

Fertility problems: assessment and treatment

[H] Surgery for hydrosalpinges before IVF

NICE guideline NGXXX

Evidence reviews underpinning recommendations 1.6.3 and 1.6.4 in the NICE guideline

September 2025

Draft for consultation

This evidence review was developed by NICE

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1 Surgery for hydrosalpinges before IVF

2 Review question

3 What is the clinical and cost effectiveness of surgery for hydrosalpinges prior to assisted
4 reproductive technology (ART), relative to standard ART without prior surgical optimisation,
5 for people with tubal disease?

6 Introduction

7 Infertility related to problems with fallopian tubes (also known as salpinges) can be caused
8 by many factors such as pelvic inflammatory disease, endometriosis and previous abdominal
9 surgery. In cases of severe tubal damage with distal occlusion, the tube becomes distended
10 with fluid – this is referred to as hydrosalpinx. The presence of hydrosalpinx is particularly
11 associated with a reduced chance of success with in vitro fertilisation (IVF), because of a
12 reduced chance of implantation of the embryo. The mechanism whereby the likelihood of
13 implantation is reduced is not established but may involve a direct mechanical effect of
14 hydrosalpinx fluid leaking into the uterine cavity.

15 A number of methods have been studied for treating hydrosalpinges in people with tubal
16 disease contemplating IVF. These include surgical removal of the fallopian tube
17 (salpingectomy), or interruption of the medial end of the tube thereby preventing access of
18 fluid to the uterine cavity (proximal tubal occlusion), and aspiration of hydrosalpingeal fluid
19 prior to embryo transfer. However, the most effective intervention for treating hydrosalpinx in
20 people with tubal disease contemplating IVF is not clear.

21 The aim of this review is to determine the effectiveness of surgery for hydrosalpinges prior to
22 ART in people with tubal disease.

23 Summary of the protocol

24 See Table 1 for a summary of the Population, Intervention, Comparison and Outcome
25 (PICO) characteristics of this review.

26 **Table 1: Summary of the protocol (PICO table)**

Population	Inclusion: <ul style="list-style-type: none"> • People with a known diagnosis of tubal disease (confirmed by diagnostic surgery, hysterosalpingogram (HSG), hysterosalpingo-contrast sonography, or ultrasound)
Intervention	Tubal surgery (unilateral or bilateral) prior to in vitro fertilisation (IVF): <ul style="list-style-type: none"> • Salpingectomy • Tubal occlusion • Aspiration of hydrosalpinx fluid • Salpingostomy
Comparison	<ul style="list-style-type: none"> • Head-to-head comparisons of different tubal surgical interventions prior to IVF • No intervention or tubal treatment prior to IVF • Non-surgical intervention prior to IVF

Outcome	<p>Critical</p> <ul style="list-style-type: none"> • Live birth per participant randomised (defined as the delivery of a live fetus after 22 completed weeks of gestational age) • Surgical complication rate per participant randomised (including intraoperative bleeding, vasomotor instability, infection, need for repeat surgery, or overall complications as defined by the study) <p>Important</p> <ul style="list-style-type: none"> • Clinical pregnancy rate per participant randomised (defined as the presence of one or more gestational sacs on ultrasound) • Multiple pregnancy rate per participant randomised (defined as number of twin, triplet or higher-order pregnancies confirmed by ultrasound or delivery) • Miscarriage rate per participant randomised (defined as the spontaneous loss of an intrauterine pregnancy prior to 22 completed weeks of gestational age) • Ectopic pregnancy rate per participant randomised (defined as pregnancy outside the uterine cavity as diagnosed by ultrasound, surgical identification or histopathology)
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1 For further details see the review protocol in appendix A.

2 **Methods and process**

3 During the development of this guideline, a recently updated published Cochrane review was
4 identified which matched the committee's intended PICO (Melo 2020). Cochrane's methods
5 are closely aligned to standard NICE methods, minor deviations (defining primary and
6 secondary outcomes as opposed to critical and important) relevant to the topic area were
7 highlighted and taken into account by the committee in discussions of the evidence. Methods
8 specific to this review question are described in the review protocol in appendix A and the
9 methods document (supplementary document 1).

10 Declarations of interest were recorded according to [NICE's conflicts of interest policy](#).

11 **Effectiveness evidence**

12 **Included studies**

13 One Cochrane review (Melo 2020) was included in this report, comparing the effectiveness
14 and safety of unilateral and/or bilateral tubal surgical intervention (salpingectomy, tubal
15 occlusion, ultrasound guided/surgical aspiration of hydrosalpinx fluid or salpingostomy) for
16 tubal disease versus any other tubal surgery, non-surgical intervention prior to IVF or no
17 intervention.

18 The review includes 11 parallel design randomised controlled trials (RCTs; An 2015,
19 Dechaud 1998, Dreyer 2016, Fouda 2011, Fouda 2015, Hammadi 2008, Kontoravdis
20 2006, Labib 2016, Moshin 2006, Strandell 1999, Vignarajan 2019) and all these studies were
21 considered in this report. Nine studies were published in full and 2 were conference abstracts
22 (Labib 2016, Moshin 2006). This review was used by the committee to make
23 recommendations as it was considered sufficiently relevant, high quality and recent.

24 The Cochrane review is summarised in Table 2 and the results of the review are summarised
25 in evidence statements in this report, however full details of the Cochrane review including
26 methods are available here: <https://doi.org/10.1002/14651858.CD002125.pub4>.

27 See the Cochrane review for the literature search strategy and study selection flow chart
28 <https://doi.org/10.1002/14651858.CD002125.pub4>.

1 Excluded studies

2 See the Cochrane review for the list of excluded studies with reasons for their exclusions:
3 <https://doi.org/10.1002/14651858.CD002125.pub4>.

4 Summary of included studies

5 Summaries of the studies that were included in this review are presented in Table 2.

6 **Table 2: Summary of included studies**

Study	Population	Comparisons	Outcomes	Comments
Melo 2020	Number of studies = 11	<u>Salpingectomy vs no tubal surgery</u>	<u>Primary outcomes</u>	<u>IVF cycles</u>
Systematic review	Number of women with tubal disease analysed = 1386 (502 underwent salpingectomy; 294 underwent tubal occlusion; 256 had transvaginal aspiration of hydrosalpingeal fluid; and 334 had no tubal surgery)	2 RCTs, N=264 (Dechaud 1998, Strandell 1999)	<ul style="list-style-type: none"> • Live birth per woman randomised, defined as the delivery of a live fetus after 22 weeks of gestational age • Surgical complication rate per woman randomised (intraoperative bleeding or vasomotor instability, infection, need for repeat surgery or overall complications as reported by trialists) 	9 RCTs reported results over 1 IVF cycle except for 2 RCTs (Dechaud 1998 and Strandell 1999)
		<u>Transvaginal aspiration of hydrosalpingeal fluid vs no aspiration</u>		<u>Time interval between surgery and IVF</u>
		2 RCTs, N=176 (Fouda 2011, Hammadih 2008)		4 RCTs reported 2-3 months (Dreyer 2016, Kontoravdis 2006, Moshin 2006, Strandell 1999), 1 RCT no later than 12 weeks (Vignarajan 2019), 1 RCT 1-17 months (Dechaud 1998)
		<u>Tubal occlusion vs salpingectomy vs no tubal surgery for hydrosalpinges</u> (3 randomisation groups)		<u>Definition of tubal disease</u>
		2 RCTs, N=209 (Kontoravdis 2006, Moshin 2006#)		All included RCTs defined tubal disease as the presence of hydrosalpinx in imaging studies or at the time of laparoscopy, except Dechaud 1998 where tubal diverticula were, in isolation or concurrently to hydrosalpinx, sufficient to make a diagnosis of tubal disease
	The upper age limit ≤41 years	<u>Tubal occlusion vs salpingectomy</u>	<u>Secondary outcomes</u>	<u>Hydrosalpinx size</u>
		3 RCTs, N=332 (Dreyer 2016, Labib 2016#, Vignarajan 2019)	<ul style="list-style-type: none"> • Clinical pregnancy rate per woman randomised, defined as the presence of one or more gestational sacs on ultrasound* • Multiple pregnancy rate per woman randomised, defined as the number of twin, triplet or higher-order pregnancies confirmed by ultrasound or delivery • Miscarriage rate per woman randomised, defined as the 	Not reported in any of the studies
		In Dreyer 2016 study tubal occlusion was done using Essure® device		
		<u>Transvaginal aspiration of hydrosalpingeal fluid vs salpingectomy</u>		
		1 RCT, N=160 (Fouda 2015)		
		<u>Transvaginal aspiration of hydrosalpinges vs no tubal surgery</u>		
		1 RCT, N=135 (An 2015)		
		No studies were		

Study	Population	Comparisons	Outcomes	Comments
		<p>identified:</p> <ul style="list-style-type: none"> that compared tubal occlusion with aspiration of hydrosalpingeal fluid, where 1 of the intervention arms underwent salpingostomy 	<p>spontaneous loss of an intrauterine pregnancy prior to 22 completed weeks of gestational age</p> <ul style="list-style-type: none"> Ectopic pregnancy rate per woman randomised, defined as pregnancy outside the uterine cavity as diagnosed by ultrasound, surgical identification or histopathology <p><u>Non-relevant secondary outcomes included in the Cochrane review</u></p> <ul style="list-style-type: none"> Mean number of oocytes retrieved per woman randomised Mean number of embryos obtained per woman randomised 	<p><u>Presence/absence of ultrasound-visible fluid in the endometrial cavity</u></p> <p>Not reported in any of the studies</p>

1 # Moshin 2006 and Labib 2016 are conference abstracts; * Ongoing and viable pregnancy rates were combined
2 with clinical pregnancy rate, as ongoing pregnancy is currently not a recognised outcome by The International
3 Glossary on Infertility and Fertility Care 2017 (Melo 2020)
4 ART: assisted reproductive technology; IVF: in vitro fertilisation; N: number (the totals for each comparison were
5 calculated by the Technical Team based on the corresponding forest plots in the Cochrane review); RCT:
6 randomised controlled trial

7 See the Cochrane review (Melo 2020) for characteristics of studies tables and forest plots:
8 <https://doi.org/10.1002/14651858.CD002125.pub4>.

9 **Summary of the evidence**

10 The results of the Cochrane review are summarised below and interpreted according to the
11 minimal important differences (MIDs) used for this guideline (see Methods supplement).

12 **Comparison 1: Tubal surgery (all methods) vs no tubal surgery**

13 **Critical outcomes**

14 There was no relevant evidence for the outcome of live birth.

15 There was no evidence of clinically important difference between salpingectomy (all
16 methods) and no tubal surgery on surgical complication rate such as conversion to
17 laparotomy (Peto OR=5.80 (95% CI 0.11 to 303.69), 1 RCT, very low quality evidence) and
18 pelvic infection (Peto OR=5.80 (95% CI 0.11 to 303.69), 1 RCT, low quality evidence).

19 **Important outcomes**

- 1 Moderate quality evidence from 4 RCTs showed a higher clinical pregnancy rate in people
2 receiving salpingectomy (all methods) relative to no tubal surgery (RR=2.02 (95%CI 1.44 to
3 2.82)).
- 4 Low quality evidence from 2 RCTs showed a higher clinical pregnancy rate in people
5 receiving tubal occlusion (all methods) relative to no tubal surgery (RR=3.21 (95% CI 1.72 to
6 5.99)). Similarly, very low quality evidence from 3 RCTs showed a higher clinical pregnancy
7 rate in people receiving transvaginal aspiration of hydrosalpingeal fluid relative to no tubal
8 surgery (RR=1.67 (95% CI 1.10 to 2.55)).
- 9 There was no evidence of clinically important difference between transvaginal aspiration of
10 hydrosalpingeal fluid and no tubal surgery on multiple pregnancy rate (Peto OR=2.15 (95%
11 CI 0.59 to 7.85), 1 RCT, very low quality evidence).
- 12 There was no clinically important difference between salpingectomy (all methods) and no
13 tubal surgery on miscarriage rate (Peto OR=0.91 (95% CI 0.33 to 2.52, 3 RCTs, low quality
14 evidence).
- 15 There was no evidence of clinically important difference between tubal occlusion (all
16 methods) (Peto OR=0.55 (95% CI 0.04 to 8.43), 1 RCT, very low quality evidence),
17 transvaginal aspiration of hydrosalpingeal fluid (Peto OR=1.27 (95% CI 0.44 to 3.66), 3
18 RCTs, very low quality evidence) and no tubal surgery on miscarriage rate.
- 19 There was also no evidence of clinically important difference between salpingectomy (all
20 methods) (Peto OR=0.29 (95% CI 0.04 to 2.11), 3 RCTs, low quality evidence), tubal
21 occlusion (all methods) (Peto OR=3.67 (95% CI 0.04 to 384.48), 1 RCT, very low quality
22 evidence), transvaginal aspiration of hydrosalpingeal fluid (Peto OR=0.59 (95% CI 0.08 to
23 4.61), 3 RCTs, very low quality evidence) and no tubal surgery on ectopic pregnancy rate.
- 24 It is reported that a subgroup analysis for women younger and older than 40 years old was
25 not possible because of a lack of data.

26 ***Comparison 2: Laparoscopic proximal tubal occlusion (all methods) vs salpingectomy***

27 **Critical outcomes**

- 28 Very low quality evidence from 1 RCT showed no clinically important difference between
29 laparoscopic proximal tubal occlusion and salpingectomy on live birth rate (RR=1.21, (95%
30 CI 0.76 to 1.95)).
- 31 There was no relevant evidence for the outcome of surgical complication rate such as wound
32 infection or pelvic infection.

33 **Important outcomes**

- 34 Very low quality evidence from 3 RCTs showed no clinically important difference between
35 laparoscopic proximal tubal occlusion and salpingectomy on clinical pregnancy rate
36 (RR=0.81 (95% CI 0.62 to 1.07)).
- 37 There was no relevant evidence for the outcome of multiple pregnancy rate.
- 38 There was no evidence of a clinically important difference between laparoscopic proximal
39 tubal occlusion and salpingectomy on ectopic pregnancy rate (Peto OR=7.39, (95% CI 0.15
40 to 372.38), 1 RCT, very low quality evidence) or on miscarriage rate (Peto OR=0.74 (95% CI
41 0.16 to 3.34), 2 RCTs, low quality evidence).
- 42 It is reported that a subgroup analysis for women younger and older than 40 years old was
43 not possible because of a lack of data.

44 ***Comparison 3: Transvaginal aspiration of hydrosalpingeal fluid vs laparoscopic*** 45 ***salpingectomy***

1 **Critical outcomes**

2 There was no relevant evidence for the outcome of live birth.

3 It is reported that there were no cases of surgical complications and therefore the effect could
4 not be estimated.

5 **Important outcomes**

6 There was no evidence of clinically important difference between transvaginal aspiration of
7 hydrosalpingeal fluid and laparoscopic salpingectomy on clinical pregnancy rate (RR=0.69
8 (95% CI 0.44 to 1.07), 1 RCT, very low quality evidence).

9 There was no relevant evidence for the outcome of multiple pregnancy rate.

10 There was no clinically important difference between transvaginal aspiration of
11 hydrosalpingeal fluid and laparoscopic salpingectomy on miscarriage rate (Peto OR=1.00
12 (95% CI 0.20 to 5.08), 1 RCT, very low quality evidence).

13 There was no evidence of clinically important difference between transvaginal aspiration of
14 hydrosalpingeal fluid and laparoscopic salpingectomy on ectopic pregnancy rate (Peto
15 OR=7.39 (95% CI 0.15 to 372.28), 1 RCT, very low quality evidence).

16 ***Comparison 4: Tubal occlusion (all methods) vs aspiration of hydrosalpingeal fluid***

17 It is reported that no relevant evidence was identified for this comparison.

18 ***Comparison 5: Laparoscopic salpingectomy vs any other method of salpingectomy***

19 It is reported that no relevant evidence was identified for this comparison.

20 ***Comparison 6: Laparoscopic tubal occlusion vs hysteroscopic tubal occlusion***

21 It is reported that no relevant evidence was identified for this comparison.

22 See the Cochrane review for summary of findings tables and full results:

23 <https://doi.org/10.1002/14651858.CD002125.pub4>.

24 **Economic evidence**

25 A total of 2,632 studies were identified in the health economic literature review for this review
26 question. After duplicates were removed, 1,911 studies were screened on title and abstract.
27 Of these studies, 2 were ordered for full text review, of which both were excluded at this
28 stage.

29 **Included studies**

30 A systematic review of the economic literature was conducted but no economic studies were
31 identified which were applicable to this review question.

32 Also see the literature search strategy in appendix B and the economic study selection flow
33 chart in appendix F.

34 **Excluded studies**

35 Economic studies not included in this review are listed, and reasons for their exclusion are
36 provided in appendix J.

1 **Economic model**

2 No economic modelling was undertaken for this review because the committee agreed that
3 other topics were higher priorities for economic evaluation.

4 **Unit costs**

Resource	Unit costs	Source
Salpingectomy (elective inpatients)	£5,853	National Schedule of NHS Costs 2021-22, Major, Laparoscopic or Endoscopic, Upper Genital Tract Procedures, with CC Score 0-1, Currency Code MA08B, Elective inpatients
Salpingectomy (day case)	£3,628	National Schedule of NHS Costs 2021-22, Major, Laparoscopic or Endoscopic, Upper Genital Tract Procedures, with CC Score 0-1, Currency Code MA08B, Day case

5 **The committee's discussion and interpretation of the evidence**

6 **The outcomes that matter most**

7 The Cochrane review's primary outcomes included live birth, which the committee agreed
8 was a critical outcome because it is the most important outcome for people with fertility
9 problems. The review also included surgical complications as a primary outcome. The
10 committee agreed that surgical complications were critical to capture because it is important
11 when discussing and deciding on treatment options that risks are considered and weighed up
12 against potential benefits.

13 The Cochrane review included clinical pregnancy as a secondary outcome. The committee
14 agreed clinical pregnancy should be an important outcome as this reflects the evidence
15 available. They were aware that pregnancy rates do not allow for differentiation between full-
16 term pregnancy and pregnancy loss, and agreed that live birth was the most important
17 outcome and clinical pregnancy only acted as a proxy for live birth in the absence of
18 available evidence.

19 Multiple pregnancy was included in the Cochrane review as a secondary outcome and the
20 committee agreed that it was important to capture this outcome as it creates a greater risk for
21 complications in pregnancy and during delivery, such as early birth.

22 The Cochrane review included miscarriage as a secondary outcome, and the committee
23 agreed this outcome was important to capture because miscarriages can be devastating for
24 people trying to have a baby.

25 Ectopic pregnancy was also included in the Cochrane review as a secondary outcome and
26 the committee agreed that it was important to assess this outcome as the risk of ectopic
27 pregnancy may be higher with interventions that leave the fallopian tube in situ compared
28 with removal of the tube.

29 All other outcomes listed in the Cochrane protocol (mean number of oocytes retrieved per
30 woman randomised and mean number of embryos obtained per woman randomised)
31 were not prioritised by the committee because it was considered they do not provide patients
32 or clinicians with meaningful additional information to inform treatment decisions about
33 management of hydrosalpinges.

34 **The quality of the evidence**

35 The quality of the evidence was assessed using GRADE methodology and ranged from very
36 low to moderate quality. However, apart from 1 moderate-quality result, the quality of the
37 evidence was very low or low.

1 All outcomes were downgraded for serious imprecision either due to a small sample size
2 from a single study, wide confidence intervals or a low number of events.

3 Where outcomes were downgraded due to risk of bias, this was mainly due to unclear
4 blinding (of participants, personnel, and outcome assessors) and attrition bias (incomplete
5 outcome data). For some of the outcomes there was also insufficient detail reported about
6 random sequence generation and allocation concealment. In terms of other potential sources
7 of bias, some studies were assessed to be at unclear risk of bias mostly because of a lack of
8 information on baseline characteristics and 1 RCT was considered at high risk of other bias
9 because of recruitment ending prematurely.

10 The Cochrane review included 2 conference abstracts (Labib 2016, Moshin 2006); however
11 additional information was obtained from the authors by the Cochrane review authors.

12 **Benefits and harms**

13 The committee discussed the relevance of the Cochrane review as the subject of this review
14 is specific to surgery for hydrosalpinges and the Cochrane review is broader covering all
15 methods of tubal surgery. However, they agreed that the Cochrane review was in line with
16 this review's PICO as it evaluated tubal surgery before IVF and usually it is surgery for
17 hydrosalpinges that is performed before IVF.

18 The committee noted the lack of evidence for the outcome of live birth, but considered the
19 Cochrane review evidence that showed some benefits of surgery for hydrosalpinges before
20 ART on clinical pregnancy rates.

21 The committee discussed moderate quality evidence for a benefit of salpingectomy before
22 IVF on clinical pregnancy, and low quality evidence showing a benefit for tubal occlusion.
23 The committee also noted that very low quality evidence showed no clinically important
24 difference between salpingectomy and laparoscopic proximal tubal occlusion on live birth
25 rate (RR 1.21 – 95% CI 0.76 – 1.95). The committee agreed that both procedures should be
26 considered as treatment options for people with hydrosalpinges undergoing assisted
27 conception. While there is evidence of effectiveness for both procedures, the committee
28 acknowledged that the evidence base for tubal occlusion is somewhat weaker. However,
29 both interventions are used in practice and tubal occlusion could be preferred as a less
30 invasive procedure of the two. The committee noted that although salpingectomy may be
31 more invasive than tubal occlusion, salpingectomy may be the preferred option for example
32 in people with hydrosalpinx and chronic pelvic pain. Based on their knowledge of a wider
33 evidence base showing that ovarian cancer often starts in the fallopian tubes, the committee
34 also highlighted potential benefits of salpingectomy in terms of long-term risk reduction and
35 emphasised that this might be another reason for preferring salpingectomy over tubal
36 occlusion.

37 The committee noted that very low quality evidence showed a higher clinical pregnancy rate
38 in people undergoing transvaginal aspiration of hydrosalpingeal fluid relative to no tubal
39 surgery. Based on their clinical knowledge and experience, the committee noted that
40 aspiration of hydrosalpinx fluid is often followed by re-accumulation over a relatively short
41 period of time and aspiration alone does not provide a permanent solution to the problem of
42 hydrosalpinx fluid gaining access to the endometrium. The committee recognised that some
43 people with hydrosalpinges may be at a high risk of complications with laparoscopic surgery,
44 for instance, in the context of previous abdominal surgery and dense adhesions. The
45 committee agreed that aspiration of hydrosalpinx fluid should be considered for these
46 patients. However, the committee noted that of the studies in the Cochrane review that
47 showed a benefit for transvaginal aspiration of hydrosalpingeal fluid, the majority (and all the
48 studies that reported timing) performed the intervention immediately following oocyte
49 retrieval, and the committee agreed that it was important to include this in the
50 recommendation to reduce the risk of re-accumulation.

1 The committee discussed that in the study by Dreyer 2016 hysteroscopic tubal occlusion was
2 performed using the Essure® device, however they agreed to disregard the evidence about
3 this device because it has been discontinued due to safety concerns and is no longer
4 available.

5 **Cost effectiveness and resource use**

6 In the absence of any included evidence or original economic analysis, the committee made
7 a qualitative assessment of the cost-effectiveness of their recommendations.

8 The committee believed that it would be cost-effective to offer salpingectomy or tubal
9 occlusion before IVF in those women with hydrosalpinx as they believed that the evidence of
10 improved clinical pregnancy rate when compared to no tubal intervention would translate into
11 higher live births and therefore reduce the need for further IVF cycles.

12 The committee acknowledged that the clinical evidence showed no clinically important
13 difference for live birth rates, however noted this was very low quality evidence. The
14 committee discussed the positive correlation between clinical pregnancy rates and live birth
15 rates which they observe in clinical practice and highlighted the evidence of a positive
16 correlation which can be found in the clinical literature for studies assessing the impact of
17 fertility interventions – where both clinical pregnancy and live birth rates are reported
18 outcomes. As seen, for example in Rao 2018. The committee also noted that this cohort of
19 people are likely to experience greater difficulty achieving a clinical pregnancy as opposed to
20 experiencing problems carrying to term due to the nature of the condition. The committee
21 also discussed that the benefits of salpingectomy were not restricted to live births as surgery
22 can also lead to improvements in chronic pelvic pain. Therefore, the committee concluded
23 that their recommendation would be cost-effective producing additional live births at an
24 acceptable cost to the NHS.

25 In addition, the committee reasoned that aspiration could be cost-effective for the NHS in
26 women at high risk of complications from laparoscopic surgery. While noting the very low
27 quality of evidence for aspiration for the treatment of hydrosalpinges, they believed that a
28 consider recommendation was appropriate because of the different trade-off between
29 benefits and risks in a sub-group with higher risks of adverse surgical outcomes. Despite its
30 lower cost the committee did not believe that aspiration would be cost-effective relative to
31 salpingectomy as, in their experience, aspiration does not provide a permanent solution
32 because of re-accumulation of fluid over time.

33 The committee stated that their recommendations were largely aligned with existing NICE
34 guidance and also reflected current practice. They therefore did not anticipate a significant
35 resource impact to the NHS.

36 **Recommendations supported by this evidence review**

37 This evidence review supports recommendations 1.6.3 and 1.6.4.

38 **References – included studies**

39 **Effectiveness**

40 **Melo 2020**

41 Melo E, Georgiou EX, Johnson N, van Voorst SF, Strandell A, Mol BWJ et al. Surgical
42 treatment for tubal disease in women due to undergo in vitro fertilisation. Cochrane Database
43 of Systematic Reviews 2020, 10(10):CD002125. doi: 10.1002/14651858.CD002125.pub4.
44 Accessed 14 December 2023

1 **Other**

2 **Rao 2018**

- 3 Rao, M., Zeng, Z. & Tang, L. Maternal physical activity before IVF/ICSI cycles improves
4 clinical pregnancy rate and live birth rate: a systematic review and meta-analysis. *Reprod*
5 *Biol Endocrinol* **16**, 11 (2018). <https://doi.org/10.1186/s12958-018-0328-z>

Appendices

Appendix A Review protocols

Review protocol for review question: What is the clinical and cost effectiveness of surgery for hydrosalpinges prior to assisted reproductive technology (ART), relative to standard ART without prior surgical optimisation, for people with tubal disease?

Table 3: Review protocol

ID	Field	Content
0.	PROSPERO registration number	N/A
1.	Review title	Clinical and cost effectiveness of surgery for hydrosalpinges prior to assisted reproductive technology (ART) for people with tubal disease
2.	Review question	What is the clinical and cost effectiveness of surgery for hydrosalpinges prior to assisted reproductive technology (ART), relative to standard ART without prior surgical optimisation, for people with tubal disease?
3.	Objective	To determine the clinical and cost effectiveness of tubal surgery as an adjunct to assisted reproductive technology (ART) for people with tubal disease
4.	Searches	No searches will be conducted. A recent Cochrane review has been identified (Melo 2020) and this will be incorporated/adopted into the guideline
5.	Condition or domain being studied	Surgical interventions for female factor fertility problems
6.	Population	Inclusion: <ul style="list-style-type: none"> • People with a known diagnosis of tubal disease (confirmed by diagnostic surgery, hysterosalpingogram (HSG), hysterosalpingo-contrast sonography, or ultrasound)
7.	Intervention	Tubal surgery (unilateral or bilateral) prior to in vitro fertilisation (IVF): <ul style="list-style-type: none"> • Salpingectomy • Tubal occlusion • Aspiration of hydrosalpinx fluid

ID	Field	Content
		<ul style="list-style-type: none"> • Salpingostomy
8.	Comparator	<ul style="list-style-type: none"> • Head-to-head comparisons of different tubal surgical interventions prior to IVF • No intervention or tubal treatment prior to IVF • Non-surgical intervention prior to IVF
9.	Types of study to be included	This guideline will incorporate/adopt the recent Cochrane review identified (Melo 2020)
10.	Other exclusion criteria	<ul style="list-style-type: none"> • N/A
11.	Context	This guidance will fully update the following NICE guideline: Fertility problems: assessment and treatment (last updated 2017; CG156)
12.	Primary outcomes (critical outcomes)	<ul style="list-style-type: none"> • Live birth per participant randomised (defined as the delivery of a live fetus after 22 completed weeks of gestational age) • Surgical complication rate per participant randomised (including intraoperative bleeding, vasomotor instability, infection, need for repeat surgery, or overall complications as defined by the study)
13.	Secondary outcomes (important outcomes)	<ul style="list-style-type: none"> • Clinical pregnancy rate per participant randomised (defined as the presence of one or more gestational sacs on ultrasound) • Multiple pregnancy rate per participant randomised (defined as number of twin, triplet or higher-order pregnancies confirmed by ultrasound or delivery) • Miscarriage rate per participant randomised (defined as the spontaneous loss of an intrauterine pregnancy prior to 22 completed weeks of gestational age) • Ectopic pregnancy rate per participant randomised (defined as pregnancy outside the uterine cavity as diagnosed by ultrasound, surgical identification or histopathology)
14.	Data extraction (selection and coding)	Analyses from the Melo 2020 Cochrane review will be presented to the committee
15.	Risk of bias (quality) assessment	Risk of bias assessments from the Melo 2020 Cochrane review will be presented to the committee
16.	Strategy for data synthesis	Analyses from the Melo 2020 Cochrane review will be presented to the committee
17.	Analysis of sub-groups	<p>In Melo 2020, planned subgroup analyses are:</p> <ul style="list-style-type: none"> • Female age: <ul style="list-style-type: none"> • <40 years • ≥40 years

ID	Field	Content																					
		Where evidence is sub-grouped the committee will consider on a case-by-case basis if separate recommendations should be made for distinct groups. Separate recommendations may be made where there is evidence of a differential effect of interventions in distinct groups. If there is a lack of evidence in one group, the committee will consider, based on their experience, whether it is reasonable to extrapolate and assume the interventions will have similar effects in that group compared with others.																					
18.	Type and method of review	<table border="1"> <tr> <td><input checked="" type="checkbox"/></td><td>Intervention</td></tr> <tr> <td><input type="checkbox"/></td><td>Diagnostic</td></tr> <tr> <td><input type="checkbox"/></td><td>Prognostic</td></tr> <tr> <td><input type="checkbox"/></td><td>Qualitative</td></tr> <tr> <td><input type="checkbox"/></td><td>Epidemiologic</td></tr> <tr> <td><input type="checkbox"/></td><td>Service Delivery</td></tr> <tr> <td><input type="checkbox"/></td><td>Other (please specify)</td></tr> </table>	<input checked="" type="checkbox"/>	Intervention	<input type="checkbox"/>	Diagnostic	<input type="checkbox"/>	Prognostic	<input type="checkbox"/>	Qualitative	<input type="checkbox"/>	Epidemiologic	<input type="checkbox"/>	Service Delivery	<input type="checkbox"/>	Other (please specify)							
<input checked="" type="checkbox"/>	Intervention																						
<input type="checkbox"/>	Diagnostic																						
<input type="checkbox"/>	Prognostic																						
<input type="checkbox"/>	Qualitative																						
<input type="checkbox"/>	Epidemiologic																						
<input type="checkbox"/>	Service Delivery																						
<input type="checkbox"/>	Other (please specify)																						
19.	Language	English																					
20.	Country	England																					
21.	Anticipated or actual start date	September 2023																					
22.	Anticipated completion date	November 2024																					
23.	Stage of review at time of this submission	<table border="1"> <thead> <tr> <th>Review stage</th><th>Started</th><th>Completed</th></tr> </thead> <tbody> <tr> <td>Preliminary searches</td><td><input type="checkbox"/></td><td><input type="checkbox"/></td></tr> <tr> <td>Piloting of the study selection process</td><td><input type="checkbox"/></td><td><input type="checkbox"/></td></tr> <tr> <td>Formal screening of search results against eligibility criteria</td><td><input type="checkbox"/></td><td><input type="checkbox"/></td></tr> <tr> <td>Data extraction</td><td><input type="checkbox"/></td><td><input type="checkbox"/></td></tr> <tr> <td>Risk of bias (quality) assessment</td><td><input type="checkbox"/></td><td><input type="checkbox"/></td></tr> <tr> <td>Data analysis</td><td><input type="checkbox"/></td><td><input type="checkbox"/></td></tr> </tbody> </table>	Review stage	Started	Completed	Preliminary searches	<input type="checkbox"/>	<input type="checkbox"/>	Piloting of the study selection process	<input type="checkbox"/>	<input type="checkbox"/>	Formal screening of search results against eligibility criteria	<input type="checkbox"/>	<input type="checkbox"/>	Data extraction	<input type="checkbox"/>	<input type="checkbox"/>	Risk of bias (quality) assessment	<input type="checkbox"/>	<input type="checkbox"/>	Data analysis	<input type="checkbox"/>	<input type="checkbox"/>
Review stage	Started	Completed																					
Preliminary searches	<input type="checkbox"/>	<input type="checkbox"/>																					
Piloting of the study selection process	<input type="checkbox"/>	<input type="checkbox"/>																					
Formal screening of search results against eligibility criteria	<input type="checkbox"/>	<input type="checkbox"/>																					
Data extraction	<input type="checkbox"/>	<input type="checkbox"/>																					
Risk of bias (quality) assessment	<input type="checkbox"/>	<input type="checkbox"/>																					
Data analysis	<input type="checkbox"/>	<input type="checkbox"/>																					

ID	Field	Content
24.	Named contact	<p>5a. Named contact Guideline development team A</p> <p>5b. Named contact e-mail FertilityProblems@nice.org.uk</p> <p>5c. Organisational affiliation of the review Guideline Development Team A, Centre for Guidelines, National Institute for Health and Care Excellence (NICE)</p>
25.	Review team members	<ul style="list-style-type: none"> • Senior Technical Analyst • Technical Analyst
26.	Funding sources/sponsor	This systematic review is being completed by the National Institute for Health and Care Excellence (NICE)
27.	Conflicts of interest	All guideline committee members and anyone who has direct input into NICE guidelines (including the evidence review team and expert witnesses) must declare any potential conflicts of interest in line with NICE's code of practice for declaring and dealing with conflicts of interest. Any relevant interests, or changes to interests, will also be declared publicly at the start of each guideline committee meeting. Before each meeting, any potential conflicts of interest will be considered by the guideline committee Chair and a senior member of the development team. Any decisions to exclude a person from all or part of a meeting will be documented. Any changes to a member's declaration of interests will be recorded in the minutes of the meeting. Declarations of interests will be published with the final guideline.
28.	Collaborators	Development of this systematic review will be overseen by an advisory committee who will use the review to inform the development of evidence-based recommendations in line with section 3 of Developing NICE guidelines: the manual . Members of the guideline committee are available on the NICE website: https://www.nice.org.uk/guidance/indevelopment/gid-ng10263
29.	Other registration details	None
30.	URL for published protocol	N/A
31.	Dissemination plans	<p>NICE may use a range of different methods to raise awareness of the guideline. These include standard approaches such as:</p> <ul style="list-style-type: none"> • notifying registered stakeholders of publication

ID	Field	Content
		<ul style="list-style-type: none"> publicising the guideline through NICE's newsletter and alerts issuing a press release or briefing as appropriate, posting news articles on the NICE website, using social media channels, and publicising the guideline within NICE.
32.	Keywords	aspiration; hydrosalpinx; hydrosalpinges; IVF; salpingectomy; tubal occlusion
33.	Details of existing review of same topic by same authors	None
34.	Current review status	<input checked="" type="checkbox"/> Ongoing
		<input type="checkbox"/> Completed but not published
		<input type="checkbox"/> Completed and published
		<input type="checkbox"/> Completed, published and being updated
		<input type="checkbox"/> Discontinued
35..	Additional information	None
36.	Details of final publication	www.nice.org.uk

1 N/A: not applicable

1 Appendix B Literature search strategies

2 Literature search strategies for review question: What is the clinical and cost
3 effectiveness of surgery for hydrosalpinges prior to assisted reproductive
4 technology (ART), relative to standard ART without prior surgical optimisation,
5 for people with tubal disease?

6 See Appendices 1 to 6 of the Cochrane review (Melo 2020):

7 <https://doi.org/10.1002/14651858.CD002125.pub4>

8 Health Economic Literature Search Strategies

9 Database: Ovid MEDLINE(R) ALL <1946 to January 16, 2024>

10 Date of last search: 17/01/2024

#	Searches
1	(in vitro fertilisation or in vitro fertilization).tw.
2	(ivf or icsi).tw.
3	intracytoplasmic sperm injection*.tw.
4	reproductive techniques, assisted/ or exp embryo transfer/ or exp fertilization in vitro/ or zygote intrafallopian transfer/
5	(ART or embryo transfer* or et).tw.
6	assisted reproducti* techn*.tw.
7	Pregnancy/
8	pregnan\$.tw.
9	Fertility/ or Infertility/ or Infertility, Female/
10	(fertil\$ or subfertil\$ or infertil\$).tw.
11	or/1-10
12	exp Fallopian Tube Diseases/
13	(Fallopian\$ adj5 Disease\$).tw.
14	Salpingectomy/ or Salpingostomy/
15	salping\$.tw.
16	Sterilization, Tubal/
17	(fallopian adj5 (ligat\$ or electrocauter\$ or occlusion\$ or occluded or block\$ or clamp\$ or factor\$ or adhesion\$)).tw.
18	(tub\$ adj5 adhesion\$).tw.
19	(tub\$ adj5 occlusion\$).tw.
20	(tub\$ adj3 disease\$).tw.
21	(tub\$ adj3 factor\$).tw.
22	(tub\$ adj5 block\$).tw.
23	(tub\$ adj5 (clamp\$ or clip\$)).tw.
24	(tub\$ adj3 electrocaut\$).tw.
25	(tub\$ adj5 Filshie\$).tw.
26	(tub\$ adj3 ligat\$).tw.
27	hydrosalpin\$.tw.
28	gynecologic surgical procedures/
29	(surg\$ adj5 tub\$).tw.
30	(surg\$ adj5 fallopian\$).tw.
31	(laparoscop\$ adj5 (tube\$ or tubal)).tw.
32	Endoscopic Ultrasound-Guided Fine Needle Aspiration/
33	(ultrasound guided adj5 aspiration\$).tw.
34	(aspirat\$ adj5 (tube\$ or tubal)).tw.
35	((tube\$ or tubal) adj5 interrupt\$).tw.
36	essure\$.tw.
37	(clip\$ adj5 fallopian\$).tw.
38	hysteroscop\$.tw.

#	Searches
39	or/12-38
40	fallopian tubes/
41	microsurgery/ or minimally invasive surgical procedures/ or Electrosurgery/ or Electrocoagulation/ or Laparoscopy/ or Laparoscopes/ or Hand-assisted laparoscopy/ or hysteroscopy/
42	40 and 41
43	fallopian tubes/su or Fallopian Tube Diseases/su or Infertility, Female/su
44	39 or 42 or 43
45	11 and 44
46	letter/
47	editorial/
48	news/
49	exp historical article/
50	Anecdotes as topic/
51	comment/
52	case reports/
53	(letter or comment*).ti.
54	or/46-53
55	randomized controlled trial/ or random*.ti,ab.
56	54 not 55
57	animals/ not humans/
58	exp Animals, Laboratory/
59	exp Animal Experimentation/
60	exp Models, Animal/
61	exp Rodentia/
62	(rat or rats or rodent* or mouse or mice).ti.
63	or/56-62
64	45 not 63
65	limit 64 to english language
66	Economics/
67	Value of life/
68	exp "Costs and Cost Analysis"/
69	exp Economics, Hospital/
70	exp Economics, Medical/
71	exp Resource Allocation/
72	Economics, Nursing/
73	Economics, Pharmaceutical/
74	exp "Fees and Charges"/
75	exp Budgets/
76	budget*.ti,ab.
77	cost*.ti,ab.
78	(economic* or pharmaco?economic*).ti,ab.
79	(price* or pricing*).ti,ab.
80	(financ* or fee or fees or expenditure* or saving*).ti,ab.
81	(value adj2 (money or monetary)).ti,ab.
82	resourc* allocat*.ti,ab.
83	(fund or funds or funding* or funded).ti,ab.
84	(ration or rations or rationing* or rationed).ti,ab.
85	ec.fs.
86	or/66-85
87	quality-adjusted life years/
88	sickness impact profile/
89	(quality adj2 (wellbeing or well being)).ti,ab.
90	sickness impact profile.ti,ab.
91	disability adjusted life.ti,ab.

#	Searches
92	(qal* or qtime* or qwb* or daly*).ti,ab.
93	(euroqol* or eq5d* or eq 5*).ti,ab.
94	(qol* or hql* or hqol* or h qol* or hrqol* or hr qol*).ti,ab.
95	(health utility* or utility score* or disutilit* or utility value*).ti,ab.
96	(hui or hui1 or hui2 or hui3).ti,ab.
97	(health* year* equivalent* or hye or hyes).ti,ab.
98	discrete choice*.ti,ab.
99	rosser.ti,ab.
100	(willingness to pay or time tradeoff or time trade off or tto or standard gamble*).ti,ab.
101	(sf36* or sf 36* or short form 36* or shortform 36* or shortform36*).ti,ab.
102	(sf20 or sf 20 or short form 20 or shortform 20 or shortform20).ti,ab.
103	(sf12* or sf 12* or short form 12* or shortform 12* or shortform12*).ti,ab.
104	(sf8* or sf 8* or short form 8* or shortform 8* or shortform8*).ti,ab.
105	(sf6* or sf 6* or short form 6* or shortform 6* or shortform6*).ti,ab.
106	or/87-105
107	65 and (86 or 106)

1 **Database: Embase <1974 to 2024 January 16>**

2 **Date of last search: 17/01/2024**

#	Searches
1	(in vitro fertilisation or in vitro fertilization).tw.
2	(ivf or icsi).tw.
3	intracytoplasmic sperm injection*.tw.
4	infertility therapy/ or exp embryo transfer/ or exp in vitro fertilization/ or zygote intrafallopian transfer/
5	(ART or embryo transfer* or et).tw.
6	assisted reproducti* techn*.tw.
7	Pregnancy/
8	pregnan\$.tw.
9	fertility/ or female fertility/ or infertility/ or female infertility/
10	(fertil\$ or subfertil\$ or infertil\$).tw.
11	or/1-10
12	exp uterine tube disease/
13	(Fallopian\$ adj5 Disease\$).tw.
14	salpingostomy/
15	salping\$.tw.
16	exp uterine tube sterilization/
17	(fallopian adj5 (ligat\$ or electrocauter\$ or occlusion\$ or occluded or block\$ or clamp\$ or factor\$ or adhesion\$)).tw.
18	(tub\$ adj5 adhesion\$).tw.
19	(tub\$ adj5 occlusion\$).tw.
20	(tub\$ adj3 disease\$).tw.
21	(tub\$ adj3 factor\$).tw.
22	(tub\$ adj5 block\$).tw.
23	(tub\$ adj5 (clamp\$ or clip\$)).tw.
24	(tub\$ adj3 electrocaut\$).tw.
25	(tub\$ adj5 Filshie\$).tw.
26	(tub\$ adj3 ligat\$).tw.
27	hydrosalpin\$.tw.
28	gynecologic surgery/ or uterine tube surgery/ or salpingoplasty/
29	(surg\$ adj5 tub\$).tw.
30	(surg\$ adj5 fallopian\$).tw.
31	(laparoscop\$ adj5 (tube\$ or tubal\$)).tw.
32	ultrasound guided fine needle aspiration/
33	(ultrasound guided adj5 aspiration\$).tw.

#	Searches
34	(aspirat\$ adj5 (tube\$ or tubal)).tw.
35	((tube\$ or tubal) adj5 interrupt\$).tw.
36	essure\$.tw.
37	(clip\$ adj5 fallopian\$).tw.
38	hysteroscop\$.tw.
39	or/12-38
40	fallopian tube/
41	Surgery/ or microsurgery/ or minimally invasive surgery/ or minimally invasive procedure/ or electrosurgery/ or electrocoagulation/ or laparoscopy/ or hand assisted laparoscopy/ or laparoscopic surgery/ or exp laparoscope/ or hysteroscopy/
42	40 and 41
43	uterine tube disease/su or female infertility/su
44	39 or 42 or 43
45	11 and 44
46	letter.pt. or letter/
47	note.pt.
48	editorial.pt.
49	case report/ or case study/
50	(letter or comment*).ti.
51	or/46-50
52	randomized controlled trial/ or random*.ti,ab.
53	51 not 52
54	animal/ not human/
55	nonhuman/
56	exp Animal Experiment/
57	exp Experimental Animal/
58	animal model/
59	exp Rodent/
60	(rat or rats or rodent* or mouse or mice).ti.
61	or/53-60
62	45 not 61
63	limit 62 to english language
64	(conference abstract* or conference review or conference paper or conference proceeding).db,pt,su.
65	63 not 64
66	health economics/
67	exp economic evaluation/
68	exp health care cost/
69	exp fee/
70	budget/
71	funding/
72	resource allocation/
73	budget*.ti,ab.
74	cost*.ti,ab.
75	(economic* or pharmaco?economic*).ti,ab.
76	(price* or pricing*).ti,ab.
77	(financ* or fee or fees or expenditure* or saving*).ti,ab.
78	(value adj2 (money or monetary)).ti,ab.
79	resourc* allocat*.ti,ab.
80	(fund or funds or funding* or funded).ti,ab.
81	(ration or rations or rationing* or rationed).ti,ab.
82	or/66-81
83	quality adjusted life year/
84	"quality of life index"/
85	short form 12/ or short form 20/ or short form 36/ or short form 8/

#	Searches
86	sickness impact profile/
87	(quality adj2 (wellbeing or well being)).ti,ab.
88	sickness impact profile.ti,ab.
89	disability adjusted life.ti,ab.
90	(qal* or qtime* or qwb* or daly*).ti,ab.
91	(euroqol* or eq5d* or eq 5*).ti,ab.
92	(qol* or hql* or hqol* or h qol* or hrqol* or hr qol*).ti,ab.
93	(health utility* or utility score* or disutilit* or utility value*).ti,ab.
94	(hui or hui1 or hui2 or hui3).ti,ab.
95	(health* year* equivalent* or hye or hyes).ti,ab.
96	discrete choice*.ti,ab.
97	rosser.ti,ab.
98	(willingness to pay or time tradeoff or time trade off or tto or standard gamble*).ti,ab.
99	(sf36* or sf 36* or short form 36* or shortform 36* or shortform36*).ti,ab.
100	(sf20 or sf 20 or short form 20 or shortform 20 or shortform20).ti,ab.
101	(sf12* or sf 12* or short form 12* or shortform 12* or shortform12*).ti,ab.
102	(sf8* or sf 8* or short form 8* or shortform 8* or shortform8*).ti,ab.
103	(sf6* or sf 6* or short form 6* or shortform 6* or shortform6*).ti,ab.
104	or/83-103
105	65 and (82 or 104)

1

2 Database: INAHTA

3 Date of last search: 17/01/2024

#	Searches
1	"in vitro fertilisation" or "in vitro fertilization"
2	ivf or icsi
3	"intracytoplasmic sperm injection" or "intracytoplasmic sperm injections"
4	"Reproductive Techniques, Assisted"[mh]
5	"Embryo Transfer"[mhe]
6	"Fertilization in Vitro"[mhe]
7	"Zygote Intrafallopian Transfer"[mh]
8	ART or "embryo transfer" or "embryo transfers" or et
9	assisted reproducti* techn*
10	"Pregnancy"[mh]
11	pregnan*
12	"Fertility"[mh]
13	"Infertility"[mh]
14	"Infertility, Female"[mh]
15	fertil* or subfertil* or infertil*
16	#15 OR #14 OR #13 OR #12 OR #11 OR #10 OR #9 OR #8 OR #7 OR #6 OR #5 OR #4 OR #3 OR #2 OR #1
17	"Fallopian Tube Diseases"[mhe]
18	(Fallopian* and Disease*)
19	"Salpingectomy"[mh]
20	"Salpingostomy"[mh]
21	salping*
22	"Sterilization, Tubal"[mh]
23	(fallopian AND (ligat* or electrocauter* or occlusion* or occluded or block* or clamp* or factor* or adhesion*))
24	(tub* and adhesion*)
25	(tub* and occlusion*)
26	(tub* and disease*)
27	(tub* and factor*)
28	(tub* and block*)

#	Searches
29	(tub* and (clamp* or clip*))
30	(tub* and electrocaut*)
31	(tub* and Filshie*)
32	(tub* and ligat*)
33	hydrosalpin*
34	"Gynecologic Surgical Procedures"[mh]
35	(surg* and tub*)
36	(surg* and fallopian*)
37	(laparoscop* and (tube* or tubal))
38	"Endoscopic Ultrasound-Guided Fine Needle Aspiration"[mh]
39	("ultrasound guided" AND aspiration*)
40	(aspirat* and (tube* or tubal))
41	((tube* or tubal) and interrupt*)
42	essure*
43	(clip* and fallopian*)
44	hysteroscop*
45	#44 OR #43 OR #42 OR #41 OR #40 OR #39 OR #38 OR #37 OR #36 OR #35 OR #34 OR #33 OR #32 OR #31 OR #30 OR #29 OR #28 OR #27 OR #26 OR #25 OR #24 OR #23 OR #22 OR #21 OR #20 OR #19 OR #18 OR #17
46	#45 AND #16
47	"Fallopian Tubes"[mh]
48	"Microsurgery"[mh]
49	"Minimally Invasive Surgical Procedures"[mh]
50	"Electrosurgery"[mh]
51	"Electrocoagulation"[mh]
52	"Laparoscopy"[mh]
53	"Laparoscopes"[mh]
54	"Hand-Assisted Laparoscopy"[mh]
55	"Hysteroscopy"[mh]
56	#55 OR #54 OR #53 OR #52 OR #51 OR #50 OR #49 OR #48
57	#56 AND #47
58	#57 OR #46

1

2 **Database: HTA via CRD**3 **Date of last search: 17/01/2024**

#	Searches
1	("in vitro fertilisation" or "in vitro fertilization")
2	(ivf or icsi)
3	(intracytoplasmic NEXT sperm NEXT injection*)
4	MESH DESCRIPTOR Reproductive Techniques, Assisted
5	MESH DESCRIPTOR Embryo Transfer EXPLODE ALL TREES
6	MESH DESCRIPTOR Fertilization in Vitro EXPLODE ALL TREES
7	MESH DESCRIPTOR Zygote Intrafallopian Transfer
8	(ART or (embryo NEXT transfer*) or et)
9	(assisted NEXT reproducti* NEXT techn*)
10	MESH DESCRIPTOR Pregnancy
11	(pregnan*)
12	MESH DESCRIPTOR Fertility
13	MESH DESCRIPTOR Infertility
14	MESH DESCRIPTOR Infertility, Female
15	(fertil* or subfertil* or infertil*)
16	#1 or #2 or #3 or #4 or #5 or #6 or #7 or #8 or #9 or #10 or #11 or #12 or #13 or #14 or #15

#	Searches
17	MESH DESCRIPTOR Fallopian Tube Diseases EXPLODE ALL TREES
18	(Fallopian* NEAR5 Disease*)
19	MESH DESCRIPTOR Salpingectomy
20	MESH DESCRIPTOR Salpingostomy
21	salping*
22	MESH DESCRIPTOR Sterilization, Tubal
23	(fallopian NEAR5 (ligat* or electrocauter* or occlusion* or occluded or block* or clamp* or factor* or adhesion*))
24	(tub* NEAR5 adhesion*)
25	(tub* NEAR5 occlusion*)
26	(tub* NEAR3 disease*)
27	(tub* NEAR3 factor*)
28	(tub* NEAR5 block*)
29	(tub* NEAR5 (clamp* or clip*))
30	(tub* NEAR3 electrocaut*)
31	(tub* NEAR5 filshie*)
32	(tub* NEAR3 ligat*)
33	hydrosalpin*
34	MESH DESCRIPTOR Gynecologic Surgical Procedures
35	(surg* NEAR5 tub*)
36	(surg* NEAR5 fallopian*)
37	(laparoscop* NEAR5 (tube* or tubal))
38	MESH DESCRIPTOR Endoscopic Ultrasound-Guided Fine Needle Aspiration
39	("ultrasound guided" NEAR5 aspiration*)
40	(aspirat* NEAR5 (tube* or tubal))
41	((tube* or tubal) NEAR5 interrupt*)
42	essure*
43	(clip* NEAR5 fallopian*)
44	hysteroscop*
45	#17 or #18 or #19 or #20 or #21 or #22 or #23 or #24 or #25 or #26 or #27 or #28 or #29 or #30 or #31 or #32 or #33 or #34 or #35 or #36 or #37 or #38 or #39 or #40 or #41 or #42 or #43 or #44
46	MESH DESCRIPTOR Fallopian Tubes
47	MESH DESCRIPTOR Microsurgery
48	MESH DESCRIPTOR Minimally Invasive Surgical Procedures
49	MESH DESCRIPTOR Electrosurgery
50	MESH DESCRIPTOR Electrocoagulation
51	MESH DESCRIPTOR Laparoscopy
52	MESH DESCRIPTOR Laparoscopes
53	MESH DESCRIPTOR Hand-Assisted Laparoscopy
54	MESH DESCRIPTOR Hysteroscopy
55	#47 or #48 or #49 or #50 or #51 or #52 or #53 or #54
56	#46 and #55
57	#45 or #56
58	#16 and #57

1

2

1 **Appendix C Effectiveness evidence study selection**

2 **Study selection for: What is the clinical and cost effectiveness of surgery for**
3 **hydrosalpinges prior to assisted reproductive technology (ART), relative to**
4 **standard ART without prior surgical optimisation, for people with tubal**
5 **disease?**

6 See results of the search – figure 1 from the Cochrane review (Melo 2020):
7 <https://doi.org/10.1002/14651858.CD002125.pub4>.

8

1 **Appendix D Characteristics of studies tables**

2 **Characteristics of studies tables for review question: What is the clinical and cost**
3 **effectiveness of surgery for hydrosalpinges prior to assisted reproductive**
4 **technology (ART), relative to standard ART without prior surgical optimisation, for**
5 **people with tubal disease?**

6 See the characteristics of included studies tables from the Cochrane review (Melo 2020):
7 <https://doi.org/10.1002/14651858.CD002125.pub4>.

1

2 **Appendix E Forest plots**

3 **Forest plots for review question: What is the clinical and cost effectiveness of**
4 **surgery for hydrosalpinges prior to assisted reproductive technology (ART),**
5 **relative to standard ART without prior surgical optimisation, for people with tubal**
6 **disease?**

7 See forest plots from the Cochrane review (Melo 2020):
8 <https://doi.org/10.1002/14651858.CD002125.pub4>.

9

1 **Appendix E Summary of findings tables**

2 **Summary of findings tables for review question: What is the clinical and cost**
3 **effectiveness of surgery for hydrosalpinges prior to assisted reproductive**
4 **technology (ART), relative to standard ART without prior surgical optimisation, for**
5 **people with tubal disease?**

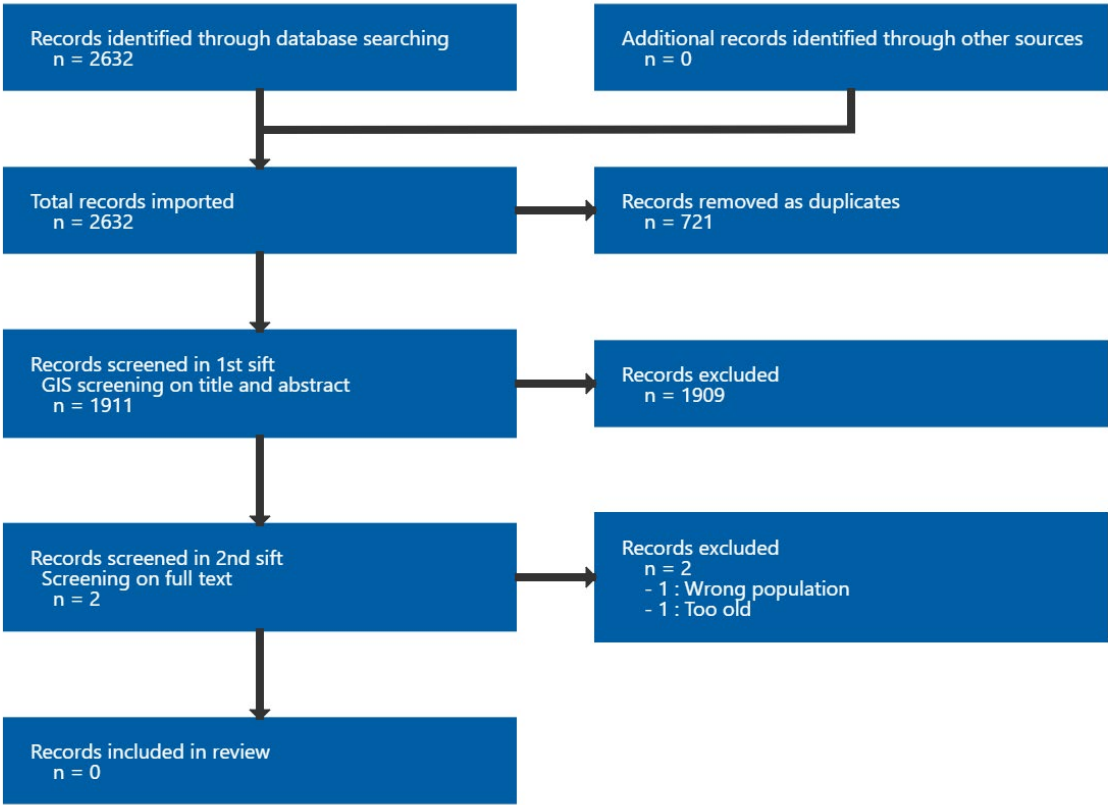
6 See the summary of findings tables from the Cochrane review (Melo 2020):
7 <https://doi.org/10.1002/14651858.CD002125.pub4>.
8

Appendix F Economic evidence study selection

Study selection for review question: What is the clinical and cost effectiveness of surgery for hydrosalpinges prior to assisted reproductive technology (ART), relative to standard ART without prior surgical optimisation, for people with tubal disease?

No economic evidence was identified which was applicable to this review question.

Figure 1: Study selection flow chart



1 **Appendix G Economic evidence tables**

2 **Economic evidence tables for review question: What is the clinical and cost**
3 **effectiveness of surgery for hydrosalpinges prior to assisted reproductive**
4 **technology (ART), relative to standard ART without prior surgical optimisation,**
5 **for people with tubal disease?**

6 No evidence was identified which was applicable to this review question.

7

8

1 **Appendix H Economic model**

2 **Economic model for review question: What is the clinical and cost**
3 **effectiveness of surgery for hydrosalpinges prior to assisted reproductive**
4 **technology (ART), relative to standard ART without prior surgical optimisation,**
5 **for people with tubal disease?**

6 No economic analysis was conducted for this review question.

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Appendix I Excluded studies

Excluded studies for review question: What is the clinical and cost effectiveness of surgery for hydrosalpinges prior to assisted reproductive technology (ART), relative to standard ART without prior surgical optimisation, for people with tubal disease?

Excluded effectiveness studies

See the characteristics of excluded studies table from the Cochrane review (Melo 2020): <https://doi.org/10.1002/14651858.CD002125.pub4>.

Excluded economic studies

Table 4: Excluded studies and reasons for their exclusion

Study	Code
Strandell, Annika; Lindhard, Anette; Eckerlund, Ingemar (2005) Cost--effectiveness analysis of salpingectomy prior to IVF, based on a randomized controlled trial. Human reproduction (Oxford, England) 20(12): 3284-92	- Too old
Verhoeve, H R, Moolenaar, L M, Hompes, P et al. (2013) Cost-effectiveness of tubal patency tests. BJOG: an international journal of obstetrics and gynaecology 120(5): 583-93	- Wrong population

1 **Appendix J Research recommendations – full details**

2 **Research recommendations for review question: What is the clinical and cost**
3 **effectiveness of surgery for hydrosalpinges prior to assisted reproductive**
4 **technology (ART), relative to standard ART without prior surgical optimisation,**
5 **for people with tubal disease?**

6 No research recommendations were made for this review question.
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