



Kunutsor, S. K., Whitehouse, M. R., Lenguerrand, E., Blom, A. W., Beswick, A. D., & INFORM Team (2016). Re-Infection Outcomes Following One- And Two-Stage Surgical Revision of Infected Knee Prosthesis: A Systematic Review and Meta-Analysis. PLoS ONE, 11(3), [e0151537]. 10.1371/journal.pone.0151537

Publisher's PDF, also known as Final Published Version

Link to published version (if available): 10.1371/journal.pone.0151537

Link to publication record in Explore Bristol Research PDF-document

# **University of Bristol - Explore Bristol Research General rights**

This document is made available in accordance with publisher policies. Please cite only the published version using the reference above. Full terms of use are available: http://www.bristol.ac.uk/pure/about/ebr-terms.html

# Take down policy

Explore Bristol Research is a digital archive and the intention is that deposited content should not be removed. However, if you believe that this version of the work breaches copyright law please contact open-access@bristol.ac.uk and include the following information in your message:

- Your contact details
- Bibliographic details for the item, including a URL
- An outline of the nature of the complaint

On receipt of your message the Open Access Team will immediately investigate your claim, make an initial judgement of the validity of the claim and, where appropriate, withdraw the item in question from public view.







Citation: Kunutsor SK, Whitehouse MR, Lenguerrand E, Blom AW, Beswick AD, INFORM Team (2016) Re-Infection Outcomes Following One-And Two-Stage Surgical Revision of Infected Knee Prosthesis: A Systematic Review and Meta-Analysis. PLoS ONE 11(3): e0151537. doi:10.1371/journal. pone.0151537

**Editor:** Heiner Baur, Bern University of Applied Sciences, SWITZERLAND

Received: November 4, 2015

Accepted: February 28, 2016

Published: March 11, 2016

Copyright: © 2016 Kunutsor et al. This is an open access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

**Data Availability Statement:** All relevant data are within the paper and its Supporting Information files.

**Funding:** This article presents independent research funded by the National Institute for Health Research (NIHR) under its Programme Grants for Applied Research program (RP-PG-1210-12005). The views expressed in this article are those of the authors and not necessarily those of the NHS, the NIHR, or the Department of Health. The funder had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

RESEARCH ARTICLE

# Re-Infection Outcomes Following One- And Two-Stage Surgical Revision of Infected Knee Prosthesis: A Systematic Review and Meta-Analysis

Setor K. Kunutsor\*, Michael R. Whitehouse, Erik Lenguerrand, Ashley W. Blom, Andrew D. Beswick, INFORM Team<sup>1</sup>

Musculoskeletal Research Unit, School of Clinical Sciences, University of Bristol, Learning & Research Building (Level 1), Southmead Hospital, Southmead Road, Bristol, BS10 5NB, United Kingdom

¶ Membership of the INFORM Team is provided in the Acknowledgments.

\* skk31@cantab.net

# **Abstract**

# **Background**

Periprosthetic joint infection (PJI) is a serious complication of total knee arthroplasty. Twostage revision is the most widely used technique and considered as the most effective for treating periprosthetic knee infection. The one-stage revision strategy is an emerging alternative option, however, its performance in comparison to the two-stage strategy is unclear. We therefore sought to ask if there was a difference in re-infection rates and other clinical outcomes when comparing the one-stage to the two-stage revision strategy.

### Objective

Our first objective was to compare re-infection (new and recurrent infections) rates for oneand two-stage revision surgery for periprosthetic knee infection. Our second objective was to compare between the two revision strategies, clinical outcomes as measured by postoperative Knee Society Knee score, Knee Society Function score, Hospital for Special Surgery knee score, WOMAC score, and range of motion.

#### Design

Systematic review and meta-analysis.

#### **Data sources**

MEDLINE, EMBASE, Web of Science, Cochrane Library, reference lists of relevant studies to August 2015, and correspondence with investigators.



**Competing Interests:** The authors have declared that no competing interests exist.

# Study selection

Longitudinal (prospective or retrospective cohort) studies conducted in generally unselected patients with periprosthetic knee infection treated exclusively by one- or two-stage revision and with re-infection outcomes reported within two years of revision surgery. No clinical trials comparing both revision strategies were identified.

#### **Review methods**

Two independent investigators extracted data and discrepancies were resolved by consensus with a third investigator. Re-infection rates from 10 one-stage studies (423 participants) and 108 two-stage studies (5,129 participants) were meta-analysed using random-effect models after arcsine transformation.

#### **Results**

The rate (95% confidence intervals) of re-infection was 7.6% (3.4–13.1) in one-stage studies. The corresponding re-infection rate for two-stage revision was 8.8% (7.2–10.6). In subgroup analyses, re-infection rates remained generally similar for several study-level and clinically relevant characteristics. Postoperative clinical outcomes of knee scores and range of motion were similar for both revision strategies.

#### Limitations

Potential bias owing to the limited number of one-stage revision studies and inability to explore heterogeneity in greater detail.

#### **Conclusions**

Available evidence from aggregate published data suggest the one-stage revision strategy may be as effective as the two-stage revision strategy in treating infected knee prostheses in generally unselected patients. Further investigation is warranted.

#### Systematic review registration

PROSPERO 2015: CRD42015017327

#### Introduction

One of the most serious complications of total knee arthroplasty (TKA) is deep periprosthetic joint infection (PJI). Cure of PJI is not possible by antibiotic treatment alone in the majority of cases and surgical treatment is required. Options include: debridement, treatment with antibiotics and retention of the prosthesis; one-stage revision; two-stage revision; resection arthroplasty; or arthrodesis or amputation as a last resort [1, 2]. Most people will receive a one-stage or two-stage revision. The two-stage revision strategy has evolved as the gold standard and preferred procedure for treating PJI of the knee [3, 4]. However, this revision strategy is associated with long periods of hospitalization, lengthy functional impairment, and high health service costs [5]. For these reasons and though still less commonly used compared to the two-stage revision strategy, the one-stage strategy is an attractive alternative option. Besides the reduced



number of surgical procedures for the patient, the one-stage procedure may be associated with a shorter overall time in hospital, reduced duration of postoperative antibiotic use, and quicker mobilization. It might also be a more cost-effective approach [5, 6]. The optimal treatment strategy for the management of deep prosthetic knee infection remains controversial.

To date, no randomized controlled trial has compared the effectiveness of the one- and twostage revision procedures. However, a number of reviews based on observational evidence have assessed re-infection outcomes (recurrent and new infections) comparing both revision strategies and have mostly reported inconclusive results. In 2009, Jamsen and colleagues in their review of a total of 31 studies, reported overall infection eradication rates of between 73 to 100% and 82 to 100% in one- and two-stage revision strategies respectively [7]. In further assessment of the clinical outcomes as measured by knee scores and range of motion, there was no significant difference between the two strategies. In another review, Romano and colleagues reported average infection eradication rates of 89.8% and 81.8% in two-stage (38 studies) and one-stage (6 studies) revisions respectively and concluded that the two-stage was associated with a higher rate of infection eradication (albeit acknowledging the limitations posed by the methodological quality of included studies) [8]. In the most recent relevant review, Masters and colleagues reviewed available evidence (58 two-stage and 5 one-stage studies) and reported re-infection rates of between 0 to 41% for two-stage studies and 0 to 11% for one-stage studies [9]. In their report, the authors concluded that there was a significantly larger evidence base for the two-stage revision strategy and recommended further work to compare the two revision strategies.

The current evidence does not conclusively support a superior revision strategy for periprosthetic knee infection. In addition, several features of these previous reviews existed which limited the generalisability and validity of the findings. First, all of the reviews were characterised by a small number of one-stage revision studies (< 10 studies), which limited reliable comparability of the two revision strategies. Second, the heterogeneous periods of follow-up for reinfection outcomes in the individual studies were considered, which did not enhance comparability and interpretation of the findings. Third, none of the reviews explored for potential sources of heterogeneity among the contributing studies. Fourth, assessment of publication bias or small study bias was not conducted. Fifth, none of the reviews conducted any subgroup analysis across important study-level and clinically relevant characteristics (e.g. geographical location, age at baseline, and quality assessment). Sixth, apart from the study by Jamsen and colleagues [7] which was limited by its narrative approach, none of the studies compared the two revision strategies using other clinical outcomes. Finally, several reports have been published on the topic since the last previous relevant review [9].

In line with the uncertain evidence and recommendations of previous reviews [7, 9], there is a need for further work to compare the effectiveness of the two revision strategies. We therefore sought to ask if there was a difference in re-infection rates and other clinical outcomes when comparing the one-stage to the two-stage revision strategy. Ideally, to compare the effectiveness of these two revision strategies will require evidence from a carefully designed randomized clinical trial; however, this is may be unlikely in the short term given the low PJI event rates recorded after TKA [10, 11]. In the absence of robust evidence from a rigorous randomized clinical trial; using a systematic meta-analytic approach, our aim was to (1) to evaluate the effectiveness of one- and two-stages revision strategies using re-infection rate (primary outcome) and other clinical outcomes as measured by postoperative Knee Society Knee score, Knee Society Function score, Hospital for Special Surgery knee score, WOMAC score, and range of motion following revision arthroplasty and (2) to compare and describe the outcomes between the two revision strategies.



#### **Methods**

# Data sources and search strategy

Our review has been registered in the PROSPERO prospective register of systematic reviews (CRD42015017327) and was conducted using a predefined protocol and in line with PRISMA (Appendix A in S1 File) and MOOSE guidelines [12, 13] (Appendix B in S1 File). We searched for longitudinal studies (retrospective, prospective, or randomized controlled trials) reporting re-infection outcomes following one- or two-stage surgical revision of infected knee prosthesis in MEDLINE, EMBASE, Web of Science, and Cochrane databases from inception up to August 2015. The search strategy used a combination of key words related to knee replacement, infection, and revision with focus on one- and two-stage surgeries. No language restrictions were employed. We complemented the search by manually scanning reference lists of identified articles and review articles for relevant publications missed by the original search. Details on our search strategy are presented in Appendix C in S1 File.

# Eligibility criteria

We included studies comprising of consecutive generally unselected patients (i.e., patients representative of the general patient population) who were treated exclusively by one-stage or two-stage revision with a follow-up duration of at least two years for re-infection outcomes (recurrent and new infections). We excluded: (i) studies that reported case series of methods in selected group of patients (such as patients with a specific infection); (ii) studies that did not include patients with less than two years of follow-up; and (iii) studies with less than 10 participants as these were more likely to be case series which did not include consecutive patients.

# Data extraction and quality assessment

After an initial screen of abstracts by one reviewer (S.K.K.), potentially relevant articles were acquired. Each article was assessed by two independent reviewers (S.K.K., A.D.B.) using the inclusion criteria and any discrepancies regarding eligibility of an article was discussed, and consensus reached with a third author (M.R.W). One author (S.K.K.) independently extracted data and performed quality assessments using a standardized data collection form. A second reviewer checked data with that in the original articles. Data were extracted on study design, year of publication, geographical location, mean baseline age, proportion of males, period of follow-up following revision surgery, type of fixation used for re-implantation used, use of and type of spacer, number of re-infection outcomes and participants, and clinical characteristics such as preoperative and postoperative Knee Society Knee Society Function score, Hospital for Special Surgery knee score, WOMAC score, and range of motion. For multiple publications involving the same study, the most comprehensive study was used. We corresponded with study investigators to provide missing information. Information on the methodological quality of included studies was assessed based on the Methodological Index for Non-Randomised Studies (MINORS), a validated instrument which is designed for assessment of methodological quality of non-randomized studies in surgery [14] and has been described previously.[15] Briefly, it uses eight pre-defined domains namely: a clearly stated aim, inclusion of consecutive patients, prospective collection of data, endpoints appropriate to the aim of the study, unbiased assessment of the study endpoint, follow-up period appropriate to the aim of the study, loss to follow-up less than 5%, and prospective calculation of the study size. For noncomparative studies, the score varies from 0 to 16, which is the global ideal score.



#### Statistical analysis

The rate of re-infection (estimated from the number of re-infections within two years of knee revision surgery/total number of participants or arthroplasties as reported) with 95% confidence intervals (CIs) was used as the summary measure and primary outcome across studies. The Freeman-Tukey variance stabilising double arcsine transformation [16] was used in estimating the rates, because the alternative use of inverse variance weight in fixed-effects metaanalysis is suboptimal when dealing with binary data with low rates. Details of the method have been reported previously [15]. To account for the effect of between-study heterogeneity anticipated, summary rates of re-infection were pooled using random effects models [17]. We also estimated 95% prediction intervals which are used to determine the degree of heterogeneity, as they provide a region in which about 95% of the true study effects are expected to be found [18, 19]. Standard chi-square tests and the I<sup>2</sup> statistic [20] were used to quantify the extent of statistical heterogeneity across studies. We explored for potential sources of heterogeneity, including study-level and clinically relevant characteristics such as geographical location, baseline mean age, use of a spacer and spacer type, type of fixation used in reimplantation (cemented or cementless), size of study, and study quality, using stratified analysis and random effects meta-regression [21, 22]. We assessed for publication bias using Egger's regression symmetry test [23]. In the presence of publication bias, we adjusted for this using the Duval and Tweedie's nonparametric trim-and-fill method [24]. Though the trim-and-fill method provides fairly conservative estimates compared to the less commonly used Copas selection model [25], we employed the trim-and-fill method as it is commonly used in meta-analysis of this nature and the Copas selection model is also cumbersome to employ and frequently encounters statistical estimation. Nonetheless, the two methods give similar point estimates [25]. Given the suggested inefficiency of Begg's funnel plots as a method for assessing publication bias in metaanalysis of proportion studies with low proportion outcomes [26], this was not employed. Due to the unsuitability of pooling data for the clinical outcomes of postoperative Knee Society Knee score, Knee Society Function score, Hospital for Special Surgery knee score, WOMAC score, and range of motion, these outcomes were compared between the two revision strategies using descriptive statistics. We employed Stata version 14 (Stata Corp, College Station, Texas, USA) for all statistical analyses. The complete dataset used for our analyses can be found in \$2 File.

#### Results

#### Study identification and selection

Our computer search and manual screening of reference lists of relevant studies identified 12,392 potentially relevant citations. After initial screening based of titles and abstracts, 244 articles remained for further evaluation. We excluded 135 articles following detailed assessments (Appendix D in <u>S1 File</u>). The remaining 109 articles were relevant to the review question and were included in the meta-analysis (<u>Fig 1</u>; <u>Table 1</u>; Table A and Appendix E in <u>S1 File</u>). Overall, there were 118 independent studies (comprising of 5,552 participants and 667 reinfection outcomes) included in the review.

#### Study characteristics and study quality

Table 1 provides summary characteristics of one- and two-stage studies included in the review. Table A in S1 File provides baseline characteristics and quality assessment scores of the individual studies. All studies were longitudinal (prospective or retrospective cohort) studies carried out in Europe (United Kingdom, Italy, Germany, France, Switzerland, Austria, Sweden, Spain,



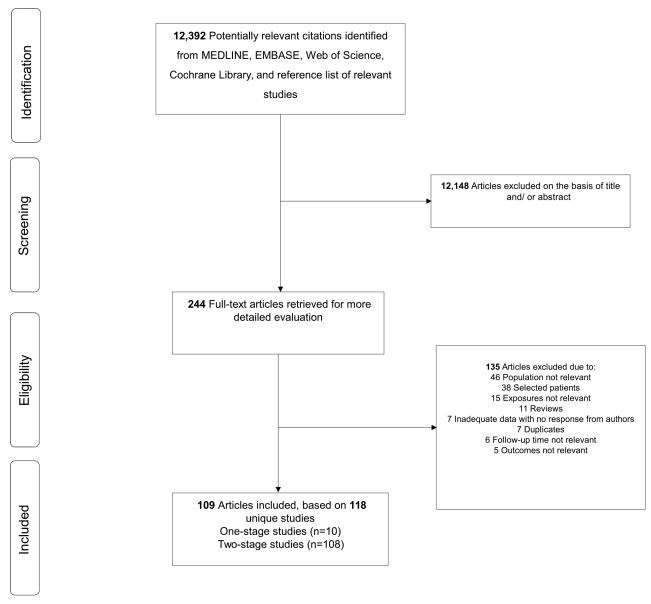


Fig 1. PRISMA Flow Diagram.

doi:10.1371/journal.pone.0151537.g001

Italy, Finland, Greece, Poland, Netherlands, Denmark, Czech Republic, Turkey, and Belgium), North America (United States of America and Canada), Asia (South Korea, India, China, and Taiwan), and the Pacific (Australia and New Zealand). No clinical trials comparing both revision strategies were identified. The methodological quality of included studies ranged from 9–15.

# One-stage revision

Ten studies comprising of 423 participants (42 re-infections) reported on re-infection outcomes in patients using the one-stage surgical revision strategy (Table 1; Table A in S1 File). The pooled random effects re-infection rate (95% CI) was 7.6% (3.4–13.1; P < 0.001) (Fig 2). The 95% prediction interval for the summary re-infection rate was 1.9 to 13.3%, suggesting



Table 1. Summary Characteristics of Included Studies.

	One-stage revision	Two-stage revision
Eligible studies		
Total number of studies included	10	108
Participants		
Total number of re-infections	42	625
Total number of participants	423	5,129
Median (IQR) age (years)	71.1 (66.8–71.8)	67.0 (65.2–70.0)
Location		
Europe	9 (401)	36 (1,181)
North America	1 (22)	52 (3,468)
Asia	-	18 (448)
Pacific	-	2 (32)
Study characteristics		
Median (IQR) follow-up (months)	52.0 (24.0-122.4)	47.0 (32.2-62.2)
Cemented reimplants	3 (97)	38 (1,349)
Use of spacers	-	88 (4,082)
Interval between stages (months)	-	3.2 (2.0-4.7)
Baseline clinical characteristics		
Median (IQR) Knee Society knee score	48.0 (32.0-55.0)	45.8 (35.1–64.0)
Median (IQR) Knee Society function score	33.0 (24.0-42.0)	28.3 (16.6–37.4)
Median (IQR) Hospital for Special Surgery knee score*	-	48.2 (42.4–57.6)
Median (IQR) WOMAC score*	-	49.5 (45.7–60.7)
Median (IQR) range of motion (degrees)	84.5 (79.0–90.0)	74.1 (64.0–83.0)

IQR = interquartile range; values are number of studies (number of participants) unless stated otherwise \*, none of the one-stage revision strategy studies reported baseline scores for these clinical characteristics

doi:10.1371/journal.pone.0151537.t001

that the true re-infection rate for any single study will usually fall within this range. The moderate heterogeneity between contributing studies ( $I^2 = 62\%$ , 95% CI: 24–81%; P = 0.005) was not explained by any study-level characteristics explored (P for meta-regression > 0.10 for each; Figure A in S1 File). The rates of re-infection did not vary significantly by levels or categories of several study-level characteristics. Heterogeneity was substantially reduced when we restricted analyses to studies of the highest quality (quality score:  $\geq 12$ ) ( $I^2 = 42\%$ , 95% CI: 0–79%; P = 0.144). Among the higher quality studies, the pooled re-infection rate (95% CI) was 6.4% (2.1–12.4; P < 0.001), which was similar to the main finding. In further exploration of heterogeneity, on exclusion of the single largest study comprising of 118 participants (with 20 reinfections) [27], heterogeneity was substantially reduced ( $I^2 = 48\%$ , 95% CI: 0–76%; P = 0.053) and the pooled re-infection rate (95% CI) was 6.2% (2.4–11.2; P < 0.001). There was no statistically significant evidence of publication bias using the Egger test (P = 0.516), consistent with the absence of selective reporting when studies were grouped by size (Figure A in S1 File).

#### Two-stage revision

Assessment of re-infection outcomes using the two-stage surgical revision strategy was reported in 108 studies involving 5,129 participants and 625 re-infection outcomes (Table 1; Table A in S1 File). The pooled re-infection rate (95% CI) was 8.8% (7.2–10.6; P < 0.001) (Fig 3). The corresponding 95% prediction interval was 7.1 to 10.6%, similar to the 95% confidence



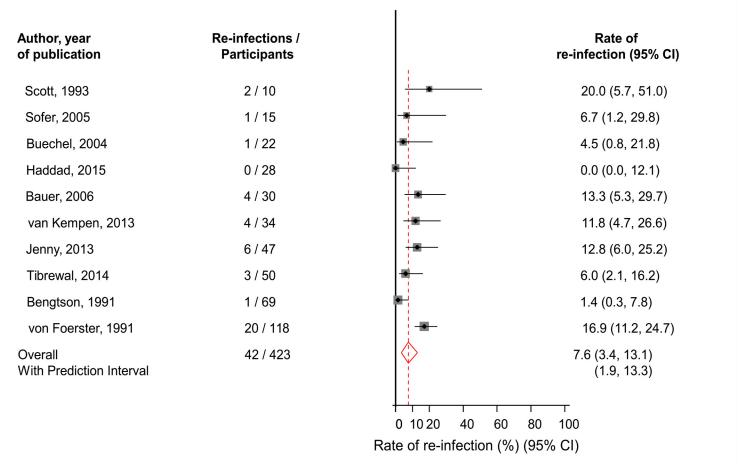


Fig 2. Rates of Re-Infection in Patients Treated by One-Stage Revision. The summary estimates presented were calculated using random effects models; CI, confidence interval (bars)

doi:10.1371/journal.pone.0151537.g002

interval. There was evidence of substantial heterogeneity between contributing studies ( $I^2 = 71\%$ , 64–76%; P < 0.001), which was unexplained by any of the study-level characteristics assessed (Figure B in S1 File). On simultaneous exclusion of the two largest studies [28, 29], the pooled re-infection rate (95% CI) was 8.6% (7.1–10.2; P < 0.001), which was similar to the main finding. The rates of re-infection were generally similar across study relevant characteristics. There was no evidence of heterogeneity ( $I^2 = 0\%$ , 0–68%; P = 0.602), when we restricted the analysis to studies of the highest quality. Among the higher quality studies, the pooled reinfection rate (95% CI) was 6.1 (3.3–9.2; P < 0.001). There was evidence of publication bias (Egger's P = 0.008) and Duval and Tweedie's trim-and-fill method resulted in no additional imputed studies (Figure C in S1 File).

### Other clinical outcomes

Clinical outcomes as measured by postoperative Knee Society Knee Society Function score, and range of motion following revision arthroplasty, were similar between both revision strategies (<u>Table 2</u>); however, only a limited number of one-stage studies reported scores for these outcomes.



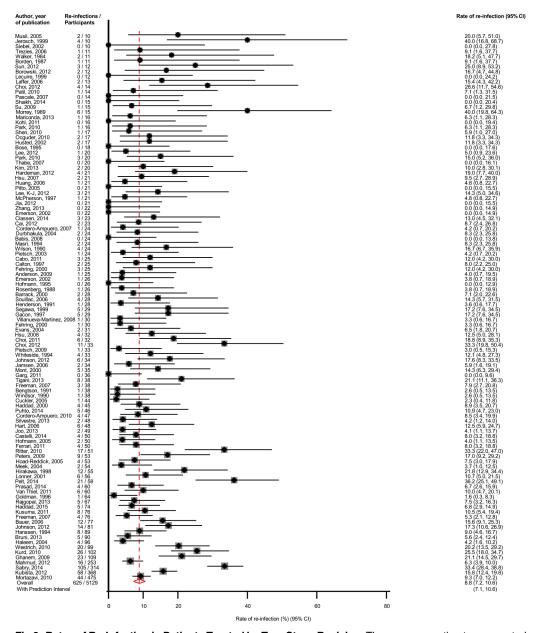


Fig 3. Rates of Re-Infection in Patients Treated by Two-Stage Revision. The summary estimates presented were calculated using random effects models; CI, confidence interval (bars)

doi:10.1371/journal.pone.0151537.g003

# **Discussion**

# Key findings

Given the uncertainty regarding the performance of the one-stage revision strategy in comparison to the two-stage strategy when treating periprosthetic knee infection, we sought to ask if there was a difference in re-infection rates and other clinical outcomes when comparing the one-stage to the two-stage revision strategy. Using a systematic and meta-analytical approach, we aimed to compare the effectiveness of the one- and two-stages revision strategies, using re-infection rate as a primary outcome and other clinical outcomes as measured by postoperative Knee Society Knee score, Knee Society Function score, Hospital for Special Surgery knee score, WOMAC score, and



Table 2. Post-Operative Cl	nical Outcomes Following O	ne- and Two-Stage Revis	ion Strategies.

	One-stage revision	Two-stage revision
Median (IQR) Knee Society knee score	80.3 (74.8–86.5)	82.1 (76.0–86.0)
Median (IQR) Knee Society function score	62.5 (61.0-78.0)	68.0 (59.4–77.9)
Median (IQR) Hospital for Special Surgery knee score*	=	82.5 (79.2-87.0)
Median (IQR) WOMAC score*	-	45.5 (30.3–66.4)
Median (IQR) range of motion (degrees)	97.5 (93.8–100.5)	97.8 (93.7–104.0)

IQR = interquartile range

doi:10.1371/journal.pone.0151537.t002

range of motion following revision arthroplasty. We found similar re-infection rates following one- or two-stage surgical revision for infected knee prostheses. For both revision strategies, our findings also showed that re-infection rates were generally similar (albeit with overlapping confidence intervals) across several study relevant characteristics. The findings for other clinical outcomes were similar for both revision strategies; however, in the context of the limited data for one-stage revision studies, these findings should be interpreted with caution

# Comparison with previous work

Some of our findings are consistent with that of previous reviews on the topic. We also provide several findings that have not been previously reported. Our findings suggest the one-stage as an equivalent revision strategy to the two-stage in terms of effectiveness for treating periprosthetic knee infection, consistent with the results of some previous reviews [7, 9]. However, in contrast to the narrative approach used by Masters et al.[9] and Jamsen et al.[7], we employed a meta-analytic approach, thereby providing more specific and reliable estimates of re-infection rates after one- and two-stage revision of infected knee prosthesis. Our results were also based on a larger number of studies (108 two-stage and 10 one-stage studies), therefore we had enhanced power to robustly demonstrate these results. For the first time, we have also shown that re-infection rates were generally similar across several study-level characteristics such as geographical location, mean baseline age, size of study, and study quality for both revision strategies. Finally, pooled estimates reported from available studies contributing data, showed that the effectiveness of the two revision strategies was similar as measured by other clinical outcomes such as knee scores and range of motion.

### Implications of our findings

Given that the control of infection and maintenance of joint function are considered as main criteria for a successful outcome following one- or two stage revision [6, 30], our findings suggest that the one-stage revision strategy may be as effective as the two-stage strategy in treating many patients with periprosthetic knee infection. The two-stage revision strategy which was pioneered and described by Insall and colleagues in 1983 [31], has for several decades been the established gold standard for treating infected knee replacements [4, 32]. Despite the high success rate in eradication of infection, the two-stage strategy frequently results in significant functional impairment and occasionally death [33]. It is also a costly process in terms of both time and money. Surgical revision of a total knee replacement has been estimated to consume considerably more resources than a primary total knee replacement [34, 35], with even more resources required for surgical revision of an infected arthroplasty [36]. The average cost to the

<sup>\*,</sup> none of the one-stage revision strategy studies reported post-operative scores for these clinical outcomes



NHS of surgical revision of an infected knee replacement is estimated to be about £30 011, which is three times more than that of an aseptic revision [37]. With increasing life expectancy, a growing healthcare burden due to osteoarthritis [38], and a predicted large rise in the numbers of primary TKA being performed, there will be a proportionate rise in the number of patients requiring revision surgery for infected knee prostheses in the coming decades [39, 40]. Within the NHS, total knee arthroplasty is one of the most common orthopaedic procedures [41]. Given the current economic climate and the financial burden associated with the two-stage revision strategy, optimisation of limited resources is of crucial importance.

With the establishment in the treatment of hip periprosthetic joint infection of the onestage revision strategy by Buchholz and colleagues over four decades ago [42], its use has become increasingly popular in recent years. The one-stage revision is widely used in infected hip arthroplasties, but has not gained the same level of momentum in the knee. However, there is an increasing enthusiasm for this procedure in some centres [6] and has been adopted in a number of small case series [43–45]. Apart from the economic benefits associated with this procedure, it has been reported that the one-stage strategy may be associated with better maintenance of joint function (allowing earlier mobility) in selected cases [5]. However, comparing the one-stage procedure in knee and hips, less favourable results have been reported in the knee [1, 46]. One-stage surgical revision has traditionally been thought to expose the patient to a higher risk of re-infection by any residual bacteria [47], given the limited opportunities for additional antimicrobial strategies associated with it. Therefore, suggestions have been made that this strategy should be used only in selected cases, such as patients with minimal bone loss, known organisms with known sensitivities, absence of a sinus tract, and non-immunocompromised patients [48, 49]. Our findings thus yield supportive evidence for the one-stage strategy as a potential alternative option among unselected patients in general.

# Strengths and limitations

The strengths of the current review deserve mention. First, given the large number of studies included (twice as many one- and two-stage studies than the most recent previous review [9]), we had enhanced power to reliably compare the effectiveness of one- and two-stage revision strategies in more detail. Second, we conducted a detailed assessment of the methodological quality of the included studies using a validated instrument for non-randomised surgical studies. Third, compared to previous reviews [7, 9], we employed a meta-analytic approach, thereby the ability to compare the effectiveness of the two revision strategies using quantitative estimates. Fourth, we compared re-infection rates among a broad range of study-level and clinically relevant characteristics. Fifth, we also compared the effectiveness of the two revision strategies using post-operative clinical outcomes such as knee scores and range of motion. Sixth, our results remained robust in several sensitivity analyses. Indeed, our confidence and estimated prediction intervals were similar. Finally, we quantified heterogeneity and explored for potential sources of bias, which included heterogeneity and small study effects.

The limitations also deserve consideration. We acknowledge the limited number of one-stage revision studies, which precluded the ability to robustly compare re-infection rates and other clinical outcomes between the two interventions. We could not conduct detailed subgroup analyses by relevant subgroups such as co-morbidities (e.g., history of diabetes) and duration of antibiotic therapy, given the limited data. We aimed to include only studies that that did not involve selected patients, however, this may not have been possible in some instances given the earlier phase of selection related to management without further replacement in some studies. Though we aimed to include studies that reported re-infection outcomes within two years of revision surgery, there is a possibility some re-infection outcomes may have occurred outside this period as some authors



did not respond to requests for further information. Our findings are comprehensive, important and timely, but should be interpreted with caution in the context of the level of evidence. Indeed, to robustly compare the effectiveness of one-stage and two-stage revision strategies, will require evidence from a carefully designed randomized clinical trial.

#### **Conclusions**

Available evidence from aggregate published data suggest the one-stage revision strategy may be as effective as the two-stage revision strategy in treating infected knee prostheses in generally unselected patients. Further investigation is warranted.

# **Supporting Information**

S1 File.

(DOC)

S2 File.

(CSV)

# **Acknowledgments**

This publication is part of the INFORM (Infection Orthopaedic Management) Programme. As such it benefits from involvement of the whole INFORM team. The INFORM team includes:

Simon Strange, Setor Kunutsor, Kirsty Garfield, Erik Lenguerrand, Rachael Gooberman-Hill, Drew Moore, Amanda Burston, Joanne Simon, Garry King, Michael Whitehouse, Vikki Wylde, Andrew Beswick, Ashley Blom, Sian Noble, Athene Lane, Fran Carroll (Musculoskeletal Research Unit, School of Clinical Sciences, University of Bristol, Southmead Hospital, Southmead Road, Bristol, BS10 5NB, UK); Jason Webb, Alasdair MacGowan (North Bristol NHS Trust, Southmead Hospital, Bristol, BS10 5NB, UK); Stephen Jones (Cardiff and Vale University Health Board, Longcross Street, Cardiff, CF24 0SZ, UK); Adrian Taylor (Oxford University Hospitals NHS Trust, John Radcliffe Hospital, Oxford OX3 9DU, UK); Paul Dieppe (University of Exeter, Medical School, Exeter, EX1 2LU, UK); Andrew Toms, Matthew Wilson (Royal Devon and Exeter NHS Foundation Trust, Newcourt House, Exeter, EX2 7JU, UK); Ian Stockley (Sheffield Teaching Hospitals NHS Trust, Northern General Hospital, Sheffield, S5 7AU, UK); Ben Burston, John-Paul Whittaker (The Robert Jones and Agnes Hunt Orthopaedic Hospital NHS Foundation Trust, Oswestry, Shropshire, SY10 7AG, UK); Tim Board (Wrightington, Wigan and Leigh NHS Foundation Trust, Apple Bridge, Wigan, Lancashire, WN6 9EP, UK); and all the surgeons and nurses from the collaborating centres.

We thank the following for readily providing additional data on request: Gorane Euba, Department of Infectious Diseases, IDIBELL, Hospital Universitari de Bellvitge, Feixa Llarga s/n. 08907, L'Hospitalet de Llobregat, Barcelona, Spain; Francois Hardeman, Orthopaedics and Traumatology, University Hospital Leuven, Leuven, Belgium; S.M.J. Mortazavi, Department of Orthopaedic Surgery, Imam University Hospital, Tehran University of Medical Sciences, End of Keshavarz Blvd, Tehran 1419733141, Iran

#### **Author Contributions**

Conceived and designed the experiments: SKK MRW EL AWB ADB INFORM Team. Performed the experiments: SKK ADB. Analyzed the data: SKK. Contributed reagents/materials/



analysis tools: SKK MRW EL AWB ADB INFORM Team. Wrote the paper: SKK MRW EL AWB ADB INFORM Team.

#### References

- Hanssen AD, Rand JA. Evaluation and treatment of infection at the site of a total hip of knee arthroplasty. J Bone Joint Surg Am. 1998; 80A(6):910–22. PMID: ISI:000074445800016.
- Matthews PC, Berendt AR, McNally MA, Byren I. Diagnosis and management of prosthetic joint infection. BMJ. 2009; 338:1378–83. doi: b1773 doi: 10.1136/bmj.b1773 PMID: ISI:000266502500001.
- 3. Antti-Poika I, Josefsson G, Konttinen Y, Lidgren L, Santavirta S, Sanzen L. Hip arthroplasty infection. Current concepts. Acta orthopaedica Scandinavica. 1990; 61(2):163–9. PMID: 2193478.
- Leone S, Borre S, Monforte A, Mordente G, Petrosillo N, Signore A, et al. Consensus document on controversial issues in the diagnosis and treatment of prosthetic joint infections. Int J Infect Dis. 2010; 14 (4):S67–77.
- Parkinson R, Kay P, Rawal A. A case for one-stage revision in infected total knee arthroplasty? Knee. 2011; 18(1):1–4. doi: 10.1016/j.knee.2010.04.008 PMID: 20719522
- Kendoff D, Gehrke T. Surgical management of periprosthetic joint infection: one-stage exchange. J Knee Surg. 2014; 27(4):273–8. doi: 10.1055/s-0034-1376882 PMID: 24854292.
- Jämsen E, Stogiannidis I, Malmivaara A, Pajamaki J, Puolakka T, Konttinen YT. Outcome of prosthesis exchange for infected knee arthroplasty: the effect of treatment approach. Acta Orthop. 2009; 80 (1):67–77. PMID: 19234888. doi: 10.1080/17453670902805064
- 8. Romano CL, Gala L, Logoluso N, Romano D, Drago L. Two-stage revision of septic knee prosthesis with articulating knee spacers yields better infection eradication rate than one-stage or two-stage revision with static spacers. Knee Surg Sports Traumatol Arthrosc. 2012; 20(12):2445–53. doi: <a href="http://dx.doi.org/10.1007/s00167-012-1885-x">http://dx.doi.org/10.1007/s00167-012-1885-x</a> PMID: 22270671.
- Masters JP, Smith NA, Foguet P, Reed M, Parsons H, Sprowson AP. A systematic review of the evidence for single stage and two stage revision of infected knee replacement. BMC musculoskeletal disorders. 2013; 14:222. doi: <a href="http://dx.doi.org/10.1186/1471-2474-14-222">http://dx.doi.org/10.1186/1471-2474-14-222</a> PMID: <a href="https://dx.doi.org/10.1186/1471-2474-14-222">23895421</a>; PubMed Central PMCID: PMCPMC3734185.
- Kurtz SM, Lau E, Schmier J, Ong KL, Zhao K, Parvizi J. Infection burden for hip and knee arthroplasty in the United States. J Arthroplasty. 2008; 23(7):984–91. doi: 10.1016/j.arth.2007.10.017 PMID: 18534466
- Tanner J, Padley W, Kiernan M, Leaper D, Norrie P, Baggott R. A benchmark too far: findings from a national survey of surgical site infection surveillance. J Hosp Infect. 2013; 83(2):87–91. doi: <a href="http://dx.doi.org/10.1016/j.jhin.2012.11.010">http://dx.doi.org/10.1016/j.jhin.2012.11.010</a> PMID: 23332352.
- Stroup DF, Berlin JA, Morton SC, Olkin I, Williamson GD, Rennie D, et al. Meta-analysis of Observational Studies in Epidemiology. JAMA: The Journal of the American Medical Association. 2000; 283 (15):2008–12. doi: <a href="https://doi.org/10.1001/jama.283.15.2008">10.1001/jama.283.15.2008</a> PubMed Central PMCID: PMC10789670. PMID: 10789670
- Moher D, Liberati A, Tetzlaff J, Altman DG. Preferred reporting items for systematic reviews and metaanalyses: the PRISMA statement. PLoS Med. 2009; 6(7):e1000097. Epub 2009/07/22. doi: 10.1371/ journal.pmed.1000097 PMID: 19621072; PubMed Central PMCID: PMC2707599.
- Slim K, Nini E, Forestier D, Kwiatkowski F, Panis Y, Chipponi J. Methodological index for non-randomized studies (MINORS): development and validation of a new instrument. ANZ J Surg. 2003; 73 (9):712–6. PMID: 12956787.
- 15. Kunutsor SK, Whitehouse MR, Blom AW, Beswick AD, Team I. Re-Infection Outcomes following Oneand Two-Stage Surgical Revision of Infected Hip Prosthesis: A Systematic Review and Meta-Analysis. PloS one. 2015; 10(9):e0139166. doi: 10.1371/journal.pone.0139166 PMID: 26407003.
- Freeman MF, Tukey JW. Transformations Related to the Angular and the Square Root. Ann Math Statist. 1950:607–11. doi: 10.1214/aoms/1177729756
- DerSimonian R, Laird N. Meta-analysis in clinical trials. Control Clin Trials. 1986; 7(3):177–88. Epub 1986/09/01. doi: 0197-2456(86)90046-2 [pii]. PMID: 3802833; PubMed Central PMCID: PMC3802833
- Riley RD, Higgins JP, Deeks JJ. Interpretation of random effects meta-analyses. BMJ. 2011; 342:d549. doi: 10.1136/bmi.d549 PMID: 21310794.
- Higgins JP, Thompson SG, Spiegelhalter DJ. A re-evaluation of random-effects meta-analysis. J R Stat Soc Ser A Stat Soc. 2009; 172(1):137–59. doi: 10.1111/j.1467-985X.2008.00552.x PMID: 19381330; PubMed Central PMCID: PMCPMC2667312.



- Higgins JP, Thompson SG, Deeks JJ, Altman DG. Measuring inconsistency in meta-analyses. BMJ. 2003; 327(7414):557–60. Epub 2003/09/06. doi: 10.1136/bmj.327.7414.557 PMID: 12958120; PubMed Central PMCID: PMC192859.
- Thompson SG, Sharp SJ. Explaining heterogeneity in meta-analysis: a comparison of methods. Stat Med. 1999; 18(20):2693–708. Epub 1999/10/16. doi: 10.1002/(SICI)1097-0258(19991030) 18:20<2693::AID-SIM235>3.0.CO;2-V [pii]. PMID: 10521860.
- 22. Harbord RM, Higgins JPT. Meta-regression in Stata. Stata Journal. 2008; 8:493–519.
- Egger M, Davey Smith G, Schneider M, Minder C. Bias in meta-analysis detected by a simple, graphical test. BMJ. 1997; 315(7109):629–34. Epub 1997/10/06. PMID: <u>9310563</u>; PubMed Central PMCID: PMC2127453.
- Duval S, Tweedie R. Trim and fill: A simple funnel-plot-based method of testing and adjusting for publication bias in meta-analysis. Biometrics. 2000; 56(2):455–63. Epub 2000/07/06. PMID: 10877304.
- Schwarzer G, Carpenter J, Rucker G. Empirical evaluation suggests Copas selection model preferable to trim-and-fill method for selection bias in meta-analysis. J Clin Epidemiol. 2010; 63(3):282–8. doi: 10.16/j.jclinepi.2009.05.008 PMID: 19836925.
- Hunter JP, Saratzis A, Sutton AJ, Boucher RH, Sayers RD, Bown MJ. In meta-analyses of proportion studies, funnel plots were found to be an inaccurate method of assessing publication bias. J Clin Epidemiol. 2014; 67(8):897–903. doi: 10.1016/j.jclinepi.2014.03.003 PMID: 24794697.
- von Foerster G, Kluber D, Kabler U. Mid- to long-term results after treatment of 118 cases of periprosthetic infections after knee joint replacement using one-stage exchange surgery. Orthopade. 1991; 20 (3):244–52. PMID: 1876406.
- Sabry FY, Buller L, Ahmed S, Klika AK, Barsoum WK. Preoperative prediction of failure following twostage revision for knee prosthetic joint infections. Journal of Arthroplasty. 2014; 29(1):115–21. doi: http://dx.doi.org/10.1016/j.arth.2013.04.016 PMID: 23707344.
- Mortazavi SM, Schwartzenberger J, Austin MS, Purtill JJ, Parvizi J. Revision total knee arthroplasty infection: incidence and predictors. Clinical Orthopaedics & Related Research. 2010; 468(8):2052–9.
   PMID: 20309657; PubMed Central PMCID: PMCPMC2895829.
- Rasul AT Jr., Tsukayama D, Gustilo RB. Effect of time of onset and depth of infection on the outcome of total knee arthroplasty infections. Clin Orthop. 1991;(273: ):98–104. PMID: 1959294.
- Insall JN, Thompson FM, Brause BD. Two-stage reimplantation for the salvage of infected total knee arthroplasty. Journal of Bone & Joint Surgery—American Volume. 1983; 65(8):1087–98. PMID: 6630253
- Lentino JR. Prosthetic joint infections: bane of orthopedists, challenge for infectious disease specialists. Clinical infectious diseases: an official publication of the Infectious Diseases Society of America. 2003; 36(9):1157–61. PMID: 12715311.
- Bernard L, Hoffmeyer P, Assal M, Vaudaux P, Schrenzel J, Lew D. Trends in the treatment of orthopaedic prosthetic infections. J Antimicrob Chemother. 2004; 53 (2):127–9. PMID: 2004089143.
- Iorio R, Healy WL, Richards JA. Comparison of the hospital cost of primary and revision total knee arthroplasty after cost containment. Orthopedics. 1999; 22(2):195–9. PMID: 10037333.
- Kurtz S, Mowat F, Ong K, Chan N, Lau E, Halpern M. Prevalence of primary and revision total hip and knee arthroplasty in the United States from 1990 through 2002. J Bone Joint Surg Am. 2005; 87 (7):1487–97. doi: 10.2106/JBJS.D.02441 PMID: 15995115.
- Oduwole KO, Molony DC, Walls RJ, Bashir SP, Mulhall KJ. Increasing financial burden of revision total knee arthroplasty. Knee Surg Sports Traumatol Arthrosc. 2010; 18(7):945–8. doi: 10.1007/s00167-010-1074-8 PMID: 20148322.
- Kallala RF, Vanhegan IS, Ibrahim MS, Sarmah S, Haddad FS. Financial analysis of revision knee surgery based on NHS tariffs and hospital costs: does it pay to provide a revision service? Bone Joint J. 2015; 97-B(2):197–201. doi: 10.1302/0301-620X.97B2.33707 PMID: 25628282.
- Cross M, Smith E, Hoy D, Nolte S, Ackerman I, Fransen M, et al. The global burden of hip and knee osteoarthritis: estimates from the global burden of disease 2010 study. Annals of the rheumatic diseases. 2014; 73(7):1323–30. doi: 10.1136/annrheumdis-2013-204763 PMID: 24553908.
- Kurtz S, Ong K, Lau E, Mowat F, Halpern M. Projections of primary and revision hip and knee arthroplasty in the United States from 2005 to 2030. J Bone Joint Surg Am. 2007; 89-A(4):780–5. doi: 10.2106/jbjs.f.00222
- 40. Patel A, Pavlou G, Mujica-Mota RE, Toms AD. The epidemiology of revision total knee and hip arthroplasty in England and Wales: a comparative analysis with projections for the United States. A study using the National Joint Registry dataset. Bone Joint J. 2015; 97-B(8):1076–81. doi: 10.1302/0301-620X.97B8.35170 PMID: 26224824.



- Health & Social Care Information Centre: Hospital Episode Statistics. <a href="http://www.hscic.gov.uk/hes">http://www.hscic.gov.uk/hes</a> (date last accessed 21 May 2015).
- **42.** Buchholz H, Elson R, Engelbrecht E, Lodenkamper H, Rottger J, Siegel A. Management of deep infection of total hip replacement. J Bone Joint Surg Am. 1981; 63-B(3):342–53.
- Bauer T, Piriou P, Lhotellier L, Leclerc P, Mamoudy P, Lortat-Jacob A. [Results of reimplantation for infected total knee arthroplasty: 107 cases]. Rev Chir Orthop Reparatrice Appar Mot. 2006; 92(7):692– 700. PMID: 17124453.
- Buechel FF, Femino FP, D'Alessio J. Primary exchange revision arthroplasty for infected total knee replacement: a long-term study. Am J Orthop. 2004; 33(4):190–8; discussion 8. PMID: <u>15132327</u>.
- **45.** Goksan SB, Freeman MA. One-stage reimplantation for infected total knee arthroplasty. Journal of Bone & Joint Surgery—British Volume. 1992; 74(1):78–82. PMID: <u>1732271</u>.
- Burnett RSJ, Kelly MA, Hanssen AD, Barrack RL. Technique and timing of two-stage exchange for infection in TKA. Clinical Orthopaedics & Related Research. 2007; 464:164–78. PMID: <u>17975376</u>.
- Buttaro MA, Pusso R, Piccaluga F. Vancomycin-supplemented impacted bone allografts in infected hip arthroplasty. Two-stage revision results. J Bone Joint Surg Am. 2005; 87-B(3):314–9. PMID: 15773637.
- 48. Gulhane S, Vanhegan IS, Haddad FS. Single stage revision: regaining momentum. J Bone Joint Surg Br. 2012; 94(11 Suppl A):120–2. doi: <a href="http://dx.doi.org/10.1302/0301-620X.94B11.30746">http://dx.doi.org/10.1302/0301-620X.94B11.30746</a> PMID: 23118398.
- 49. Vanhegan IS, Morgan-Jones R, Barrett DS, Haddad FS. Developing a strategy to treat established infection in total knee replacement: a review of the latest evidence and clinical practice. J Bone Joint Surg Br. 2012; 94(7):875–81. doi: http://dx.doi.org/10.1302/0301-620X.94B7.28710 PMID: 22733939.