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Abu Al-Rub, Z, Lamb, JN orcid.org/0000-0002-0166-9406, West, RM orcid.org/0000-0001-7305-3654 et al. (3 more authors) (2020) Survivorship of fixed vs mobile bearing unicompartmental knee replacement: A systematic review and meta-analysis of sixty-four studies and National Joint Registries. *The Knee*, 27 (5). pp. 1635-1644. ISSN 0968-0160

<https://doi.org/10.1016/j.knee.2020.09.004>

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1 Survivorship of Fixed Vs Mobile Bearing
2 Unicompartamental Knee Replacement: A Systematic
3 Review and Meta-analysis of Sixty-Four Studies and
4 National Joint Registries

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9

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17

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19 **Conflict of Interest:** JL is supported by the NIHR Biomedical Research Centre at the Leeds Teaching Hospitals
20 NHS Trust and the University of Leeds.

21 HGP is a paid consultant for Bristol Myers Squibb, Depuy Synthes, JRI Orthopaedics, Kennedys Law, Medacta

22 Int, Meril Life, Smith & Nephew and Zimmer Biomet. He has received institutional research grants

23 from Charnley Trust, Depuy Synthes, Glaxo Smith Kline, NIHR, Versus Arthritis and Zimmer Biomet.

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25

26

27 **Funding:** The study is funded by departmental funds held at the University of Leeds, England and Smith and
28 Nephew.

29

30 **Acknowledgment:** *Professor Pandit is a National Institute for Health Research (NIHR) Senior Investigator.*

31 *The views expressed in this article are those of the author(s) and not necessarily those of the NIHR, or the*

32 *Department of Health and Social Care.*

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34 **Word count (Abstract + Manuscript) = 2859 words.**

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58 **Abstract**

59 **Background:** Unicompartmental knee replacement (UKR) prostheses can use fixed (FB) or mobile bearing
60 (MB) constructs. We compared survivorship and failure modes of both designs.

61 **Methods:** The inclusion criteria were studies published between 2005-2020 with minimum average follow-up
62 of 5 years reporting the survival and/or number of revisions of specific designs in medial and lateral UKR.
63 Pooled rate of revision per 100 patient years (PTIR) was estimated using a random effects model.

64 **Results:** 70 cohorts of 17 405 UKRs with weighted mean follow-up of 7.3 years (0.1-29.4 years) were included.
65 A total of 170 923 UKRs were identified in registry reports at a weighted mean implant survival time of 15.4
66 years. PTIR in MB UKR versus FB UKR was similar [1.45 Vs 1.40, (p = 0.8)].

67 In cohort studies, the overall PTIR for MB was also similar to FB [1.03 Vs 0.78, (p= 0.1)]. For medial UKR, the
68 PTIR for MB was marginally greater but not significantly different to FB [0.96 Vs 0.81, (p= 0.3)], whilst for
69 lateral UKR, the PTIR for MB was significantly worse than for FB [(2.20 Vs 0.72, (p<0.01)]. Polyethylene wear
70 is more common in FB implants, whilst MB implants are revised more often for bearing dislocation.

71 **Conclusion:** Overall implant survival in mid- to long-term studies is similar for MB versus FB medial UKR.
72 MB have a four-fold higher risk of revision in comparison to FB when used for lateral UKR.

73

74 **Keywords:** Unicompartmental knee; Revision; Survivorship; Mobile; Fixed; Arthroplasty

75 **1. Introduction**

76 Unicompartmental knee replacement (UKR) is deemed to be appropriate for about one in four osteoarthritic
77 knees needing replacement [1]. UKR is associated with faster recovery, better function and lower morbidity and
78 mortality [2–4]. However, the UKR usage has not exceeded 5 to 10% [5]. Many surgeons are reluctant to
79 perform UKR where indications allow because of higher revision rates than total knee replacement [5].

80

81 A UKR can be performed with either a mobile bearing (MB) or a fixed (FB). Proponents of the MB construct
82 claim a reduction in polyethylene wear in comparison to FB constructs. MB UKR can be more technically
83 challenging with suboptimal outcomes, and higher rate of complications like bearing dislocation [6]. Implants
84 which reduce the technical demands on surgeons may reduce poor outcomes and encourage surgeons to implant
85 UKRs when appropriate. FB UKR utilisation has increased from 30% in 2010, to 49% in 2018 [5]. Adoption of
86 FB designs may be due to a perceived reduction in technical difficulty, reduced bearing dislocation and
87 improved wear of modern polyethylene [7].

88

89 Outcomes and survivorship of both FB and MB constructs are excellent but there are significant differences in
90 the revision rates between cohort studies and in registry data [5,8–10]. Over the past decade, systematic reviews
91 and meta-analyses attempted to address the debate of superiority between the two constructs. Three reviews
92 included RCTs and non-randomised comparative studies analysed 5,10, and 15 studies, respectively [11–13].
93 Other reviews focused on either medial or lateral only UKRs with limited study inclusion potentially limiting
94 the usefulness of their findings [14–19]. Therefore, the aim of this study was to address the issue of limited
95 inclusion of studies and registries in previous reviews and complement their findings through a comprehensive
96 systematic review and meta-analysis of the implant survival in mobile versus fixed bearing UKR for medial as
97 well as lateral constructs, using all available cohort studies with long-term follow up and registry data with
98 minimal restriction from the past 15 years.

99

100 **2. Material and methods**

101 *2.1 Data sources*

102 This analysis was performed in two parts. Firstly, a systematic review and meta-analysis of cohort studies
103 reporting revisions for any cause for MB or FB UKR in either the medial or lateral tibiofemoral joint, over an

104 average follow up of at least five years was performed. Secondly, a meta-analysis of registry data was
105 performed for UKRs where survival was reported separately for either FB or MB UKRs at or beyond least five
106 years.

107

108 ***2.2 Systematic review and meta-analysis of cohort studies***

109 The study methodology was peer reviewed and registered on the PROSPERO (id: CRD42020167444). The
110 literature search was conducted using the online databases Medline and EMBASE. Articles were identified
111 using a combination of keyword searches describing unicompartmental knee replacement, survival and revision.
112 Results were combined with searches for Mesh terms (Appendix 1). Citation searching was performed for all
113 full text manuscripts to identify manuscripts which were not found in initial searches.

114

115 Inclusion criteria for cohort studies included: Articles written in English language, available in full text,
116 published between 2005 and 2020, human studies, with greater or equal to 5 years of average follow-up and
117 reporting the survival and/or number of joints revised of specific UKR (mobile or fixed for either medial or
118 lateral unicompartmental replacement). Revision was defined as revision of any part of the UKR construct. We
119 excluded conference abstracts, manuscripts which reported on the same cohort twice, systematic reviews and
120 registry studies to prevent duplication of observation.

121 Abstracts and the full texts were screened by two authors (JL and ZA) independently and disagreements at each
122 stage were settled by consensus. Risk of bias/quality of studies was assessed using criteria developed by Wyld
123 et al. independently by two authors (JL and ZA) [20]. Where available extracted data included: Title, authors,
124 year of publication, prostheses used (brand and model), reconstructed tibiofemoral joint (medial or lateral),
125 number of female patients, average age of cohort, indication for UKR (osteoarthritis (OA), spontaneous
126 osteonecrosis of the knee (SONK), post traumatic osteoarthritis and other), number of UKRs, number of
127 revisions, survival of cohort (with confidence intervals), revision indications and loss to follow up. Data was
128 extracted by one author (ZA) and a random selection of 10% of screened records were double checked by a
129 second reviewer (JL).

130

131 ***2.3 Meta-analysis of registry reported data***

132 Registry data was obtained through manual searching of all publicly accessible arthroplasty registers. Survival
133 rates for MB and FB UKR were separately reported by four arthroplasty registries and data collected included:

134 Date of report, country of report, prostheses used (brand and model), number of female patients, average age of
135 cohort, number of UKRs, number of revisions, estimated proportion of cohort surviving and time point of
136 survival estimate.

137

138 **2.4 Statistical analysis**

139 The primary exposure was the UKR construct, and the primary outcome measure was all-cause revision of any
140 part of the construct. Study and patient level statistics were estimated using mean values weighted by number of
141 cases. Patient time (PT) was approximated using the product of number of cases in follow-up and average
142 follow-up time in centuries. Patient time incidence rate of revision per 100 years (PTIR) was estimated by
143 dividing the patient time by the number of revisions. For registry reported data, PTIR was derived from the
144 Kaplan-Meier estimates of survival for mobile and fixed bearing UKRs separately. PT was estimated using the
145 product of number of UKRs and the survival time in centuries. PTIR was estimated by dividing PT by the
146 product of the number of UKRs and the proportion revised. PTIR estimates were pooled using a random effects
147 model with a comparison of subgroups based on bearing fixation (FB versus MB UKRs). The random effects
148 method used an inverse-variance method, DerSimonian-Laird estimator for τ^2 , Jackson method for confidence
149 interval of τ^2 and τ and the results are given on a log transformed scale. All data analysis was completed
150 using R (version 3.6.2, Vienna Austria, 2019).

151

152 **3. Results**

153

154 **3.1 Cohort studies:**

155 Our search resulted in 213 references from database and citation searches. After title and abstract screening, 99
156 records were excluded and 114 manuscripts underwent full text review (figure 1).

157

158 Full text review identified 70 cohorts (61 medial UKR and nine lateral UKR) in 64 studies which were included
159 in further analysis. The cohorts included a total of 17 405 cases, which were comprised of 16 619 medial
160 compartment UKR and 786 lateral compartment UKR. Weighted mean follow-up was 7.3 years and ranged
161 from 0.1 to 29.4 years after implantation (table 1). Patient level characteristics derived from reported data are
162 displayed in table 1.

163 Meta-analysis of PTIR revealed an overall rate of revision per 100 patient years (95% CI) was 0.8 (0.6 to 1) for
164 FB UKR and 1.0 (0.8 to 1.3) for MB UKR (test for sub-group differences, $p=0.1$). When analysing by subgroup
165 of constructs on the medial and lateral condyles the rate of revision per 100 patient years (95% CI) was 1 (0.8 to
166 1.1) for medial UKR and 1.3 (0.8 to 2.1) for lateral UKR (test for sub-group differences, $p=0.2$).
167 For medial UKR the PTIR for MBs UKR was higher than that for FB UKR but the difference did not reach
168 significance (0.96 [0.8 to 1.2] versus 0.8 [0.6 to 1.1], $p=0.3$, figure 2).
169 For lateral UKR the PTIR for MBs was significantly greater than for FBs (2.2 [1.3 to 3.6] versus 0.7 [0.4 to 1.2]
170 $p<0.01$, figure 3).
171 Revision indications were reported in 60 of 61 medial UKR cohorts and 6 of 9 lateral UKR cohorts (table 2).
172 The quality of cohort studies was generally low with a majority of studies reporting only from a single centre
173 without any form of adjustment for confounding factors (figure 4).

174

175 **3.2 Registry data**

176 Our search of public reports from national and international registries yielded results for 176 412 UKRs,
177 comprising of 75 625 FB UKRs and 100 787 MB UKRs. Weighted mean time point for survival estimates was
178 15.3 years.

179 Pooled PTIR derived from survival estimates reported by registries demonstrated a marginally increased PTIR
180 (95% CI) for revision in MB UKR versus FB UKR, (1.44 [1.2 to 1.7] versus 1.37 [1.1 to 1.7] respectively, $p=$
181 0.75, Figure 5). The registry data does not differentiate between medial and lateral UKR and therefore sub-
182 group analysis could not be performed.

183 Indications for revision in all UKRs combined were reported in all registries but subgroup analysis according to
184 the mobility of bearing surface was only provided in the National Joint Registry of England, Wales, Northern
185 Ireland and the Isle of Man, 16th report (table 3)[5] .

186

187 **4. Discussion**

188

189 This comprehensive review of fixed versus mobile bearings used in medial or lateral UKRs has shown a
190 marginally better implant survival for FB UKR over MB UKR in cohort as well as registry data. These
191 differences are more apparent in the lateral UKR (nearly four-fold worse for mobile UKR). Progression of

192 arthritis is the commonest cause of revision of a UKR. This study is the first to integrate large amounts of
193 cohort and registry data to evaluate the survivorship of the two available bearing designs for UKRs in
194 both the medial and lateral compartments.

195
196 Results extracted from manuscripts and registry reports did not demonstrate a statistically significant difference
197 between FB and MB UKR overall, which is similar to results reported from other reviews [11,12,21]. Overall,
198 there was a small increased risk of revision associated with MB implants as compared to FB implants, although
199 the difference is too small to make a meaningful statistical difference. Given that the numbers of lateral UKR
200 are far less than medial UKR, the results are heavily skewed by medial UKR outcomes. It is up to the surgeon to
201 choose a mode of bearing fixation for both medial and lateral UKR with which they can safely rebalance the
202 reconstructed knee.

203
204 There was no significant difference in the rate of revision between FB and MB for medial UKR. Cohort studies
205 demonstrated that bearing dislocation and bearing breakage as a failure mechanism are unique in the MB group
206 whilst bearing wear is more common in the FB group. The higher bearing wear seen in FBs is compatible with
207 previously published literature showing a higher mean penetrative wear rate 0.15 vs 0.04 mm/year in fixed
208 versus mobile bearing UKRs [22,23]. In anteromedial OA of the knee, patients typically bear more weight
209 through the medial tibio-femoral joint which has greater constraint in normal walking than the lateral tibio-
210 femoral joint. This can increase the loading of the reconstructed medial UKR and increase the likelihood of
211 bearing wear. Therefore, any potential benefit of bearing stability might be offset by the increased risk of wear
212 with medial FB UKR, particularly where older high wear polyethylene bearings are used [24].

213
214 For lateral UKR, there was an approximately four-fold risk of revision associated with the use of MB UKR
215 versus FB UKR, which is agreement with a previous review [14]. The lateral condylar joint is less constrained
216 which increases the underlying risk of bearing dislocation for MB UKR. Lateral UKR performed for valgus
217 knee OA may bear a larger proportion of body weight through the lateral versus medial condylar joint.
218 However, there were no reported cases of revision of lateral UKR (fixed or mobile) due to bearing wear [25,26].
219 It appears that the FB construct maintains the benefits of a stable bearing without any reciprocal increase in
220 wear.

221

222 Analysis of revision indications provide further insight into the potential mechanism of failure for fixed or
223 mobile constructs. By definition, bearing dislocation is only confined to mobile constructs and therefore not
224 possible to compare the risk, whilst bearing breakage is significantly higher in the mobile cohort and may be
225 related to the bearing thickness. In comparison, FB wear is significantly higher than MB potentially representing
226 the price paid by the FB constructs in exchange for reduced risk of bearing dislocation and bearing breakage.

227

228 Progression of arthritis is a dominant indication for revision for both FB and MB UKRs, an effect seen in the
229 reported revision indications in cohort studies and in results reported by the NJR (National Joint Registry for
230 England, Wales, Northern Ireland and the Isle of Man). The ability of the surgeon to correctly balance the knee
231 and prevent overloading of the contralateral compartment is likely to be a key factor. Therefore, this skill
232 becomes more challenging in the MB UKR when the surgeon must also carefully cut and select the appropriate
233 bearing size for the reconstructed condyle to reduce the risk of dislocation, particularly following lateral MB
234 UKR. As such, reduction of bearing instability and surgical complexity without excessive bearing wear may
235 potentially be achieved by using a FB UKR construct, particularly when used in the lateral compartment or
236 when low wear polyethylene bearings are used.

237

238 There is a noticeable difference in the reported indications for revision between the registry reports and the
239 results of this meta-analysis. For example, bearing dislocation as an indication for revision was less than half the
240 frequency reported in cohort studies, whilst the incidence of aseptic loosening and lysis was significantly higher
241 for registry reported UKRs. This can possibly be explained by the fact that bearing dislocation treated with
242 bearing exchange may not be recorded consistently in registry data collection methods. Furthermore, it is more
243 likely that revisions involving removal or exchange of tibial or femoral component, in a planned setting where
244 data collection techniques are readily used, will be more accurately recorded. Those differences are likely to
245 reflect the constraints of the literature and registries to accurately report the true indication for revisions and
246 may account for those findings.

247

248 There is also a prominent difference in the revision rates between different designs and different registry
249 datasets. Those differences are likely to be multi-factorial and may be related to patient, surgical, implant
250 and registry factors. Patient factors may include differing levels of activity, comorbidity, weight, height
251 and baseline grade of arthritis [27–29] . Surgical factors include surgeon’s annual caseload, patient

252 selection, misinterpretation of radiographs and threshold for revision [27–32]. Implant factors include
253 variation in implant design along with variation described in bearing mobility [33]. Registry factors may
254 include reporting method, definition of revision procedures and data completeness.

255 *4.1 Study Limitations*

256

257 Our meta-analysis is based on observational data subject to a majority of low study quality, large statistical
258 heterogeneity because of their inherent biases, and design differences that meta-analysis cannot overcome. Four
259 studies reported results from multi-centre patients, whilst the rest of studies included in this analysis were
260 single-centre and/or single surgeon cohorts [10,27–29]. Two randomised prospective studies were identified but
261 didn't meet our inclusion criteria due to either short-term follow up and/or not reporting on survival of implant
262 construct specifically [30,31]. Our inclusion criteria were set to reduce the bias of shorter follow up studies
263 which may report early complication such as bearing dislocation. Those criteria may unintentionally reduce the
264 number of cases from studies reporting survival of the most recently produced implants and approaches.

265 Estimated PTIR in the study assumes that the revision rate is relatively constant over different periods of patient
266 time, which may not accurately reflect the true risk of revision for any particular patient. PTIR estimates derived
267 from average follow up is a useful method but may have limited precision, particularly with abnormal
268 distributions of follow up time in the reported studies. Lateral UKR is an uncommon indication for UKR which
269 reduced the number of cohorts available limiting our sample sizes in the pooled analysis. This is reflected in the
270 confidence limits of our estimates. Implants used in this analysis may have used polyethylene with higher wear
271 rates, suboptimal fixation methods, and outdated technology which is no longer commonly used. No adjustment
272 for those limitations is possible due to the lack of detail given in the studies. As such, the results in this study
273 may not accurately represent the benefits provided by recent or future technological advancements. Further
274 insights into the performance of modern implants may be better understood using robust analysis of registry
275 data or well-designed prospective trials. Joint registry data has the advantages of large sample size, usage by
276 multiple surgeons in different settings and may be more generalisable in comparison to cohort study data.
277 However, it does not differentiate between medial and lateral UKRs and does not take into consideration the
278 learning curve and/or indication for primary surgery or revision surgery.

279

280 **5. Conclusions**

281

282 This up to date and comprehensive study demonstrates a comparable incidence of all cause revision between
283 fixed and mobile bearing UKR. Subgroup analysis demonstrated a four-fold increase in risk of all cause revision
284 associated with mobile UKR when used for lateral condylar reconstruction in comparison to fixed bearing
285 UKRs. Failure of prosthesis is multifactorial and may be determined by factors other than bearing stability and
286 risk of bearing dislocation. Future well-designed randomised controlled trial with standardised patient
287 selection criteria to assess the long-term survival of fixed versus mobile bearing would be the ideal way to
288 verify the results of the current available literature regarding the two constructs.

289

290

291 **6. Contribution of Authors:**

292 **1- ZA: Data retrieval, manuscript preparation and editing.**

293 **2- JL: Study design, Literature search, statistical analysis, manuscript preparation and editing.**

294 **3- RMW: Statistical methods and analysis**

295 **4- XY: Manuscript editing**

296 **5- YH: Manuscript editing**

297 **6- HGP: Study design, manuscript editing.**

298

299 **7. Declaration**

300 **7.1 Conflict of Interest:** JL is supported by the NIHR Biomedical Research Centre at the Leeds Teaching
301 Hospitals NHS Trust and the University of Leeds.

302 HGP is a paid consultant for Bristol Myers Squibb, Depuy Synthes, JRI Orthopaedics, Kennedys Law, Medacta
303 Int, Meril Life, Smith & Nephew and Zimmer Biomet. He has received institutional research grants
304 from Charnley Trust, Depuy Synthes, Glaxo Smith Kline, NIHR, Versus Arthritis and Zimmer Biomet.

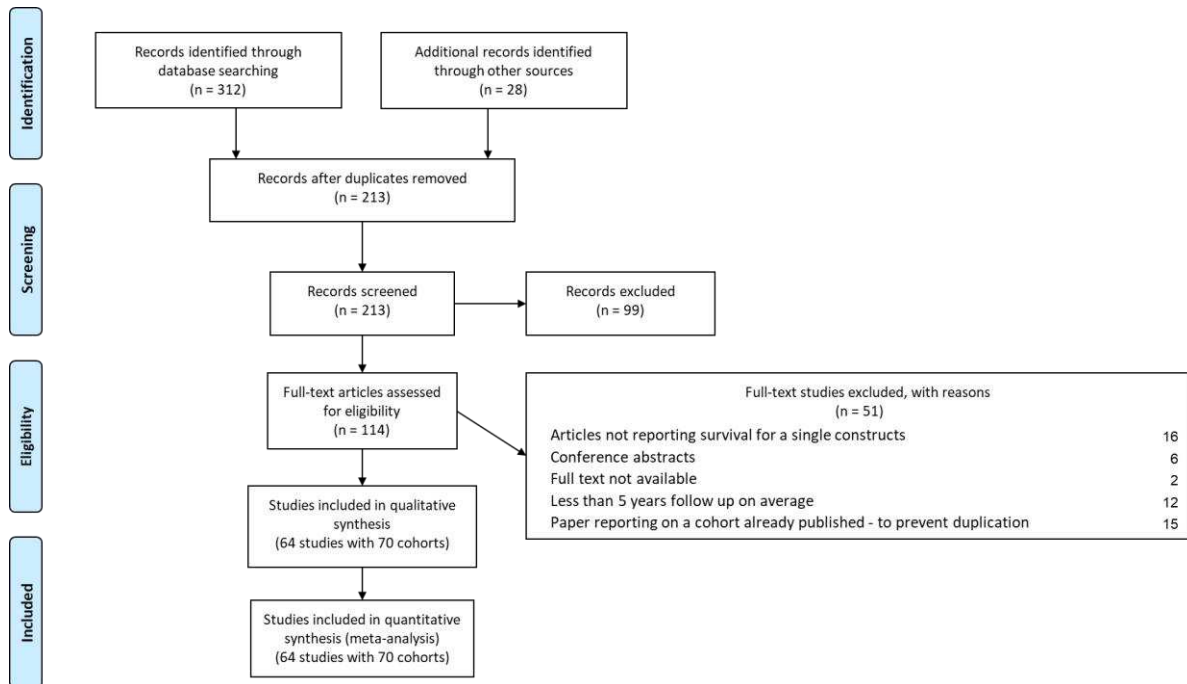
305

306 **7.2 Acknowledgment:** *Professor Pandit is a National Institute for Health Research (NIHR) Senior Investigator.*
307 *The views expressed in this article are those of the author(s) and not necessarily those of the NIHR, or the*
308 *Department of Health and Social Care.*

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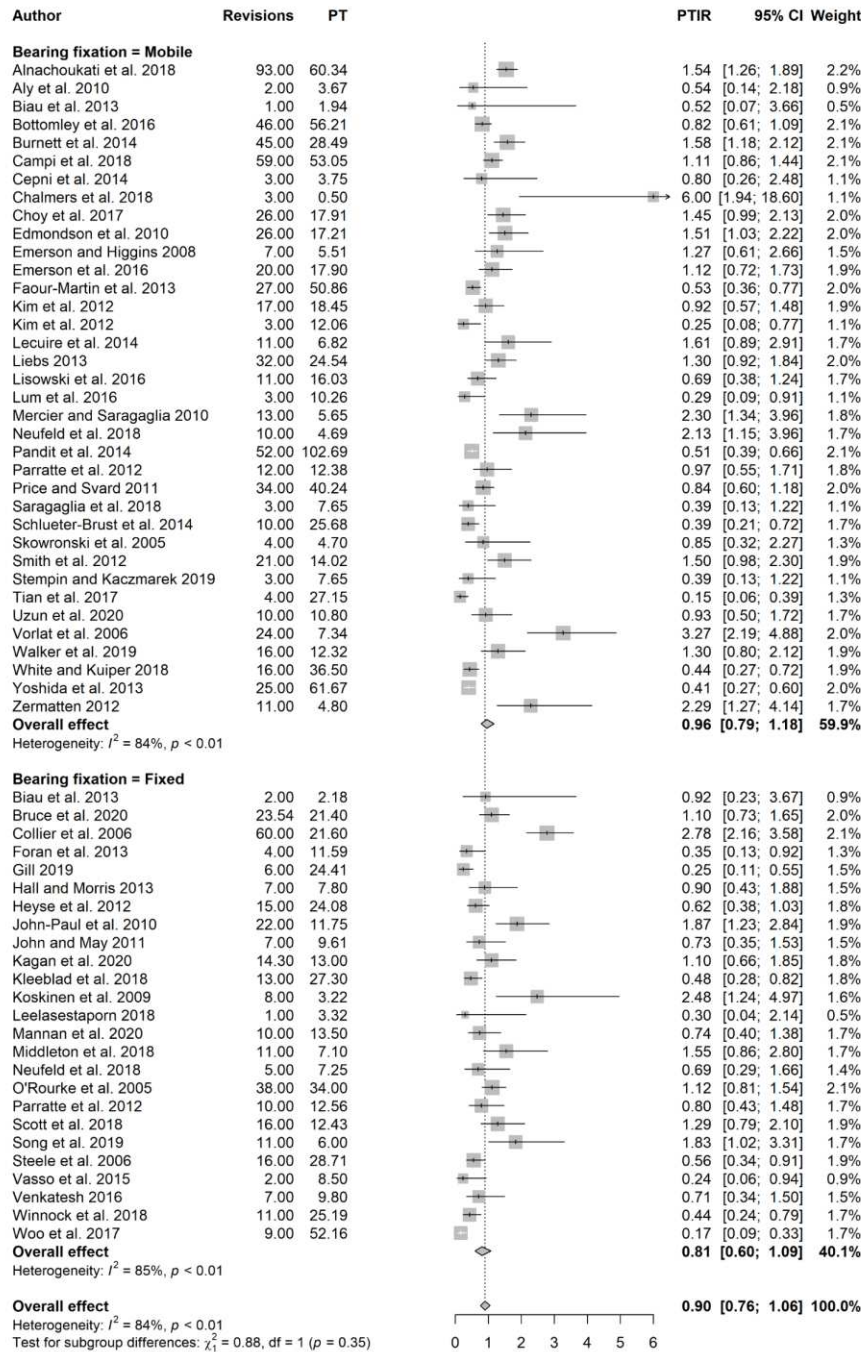
310 **7.3 Funding:** The study is funded by departmental funds held at the University of Leeds, England and Smith
311 and Nephew.

312 **Figure 1.** PRISMA flow chart summarising data collection for cohorts reported in peer reviewed manuscripts.



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331 **Figure 2.** Forest plot demonstrating pooled patient-time incidence rates of revision per 100 years (PTIR) using a
 332 random effects model for all medial UKR cohorts. Incidence rates for fixed and mobile bearing constructs are
 333 reported separately. Note: PT indicates Patient time in centuries, CI indicates confidence interval.

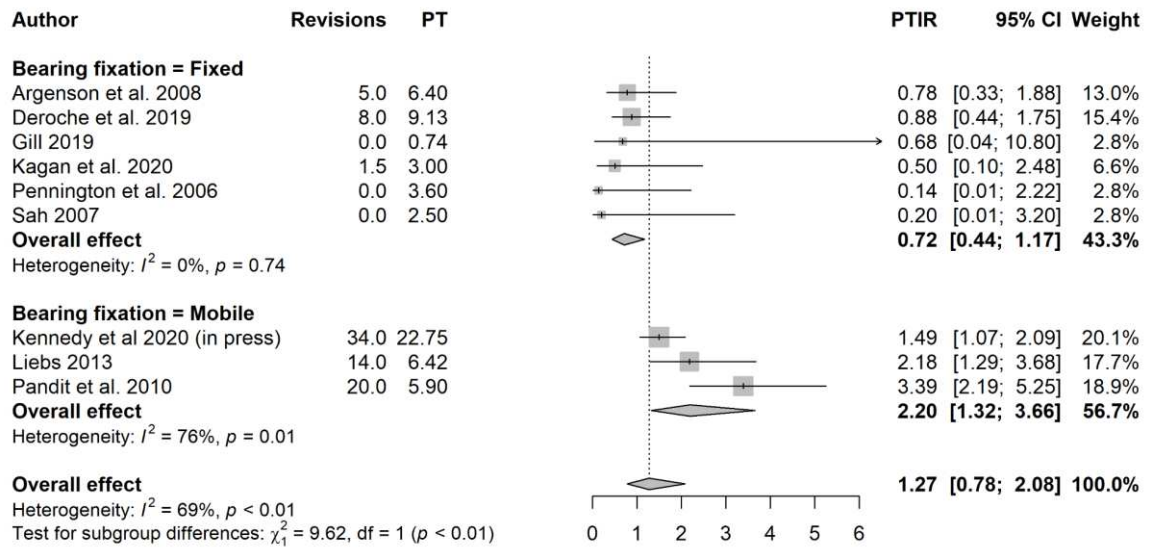


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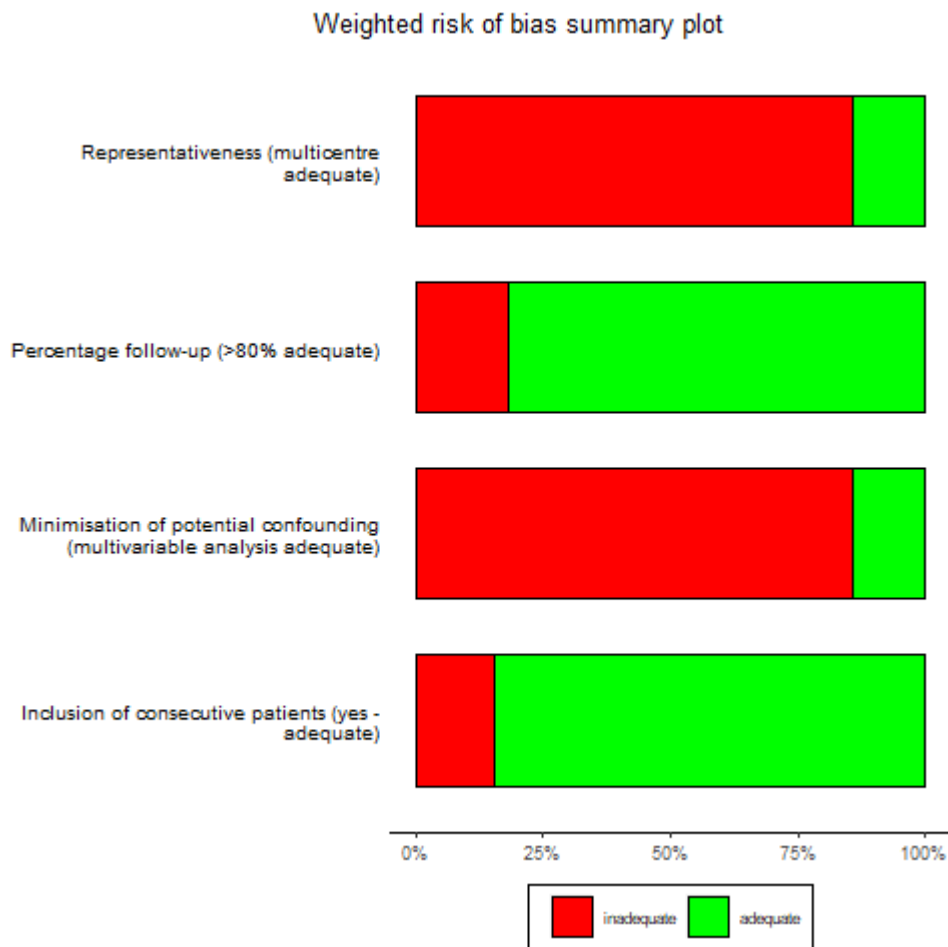
337 **Figure 3.** Forest plot demonstrating pooled patient-time incidence rates of revision per 100 years (PTIR) using a
 338 random effects model for all lateral UKR cohorts. Incidence rates for fixed and mobile bearing constructs are
 339 reported separately. Note: PT indicates Patient time in centuries, CI indicates confidence interval.



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356 **Figure 4.** Plot demonstrating the quality assessment of included studies.

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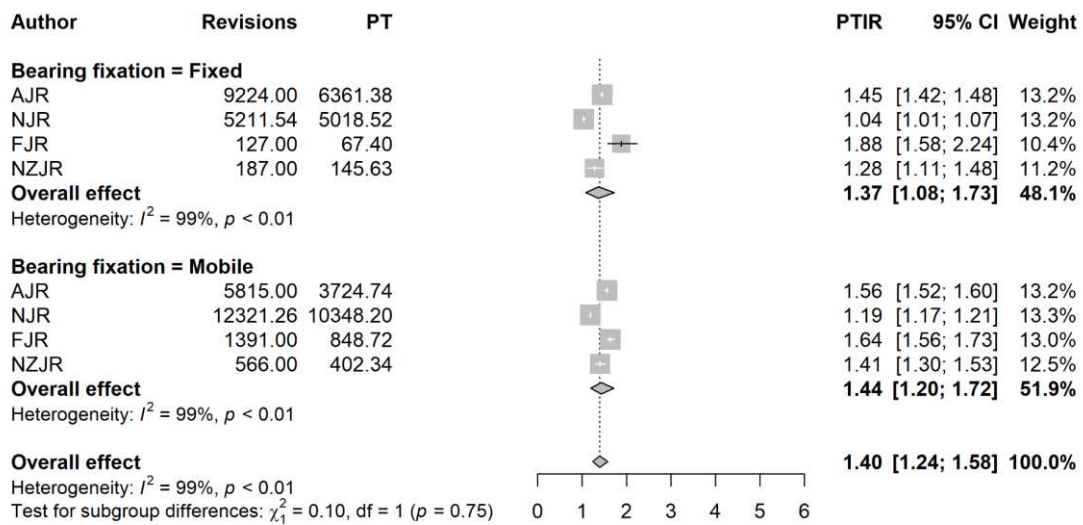
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369 **Figure 5.** Forest plot demonstrating pooled patient-time incidence rates for revision per 100 years (PTIR) using
 370 a random effects model for all registry reported UKR cohorts. Incidence rates for fixed and mobile bearing
 371 constructs are reported separately.
 372 PT: Patient time in centuries, CI: confidence interval, NJR: National Joint Registry for England, Wales, North
 373 of Ireland and Isle of Man, FJR: Finnish Joint Registry, AJR: Australian Orthopaedic Association National Joint
 374 Replacement Registry, NZJR: New Zealand Joint Registry.



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Table 1. Summary of study and patient characteristics

| | Lateral UKR | | Medial UKR | |
|--------------------------------|--------------------|----------------|-------------------|----------------|
| | Fixed bearing | Mobile bearing | Fixed bearing | Mobile bearing |
| Number of UKR | 215 | 571 | 4982 | 11637 |
| Year of publication | 2006-2020 | 2010-2013 | 2005-2020 | 2005-2020 |
| Patient characteristics | | | | |
| Mean age (years) | 64.0 | 65.8 | 64.9 | 67.1 |
| Female patients (%) | 67.7 | 64.8 | 47.3 | 54.7 |
| Indications | | | | |
| Osteoarthritis (%) | 81.4 | 98.8 | 91.8 | 95.1 |
| SONK (%) | 2.6 | 1.1 | 3.1 | 2.1 |
| Post-Traumatic OA (%) | 16.0 | 0.0 | 0.9 | 0.3 |
| Other (%) | 0.0 | 0.0 | 4.3 | 2.2 |
| Mean follow up | 11.6 | 6.4 | 7.6 | 7.1 |
| Follow up range | 1.5-24.0 | 2.1-9.8 | 1-29.4 | 0.1-22.0 |

Note: Means weighted by number in study, where reported. UKR indicates unicompartmental knee replacement, SONK spontaneous osteonecrosis of the knee and OA osteoarthritis

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Table 2. Summary of cases revised for each unicompartmental knee replacement construct

| Revision indication | Lateral UKR | | Medial UKR | |
|--|--------------|---------------|--------------|---------------|
| | <i>fixed</i> | <i>mobile</i> | <i>fixed</i> | <i>mobile</i> |
| | n (%) | n (%) | n (%) | n (%) |
| Arthrofibrosis | 0 (0) | 1 (1.5) | 1 (0.3) | 4 (0.6) |
| Bearing breakage | 0 (0) | 0 (0) | 1 (0.3) | 10 (1.5) |
| Bearing dislocation | 0 (0) | 23 (33.8) | 0 (0) | 121 (17.6) |
| Bearing wear | 0 (0) | 0 (0) | 73 (22.1) | 7 (1) |
| Both or unspecified components loosening | 0 (0) | 6 (8.8) | 39 (11.8) | 37 (5.4) |
| Component malposition | 0 (0) | 0 (0) | 3 (0.9) | 3 (0.4) |
| Femoral component loosening | 0 (0) | 1 (1.5) | 8 (2.4) | 42 (6.1) |
| Peri-prosthetic Fracture | 0 (0) | 4 (5.9) | 5 (1.5) | 16 (2.3) |
| Haemarthrosis | 0 (0) | 3 (4.4) | 0 (0) | 8 (1.2) |
| Impingement | 0 (0) | 2 (2.9) | 0 (0) | 6 (0.9) |
| Infection | 0 (0) | 7 (10.3) | 18 (5.4) | 45 (6.6) |
| Instability | 0 (0) | 0 (0) | 4 (1.2) | 12 (1.7) |
| Pain | 0 (0) | 1 (1.5) | 36 (10.9) | 45 (6.6) |
| Progression of OA | 12 (80) | 18 (26.5) | 92 (27.8) | 210 (30.6) |
| Synovitis | 0 (0) | 1 (1.5) | 1 (0.3) | 3 (0.4) |
| Tibial component loosening | 1 (6.7) | 1 (1.5) | 41 (12.4) | 86 (12.5) |
| Unknown | 2 (13.3) | 0 (0) | 9 (2.7) | 31 (4.5) |

Note: Counts are displayed with percentage of all extracted reasons for revision in each column in parentheses. UKR

indicates unicompartmental knee replacement, OA osteoarthritis

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Table 3. Patient time incidence rate estimates (per 100 years) of revision indications for unicompartmental knee replacement constructs in the National Joint Registry of England, Wales, Northern Ireland and the Isle of Man, 16th report

| | All unicompartmental | | Fixed bearing | | Mobile bearing | |
|---------------------------|----------------------|-------------|---------------|-------------|----------------|-------------|
| | PTIR | % | PTIR | % | PTIR | % |
| Pain | 0.24 | 15.8 | 0.25 | 16.1 | 0.24 | 15.6 |
| Dislocation / subluxation | 0.08 | 4.9 | 0.01 | 0.8 | 0.10 | 6.7 |
| Infection | 0.05 | 3.4 | 0.06 | 3.8 | 0.05 | 3.2 |
| Aseptic loosening / lysis | 0.33 | 21.5 | 0.31 | 20.0 | 0.34 | 22.2 |
| Peri-prosthetic fracture | 0.03 | 1.8 | 0.02 | 1.5 | 0.03 | 1.8 |
| Implant wear | 0.11 | 7.3 | 0.10 | 6.5 | 0.12 | 7.7 |
| Instability | 0.10 | 6.4 | 0.07 | 4.7 | 0.11 | 7.1 |
| Malalignment | 0.06 | 4.2 | 0.06 | 3.6 | 0.07 | 4.3 |
| Other indication | 0.17 | 11.1 | 0.12 | 7.8 | 0.19 | 12.5 |
| Stiffness | 0.02 | 1.4 | 0.03 | 1.6 | 0.02 | 1.2 |
| Progress | 0.34 | 22.2 | 0.32 | 20.9 | 0.35 | 22.8 |

Note: PTIR indicates Patient time incidence rate (per 100 patient years). Estimates are derived from different periods of patient time exposure and multiple indications may exist for a single revision.

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