by the study

Quality of life ; Cancer diagnoses in iodine uptake glands ; Cancer diagnoses in non-iodine uptake glands ; Infertility

1 Appendix E: Forest plots

2 E.1 Drugs vs Surgery vs Radioactive iodine

3 E.1.1 Radioactive iodine vs antithyroid drugs, adults with Graves' disease, first line 4 treatment

Figure 3: Mortality

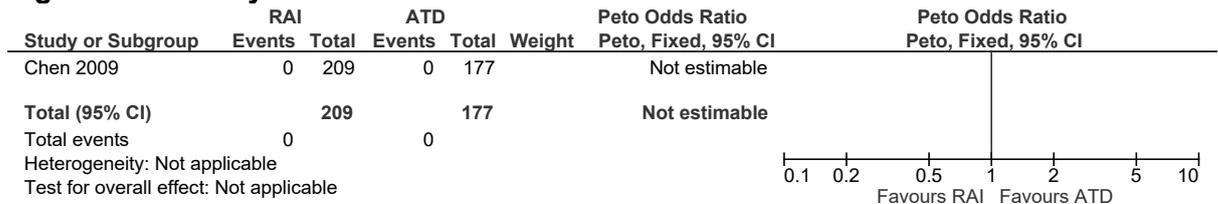
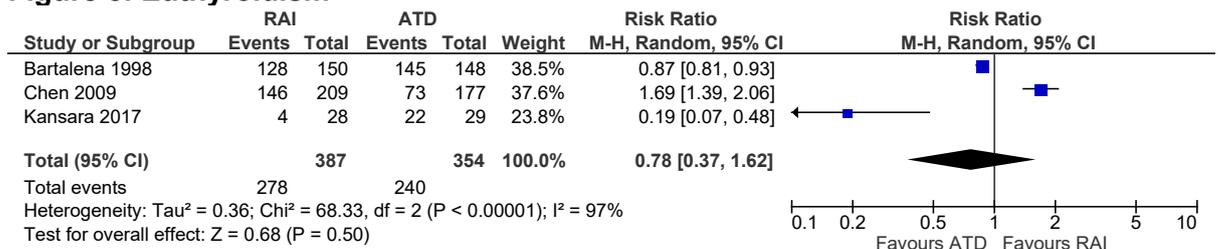


Figure 4: Ophthalmopathy (new or worsening cases)



5

Figure 5: Euthyroidism



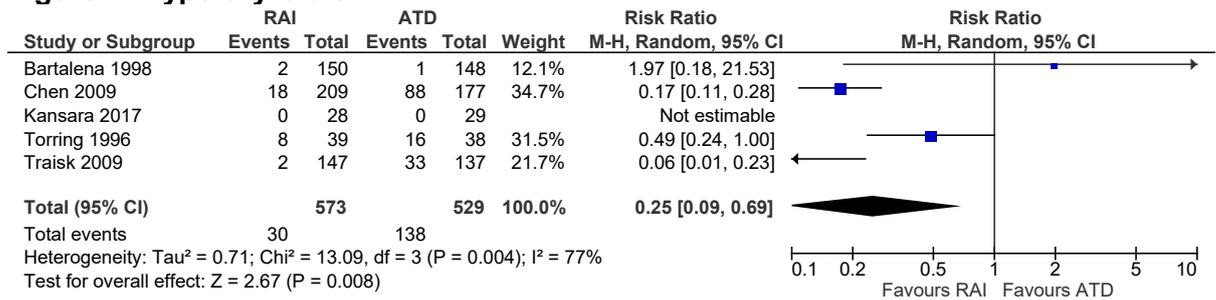
6

Figure 6: Hypothyroidism



7

Figure 7: Hyperthyroidism



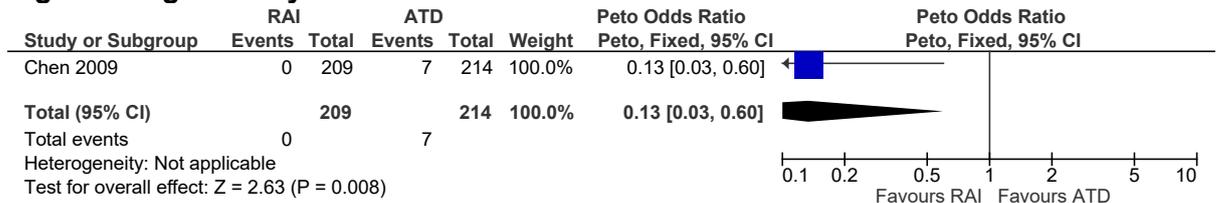
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Figure 8: Osteoporosis



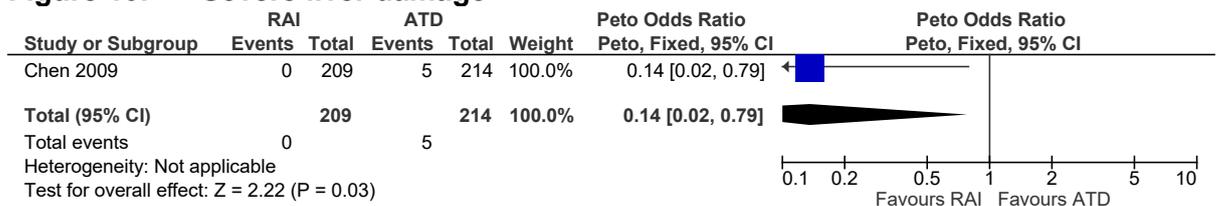
2

Figure 9: Agranulocytosis



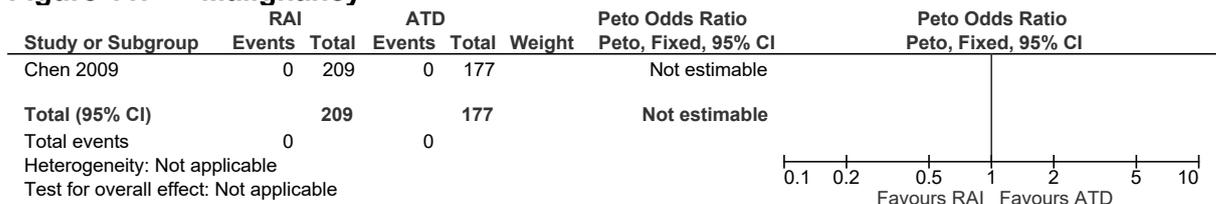
3

Figure 10: Severe liver damage



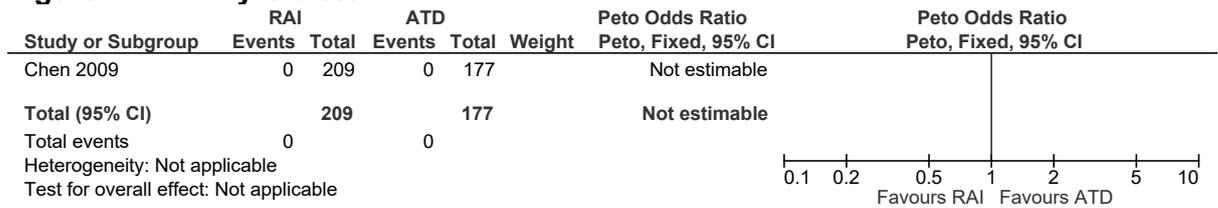
4

Figure 11: Malignancy



5

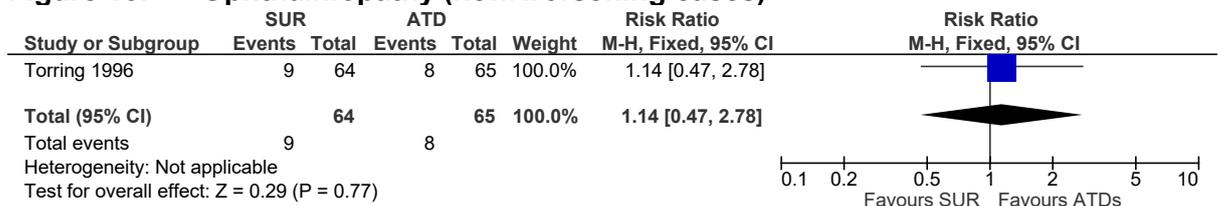
Figure 12: Thyroid storm



1

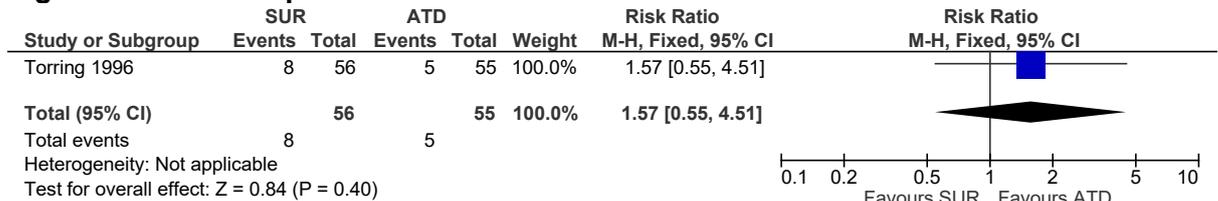
2 E.1.2 Surgery vs antithyroid drugs, adults with Graves' disease, first line treatment

Figure 13: Ophthalmopathy (new/worsening cases)



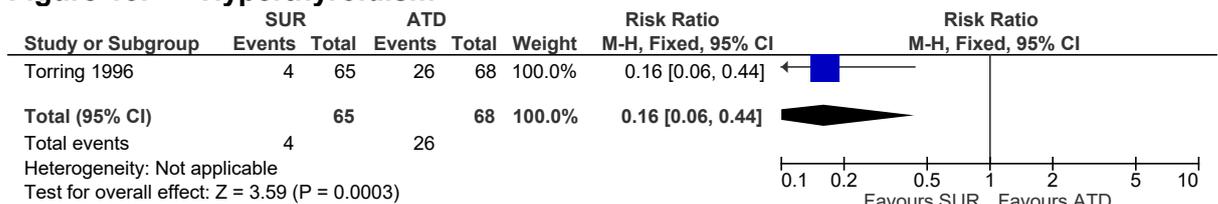
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Figure 14: Osteoporosis



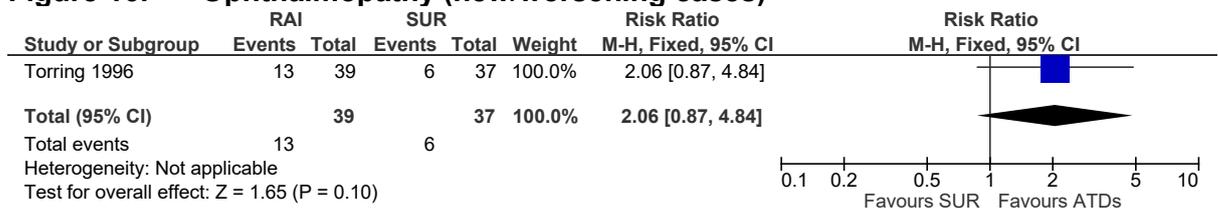
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Figure 15: Hyperthyroidism



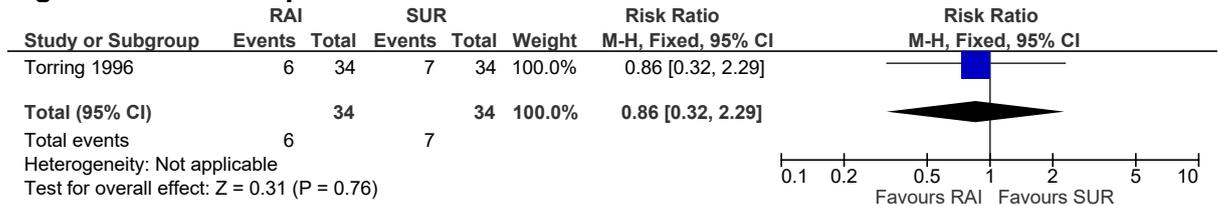
5 E.1.3 Radioactive iodine vs surgery, adults with Graves' disease, first line treatment

Figure 16: Ophthalmopathy (new/worsening cases)



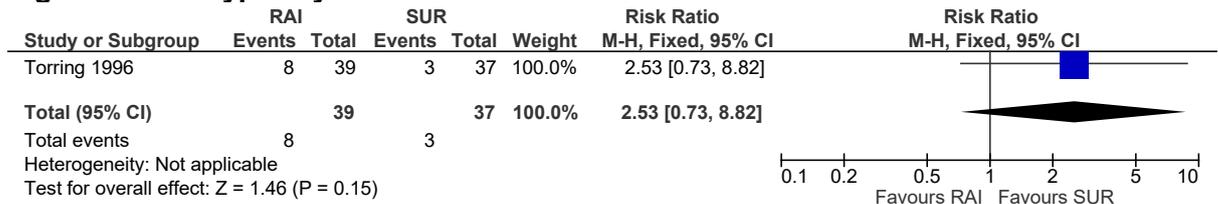
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Figure 17: Osteoporosis



2

Figure 18: Hyperthyroidism



3 E.1.4 Radioactive iodine vs antithyroid drugs, adults with Graves' disease, second line treatment

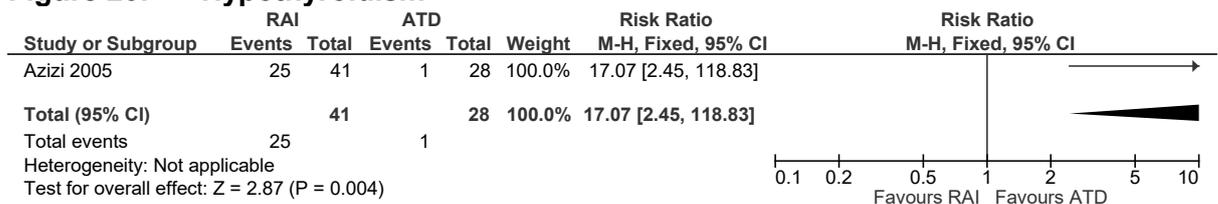
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Figure 19: Euthyroidism



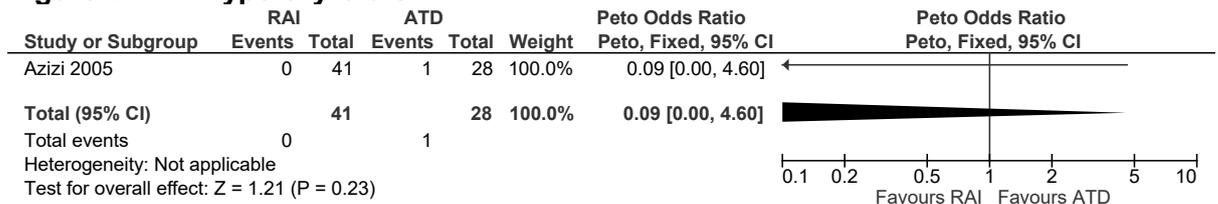
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Figure 20: Hypothyroidism



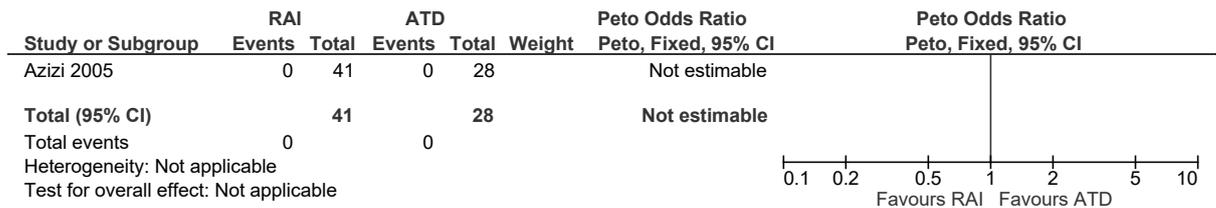
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Figure 21: Hyperthyroidism



7

Figure 22: Agranulocytosis



1 E.2 Forest plots: Radioactive Iodine safety

2 E.2.1 Radioactive iodine vs surgery

Figure 23: Total cancer diagnoses (RR)

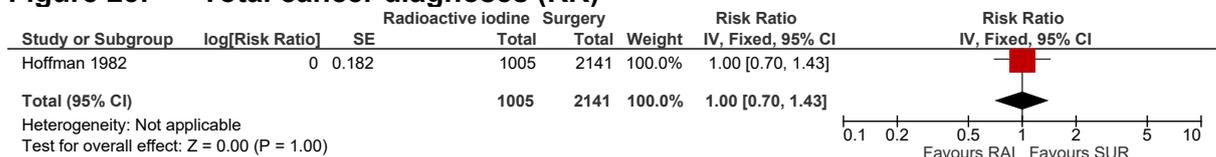
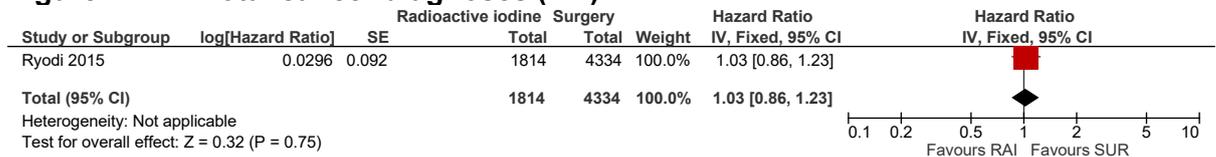
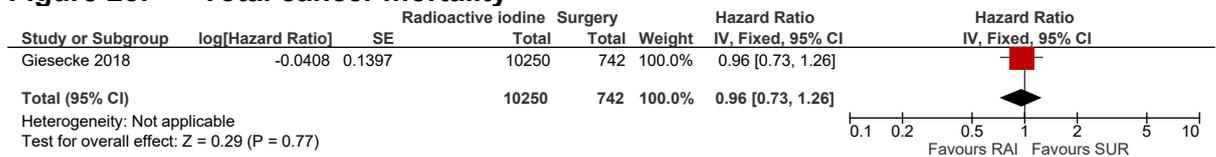


Figure 24: Total cancer diagnoses (HR)



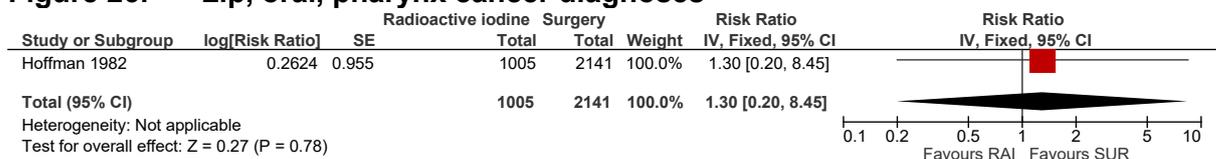
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Figure 25: Total cancer mortality



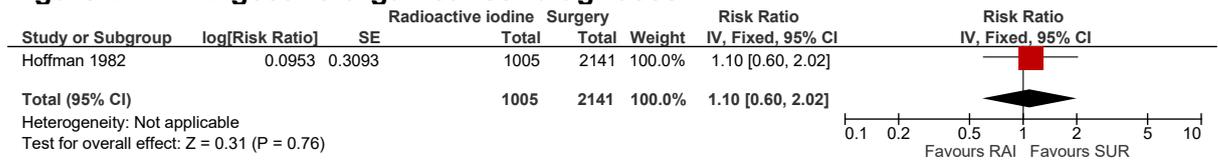
4

Figure 26: Lip, oral, pharynx cancer diagnoses



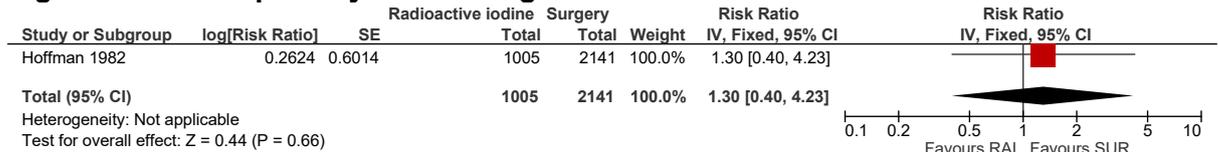
5

Figure 27: Digestive organ cancer diagnoses



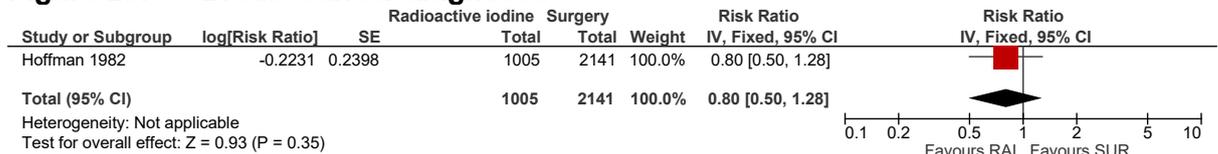
1

Figure 28: Respiratory cancer diagnoses



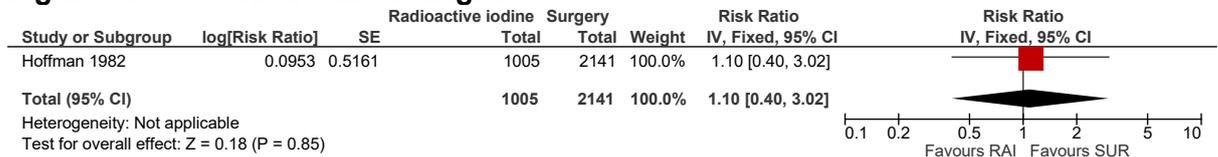
2

Figure 29: Breast cancer diagnoses



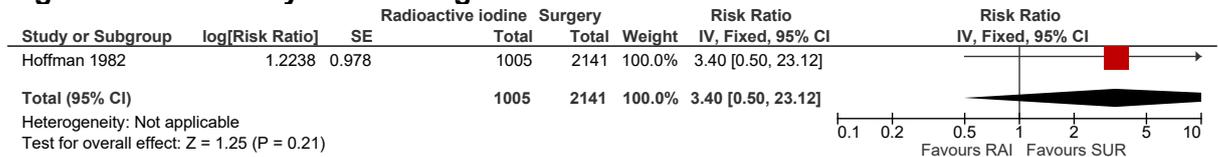
3

Figure 30: Genital cancer diagnoses



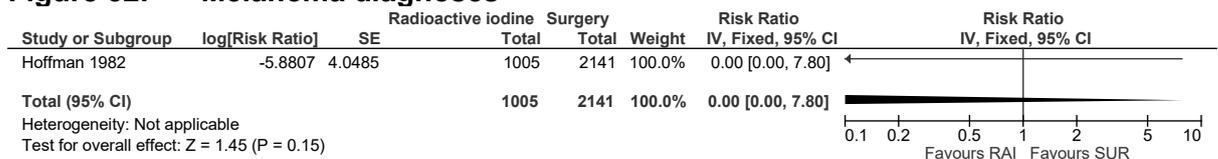
4

Figure 31: Kidney cancer diagnoses



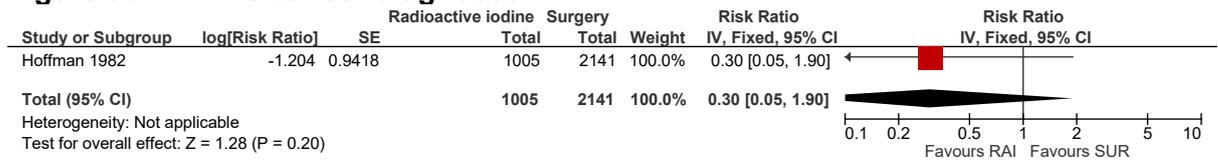
5

Figure 32: Melanoma diagnoses



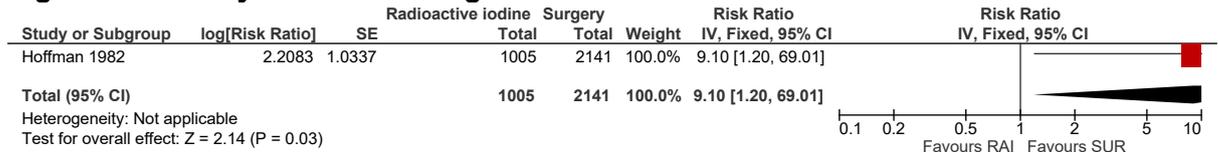
6

Figure 33: CNS cancer diagnoses



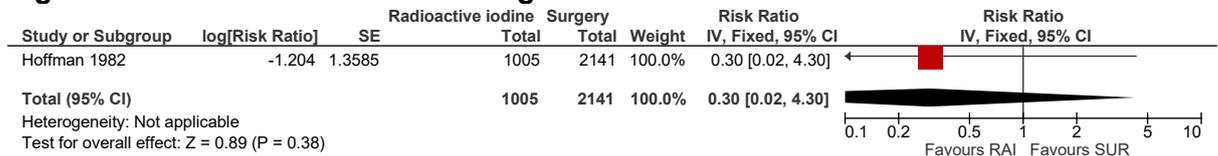
1

Figure 34: Thyroid cancer diagnoses



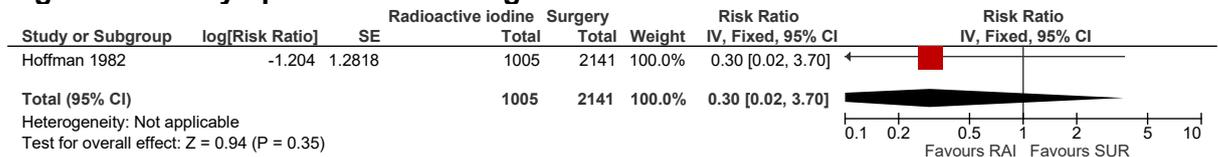
2

Figure 35: Other solid tumour diagnoses



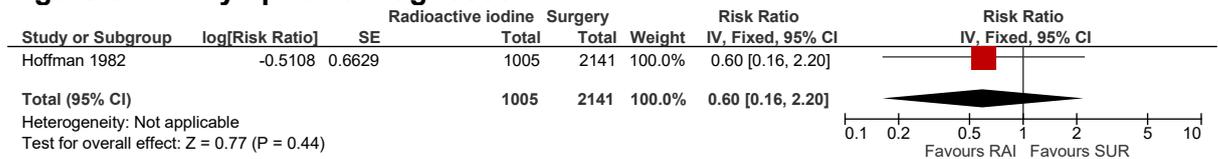
3

Figure 36: Lymphatic cancer diagnoses



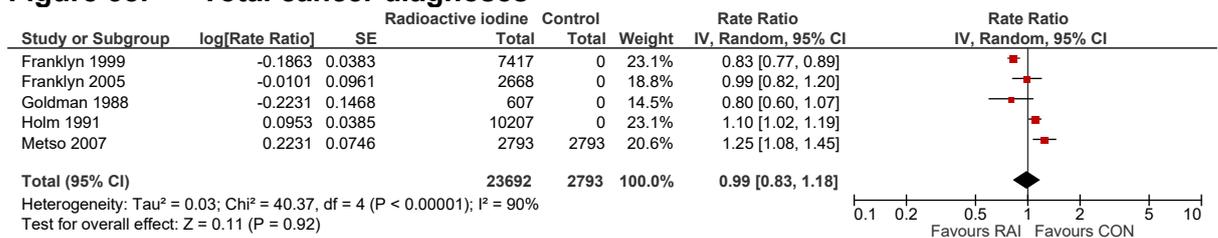
4

Figure 37: Lymphoma diagnoses



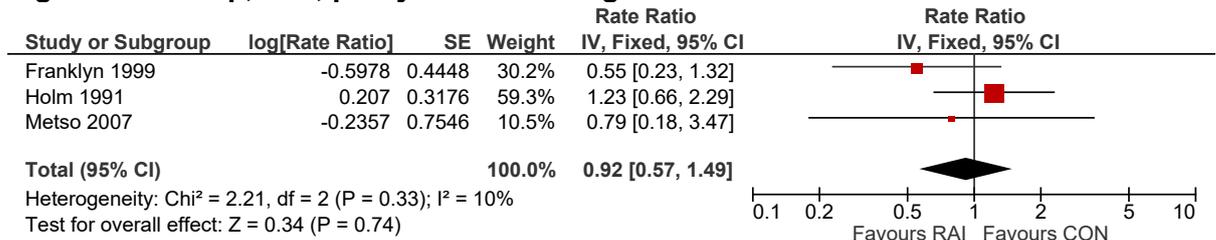
5 E.2.2 Radioactive iodine vs general population

Figure 38: Total cancer diagnoses



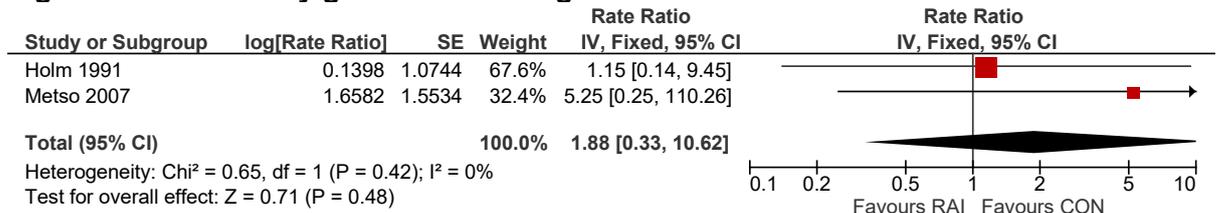
1

Figure 39: Lip, oral, pharynx cancer diagnoses



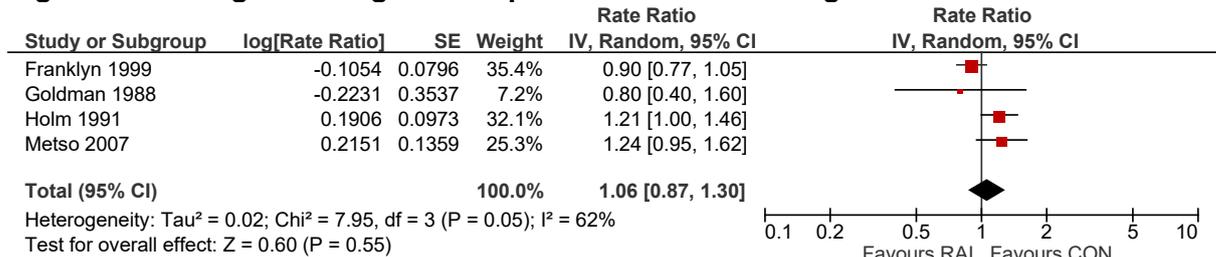
2

Figure 40: Salivary gland cancer diagnoses



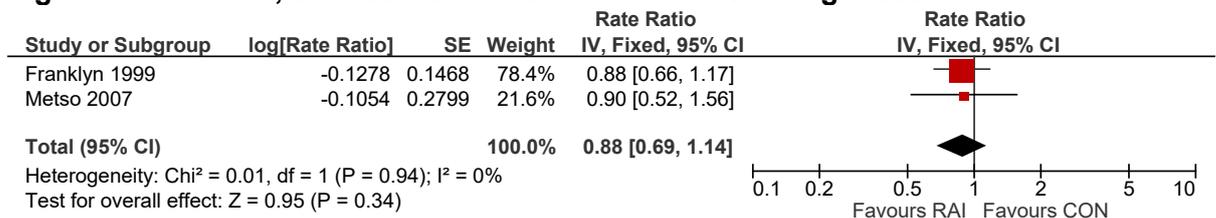
3

Figure 41: Digestive organs and peritoneum cancer diagnoses



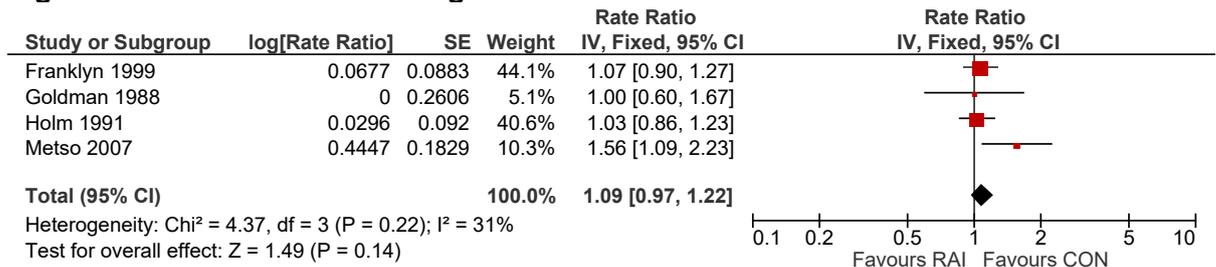
4

Figure 42: Bone, connective tissue and skin cancer diagnoses



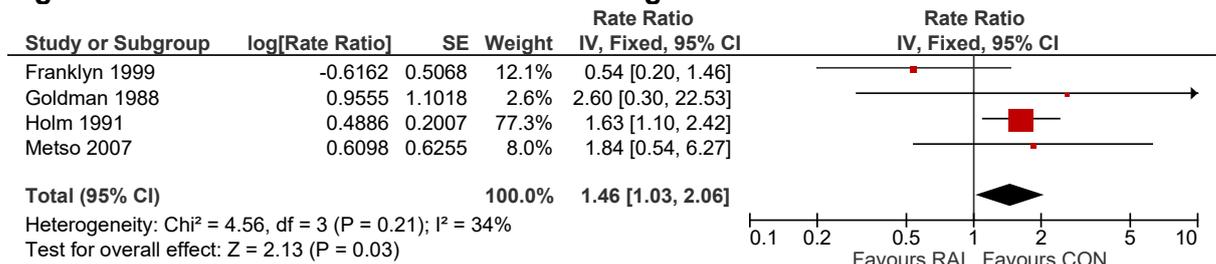
5

Figure 43: Breast cancer diagnoses



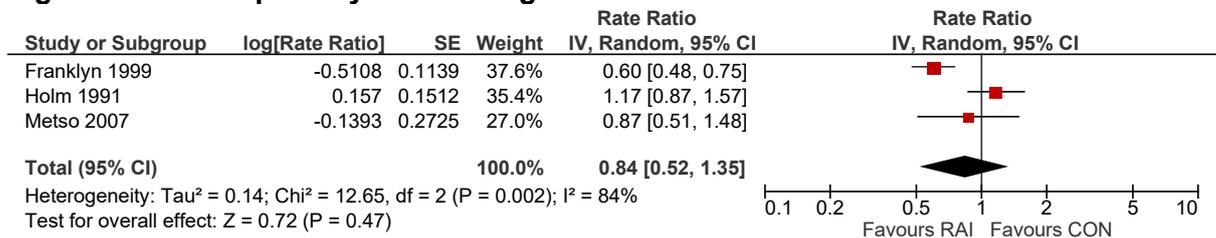
1

Figure 44: Brain and other CNS cancer diagnoses



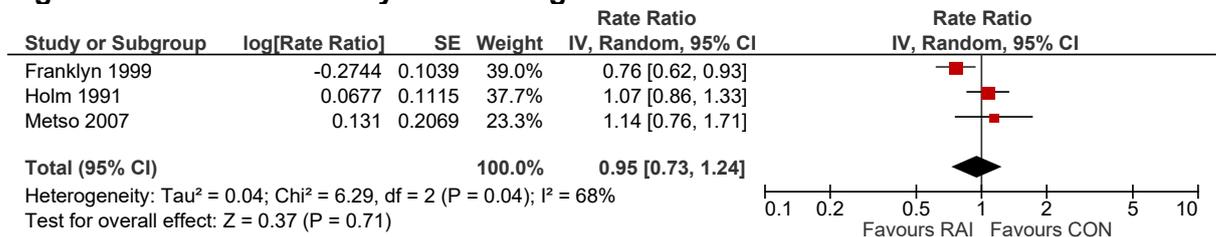
2

Figure 45: Respiratory cancer diagnoses



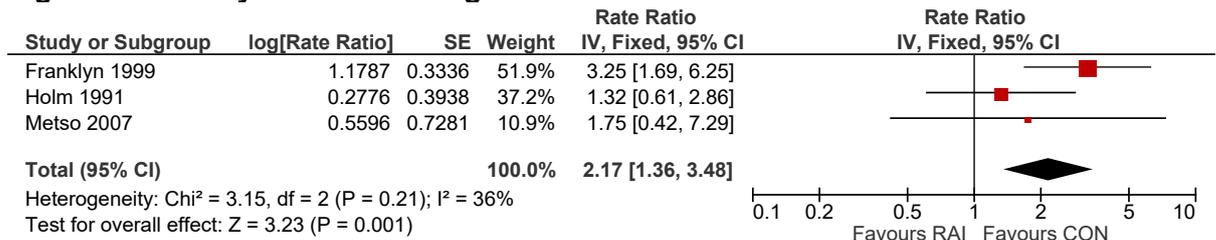
3

Figure 46: Genitourinary cancer diagnoses



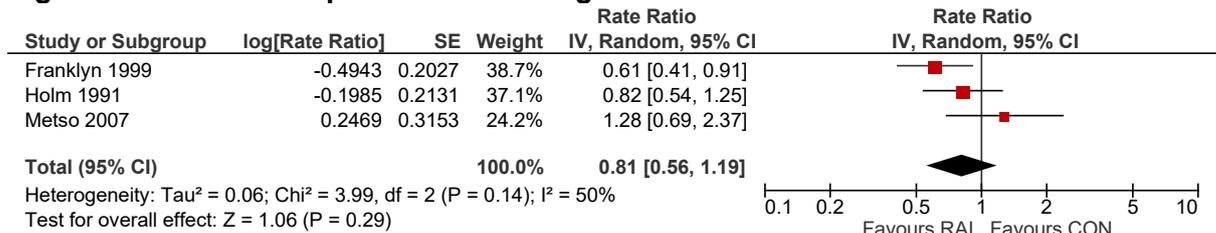
4

Figure 47: Thyroid cancer diagnoses



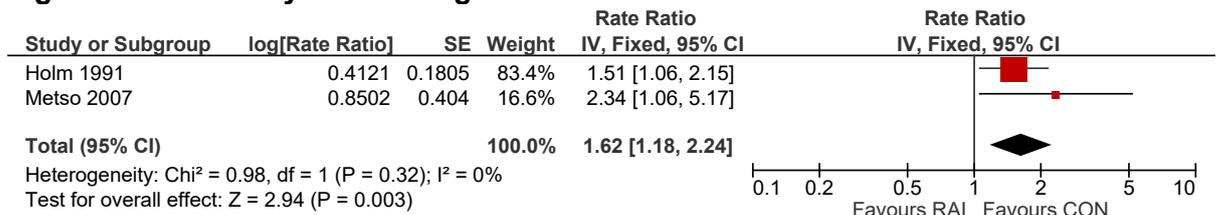
1

Figure 48: Haematopoietic cancer diagnoses



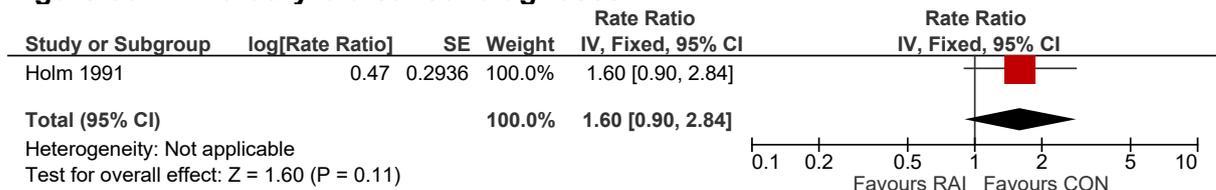
2

Figure 49: Kidney cancer diagnoses



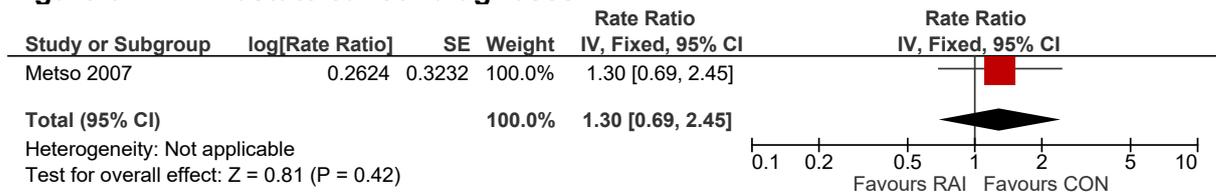
3

Figure 50: Parathyroid cancer diagnoses



4

Figure 51: Prostate cancer diagnoses



5

6

1

2

Appendix F: GRADE tables

F.1 Drugs vs Surgery vs Radioactive Iodine

Table 18: Clinical evidence profile: Radioactive iodine vs antithyroid drugs, Graves' disease, first line treatment

Quality assessment							No of patients		Effect		Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	RAI	ATD	Relative (95% CI)	Absolute		
Mortality (follow-up 9 years)												
1	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	0/209 (0%)	0%	-	not estimable ⁵	⊕⊕○○ LOW	CRITICAL
Ophthalmopathy (new/worsening cases) (follow-up 1-9 years)												
4	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	none	125/487 (25.7%)	10.3%	RR 2.17 (1.64 to 2.88)	121 more per 1000 (from 66 more to 194 more)	⊕⊕⊕○ MODERATE	IMPORTANT
Euthyroidism (follow-up 1-9 years; assessed with: (at end of follow-up))												
3	randomised trials	serious ¹	very serious ³	no serious indirectness	no serious imprecision	none	278/387 (71.8%)	75.9%	RR 0.78 (0.37 to 1.62)	167 fewer per 1000 (from 478 fewer to 471 more)	⊕○○○ VERY LOW	IMPORTANT
Hypothyroidism (follow-up 1-9 years; assessed with: (at end of follow-up))												
3	randomised	serious ¹	no serious	no serious	no serious	none	63/387	3.4%	RR 5.89 (3.12	166 more per 1000 (from	⊕⊕⊕○	IMPORTANT

	trials		inconsistency	indirectness	imprecision		(16.3%)		to 11.11)	72 more to 344 more)	MODERATE	
Hyperthyroidism (persistence/recurrence) (follow-up 1-9 years)												
5	randomised trials	serious ¹	serious ³	no serious indirectness	no serious imprecision	none	30/573 (5.2%)	24.1%	RR 0.25 (0.09 to 0.69)	181 fewer per 1000 (from 75 fewer to 219 fewer)	⊕⊕⊕⊕ LOW	IMPORTANT
Osteoporosis (follow-up 3 years)												
1	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	very serious ⁴	none	6/34 (17.6%)	13.9%	RR 1.27 (0.43 to 3.78)	38 more per 1000 (from 79 fewer to 386 more)	⊕⊕⊕⊕ VERY LOW	IMPORTANT
Agranulocytosis (follow-up 9 years)												
1	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	0/209 (0%)	3.3%	Peto OR 0.13 (0.03 to 0.6)	29 fewer per 1000 (from 13 fewer to 32 fewer)	⊕⊕⊕⊕ LOW	IMPORTANT
Severe liver damage (follow-up 9 years)												
1	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	serious ⁴	none	0/209 (0%)	2.3%	OR 0.14 (0.02 to 0.79)	20 fewer per 1000 (from 5 fewer to 23 fewer)	⊕⊕⊕⊕ LOW	IMPORTANT
Malignancy (follow-up 9 years)												
1	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	0/209 (0%)	0%	-	not estimable ⁵	⊕⊕⊕⊕ LOW	IMPORTANT
Thyroid storm (follow-up 9 years)												
1	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	0/209 (0%)	0%	-	not estimable ⁵	⊕⊕⊕⊕ LOW	IMPORTANT

- ¹ Downgraded by 1 increment if the majority of the evidence was at high risk of bias, and downgraded by 2 increments if the majority of the evidence was at very high risk of bias
- ² Downgraded by 1 increment as zero events in at least one arm
- ³ Downgraded by 1 or 2 increments because the point estimate and or the confidence intervals varied widely across studies, unexplained by subgroup analysis
- ⁴ Downgraded by 1 increment if the confidence interval crossed one MID or by 2 increments if the confidence interval crossed both MIDs
- ⁵ Zero events in both arms

Table 19: Clinical evidence profile: Surgery vs antithyroid drugs, Graves' disease, first line treatment

Quality assessment							No of patients		Effect		Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	SUR	ATD	Relative (95% CI)	Absolute		
Ophthalmopathy (follow-up 4 years; assessed with: (new/worsening cases))												
1	randomised trials	no serious risk of bias	no serious inconsistency	no serious indirectness	very serious ¹	none	9/64 (14.1%)	12.3%	RR 1.14 (0.47 to 2.78)	17 more per 1000 (from 65 fewer to 219 more)	⊕⊕⊕⊕ LOW	IMPORTANT
Osteoporosis (follow-up 14-21 years)												
1	randomised trials	serious ²	no serious inconsistency	no serious indirectness	very serious ¹	none	8/56 (14.3%)	9.1%	RR 1.57 (0.55 to 4.51)	52 more per 1000 (from 41 fewer to 319 more)	⊕⊕⊕⊕ VERY LOW	IMPORTANT
Hyperthyroidism (follow-up 4 years; assessed with: (persistence/recurrence))												
1	randomised trials	no serious risk of bias	no serious inconsistency	no serious indirectness	no serious imprecision	none	4/65 (6.2%)	38.2%	RR 0.16 (0.06 to 0.44)	321 fewer per 1000 (from 214 fewer to 359 fewer)	⊕⊕⊕⊕ HIGH	IMPORTANT

¹ Downgraded by 1 increment if the confidence interval crossed one MID or by 2 increments if the confidence interval crossed both MIDs ² Downgraded by 1 increment if the majority of the evidence was at high risk of bias, and downgraded by 2 increments if the majority of the evidence was at very high risk of bias

Table 20: Clinical evidence profile: Radioactive iodine vs surgery, Graves' disease, first line treatment

Quality assessment							No of patients		Effect		Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	RAI	Control	Relative (95% CI)	Absolute		
Ophthalmopathy (follow-up 4 years; assessed with: (new/worsening cases))												
1	randomised trials	no serious risk of bias	no serious inconsistency	no serious indirectness	serious ¹	none	13/39 (33.3%)	16.2%	RR 2.06 (0.87 to 4.84)	172 more per 1000 (from 21 fewer to 622 more)	⊕⊕⊕○ MODERATE	IMPORTANT
Osteoporosis (follow-up 14-21 years)												
1	randomised trials	serious ²	no serious inconsistency	no serious indirectness	very serious ¹	none	6/34 (17.6%)	20.6%	RR 0.86 (0.32 to 2.29)	29 fewer per 1000 (from 140 fewer to 266 more)	⊕○○○ VERY LOW	IMPORTANT
Hyperthyroidism (persistence/recurrence) (follow-up 4 years)												
1	randomised trials	no serious risk of bias	no serious inconsistency	no serious indirectness	very serious ¹	none	8/39 (20.5%)	8.1%	RR 2.53 (0.73 to 8.82)	124 more per 1000 (from 22 fewer to 633 more)	⊕⊕○○ LOW	IMPORTANT

¹ Downgraded by 1 increment if the confidence interval crossed one MID or by 2 increments if the confidence interval crossed both MIDs

² Downgraded by 1 increment if the majority of the evidence was at high risk of bias, and downgraded by 2 increments if the majority of the evidence was at very high risk of bias

Table 21: Clinical evidence profile: Radioactive iodine vs antithyroid drugs, Graves' disease, second line treatment

Quality assessment							No of patients		Effect		Quality	Importance
No of	Design	Risk of	Inconsistency	Indirectness	Imprecision	Other	RAI	ATD	Relative	Absolute		

studies		bias				considerations			(95% CI)			
Euthyroidism (follow-up 10 years; assessed with: (at end of follow-up))												
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	none	16/41 (39%)	92.9%	RR 0.42 (0.28 to 0.62)	539 fewer per 1000 (from 353 fewer to 669 fewer)	⊕⊕⊕⊕ LOW	IMPORTANT
Hypothyroidism (follow-up 10 years; assessed with: (at end of follow-up))												
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	none	25/41 (61%)	3.6%	RR 17.07 (2.45 to 118.83)	579 more per 1000 (from 52 more to 1000 more)	⊕⊕⊕⊕ LOW	IMPORTANT
Hyperthyroidism (follow-up 10 years; assessed with: (at end of follow-up))												
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	very serious ¹	none	0/41 (0%)	3.6%	OR 0.09 (0 to 4.6)	33 fewer per 1000 (from 36 fewer to 111 more)	⊕⊕⊕⊕ VERY LOW	IMPORTANT
Agranulocytosis (follow-up 10 years)												
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	0/41 (0%)	0%	-	not estimable ³	⊕⊕⊕⊕ VERY LOW	IMPORTANT

¹ Downgraded by 1 increment if the majority of the evidence was at high risk of bias, and downgraded by 2 increments if the majority of the evidence was at very high risk of bias

² Downgraded by 1 increment as at least one arm with zero events

³ zero events in both arms

F.2 Radioactive iodine safety

Table 22: Clinical evidence profile: radioactive iodine vs surgery

Quality assessment	No of patients	Effect	Quality	Importance
--------------------	----------------	--------	---------	------------

No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Radioactive iodine	Surgery	Relative (95% CI)	Absolute		
Total cancer diagnoses (RR) (follow-up median 15 years)												
1	observational studies	serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	1005	11.5%	RR 1.00 (0.7 to 1.43)	0 fewer per 1000 (from 35 fewer to 49 more)	⊕○○○ VERY LOW	CRITICAL
Total cancer diagnoses (HR) (follow-up median 10 years)												
1	observational studies	very serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	none	1814	0% ³	HR 1.02 (0.86 to 1.23)	Not estimable	⊕○○○ VERY LOW	CRITICAL
Total cancer mortality (follow-up median 16.3 years)												
1	observational studies	no serious risk of bias ¹	no serious inconsistency	no serious indirectness	very serious ²	none	10250	0% ³	HR 0.96 (0.73 to 1.26)	Not estimable	⊕○○○ VERY LOW	CRITICAL
Lip, oral, pharynx cancer diagnoses (follow-up median 15 years)												
1	observational studies	serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	1005	0.37%	RR 1.3 (0.2 to 8.45)	1 more per 1000 (from 3 fewer to 28 more)	⊕○○○ VERY LOW	CRITICAL
Digestive organ and peritoneum cancer diagnoses (follow-up median 15 years)												
1	observational studies	serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	1005	2.38%	RR 1.1 (0.6 to 2.02)	2 more per 1000 (from 10 fewer to 24 more)	⊕○○○ VERY LOW	CRITICAL
Respiratory cancer diagnoses (follow-up median 15 years)												
1	observational studies	serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	1005	0.7%	RR 1.3 (0.4 to 4.23)	2 more per 1000 (from 4 fewer to 23 more)	⊕○○○ VERY LOW	CRITICAL
Breast cancer diagnoses (follow-up median 15 years)												

1	observational studies	serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	1005	3.41%	RR 0.8 (0.5 to 1.28)	7 fewer per 1000 (from 17 fewer to 10 more)	⊕○○○ VERY LOW	CRITICAL
Genital cancer diagnoses (follow-up 15 years)												
1	observational studies	serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	1005	2.1%	RR 1.1 (0.4 to 3.02)	2 more per 1000 (from 13 fewer to 42 more)	⊕○○○ VERY LOW	CRITICAL
Kidney and bladder cancer diagnoses (follow-up median 15 years)												
1	observational studies	serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	1005	0.19%	RR 3.4 (0.5 to 23.12)	5 more per 1000 (from 1 fewer to 42 more)	⊕○○○ VERY LOW	CRITICAL
Melanoma cancer diagnoses (follow-up median 15 years)												
1	observational studies	serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	1005	0.05%	RR 0 (0 to 7.8)	1 fewer per 1000 (from 1 fewer to 3 more)	⊕○○○ VERY LOW	CRITICAL
CNS cancer diagnoses (follow-up median 15 years)												
1	observational studies	serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	1005	0.28%	RR 0.3 (0.05 to 1.9)	2 fewer per 1000 (from 3 fewer to 3 more)	⊕○○○ VERY LOW	CRITICAL
Thyroid cancer diagnoses (follow-up median 15 years)												
1	observational studies	serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	1005	0.05%	RR 9.1 (1.2 to 69.01)	4 more per 1000 (from 0 more to 34 more)	⊕○○○ VERY LOW	CRITICAL
Other solid tumour cancer diagnoses (follow-up median 15 years)												
1	observational studies	serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	1005	0.28%	RR 0.3 (0.02 to 4.3)	2 fewer per 1000 (from 3 fewer to 9 more)	⊕○○○ VERY LOW	CRITICAL
Lymphatic cancer diagnoses (follow-up median 15 years)												
1	observational	serious ¹	no serious	no serious	very serious ²	none	1005	0.33%	RR 0.3 (0.02	2 fewer per 1000	⊕○○○	CRITICAL

	studies		inconsistency	indirectness					to 3.7)	(from 3 fewer to 9 more)	VERY LOW	
Leukaemia diagnoses (follow-up median 15 years)												
1	observational studies	serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	1005	0.47%	RR 0.6 (0.16 to 2.2)	2 fewer per 1000 (from 4 fewer to 6 more)	⊕○○○ VERY LOW	CRITICAL

¹ Default starting quality of low overall due to selection bias in non-randomised studies. Downgraded further for risk of bias if the majority of evidence was at additional risk of bias, either once if high risk of bias or twice if very high risk of bias

² Downgraded by 1 increment if the confidence interval crossed one MID or by 2 increments if the confidence interval crossed both MIDs

³ No control group risk provided

Table 23: Clinical evidence profile: radioactive iodine vs general population

Quality assessment							No of patients		Effect		Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Radioactive iodine	General population	Relative (95% CI)	Absolute		
Total cancer diagnoses (follow-up 5-17 years)												
5	observational studies	serious ¹	serious ²	no serious indirectness	no serious imprecision	none	23692	7.4%	RR 0.99 (0.83 to 1.18)	1 fewer per 1000 (from 13 fewer to 13 more)	⊕○○○ VERY LOW	CRITICAL
Lip, oral, pharynx cancer diagnoses (follow-up 5-15 years)												
3	observational studies	serious ¹	no serious inconsistency	no serious indirectness	very serious ³	none	20417	0.1%	RR 0.92 (0.57 to 1.49)	0 fewer per 1000 (from 0 fewer to 0 more)	⊕○○○ VERY LOW	CRITICAL
Salivary gland cancer diagnoses (follow-up 10-15 years)												
2	observational studies	serious ¹	no serious inconsistency	no serious indirectness	very serious ³	none	13000	0.01%	RR 1.88 (0.33 to 10.62)	0 more per 1000 (from 0 fewer to 1 more)	⊕○○○ VERY LOW	CRITICAL

Digestive organs and peritoneum cancer diagnoses (follow-up 5-17 years)												
4	observational studies	serious ¹	serious ²	no serious indirectness	no serious imprecision	none	21024	2.7%	RR 1.06(0.87 to 1.30)	2 more per 1000 (from 4 fewer to 8 more)	⊕○○○ VERY LOW	CRITICAL
Bone, connective tissue and skin cancer diagnoses (follow-up 5-10 years)												
2	observational studies	serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	10210	1.3%	RR 0.88 (0.69 to 1.14)	2 fewer per 1000 (from 4 fewer to 2 more)	⊕○○○ VERY LOW	CRITICAL
Breast cancer diagnoses (follow-up 5-17 years)												
4	observational studies	serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	none	21024	1.7%	RR 1.09 (0.97 to 1.22)	2 more per 1000 (from 1 fewer to 4 more)	⊕○○○ VERY LOW	CRITICAL
Brain and other CNS cancer diagnoses (follow-up 5-17 years)												
4	observational studies	serious ¹	no serious inconsistency	no serious indirectness	serious ³	none	21024	0.3%	RR 1.46 (1.03 to 2.06)	1 more per 1000 (from 0 more to 3 more)	⊕○○○ VERY LOW	CRITICAL
Respiratory cancer diagnoses (follow-up 5-17 years)												
3	observational studies	serious ¹	serious ²	no serious indirectness	serious ³	none	20417	0.9%	RR 0.84 (0.52 to 1.35)	1 fewer per 1000 (from 4 fewer to 3 more)	⊕○○○ VERY LOW	CRITICAL
Genitourinary cancer diagnoses (follow-up 5-17 years)												
3	observational studies	serious ¹	serious ²	no serious indirectness	serious ³	none	20417	1.6%	RR 0.95 (0.73 to 1.24)	1 fewer per 1000 (from 4 fewer to 4 more)	⊕○○○ VERY LOW	CRITICAL
Thyroid cancer diagnoses (follow-up 5-17 years)												
3	observational studies	serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	none	20417	0.1%	RR 2.17 (1.36 to 3.48)	1 more per 1000 (from 0 more to 2 more)	⊕○○○ VERY LOW	CRITICAL
Haematopoietic cancer diagnoses (follow-up 5-17 years)												

3	observational studies	serious ¹	serious ²	no serious indirectness	serious ³	none	20417	0.5%	RR 0.81 (0.56 to 1.19)	1 fewer per 1000 (from 2 fewer to 0 more)	⊕000 VERY LOW	CRITICAL
Kidney cancer diagnoses (follow-up 10-15 years)												
2	observational studies	serious ¹	no serious inconsistency	no serious indirectness	serious ³	none	13000	0.4%	RR 1.62 (1.18 to 2.24)	2 more per 1000 (from 1 more to 5 more)	⊕000 VERY LOW	CRITICAL
Parathyroid cancer diagnoses (follow-up 15 years)												
1	observational studies	serious ¹	no serious inconsistency	no serious indirectness	serious ³	none	10207	0.22%	RR 1.6 (0.9 to 2.84)	1 more per 1000 (from 0 fewer to 4 more)	⊕000 VERY LOW	CRITICAL
Prostate cancer diagnoses (follow-up 10 years)												
1	observational studies	serious ¹	no serious inconsistency	no serious indirectness	very serious ³	none	2793	3.7%	RR 1.3 (0.69 to 2.45)	11 more per 1000 (from 11 fewer to 54 more)	⊕000 VERY LOW	CRITICAL

¹ Default starting quality of low overall due to selection bias in non-randomised studies. Downgraded further for risk of bias if the majority of evidence was at additional risk of bias, either once if high risk of bias or twice if very high risk of bias

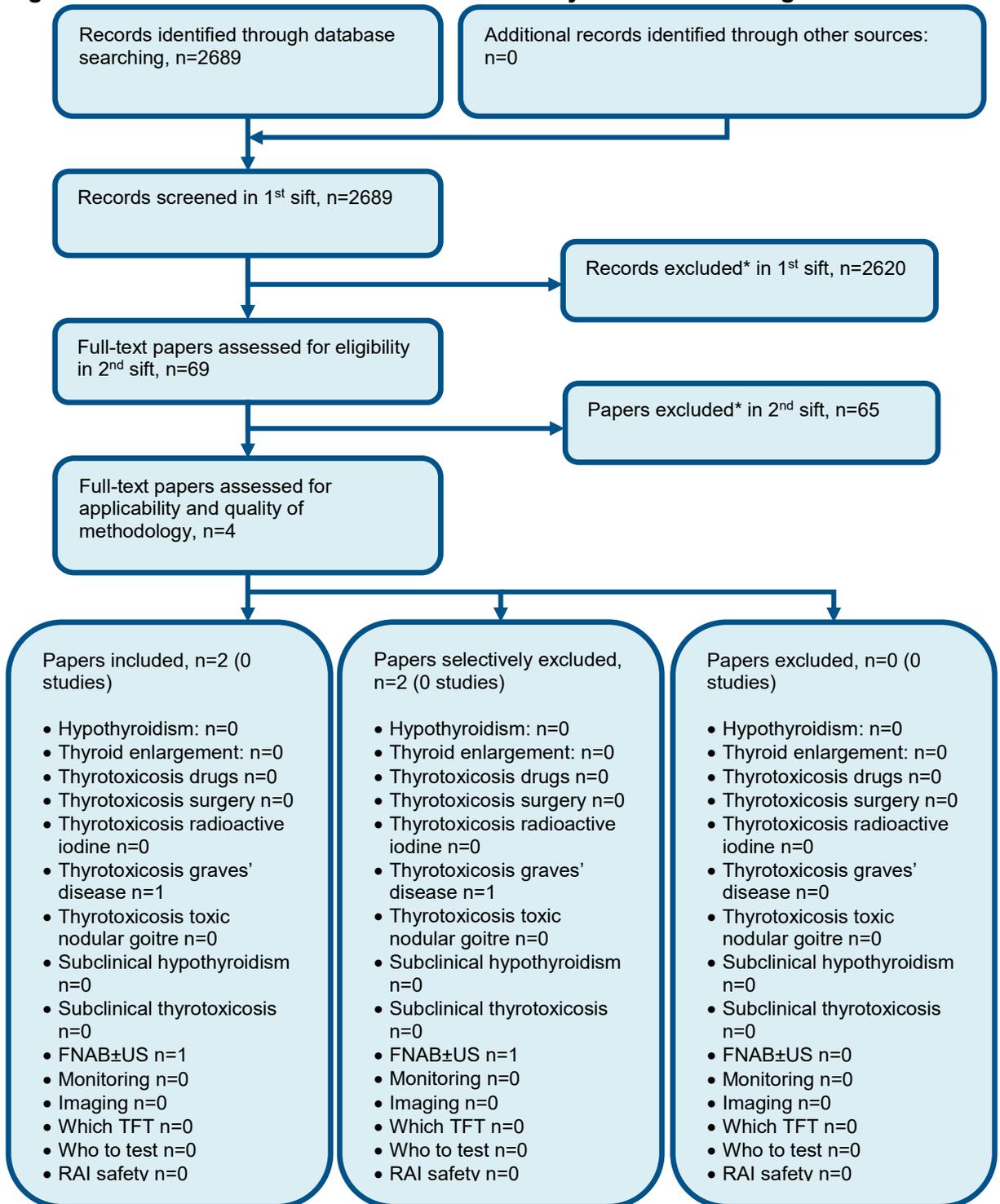
² Downgraded by 1 or 2 increments because the point estimate and or the confidence intervals varied widely across studies, unexplained by subgroup analysis

³ Downgraded by 1 increment if the confidence interval crossed one MID or by 2 increments if the confidence interval crossed both MIDs

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Appendix G: Health economic evidence selection

Figure 52: Flow chart of health economic study selection for the guideline



* Non-relevant population, intervention, comparison, design or setting; non-English language

3

Appendix H: Health economic evidence tables

H.1 Drugs vs Surgery vs Radioactive iodine

Study	Donovan et al, 2016 ³⁵			
Study details	Population & interventions	Costs	Health outcomes	Cost effectiveness
<p>Economic analysis: CUA (health outcome: QALYs)</p> <p>Study design: Deterministic decision analytic model</p> <p>Approach to analysis: Markov model, cyclical and tracks key clinical options and outcomes of persons with Graves' disease following each of the 3 interventions. 3-monthly cycles.</p> <p>Perspective: UK NHS</p> <p>Time Horizon: lifetime</p> <p>Treatment effect duration: NR</p> <p>Discounting: Costs: 3.5% ; Outcomes: 3.5%</p>	<p>Population: People diagnosed with Graves' disease.</p> <p>Cohort settings: Start age: 40 years old women</p> <p>Intervention 1: Radioactive iodine (RAI)</p> <p>Intervention 2: Antithyroid drug (ATD) (carbimazole 5mg).</p> <p>Intervention 3: Total thyroidectomy (TT).</p>	<p>Total costs (mean per patient): Intervention 1: £5,425 Intervention 2: £16,866 Intervention 3: £7,115</p> <p>Incremental (2-1):£11,441 (95% CI: NR; p=NR) Incremental (3-1): £1,690 (95% CI: NR; p=NR) Incremental (3-2): saves £9,751 (95% CI: NR; p=NR)</p> <p>Currency & cost year: 2015 UK pounds</p> <p>Cost components incorporated:</p> <ul style="list-style-type: none"> • Long-term costs of medications • medical practitioner visits • pathology tests 	<p>QALYs (mean per patient): Intervention 1: 34.73 Intervention 2: 35.17 Intervention 3: 33.93</p> <p>Incremental (2-1): 0.44 (95% CI: NR; p=NR) Incremental (3-1): -0.8 (95% CI: NR; p=NR) Incremental (3-2): -1.24 (95% CI: NR; p=NR)</p>	<p>Full incremental analysis: RAI dominated TT.</p> <p>At cost effectiveness threshold of £20,000 per QALY gained, RAI is cost-effective compared to ATD, while at a cost effectiveness threshold of £30,000 per QALY-gained; ATD is the cost-effective alternative, (ICER £26,279 per QALY-gained) compared to RAI.</p> <p>Analysis of uncertainty: One-way sensitivity analyses, where the value of a single parameter is changed across range of values with ICER values calculated. Transition probabilities ranges were based on 95% CI from published literature. Cost were varied from 50% to 150% depending on base case values. Results from these sensitivity analyses showed that ATD was a cost-effective alternative to RAI in most sensitivity analyses (calculated ICER remained below the £30,000 threshold). RAI was dominant over TT in all sensitivity</p>

		associated with treatments and their complications.		analyses of all parameters assessed.
Data sources				
<p>Health outcomes: Effectiveness data for the three interventions were based on a literature review that identified rates of efficacy, relapse, complications and HRQoL values associated with each treatment option. Some assumptions were made e.g. the failure rate with ATD and the incidence of hypothyroid post third dose of RAI. Quality-of-life weights: Effectiveness was evaluated by using the HRQoL estimates (health utilities) from published data, Euro-Qol – 5 dimensions or SF-36 values mapped to EQ-5D. Some of the values were also based on expert opinion using Delphi methodology. Cost sources: Unit costs based on 2015 UK sources (BNF, National Tariff). Where unit costs were not available, estimates were obtained from published literature or currency conversion.</p>				
Comments				
<p>Source of funding: An NHMRC early career fellowship (APPP1092153) support. Limitations: The estimates of relative treatment effects are not based on met-analysis of all the available evidence. Some costs have been based on the national tariff Payment System and maybe overestimated. The model has not been run probabilistically, to adequately assess parameter uncertainty.</p>				
<p>Overall applicability:^(c) Directly applicable Overall quality:^(d) Minor limitations</p>				

Abbreviations: CI: 95% confidence interval; CUA: cost–utility analysis; EQ-5D: Euro-qol 5 dimensions (scale: 0.0 [death] to 1.0 [full health], negative values mean worse than death); ICER: incremental cost-effectiveness ratio; NR: not reported; QALYs: quality-adjusted life years

(a) Directly applicable / Partially applicable / Not applicable

(b) Minor limitations / Potentially serious limitations / Very serious limitations

H.2 Radioactive iodine safety

None

1 **Appendix I: Health economic analysis**

2 **I.1 Drugs vs Surgery vs Radioactive Iodine**

3 None

4 **I.2 Radioactive iodine safety**

5 None

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Appendix J: Excluded studies

J.1 Excluded clinical studies

J.1.1 Drugs vs Surgery vs Radioactive Iodine

Table 24: Studies excluded from the clinical review

Study	Exclusion reason
Abraham 2010 ⁴	Systematic review is not relevant to review question or unclear PICO
Abraham-nordling 2007 ²	No usable outcomes
Allannic 1990 ⁵	Incorrect interventions
Andrade 1999 ⁶	Less than minimum duration
Andrade 2001 ⁷	Incorrect interventions
Andrade 2004 ⁸	Incorrect interventions
Azizi 2012 ¹²	Wrong study design
Azizi 2018 ¹⁰	NRS where RCTs are available
Barczynski 2012 ¹³	Incorrect interventions
Barczynski 2010 ¹⁴	Abstract only
Barczynski 2018 ¹⁵	Incorrect interventions
Benker 1995 ¹⁸	Incorrect interventions
Benker 1998 ¹⁷	Incorrect interventions
Bonnema 2003 ¹⁹	Incorrect interventions
Bonnema 2004 ²⁰	Incorrect interventions
Bonnema 2011 ²¹	Inappropriate comparison
Braga 2002 ²²	Less than minimum duration
Burch 2001 ²³	No usable outcomes
Buscemi 2007 ²⁴	Not guideline condition
Canto 2016 ²⁵	Incorrect interventions
Chen 2011 ²⁹	Inappropriate comparison
Chen 2014 ³⁰	No additional outcomes to those reported elsewhere
Chi 2005 ³¹	Inappropriate comparison
Connell 1987 ³²	No usable outcomes
De Luca 2018 ³³	SR, checked for references
Edmonds 1994 ³⁶	Incorrect interventions
Esfahani 2005 ³⁷	Inappropriate comparison
García-mayor 1992 ⁴¹	Incorrect interventions
Glinoeer 2001 ⁴³	Incorrect interventions
Goni iriarte 1995 ⁴⁵	Not in English
Grebe 1998 ⁴⁶	Incorrect interventions
Hamide 2014 ⁵²	NRS where RCTs are available
Hashizume 1991 ⁵³	NRS without adequate adjustment
He 2004 ⁵⁴	Incorrect interventions
Hoermann 2002 ⁵⁶	Incorrect interventions
Homsanit 2001 ⁶³	Incorrect interventions
Howarth 2001 ⁶⁴	Incorrect interventions

Study	Exclusion reason
Jaiswal 2014 ⁶⁵	Incorrect interventions
Järhult 2005 ⁶⁶	Incorrect interventions
Jorde 1995 ⁶⁷	Incorrect interventions
Kallner 1996 ⁶⁹	Incorrect interventions
Kung 1995 ⁷¹	Incorrect interventions
Leclere 1994 ⁷²	Not in English
Leslie 2003 ⁷³	Incorrect interventions
Leung 2017 ⁷⁴	SR, checked for references
Li 2016 ⁷⁵	SR, checked for references
Liu 2015 ⁷⁷	Incorrect interventions
Liu 2017 ⁷⁶	Incorrect interventions
Ljunggren 1998 ⁷⁸	No usable outcomes
Lucas 1997 ⁷⁹	Incorrect interventions
Ma 2008 ⁸⁰	SR, checked for references
Ma 2016 ⁸¹	SR checked for references
Marcocci 1989 ⁸²	Incorrect interventions
Mashio 1997 ⁸³	Inappropriate comparison
Mastorakos 2003 ⁸⁴	Incorrect interventions
Maugendre 1999 ⁸⁵	Incorrect interventions
Mciver 1996 ⁸⁶	Incorrect interventions
Menconi 2007 ⁸⁸	No usable outcomes
Miranda-padua 2014 ⁹⁴	Incorrect interventions
Müller 2001 ⁹⁵	Inappropriate comparison
Nakamura 2007 ⁹⁶	Incorrect interventions
Nedrebo 2002 ⁹⁸	Incorrect interventions
Noh 2015 ⁹⁹	Incorrect interventions
Orsini 2012 ¹⁰⁰	Inappropriate comparison
Peixoto 2006 ¹⁰²	Incorrect interventions
Peters 1995 ¹⁰³	Incorrect interventions
Peters 1996 ¹⁰⁴	No usable outcomes
Peters 1997 ¹⁰⁵	Incorrect interventions
Pfeilschifter 1997 ¹⁰⁶	Inappropriate comparison
Pirnat 2011 ¹⁰⁷	Incorrect interventions
Pusuwan 2011 ¹⁰⁸	Inappropriate comparison
Raber 2000 ¹⁰⁹	Incorrect interventions
Reinwein 1993 ¹¹⁰	Inappropriate comparison
Rittmaster 1998 ¹¹¹	Incorrect interventions
Rokni 2014 ¹¹²	SR checked for references
Romaldini 1983 ¹¹³	Incorrect interventions
Santos 2004 ¹¹⁶	NRS without adequate adjustment
Santos 2012 ¹¹⁷	Inappropriate comparison
Sapienza 2015 ¹¹⁸	Inappropriate comparison
Schneider 2005 ¹¹⁹	Inappropriate comparison
Singhal 2014 ¹²²	Withdrawn Cochrane review
Taieb 2016 ¹²³	Incorrect interventions

Study	Exclusion reason
Thientunyakit 2010 ¹²⁵	Inappropriate comparison
Tian 2001 ¹²⁶	Not in English
Unalp 2009 ¹²⁹	No usable outcomes
Walter 2006 ¹³¹	NRS without adequate adjustment
Wang 2016 ¹³²	SR, checked for references
Weetman 1994 ¹³³	Incorrect interventions
Witte 2000 ¹³⁴	Incorrect interventions
Yousefi 2011 ¹³⁵	Not in English
Yuan 2017 ¹³⁶	SR, checked for references

1 J.1.2 Radioactive Iodine safety

2 **Table 25: Studies excluded from the clinical review**

Study	Exclusion reason
Angusti 2000 ⁹	No usable outcomes
Cevallos 1974 ²⁶	Not minimum sample size
Chao 2009 ²⁷	SR, references checked
Franklyn 1998 ³⁹	No usable outcomes
Hall 1992 ⁴⁷	No usable outcomes
Hall 1992 ⁴⁸	Majority of radioactive iodine exposure not therapeutic
Hall 1993 ⁵¹	No usable outcomes
Hall 1995 ⁴⁹	Outcomes reported elsewhere and included
Hall 1997 ⁵⁰	Non-systematic review
Hieu 2012 ⁵⁵	SR, references checked
Hoffman 1982 ⁵⁸	No usable outcomes
Hoffman 1983 ⁵⁷	Outcomes reported elsewhere and included
Holm 1980 ⁶¹	Outcomes reported elsewhere and included
Holm 2006 ⁶⁰	Non-systematic review
Journy 2017 ⁶⁸	Inappropriate population
Mctiernan 1984 ⁸⁷	Inappropriate study design
Metso 2004 ⁹³	No usable outcomes
Metso 2007 ⁸⁹	Erratum, not relevant
Metso 2007 ⁹¹	Erratum, not relevant
Metso 2007 ⁹²	No usable outcomes
Ron 1998 ¹¹⁴	No usable outcomes
Singer 2001 ¹²⁰	Commentary only
Singer 2001 ¹²¹	No usable outcomes
Verburg 2011 ¹³⁰	SR, references checked

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4 J.2 Excluded health economic studies

5 J.2.1 Drugs vs Surgery vs Radioactive Iodine

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Study	Exclusion reason
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Study	Exclusion reason
Patel ¹⁰¹	Not applicable, resource use and cost data from 2002

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2 **J.2.2 Radioactive Iodine safety**

3 None

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Appendix K: Research recommendations

K.1 Research question: Are there subgroups of people with Graves' thyrotoxicosis who respond particularly well to antithyroid drugs?

Why this is important:

Antithyroid drugs (ATDs) are commonly used for treatment of Graves' disease. With a 12-18 months course of ATDs, there is about 50% chance for peoples with Graves' disease achieving a long-term remission. Previous studies have suggested patients with certain clinical characteristics are more likely to relapse following ATD. These characteristics, variably suggested by different studies, include male sex, young age, cigarette smoking, presence of large goitre, high levels of thyroid hormones at the time of diagnosis and high titres of TSH-receptor antibodies. However, most of these studies are small and retrospective in design, and these findings need confirmation by large prospective multi-centre studies. If the findings are confirmed, it will allow clinicians to stratify patients with Graves' hyperthyroidism who are unlikely to remain in remission following a course of ATD and offer early definitive treatments such as radioactive iodine or thyroidectomy.

Within the present guideline, the committee agreed that radioactive iodine should constitute the first line treatment option for adults with thyrotoxicosis/hyperthyroidism/Graves' disease according to both clinical and cost-effectiveness, but that for people in whom ATDs are particularly likely to achieve remission the, need for definitive treatment might be less. These were hypothesised by the committee to be people with milder, predominantly T3 thyrotoxicosis. However, no evidence was currently identified about any specific group of people who are likely to respond particularly well to ATDs. Further research is required to allow us to identify those people and allow clinicians to stratify patients with Graves' disease who are likely to remain in remission following the course of ATDs and avoid offering them a definitive treatment such as radioactive iodine or thyroidectomy.

Criteria for selecting high-priority research recommendations:

PICO question	<p>Population: People with Graves' disease who are being treated with an antithyroid drug (ATD)</p> <p>Indicator: Absence of goitre, absence of thyroid eye disease, low titres of TSH receptor antibodies, low tires of free thyroid hormone levels at diagnosis, non-smoking mild thyrotoxicosis/ Graves' disease, T3 thyrotoxicosis...</p> <p>Comparator: Presence of goitre, presence of thyroid eye disease, high titres of TSH receptor antibodies, high tires of free thyroid hormone levels at diagnosis, smoking, non-mild Grave's disease...</p> <p>Outcome(s): hyperthyroidism relapse rate</p>
Importance to patients or the population	<p>This research will help to ascertain if simple clinical characteristics are useful in predicting the achievement of remission following a course of ATD. This will enable clinicians to stratify people with Graves' disease who are likely to achieve long-term remission after a course of ATD and those who are not and 'provide particular</p>

	groups of people with the treatment they are most likely to benefit from' OR 'avoid definitive treatment for people who do not need it'.
Relevance to NICE guidance	This research will allow future guidelines to clearly recommend which people with Graves' disease should be offered ATDs as first line treatment instead of definitive treatment with radioactive iodine or thyroidectomy.
Relevance to the NHS	This research will provide clear evidence of the potential subgroup(s) of people with Graves' disease that could effectively be treated with an ATD. This will allow the identification of people who are likely to achieve long-term remission with a course of ATD, and avoid offering them definitive treatment in early course of the disease.
National priorities	Hyperthyroidism comes under the long-term condition directorate in the UK.
Current evidence base	Several retrospective single site studies have suggested various clinical characteristics, such as the presence of large goitre, high titres of free thyroid hormones at presentation, high titres of TSH receptor antibodies and smoking status are associated with the risk of relapse following a course of ATDs in patients with Graves' hyperthyroidism. However, no evidence about groups of people likely to respond particularly well to ATDs has been identified.
Equality	This recommendation is unlikely to impact on equality issues.
Study design	A multi-centre prospective observational study.
Feasibility	As Graves' disease is common, and ATDs are widely used in the UK for the treatment of Graves' disease, a multi-centre prospective observational study is feasible. A key challenge will be differences in clinical practice, in terms of regimes and duration of ATD, between different centres in the UK.
Other comments	
Importance	Medium: the research is relevant to the recommendations in the guideline, but the research recommendations are not key to future updates.

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3 **K2. Research question: What is the long-term clinical and cost effectiveness,**
4 **including safety, of radioactive iodine for thyrotoxicosis?**

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6 **Why this is important:**

7 Radioactive iodine (I-131 NaI) is used to treat benign thyroid disease in approximately
8 10,000 patients in the UK each year by delivering absorbed doses (radiation) preferentially to
9 the thyroid. This treatment is used globally with and is considered clinically effective. Despite
10 the large number of patients treated with radioactive iodine over the past 50 years, there are
11 still questions concerning the medium and longer-term effects and in particular the potential
12 impact of exposure to low doses of radiation. A registry would enable the long-term effects of

1 radiation to be recorded and in time would provide definitive answers regarding the
2 association of radiation and its specific dose with medium and long term risks. This would
3 inform treatment protocols and would potentially provide reassurance to patients.

4 **Criteria for selecting high-priority research recommendations:**

PICO question	Population: Patients receiving radio-iodine for benign thyroid disease Intervention(s): Radio-iodine (RAI) therapy Comparison: The population not receiving RAI therapy Outcome(s): Neoplasia, fertility, quality of life, morbidity, death
Importance to patients or the population	The registry will be used to develop and refine our understanding of the risks and benefits associated with RAI therapy. This would help patients to make informed choices and place the risks / benefits of RAI into context.
Relevance to NICE guidance	Registry development would help to establish the role of RAI in the management of benign thyroid disease.
Relevance to the NHS	RAI therapy is relatively cost-effective when compared to interventions like surgery. There may be associated benefits from a financial and resource perspective.
National priorities	The NHS Five Year Forward View (2014) aims to address variations in treatment and outcomes. A register of treatments and outcomes would enable this information to be collected.
Current evidence base	RAI safety review did not support an association of RAI with increased risk of malignancy, however results have been largely based on older studies using lower RAI doses than those currently used in the UK. The general public and health professionals are not clear about the risks and benefits of RAI therapy. A registry documenting outcomes following RAI treatment in the medium and long term according to current practice will provide greater clarity in this area.
Equality	A registry might be of particular benefit in the context of young people receiving RAI who statistically will have more life-years ahead of them and an increased theoretical risk of health issues such as neoplasia as a result.
Study design	A central registry of all patients receiving RAI would be established. The key national bodies including those in the field of medical physics would agree to submit data on a regular basis. This data could be linked at national level to cancer registries / cause of death and patients will be asked to consent to being contacted about studies in areas such as QOL at a later stage. In the absence of consent, anonymised data will still be linked to long term morbidity / mortality data.
Feasibility	Collecting this data is a long-term project but is relatively inexpensive. A key issue would be to ensure high ascertainment.
Other comments	Radioactive iodine treatment for benign thyroid disease is performed widely. A registry could therefore have international impact.
Importance	<ul style="list-style-type: none"> The research is important to quantify the risks associated with RAI therapy in greater detail.

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