# National Institute for Health and Care Excellence

Draft for consultation

# Joint replacement (primary): hip, knee and shoulder

[J] Evidence review for wrong implant selection

NICE guideline

Intervention evidence review

October 2019

**Draft for Consultation** 

This evidence review was developed by the National Guideline Centre, hosted by the Royal College of Physicians



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#### **ISBN**

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## 1 1 Wrong implant selection

- 1.1 2 Review question: What interventions would reduce the
  - 3 number of intraoperative implant selection errors,
  - 4 including systems and processes for selection, in adults
  - 5 having primary elective joint replacement?

#### 1.2 6 Introduction

- 7 Wrong implant selection for primary elective joint replacement refers to a situation when the
- 8 prosthesis implanted into a patient by the surgeon is the incorrect size, wrong side or where
- 9 parts of the prosthesis used are incompatible with each other. This is often termed a
- 10 'mismatch' and is a rare event. Sometimes this is recognised by the surgeon and operating
- 11 theatre team and can be addressed at the time of the person's replacement, but often it
- 12 requires a revision. If not recognised at the time, the situation is picked up by the National
- 13 Joint Register as a 'Never Event' and the person informed.
- 14 This review seeks to establish what systems or processes could be implemented to prevent
- 15 any never events in relation to wrong implant selection.

16

#### 1.3<sub>17</sub> PICO table

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

#### 19 Table 1: PICO characteristics of review question

Population	Adults having primary elective joint replacement.
	Indirect populations that will be considered:  People having a pacemaker fitted  People having maxillofacial implant surgery  People having ocular prosthesis surgery
Intervention(s)	Interventions to reduce incorrect implant use. For example:  Clearer labelling on the implant packaging  Regimen for implant verification  Use of new technology  Colour coding  Shared learning/training  Scan for safety  Unique device identifiers
Comparison(s)	Usual care
Outcomes	<ul> <li>Critical</li> <li>Incorrect implant use (dichotomous)</li> <li>Revision rate</li> <li>Revision surgery (time to event)</li> <li>Mortality: life expectancy (time to event)</li> <li>Mortality: 30 day (dichotomous)</li> <li>Quality of life (continuous)</li> <li>Important</li> </ul>



- Hospital readmission (dichotomous)
- Length of stay (continuous)
- Enhanced follow up recommend blood tests, cross sectional imaging (dichotomous)

Randomised controlled trials

#### 1.4 1 Clinical evidence

#### 1.4.1 2 Included studies

- 3 A search was conducted for randomised trials comparing interventions to reduce wrong
- 4 implant selection with usual care, and no relevant clinical studies were identified.
- 5 See also the study selection flow chart in Appendix C:, study evidence tables in Appendix D:,
- 6 forest plots in Appendix E: and GRADE tables in Appendix H:.

#### 1.4.2 7 Excluded studies

8 See the excluded studies list in **Table 6**.

#### 1.5 9 Economic evidence

#### 1.5.110 Included studies

11 No health economic studies were included.

#### 1.5.2 Excluded studies

- 13 No relevant health economic studies were excluded due to assessment of limited
- 14 applicability or methodological limitations.
- 15 See also the health economic study selection flow chart in Appendix G:.

#### 1.6<sub>16</sub> Evidence statements

#### 1.6.117 Clinical evidence statements

18 No relevant published evidence was identified.

#### 1.6.29 Health economic evidence statements

20 No relevant economic evaluations were identified.

21

#### 1.7<sub>22</sub> The committee's discussion of the evidence

#### 1.7.123 Interpreting the evidence

#### 1.7.1.24 The outcomes that matter most

- 25 The critical outcomes were incorrect implant use, revision rate, revision surgery, mortality
- 26 and quality of life. The most critical outcome was incorrect implant selection and use as this
- 27 is the issue this evidence review seeks to address. The other outcomes are thought likely to
- 28 be affected by incorrect implant use such as earlier revision surgery and reduced quality of

- 1 life that might be associated with revision surgery or a joint replacement that is not
- 2 functioning as well as it could if the correct implant was used. The important outcomes were
- 3 hospital readmission, length of stay and enhanced follow up.

#### 1.7.1.2 5 The quality of the evidence

6 No evidence was found comparing interventions to reduce incorrect implant use with usual 7 care.

8

#### 1.7.1.3 9 Benefits and harms

- 10 The committee agreed that wrong implant selection and implantation is a very rare event but
- 11 the implications of such an error are potentially significant for the person who has undergone
- 12 joint replacement surgery. These are often referred to as "never events" because they are
- 13 viewed as preventable and caused by human and process error. It was noted that this is an
- 14 issue that exists in a wider context than simply orthopaedic surgery and indeed the evidence
- 15 review was expanded to look for studies in people who are having other implant surgery such
- 16 as having a pacemaker fitted, maxillofacial implant surgery, or ocular prosthesis surgery.
- 17 How implant selection errors can occur was discussed by the committee. The implant should
- 18 be ultimately checked by the surgeon before implantation, after initial checks by the scrub
- 19 nurse, runner, and sometimes by an industry representative supporting the case. However
- 20 despite multiple parties checking the prosthesis, implant selection errors still occur. The
- 21 committee agreed that there is unlikely to be a single intervention that will solve this and it
- 22 should be approached from multiple angles.
- 23 The committee discussed the work that has been done to reduce or eradicate implant
- 24 selection errors. Two national safety initiatives have provided guidance in this area. The
- 25 National Safety Standards for Invasive Procedures<sup>48</sup> (NatSSIPs), an NHS Improvement
- 26 initiative to reduce the number of patient safety incidents related to invasive procedures in
- 27 which surgical never events could occur. Recommendations have been for prosthesis
- 28 verification before the procedure, during the procedure, and after the procedure. The
- 29 Healthcare Safety Investigation Branch (HSIB) published a report on its investigation into the
- 30 implantation of wrong prostheses during joint replacement surgery. 31 This made 5 safety
- 31 recommendations aimed at reducing wrong implant selection.
- 32 Current joint replacement surgery practice requires the use of the WHO surgical safety
- 33 checklist to address local and national safety data which could include Never Events and
- 34 National Patient Safety Agency (NPSA) alerts.
- 35 The evidence review found no relevant evidence, resulting in the committee using their
- 36 experience and knowledge of current evidence to make consensus recommendations to
- 37 reduce implant selection errors. They spoke of the importance of stop moments during
- 38 surgery. This is where other actions are ceased by all staff and the details of the prosthesis
- 39 components are checked so everyone agrees the correct prostheses are going to be
- 40 implanted. The committee believe these time outs are common practice but are uncertain it is
- 41 universal and stated how important it is to formally undertake them.
- 42 The 2<sup>nd</sup> recommendation is to use real time data entry before implantation using a system
- 43 that will highlight some instances of wrong selection. The committee spoke about NJR Data
- 44 collection forms that are normally completed after the surgery is completed. All hip and knee
- 45 joint replacement operations undertaken in the UK are put into this database. When these
- 46 data collection forms are inputted onto the NJR database incorrect combinations of implants
- 47 are highlighted. However at this point the joint replacement operation has already occurred

- 1 and any changes would require revision surgery. The committee considered the advantage
- 2 of detecting incorrect combinations of implants intraoperatively so they could be prevented
- 3 from occurring and made a recommendation to consider this.
- 4 The committee were aware that undertaking this intraoperative data entry has process
- 5 implications for the orthopaedic centres undertaking the surgery. The committee were keen
- 6 not to be prescriptive in terms of the specific system used whether it is the NJR database or
- 7 an alternative system or how the data is entered, the method of data entry be it barcode,
- 8 RFID or manual entry, or indeed the person entering the data be it OR staff or a HCA data
- 9 collection clerk. The key result is "real time" warnings of wrong implant selection that can be
- 10 acted on prior to implantation.
- 11 The committee agreed that there could be technological solutions to supplement current
- 12 manual checks that may help reduce errors and therefore the made a research
- 13 recommendation.

#### 1.7.215 Cost effectiveness and resource use

- 16 No economic evidence was found for this review question. The first recommendation would
- 17 not represent any significant additional use of resources. An intraoperative pause to check
- 18 implant details and compatibility would not require additional personal and could also be
- 19 conducted in a matter of seconds.
- 20 It was noted there may be economic implications to entering data intraoperatively, depending
- 21 on how this is done. One option would be to install scanners that can inform the surgeons if
- 22 the correct components are being used. This option is likely to have a large resource impact
- 23 as there is national variability in terms of technology and IT infrastructure in hospitals.
- 24 Therefore if technological investments were required for real time scanning, the costs
- 25 incurred for some hospitals may be more than for others. There is also the option of having
- 26 this check done manually. Component details are already entered into the NJR manually
- 27 post-operatively for all procedures, however, it could be possible to move or copy this
- 28 process during the intraoperative period. This would represent less of a resource impact and
- 29 is better than no check; however, the implementation of technology would be the safest
- 30 choice.
- 31 Given there was no clinical or cost effectiveness evidence, the committee did not specify how
- 32 the real time scanning should be done.

33

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## 1 Appendices

## 2 Appendix A: Review protocols

3 Table 2: Review protocol: Wrong implant selection

ıα	ible 2. Review protocol. Wrong implant selection	
ID	Field	Content
0.	PROSPERO registration number	Not registered
1.	Review title	Reducing wrong implant selection during joint replacement
2.	Review question	What interventions would reduce the number of intraoperative implant selection errors, including systems and processes for selection, in adults having primary elective joint replacement?
3.	Objective	Surgical placement of the wrong implant or prosthesis where the implant/prosthesis placed in the patient is other than that specified in the surgical plan. This review seeks to find systems, processes or methods that will reduce these errors.
4.	Searches	The following databases will be searched: Cochrane Central Register of Controlled Trials (CENTRAL) Cochrane Database of Systematic Reviews (CDSR) Embase MEDLINE  Searches will be restricted by: English language Human studies Letters and comments are excluded.  Other searches: Inclusion lists of relevant systematic reviews will be checked by the reviewer.  The searches may be re-run 6 weeks before final committee meeting and further studies retrieved for inclusion if relevant.  The full search strategies will be published in the final review.
5.	Condition or domain being studied	Methods to reduce wrong implant selection during joint replacement

ID	Field	Content
6.	Population	<ul> <li>Adults having primary elective joint replacement.</li> <li>Indirect populations that will be considered:</li> <li>People having a pacemaker fitted</li> <li>People having maxillofacial implant surgery</li> <li>People having ocular prosthesis surgery</li> <li>Include mixed studies with adults having primary elective joint replacement and:</li> <li>Adults having joint replacement as immediate treatment following fracture.</li> <li>Adults having revision joint replacement.</li> <li>Adults having joint replacement as treatment for primary or secondary cancer affecting the bones.</li> <li>Exclusion:</li> <li>N/A</li> </ul>
7.	Intervention/Exposure/T est	Interventions to reduce incorrect implant use. For example: Clearer labelling on the implant packaging Regimen for implant verification Use of new technology Colour coding Shared learning/training Scan for safety Unique device identifiers
8.	Comparator/Reference standard/Confounding factors	Usual care
9.	Types of study to be included	Randomised controlled trials
10.	Other exclusion criteria	Non-English language studies.  Abstracts will be excluded as it is expected there will be sufficient full text published studies available.
11.	Context	N/A

ID	Field	Content
12.	Primary outcomes (critical outcomes)	Incorrect implant use (dichotomous) Revision rate Revision surgery (time to event) Mortality: life expectancy (time to event) Mortality: 30 day (dichotomous) Quality of life (continuous)
13.	Secondary outcomes (important outcomes)	Hospital readmission (dichotomous) Length of stay (continuous) Enhanced follow up – recommend blood tests, cross sectional imaging (dichotomous)
14.	Data extraction (selection and coding)	EndNote will be used for reference management, sifting, citations and bibliographies. Titles and/or abstracts of studies retrieved using the search strategy and those from additional sources will be screened for inclusion.  The full text of potentially eligible studies will be retrieved and will be assessed for eligibility in line with the criteria outlined above.  10% of the abstracts will be reviewed by two reviewers, with any disagreements resolved by discussion or, if necessary, a third independent reviewer.  An in-house developed database; EviBase, will be used for data extraction. A standardised form is followed to extract data from studies (see Developing NICE guidelines: the manual section 6.4) and for undertaking assessment of study quality. Summary evidence tables will be produced including information on: study setting; study population and participant demographics and baseline characteristics; details of the intervention and control interventions; study methodology' recruitment and missing data rates; outcomes and times of measurement; critical appraisal ratings.  A second reviewer will quality assure the extracted data. Discrepancies will be identified and resolved through discussion (with a third reviewer where necessary).
15.	Risk of bias (quality) assessment	Risk of bias will be assessed using the appropriate checklist as described in Developing NICE guidelines: the manual. For Intervention reviews the following checklist will be used according to study design being assessed:  Systematic reviews: Risk of Bias in Systematic Reviews (ROBIS)  Randomised Controlled Trial: Cochrane RoB (2.0)  Disagreements between the review authors over the risk of bias in particular studies will be resolved by discussion, with involvement of a third review author where necessary.
16.	Strategy for data	Where possible, data will be meta-analysed. Pairwise meta-analyses will be performed using Cochrane Review Manager

ID	Field	Content	
	synthesis	with weighted mean differences for continuous outcomes a confidence intervals will be calculated for each outcome. Heterogeneity between the studies in effect measures will	h of the outcomes stated above. A fixed effect meta-analysis, and risk ratios for binary outcomes will be used, and 95% be assessed using the I <sup>2</sup> statistic and visually inspected. We will be the terogeneity. Sensitivity analyses will be conducted based
			o explore the heterogeneity in effect estimates. If this does not
GRADE pro will be used to assess the quality of each outcome, taking into account analysis results. The 4 main quality elements (risk of bias, indirectness, inconsistent for each outcome.			
			nildren aged under 12, it will be included if the majority of the ss if the overlap into those aged less than 12 is greater than
	Publication bias is tested for when there are more than 5 studies for an outcome.  Other bias will only be taken into consideration in the quality assessment if it is apparent.		
		Where meta-analysis is not possible, data will be presented	d and quality assessed individually per outcome.
		If sufficient data is available to make a network of treatmer	nts, WinBUGS will be used for network meta-analysis.
17.	Analysis of sub-groups	Site of joint replacement: knee shoulder hip	
18.	Type and method of		Intervention
	review		Diagnostic
			Prognostic
			Qualitative
			Epidemiologic

ID	Field	Content			
			Service Delivery		
			Other (please spe	ecify)	
19.	Language	English			
20.	Country	England			
21.	Anticipated or actual start date	25/04/19			
22.	Anticipated completion date	20/03/20			
23.	Stage of review at time	Review stage	;	Started	Completed
	of this submission	Preliminary searches			▼
		Piloting of the study selection process			V
		Formal screening of search results against eligibility criteri	a		✓
		Data extraction			✓
		Risk of bias (quality) assessment			✓
		Data analysis			✓
24.	Named contact	<ul> <li>5a. Named contact National Guideline Centre</li> <li>5b Named contact e-mail TBC</li> <li>5e Organisational affiliation of the review National Institute for Health and Care Excellence (NICE) and the National Guideline Centre</li> </ul>			
25.	Review team members	From the National Guideline Centre:  Mr Carlos Sharpin [Guideline lead]  Mr Alex Allen [Senior Systematic Reviewer]			

ID	Field	Content	
		Ms Rafina Yarde [Systematic reviewer] Mr Robert King [Health economist] Ms Agnès Cuyàs [Information specialist] Ms Eleanor Priestnall [Project Manager]	
26.	Funding sources/sponsor	This systematic review is being completed by the National	Il Guideline Centre which receives funding from NICE.
27.	Conflicts of interest	team and expert witnesses) must declare any potential codeclaring and dealing with conflicts of interest. Any relevan publicly at the start of each guideline committee meeting, considered by the guideline committee Chair and a senior	Before each meeting, any potential conflicts of interest will be remember of the development team. Any decisions to exclude a Any changes to a member's declaration of interests will be
28.	Collaborators		y an advisory committee who will use the review to inform the with section 3 of Developing NICE guidelines: the manual. NICE website: [NICE guideline webpage].
29.	Other registration details		
30.	Reference/URL for published protocol		
31.	Dissemination plans	NICE may use a range of different methods to raise awar as: notifying registered stakeholders of publication publicising the guideline through NICE's newsletter and a issuing a press release or briefing as appropriate, posting channels, and publicising the guideline within NICE.	
32.	Keywords	Wrong implant, prosthesis, joint replacement, clear labelli	ng, checking, data entry
33.	Details of existing review of same topic by same authors	N/A	
34.	Current review status		Ongoing
			Completed but not published

ID	Field	Content	
			Completed and published
			Completed, published and being updated
			Discontinued
35	Additional information	N/A	
36.	Details of final publication	www.nice.org.uk	

#### 1 Table 3: Health economic review protocol

	attri economic review protocor
Review question	All questions – health economic evidence
Objectives	To identify health economic studies relevant to any of the review questions.
Search criteria	<ul> <li>Populations, interventions and comparators must be as specified in the clinical review protocol above.</li> </ul>
	<ul> <li>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).</li> </ul>
	<ul> <li>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.)</li> </ul>
	<ul> <li>Unpublished reports will not be considered unless submitted as part of a call for evidence.</li> </ul>
_	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	Studies not meeting any of the search criteria above will be excluded. Studies published before 2003, abstract-only studies and studies from low or middle-income countries (e.g. most non-OECD countries) or the USA will also be excluded.
	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).
	Inclusion and exclusion criteria
	<ul> <li>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.</li> </ul>
	<ul> <li>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.</li> </ul>
	<ul> <li>If a study is rated as 'Partially applicable', with 'Potentially serious limitations' or both then there is discretion over whether it should be included.</li> </ul>
	Where there is discretion
	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.
	The health economist will be guided by the following hierarchies.  Setting:  A LIK NHS (most applicable)
	<ul> <li>UK NHS (most applicable).</li> <li>OECD countries with predominantly public health insurance systems (for example, France, Germany, Sweden).</li> </ul>
	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

- 2 The literature searches for this review are detailed below and complied with the methodology
- 3 outlined in Developing NICE guidelines: the manual. 46
- 4 For more detailed information, please see the Methodology Review.

#### **B.1**<sup>5</sup> Clinical search literature search strategy

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

#### 11 Table 4: Database date parameters and filters used

Database	Dates searched	Search filter used
Medline (OVID)	1946 – 01 May 2019	Exclusions Randomised controlled trials Systematic review studies Observational studies
Embase (OVID)	1974 – 01 May 2019	Exclusions Randomised controlled trials Systematic review studies Observational studies
The Cochrane Library (Wiley)	Cochrane Reviews to 2019 Issue 5 of 12 CENTRAL to 2019 Issue 5 of 12	None

#### 12 Medline (Ovid) search terms

1.	arthroplasty/ or arthroplasty, replacement/ or arthroplasty, replacement, hip/ or arthroplasty, replacement, knee/ or arthroplasty, replacement, shoulder/ or hemiarthroplasty/ or exp Maxillofacial Prosthesis Implantation/
2.	joint prosthesis/ or hip prosthesis/ or knee prosthesis/ or shoulder prosthesis/ or eye, artificial/ or exp maxillofacial prosthesis/
3.	((joint* or knee* or shoulder* or hip* or ocular* or maxillofacial*) adj5 (surger* or replace* or prosthe* or endoprosthe* or implant* or artificial or arthroplast* or hemiarthroplast*)).ti,ab.
4.	((femoral or patellar or polyethylene or tibial or humeral or glenoid) adj2 component*).ti,ab.
5.	(((femoral or humeral) adj2 (steam or head)) or (acetabular or liner)).ti,ab.
6.	Pacemaker, Artificial/ or Cardiac Pacing, Artificial/
7.	(pacemaker* adj3 (implant* or fit* or surg*)).ti,ab.
8.	or/1-7
9.	letter/
10.	editorial/
11.	news/
12.	exp historical article/
13.	Anecdotes as Topic/
14.	comment/

15.	case report/
16.	(letter or comment*).ti.
17.	or/9-16
18.	randomized controlled trial/ or random*.ti,ab.
19.	17 not 18
20.	animals/ not humans/
21.	exp Animals, Laboratory/
22.	exp Animal Experimentation/
23.	exp Models, Animal/
24.	exp Rodentia/
25.	(rat or rats or mouse or mice).ti.
26.	or/19-25
27.	8 not 26
28.	limit 27 to English language
29.	"Prostheses and Implants"/ae, st [Adverse Effects, Standards]
30.	(wrong* or incorrect* or erroneous or error* or mistak* or unsuitabl* or incompatibl*).ti,ab.
31.	medical errors/ or diagnostic errors/
32.	exp Malpractice/
33.	(malpractice* or negligen*).ti,ab.
34.	(confus* or disorganiz* or disorganis* or distract*).ti,ab.
35.	((never or sentinel) adj4 event*).ti,ab.
36.	or/29-35
37.	28 and 36
38.	randomized controlled trial.pt.
39.	controlled clinical trial.pt.
40.	randomi#ed.ti,ab.
41.	placebo.ab.
42.	randomly.ti,ab.
43.	Clinical Trials as topic.sh.
44.	trial.ti.
45.	or/38-44
46.	Meta-Analysis/
47.	exp Meta-Analysis as Topic/
48.	(meta analy* or metanaly* or metaanaly* or meta regression).ti,ab.
49.	((systematic* or evidence*) adj3 (review* or overview*)).ti,ab.
50.	(reference list* or bibliograph* or hand search* or manual search* or relevant journals).ab.
51.	(search strategy or search criteria or systematic search or study selection or data extraction).ab.
52.	(search* adj4 literature).ab.
53.	(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.
54.	cochrane.jw.
55.	((multiple treatment* or indirect or mixed) adj2 comparison*).ti,ab.
56.	or/46-55

57.	Epidemiologic studies/
58.	Observational study/
59.	exp Cohort studies/
60.	(cohort adj (study or studies or analys* or data)).ti,ab.
61.	((follow up or observational or uncontrolled or non randomi#ed or epidemiologic*) adj (study or studies or data)).ti,ab.
62.	((longitudinal or retrospective or prospective or cross sectional) and (study or studies or review or analys* or cohort* or data)).ti,ab.
63.	Controlled Before-After Studies/
64.	Historically Controlled Study/
65.	Interrupted Time Series Analysis/
66.	(before adj2 after adj2 (study or studies or data)).ti,ab.
67.	or/57-66
68.	exp case control study/
69.	case control*.ti,ab.
70.	or/68-69
71.	67 or 70
72.	Cross-sectional studies/
73.	(cross sectional and (study or studies or review or analys* or cohort* or data)).ti,ab.
74.	or/72-73
75.	67 or 74
76.	67 or 70 or 74
77.	37 and (45 or 56 or 76)

#### 1 Embase (Ovid) search terms

1.	*arthroplasty/ or *replacement arthroplasty/ or *hip replacement/ or *knee replacement/ or *shoulder replacement/ or *hemiarthroplasty/ or *maxillofacial implant/
2.	*joint prosthesis/ or *hip prosthesis/ or *knee prosthesis/ or *shoulder prosthesis/ or *maxillofacial prosthesis/
3.	((joint* or knee* or shoulder* or hip* or ocular* or maxillofacial*) adj5 (surger* or replace* or prosthe* or endoprosthe* or implant* or artificial or arthroplast* or hemiarthroplast*)).ti,ab.
4.	((femoral or patellar or polyethylene or tibial or humeral or glenoid) adj2 component*).ti,ab.
5.	(((femoral or humeral) adj2 (steam or head)) or (acetabular or liner)).ti,ab.
6.	*artificial heart pacemaker/ or *heart pacing/
7.	(pacemaker* adj3 (implant* or fit* or surg*)).ti,ab.
8.	or/1-7
9.	letter.pt. or letter/
10.	note.pt.
11.	editorial.pt.
12.	case report/ or case study/
13.	(letter or comment*).ti.
14.	or/9-13
15.	randomized controlled trial/ or random*.ti,ab.
16.	14 not 15
17.	animal/ not human/
18.	nonhuman/
	·

19.	ovn Animal Evnoriment/
	exp Animal Experiment/
20.	exp Experimental Animal/
21.	animal model/
22.	exp Rodent/
23.	(rat or rats or mouse or mice).ti.
24.	or/16-23
25.	8 not 24
26.	limit 25 to English language
27.	(wrong* or incorrect* or erroneous or error* or mistak* or unsuitabl* or incompatibl*).ti,ab.
28.	*medical error/ or *diagnostic error/
29.	*malpractice/
30.	(malpractice* or negligen*).ti,ab.
31.	(confus* or disorganiz* or disorganis* or distract*).ti,ab.
32.	((never or sentinel) adj4 event*).ti,ab.
33.	or/27-32
34.	26 and 33
35.	random*.ti,ab.
36.	factorial*.ti,ab.
37.	(crossover* or cross over*).ti,ab.
38.	((doubl* or singl*) adj blind*).ti,ab.
39.	(assign* or allocat* or volunteer* or placebo*).ti,ab.
40.	crossover procedure/
41.	single blind procedure/
42.	randomized controlled trial/
43.	double blind procedure/
44.	or/35-43
45.	systematic review/
46.	meta-analysis/
47.	(meta analy* or metanaly* or metanaly* or meta regression).ti,ab.
48.	((systematic* or evidence*) adj3 (review* or overview*)).ti,ab.
49.	((systematic of evidence / adjo (review of overview /).ti,ab.  (reference list* or bibliograph* or hand search* or manual search* or relevant
45.	journals).ab.
50.	(search strategy or search criteria or systematic search or study selection or data extraction).ab.
51.	(search* adj4 literature).ab.
52.	(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.
53.	cochrane.jw.
54.	((multiple treatment* or indirect or mixed) adj2 comparison*).ti,ab.
55.	or/45-54
56.	Clinical study/
57.	Observational study/
58.	family study/
59.	longitudinal study/
60.	retrospective study/
	- Chaspoonto Guayi

61.	prospective study/
62.	cohort analysis/
63.	follow-up/
64.	cohort*.ti,ab.
65.	63 and 64
66.	(cohort adj (study or studies or analys* or data)).ti,ab.
67.	((follow up or observational or uncontrolled or non randomi#ed or epidemiologic*) adj (study or studies or data)).ti,ab.
68.	((longitudinal or retrospective or prospective or cross sectional) and (study or studies or review or analys* or cohort* or data)).ti,ab.
69.	(before adj2 after adj2 (study or studies or data)).ti,ab.
70.	or/56-62,65-69
71.	exp case control study/
72.	case control*.ti,ab.
73.	or/71-72
74.	70 or 73
75.	cross-sectional study/
76.	(cross sectional and (study or studies or review or analys* or cohort* or data)).ti,ab.
77.	or/75-76
78.	70 or 77
79.	70 or 73 or 77
80.	34 and (44 or 55 or 79)

#### 1 Cochrane Library (Wiley) search terms

#1.	MeSH descriptor: [Arthroplasty] this term only
#2.	MeSH descriptor: [Arthroplasty, Replacement] this term only
#3.	MeSH descriptor: [Arthroplasty, Replacement, Hip] this term only
#4.	MeSH descriptor: [Arthroplasty, Replacement, Knee] this term only
#5.	MeSH descriptor: [Arthroplasty, Replacement, Shoulder] this term only
#6.	MeSH descriptor: [Hemiarthroplasty] this term only
#7.	MeSH descriptor: [Maxillofacial Prosthesis Implantation] explode all trees
#8.	(or #1-#7)
#9.	MeSH descriptor: [Joint Prosthesis] this term only
#10.	MeSH descriptor: [Hip Prosthesis] this term only
#11.	MeSH descriptor: [Knee Prosthesis] this term only
#12.	MeSH descriptor: [Shoulder Prosthesis] this term only
#13.	MeSH descriptor: [Maxillofacial Prosthesis] explode all trees
#14.	(or #9-#13)
#15.	((joint* or knee* or shoulder* or hip* or ocular* or maxillofacial*) near/5 (surger* or replace* or prosthe* or endoprosthe* or implant* or artificial or arthroplast* or hemiarthroplast*)):ti,ab
#16.	(((femoral or humeral) near/2 (steam or head)) or (acetabular or liner)):ti,ab
#17.	((femoral or patellar or polyethylene or tibial or humeral or glenoid) near/2 component*):ti,ab
#18.	MeSH descriptor: [Pacemaker, Artificial] this term only
#19.	MeSH descriptor: [Cardiac Pacing, Artificial] this term only
#20.	(pacemaker* near/3 (implant* or fit* or surg*)):ti,ab

#21.	(or #15-#20)
#22.	(or #8, #14, #21)
#23.	(wrong* or incorrect* or erroneous or error* or mistak* or unsuitabl* or incompatibl*):ti,ab
#24.	MeSH descriptor: [Medical Errors] this term only
#25.	MeSH descriptor: [Diagnostic Errors] this term only
#26.	MeSH descriptor: [Malpractice] this term only
#27.	(malpractice* or negligen*):ti,ab
#28.	(confus* or disorganiz* or disorganis* or distract*):ti,ab
#29.	((never or sentinel) near/4 event*):ti,ab
#30.	(or #23-#29)
#31.	#22 AND #30

#### **B.21** Health Economics literature search strategy

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

#### 8 Table 5: Database date parameters and filters used

Database	Dates searched	Search filter used
Medline	2014 – 01 May 2019	Exclusions Health economics studies
Embase	2014 – 01 May 2019	Exclusions Health economics studies
Centre for Research and Dissemination (CRD)	HTA - Inception – 01 May 2019 NHSEED - Inception to March 2015	None

#### 9 Medline (Ovid) search terms

1.	arthroplasty/ or arthroplasty, replacement/ or arthroplasty, replacement, hip/ or arthroplasty, replacement, knee/ or arthroplasty, replacement, shoulder/ or hemiarthroplasty/
2.	joint prosthesis/ or hip prosthesis/ or knee prosthesis/ or shoulder prosthesis/
3.	((joint* or knee* or shoulder* or hip*) adj5 (surger* or replace* or prosthe* or endoprosthe* or implant* or artificial or arthroplast* or hemiarthroplast*)).ti,ab.
4.	or/1-3
5.	letter/
6.	editorial/
7.	news/
8.	exp historical article/
9.	Anecdotes as Topic/
10.	comment/
11.	case report/
12.	(letter or comment*).ti.

13.	or/5-12
14.	randomized controlled trial/ or random*.ti,ab.
15.	13 not 14
16.	animals/ not humans/
17.	exp Animals, Laboratory/
18.	exp Animal Experimentation/
19.	exp Models, Animal/
20.	exp Rodentia/
21.	(rat or rats or mouse or mice).ti.
22.	or/15-21
23.	4 not 22
24.	limit 23 to English language
25.	Economics/
26.	Value of life/
27.	exp "Costs and Cost Analysis"/
28.	exp Economics, Hospital/
29.	exp Economics, Medical/
30.	Economics, Nursing/
31.	Economics, Pharmaceutical/
32.	exp "Fees and Charges"/
33.	exp Budgets/
34.	budget*.ti,ab.
35.	cost*.ti.
36.	(economic* or pharmaco?economic*).ti.
37.	(price* or pricing*).ti,ab.
38.	(cost* adj2 (effective* or utilit* or benefit* or minimi* or unit* or estimat* or variable*)).ab.
39.	(financ* or fee or fees).ti,ab.
40.	(value adj2 (money or monetary)).ti,ab.
41.	or/25-40
42.	24 and 41

#### 1 Embase (Ovid) search terms

*arthroplasty/ or *replacement arthroplasty/ or *hip replacement/ or *knee replacement/ or *shoulder replacement/ or *hemiarthroplasty/
*joint prosthesis/ or *hip prosthesis/ or *knee prosthesis/ or *shoulder prosthesis/
((joint* or knee* or shoulder* or hip*) adj5 (surger* or replace* or prosthe* or endoprosthe* or implant* or artificial or arthroplast* or hemiarthroplast*)).ti,ab.
or/1-3
letter.pt. or letter/
note.pt.
editorial.pt.
case report/ or case study/
(letter or comment*).ti.
or/5-9

11.	randomized controlled trial/ or random*.ti,ab.
12.	10 not 11
13.	animal/ not human/
14.	nonhuman/
15.	exp Animal Experiment/
16.	exp Experimental Animal/
17.	animal model/
18.	exp Rodent/
19.	(rat or rats or mouse or mice).ti.
20.	or/12-19
21.	4 not 20
22.	limit 21 to English language
23.	health economics/
24.	exp economic evaluation/
25.	exp health care cost/
26.	exp fee/
27.	budget/
28.	funding/
29.	budget*.ti,ab.
30.	cost*.ti.
31.	(economic* or pharmaco?economic*).ti.
32.	(price* or pricing*).ti,ab.
33.	(cost* adj2 (effective* or utilit* or benefit* or minimi* or unit* or estimat* or variable*)).ab.
34.	(financ* or fee or fees).ti,ab.
35.	(value adj2 (money or monetary)).ti,ab.
36.	or/23-35
37.	22 and 36

#### 1 NHS EED and HTA (CRD) search terms

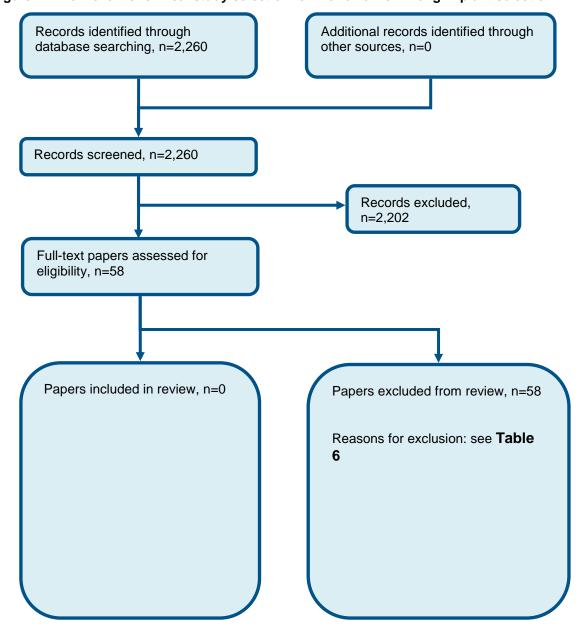
#1.	MeSH DESCRIPTOR arthroplasty
#2.	MeSH DESCRIPTOR arthroplasty, replacement
#3.	MeSH DESCRIPTOR arthroplasty, replacement, hip
#4.	MeSH DESCRIPTOR arthroplasty, replacement, knee
#5.	MeSH DESCRIPTOR arthroplasty, replacement, shoulder
#6.	MeSH DESCRIPTOR hemiarthroplasty
#7.	MeSH DESCRIPTOR joint prosthesis
#8.	MeSH DESCRIPTOR hip prosthesis
#9.	MeSH DESCRIPTOR knee prosthesis
#10.	MeSH DESCRIPTOR shoulder prosthesis
#11.	(((joint* or knee* or shoulder* or hip*) adj5 (surger* or replace* or prosthe* or endoprosthe* or implant* or artificial or arthroplast* or hemiarthroplast*)))
#12.	(#1 OR #2 OR #3 OR #4 OR #5 OR #6 OR #7 OR #8 OR #9 OR #10 OR #11) IN NHSEED

#13. (#1 OR #2 OR #3 OR #4 OR #5 OR #6 OR #7 OR #8 OR #9 OR #10 OR #11) IN HTA

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## 2 Appendix C: Clinical evidence selection

Figure 1: Flow chart of clinical study selection for the review of wrong implant selection



## <sup>1</sup> Appendix D: Clinical evidence tables

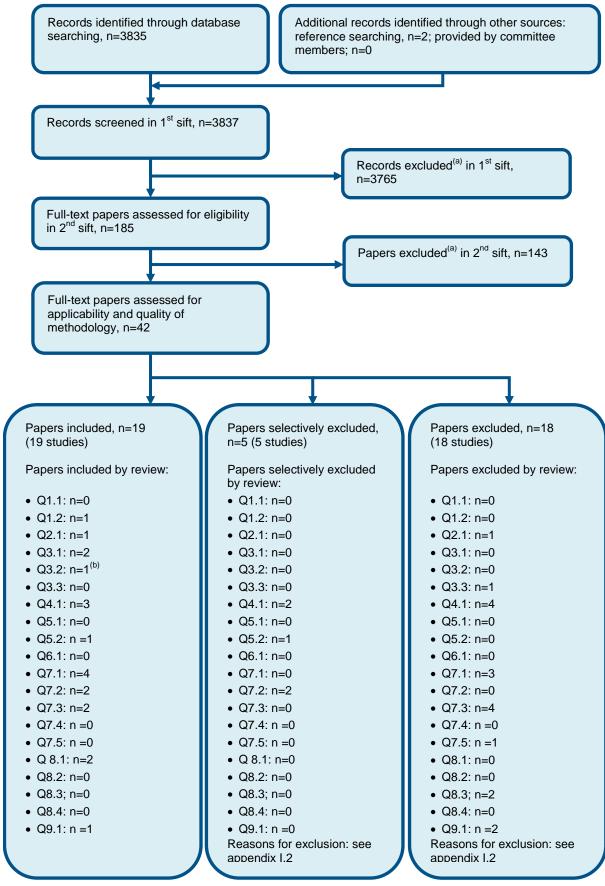
## **Appendix E: Forest plots**

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## <sup>1</sup> Appendix F: GRADE tables

## Appendix G: Health economic evidenceselection

Figure 2: Flow chart of health economic study selection for the guideline



- a) Non-relevant population, intervention, comparison, design or setting; non-English language
- b) One study was applicable to both Q3.1 and Q3.2

## <sup>1</sup> Appendix H: Health economic evidence tables

- 2 No health economic studies were included in this review
- 3
- 4

## 2 Appendix I: Excluded studies

#### I.13 Excluded clinical studies

#### 4 Table 6: Studies excluded from the clinical review

Table 0. Studies excluded from the chilical	ICAICAA
Actrn 2016 <sup>1</sup>	Trial web page
Al-Bataineh 2010 <sup>2</sup>	Incorrect study design, incorrect intervention
Asada 2014 <sup>3</sup>	Incorrect study design
Ast 2019 <sup>4</sup>	Incorrect study design
Bai 2014 <sup>5</sup>	Incorrect study design
Ballas 2013 <sup>6</sup>	Incorrect study design, incorrect comparisons
Bandeira 2018 <sup>7</sup>	Incorrect study design
Bargar 1998 <sup>8</sup>	Incorrect intervention
Bell 2016 <sup>9</sup>	Incorrect intervention
Bellemans 2007 <sup>10</sup>	Incorrect study design
Benedek 1984 <sup>11</sup>	Incorrect study design, incorrect intervention
Bernstein 1986 <sup>12</sup>	Incorrect study design
Bjorkenheim 2004 <sup>13</sup>	Incorrect study design
Bove 2010 <sup>14</sup>	Incorrect study design
Brandicourt 2017 <sup>15</sup>	Incorrect study design
Buchbender 2013 <sup>16</sup>	Incorrect study design, incorrect intervention, incorrect population
Cobb 2006 <sup>17</sup>	Incorrect intervention
Elmallah 2015 <sup>18</sup>	Incorrect comparisons
Fu 2018 <sup>19</sup>	Systematic review; references individually checked
Gan 2015 <sup>20</sup>	Incorrect intervention
Gandhi 2016 <sup>21</sup>	Incorrect study design
Gauci 2016 <sup>22</sup>	Incorrect study design
Goebel 2005 <sup>23</sup>	Not in English
Hafez 2006 <sup>24</sup>	Incorrect study population
Hampp 2019 <sup>25</sup>	Incorrect study population
Harrison 2010 <sup>26</sup>	Incorrect study design
Hashemian 2018 <sup>27</sup>	Incorrect study design
Hassan 1998 <sup>28</sup>	Incorrect study design
Hassanein 2017 <sup>29</sup>	Incorrect study design, incorrect population
Hayward 2015 <sup>30</sup>	Incorrect intervention, incorrect study design
Hoenecke 2010 <sup>32</sup>	Incorrect study design
Holt 1986 <sup>33</sup>	Incorrect comparisons
Hourlier 2014 <sup>34</sup>	Incorrect intervention
Isrctn 2014 <sup>35</sup>	Trial web page
Issa 2013 <sup>36</sup>	Incorrect study design
Jacobs 2002 <sup>37</sup>	Incorrect comparisons
Jacquot 2018 <sup>38</sup>	Incorrect comparisons

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Knafo 2019 <sup>39</sup>	Incorrect intervention
Lin 2011 <sup>40</sup>	Incorrect intervention
Liow 2014 <sup>41</sup>	Incorrect study design
Martelli 2000 <sup>42</sup>	Incorrect study design
Marx 2006 <sup>43</sup>	Incorrect study design
Michaels 2007 <sup>44</sup>	Incorrect study design
Naqvi 2016 <sup>45</sup>	Incorrect study design
Nguyen 2009 <sup>47</sup>	Incorrect population
Ozsoy 2009 <sup>49</sup>	Incorrect comparisons
Pagkalos 2014 <sup>50</sup>	Incorrect study design
Parsonnet 1974 <sup>52</sup>	Incorrect study design, incorrect intervention
Parsonnet 1975 <sup>51</sup>	Incorrect comparisons
Riddick 2014 <sup>53</sup>	Incorrect study design
Rodriguez 2005 <sup>54</sup>	Incorrect study design
Schulz 2009 <sup>55</sup>	Incorrect study design
Sendtner 2011 <sup>56</sup>	Incorrect intervention
Seyler 2008 <sup>57</sup>	Incorrect study design
Steppacher 2011 <sup>58</sup>	Incorrect study design
Stockl 2004 <sup>59</sup>	Incorrect intervention
Strik 2016 <sup>60</sup>	Incorrect study design, incorrect comparisons
Wirtz 1999 <sup>61</sup>	Incorrect study population

#### I.22 Excluded health economic studies

#### 3 Table 7: Studies excluded from the health economic review

Reference	Reason for exclusion
No studies	

## Appendix J: Research recommendations

#### J.12 Avoiding implant selection errors

- 3 Research question: What is the most effective technological solution for minimising
- 4 wrong implant selection during joint replacement surgery?
- 5 Why this is important:
- 6 The committee agreed that wrong implant selection and implantation is a very rare event but
- 7 the implications of such an error are potentially significant for the person who has undergone
- 8 joint replacement surgery. These are often referred to as "never events" because they are
- 9 viewed as preventable and caused by human and process error.

PICO question	Population: Adults undergoing primary hip, knee or shoulder joint replacement surgery Intervention(s): technological solution to prevent implant selection errors Comparison: Usual care Outcome(s): Implant selection errors and the resulting reduction in quality of life and economic costs
Study design	Randomised controlled trial
Other details	Two national safety initiatives have been asked to provide guidance in this area. The National Safety Standards for Invasive Procedures (NatSSIPs) and the Healthcare Safety Investigation Branch (HSIB). NatSSIPs is an NHS Improvement initiative that aims to reduce the number of patient safety incidents related to invasive procedures in which surgical never events could occur.