The influence of preoperative determinants on quality of life, functioning and pain after total knee and hip replacement: a pooled analysis of Dutch cohorts

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Abstract

Background: Previous research has identified preoperative determinants that predict health related quality of life (HRQoL), functioning and pain after total knee or hip arthroplasty (TKA/THA), but these differed between studies and had opposite directions. This may be due to lack of power and not adjusting for confounders. The present study aims to identify the preoperative determinants that influence health related quality of life (HRQoL), functioning and pain after total knee or hip arthroplasty (TKA/THA).

Methods: We pooled individual patient from 20 cohorts with OA patients data (n = 1783 TKA and n = 2400 THA) in the Netherlands. We examined the influence of age, gender, BMI and preoperative values of HRQoL, functioning and pain on postoperative status and total improvement. Linear mixed models were used to estimate the effect of each preoperative variable on a particular outcome for each cohort separately. These effects were pooled across cohorts using a random effects model.

Results: For each increase in preoperative point in HRQoL, the postoperative HRQoL increased by 0.51 points in TKA and 0.37 points in THA (SF-36 scale). Similarly, each point increase in preoperative functioning, resulted in a higher postoperative functioning of 0.31 (TKA) and 0.21 (THA) points (KOOS/HOOS-ADL scale). For pain this was 0.18 (TKA) and 0.15 (THA) points higher (KOOS/HOOS-pain scale) (higher means less pain). Even though patients with better preoperative values achieved better postoperative outcomes, their improvement was smaller. Women and patients with a higher BMI had more pain after a TKA and THA. Higher age and higher BMI was associated with lower postoperative HRQoL and functioning and more pain after a THA.

Conclusions: Patients with a better preoperative health status have better outcomes, but less improvement. Even though the independent effects may seem small, combined results of preoperative variables may result in larger effects on postoperative outcomes.

Keywords: Total hip replacement, Total knee replacement, Preoperative, Postoperative, Quality of life, Functioning, Pain

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Background
Total knee or hip arthroplasty (TKA/THA) is an effective treatment for most individuals who suffer from pain and loss of function due to end stage symptomatic osteoarthritis of the hip and knee (OA). In 2010, 109 and 153 patients per 100,000 persons received a TKA or THA respectively in Europe [1]. The development and progression of OA are strongly influenced by age and obesity and both occur more frequently in women [1]. Parallel to the rising prevalence of knee and hip OA, due to an ageing society and obesity, surgery rates are rising as well [2–4].

TKA and THA should not be performed too early since revision rates are higher in younger patients and the length of life of a prosthesis is limited [5]. On the other hand performing a surgery earlier gives more years of productive quality-adjusted life years (QALY’s). However, outcomes after revision surgery are generally worse compared to primary surgery [6]. Current practice shows that preoperative disease severity varies largely among centers and countries [7, 8], suggesting differences in timing. In addition, about 10–20% of the patients is not satisfied after primary TKA/THA [9–12], possibly caused by unmet expectations of patients or due to suboptimal timing of surgery.

Previous research has identified preoperative determinants that influence outcomes, but these differed between studies and had opposite directions [13]. This may be due to lack of power so that some studies did not find any effect, while other studies did not adjust for confounders. In addition, most registries collect a minimal data set [14] e.g. only the VAS scale for pain. Therefore, pooling the data from available cohort studies may provide more reliable evidence on which determinants influence the outcome after TKA/THA because of the larger sample size than separate studies and a more comprehensive set of questionnaires with the ability to measure each outcome more reliably compared to registry studies.

Objective
The present study aims to examine the independent effect of several preoperative determinants for outcomes after TKA or THA by pooling individual patient data from available prospective cohorts in the Netherlands.

Methods
Study design and setting
The ARGON-OPTIMA (Outcome Predictors for TIMing of ArthropLasty) study is part of the ARGON program (Arthritis Research Group Orthopaedics in The Netherlands). Within this study, we pooled individual patient data from all available prospective TKA/THA cohorts in the Netherlands. All orthopaedic clinics in The Netherlands were invited to participate and submit data. We included prospective cohorts among patients with primary OA who underwent TKA or THA, with at least one preoperative and one postoperative measurement on functional or clinical outcomes and a follow-up of at least one year. Cohorts regarding metal-on-metal (MoM) prostheses were excluded, since these are not recommended in current guidelines in The Netherlands.

Participants
Twenty hospitals submitted data and 20 cohorts from 11 hospitals were included. Nine hospitals were excluded because they did not meet the inclusion criteria. Of the included cohorts, 8 cohorts included 1783 knee OA patients undergoing primary TKA and 12 cohorts included 2400 hip OA patients undergoing primary THA. Table 1 shows the characteristics of patients per cohort.

Preoperative determinants
The assessed preoperative determinants were age, gender and BMIs. Furthermore, we examined the influence of preoperative health related quality of life (HRQoL), functioning and pain.

Postoperative outcomes
We studied the effect on the absolute level of the postoperative outcome, but also on the extent of improvement to assess which patients would benefit most from change in health related quality of life (HRQoL), functioning and pain.

Standardization
Since different cohorts used different questionnaires, these were standardized to compare the same domains across different questionnaires. Furthermore, multiple questionnaires were sometimes used to measure the same domain within a cohort. As each patient should be included only once for each domain, we ordered questionnaires in their ability to measure each outcome reliably. This was done during an ARGON consortium meeting. A group of experts within the ARGON consortium discussed about the ordering of questionnaires until consensus was reach. The following main points were taken into concern: is it a general or disease specific questionnaire, how many items are used to calculate the composite score, and is it a common used questionnaire in the Netherlands.

Only the highest rated questionnaire in each dataset was included. The following ordering was used:
For each patient we calculated the standardized score at each time point using the following formula (functioning as example):

\[
\text{Standardized Functioning score for patient X at time point } t = \frac{\text{functioning score for patient X in cohort Y at time point } t - \text{preoperative mean of functioning among patients in cohort Y}}{\text{preoperative SD of functioning}}
\]

Some questionnaires differed in the direction of the scale e.g. on the VAS pain scale, lower scores mean less pain whereas lower scores mean more pain on the HOOS/KOOS subscale pain. The direction of all scales were recoded so that higher scores referred to better values.

**Statistical analysis**

Data of TKA and THA were analyzed separately. As a first step, linear mixed models (LMM) were used to estimate the influence of each preoperative variable on each major outcome for each cohort separately, adjusted for the other variables. As determinants were included in the fixed part of the LMM: the standardized preoperative score (HRQoL, functioning and pain), age, sex, BMI and follow-up time. Interaction terms were fitted between the determinants and follow-up time. In the LMM the patients were specified as the subjects, with an unstructured covariance matrix. This
was done for each standardized postoperative outcome. In the second step, the regression coefficients from all cohorts were pooled using a random effects model to obtain one pooled estimate for each preoperative variable and outcome. Given the pooled estimates of the impact of preoperative status on postoperative status, we can also determine the total improvement (postoperative minus the preoperative status). If patients would have the same amount of improvement, 1 point higher in preoperative status would result in a postoperative status of 1 point higher. So if the increase in postoperative status is < 1 (e.g. 0.4), this means that the improvement is 0.6 points smaller for every point increase in preoperative status.

Given that preoperative scores were standardized, the pooled regression coefficient should be interpreted as the number of standard deviations that an outcome will change, per point increase in the preoperative variable. For example looking at the effect of age on postoperative functioning with a standardized regression coefficient of 0.2 and the preoperative SD of functioning is 7, this means that one year increase in age is estimated to increase the postoperative functioning by: 0.2*7. To facilitate interpretation of the pooled standardized regression coefficients of age, BMI and gender, we transformed standardized regression coefficients back to a 0–100 scale (e.g. HOOS, SF-36), using the preoperative values. Most effects were small and homogeneous. For example looking at the effect of age on postoperative functioning with a standardized regression coefficient of 0.2 and the preoperative status on postoperative status, we can also determine the total improvement (postoperative minus the preoperative status) of the study participants.

**Assessment of heterogeneity**

The I^2 statistic was used to quantify heterogeneity between cohorts. This can be interpreted as the percentage of total variability in a set of effect sizes due to between-studies variability. We considered results as heterogeneous when I^2 was 50% or greater [17].

**Ethics approval and consent to participate**

The Medical Ethical Committee of the Leiden University Medical Center (CME P15.043/SH/sh) confirmed on February 13 2015 that ethical approval for this type of study is not required under the Dutch Medical Research (Human Subjects) Act. The hospitals that supplied anonymous data obtained written informed consent from the study participants.

**Results**

**Age, gender and BMI**

Table 2 shows the pooled estimates of the effect of age, gender and BMI on outcomes as well as the transformed coefficients (95% CI).

| Arthroplasty | Patients characteristic | Outcome | Cohorts (n) | Patients (n) | Standardized regression coefficients (95% CI) | Pooled estimate (0–100 scale) | I^2 (%) |
|-------------|------------------------|---------|-------------|-------------|----------------------------------------------|------------------------------|-------|
| TKA^a       | Age                    | HRQoL^d | 4           | 774         | 0.00 (−0.00, 0.01)                              | 0.00                          | 0.0   |
|             | Functioning            | 6       | 1021        |             | −0.01 (−0.01, 0.00)                             | −0.18                        | 0.0   |
|             | Pain                   | 6       | 1102        |             | 0.01 (−0.00, 0.02)                              | 0.16                         | 47.0  |
| Gender (women) | HRQoL        | 4       | 774         |             | −0.05 (−0.23, 0.13)                             | −0.38                        | 0.0   |
|             | Functioning            | 6       | 1021        |             | −0.24 (−0.50, 0.01)                             | −4.12                        | 53.6  |
|             | Pain                   | 6       | 1102        |             | −0.25 (−0.50, −0.01)                            | −3.92                        | 50.5  |
| BMI^b       | HRQoL                  | 4       | 774         |             | −0.02 (−0.06, 0.02)                             | −0.23                        | 76.1  |
|             | Functioning            | 6       | 1021        |             | −0.01 (−0.05, 0.02)                             | −0.18                        | 62.5  |
|             | Pain                   | 6       | 1102        |             | −0.03 (−0.05, −0.01)                            | −0.47                        | 13.1  |
| THA^b       | Age                    | HRQoL    | 8           | 1436        | −0.01 (−0.02, −0.01)                             | −0.08                        | 0.0   |
|             | Functioning            | 10      | 1271        |             | −0.02 (−0.02, −0.01)                             | −0.33                        | 0.0   |
|             | Pain                   | 10      | 1492        |             | −0.01 (−0.01, 0.00)                             | −0.18                        | 0.0   |
| Gender (women) | HRQoL        | 8       | 1436        |             | −0.10 (−0.22, 0.01)                             | −0.78                        | 0.0   |
|             | Functioning            | 10      | 1271        |             | −0.11 (−0.22, 0.01)                             | −1.95                        | 10.9  |
|             | Pain                   | 10      | 1492        |             | −0.11 (−0.21, −0.00)                            | −2.00                        | 0.0   |
| BMI         | HRQoL                  | 8       | 1436        |             | −0.03 (−0.04, −0.01)                             | −0.23                        | 0.0   |
|             | Functioning            | 10      | 1271        |             | −0.02 (−0.04, −0.01)                             | −0.35                        | 0.0   |
|             | Pain                   | 10      | 1492        |             | −0.02 (−0.03, −0.00)                             | −0.36                        | 0.0   |

^a TKA Total Knee Arthroplasty, ^b THA Total Hip Arthroplasty, ^c BMI Body Mass Index, ^d HRQoL Health Related Quality of Life
An increase in BMI with one point, resulted in more postoperative pain (0.5 points on a 0–100 scale). For THA, age and BMI were significantly associated with HRQoL, functioning and pain. One year increase in age decreased postoperative functioning by 0.3 point on a 0–100 scale. Furthermore, women perceived more pain after a THA (2 points on a 0–100 scale).

**Health related quality of life**

Four cohorts examined the effect of preoperative HRQoL on postoperative HRQoL in 760 patients after TKA (Fig. 1). Eight cohorts examined this effect in 1436 patients with a THA (Fig. 2). A significant positive effect of preoperative HRQoL was found of 0.51 (95% CI 0.32 to 0.71) for patients after TKA and 0.37 (95% CI 0.21 to 0.53) after THA. This means that a patient with 1 point higher preoperative HRQoL on average achieves a 0.51 point (TKA) and 0.37 point (THA) higher postoperative HRQoL on the SF-36 scale. At the same time, if patients with a 1 point higher preoperative HRQoL reach a 0.51 point higher postoperative HRQoL after TKA, this also means that their improvement is 0.49 (0.51–1) points less. For THA this implies 0.63 (0.37–1) points less improvement postoperative. The results were heterogeneous, meaning that included cohorts differed with respect to the estimated effect for either TKA or THA.

**Functioning**

Six cohorts examined the effect of preoperative functioning on postoperative functioning in 1221 patients with a TKA (Fig. 3) and 10 cohorts examined this effect in 1271 patients with a THA (Fig. 4). We found a significant positive effect of 0.31 (95% CI 0.23 to 0.39) for TKA and 0.21 (95% CI 0.16 to 0.26) for THA. This means that a patient with a 1 point higher preoperative functioning on average achieves a 0.31 points higher postoperative functioning on the KOOS scale (TKA) and 0.21 points of the HOOS scale (THA). At the same time this means that these patients have a 0.69 and 0.79 point less improvement for TKA and THA respectively for every 1 point higher on preoperative functioning. The results were homogeneous meaning that the estimated effects did not differ between cohorts.

**Pain**

Six cohorts examined the effect of preoperative pain on postoperative pain in 1102 TKA patients (Fig. 5) and 12 cohorts examined this effect in 1492 THA patients (Fig. 6). We found that every point increase in preoperative pain (i.e. less pain) was associated with 0.18 (95% CI 0.11 to 0.26) point increase in postoperative pain after a TKA and 0.15 (95% CI 0.08 to 0.21) after a THA. This also means that patients with less preoperative pain improve 0.82 points less after TKA and 0.85 points less after THA. The results were homogeneous meaning that the estimated effects did not differ between cohorts.

**Combined results**

Even though the independent effect of one variable may be small, the combined effect of different determinants may result in clinically relevant differences. Table 3 shows some hypothetical scenarios in which several determinants are combined. The first scenario is that a patient first loses some weight and reduces the BMI with 5 points to improve the postoperative functioning after THA. This takes some time (e.g. 5 years) and a higher age decreases the postoperative functioning. Suppose that due to the weight loss the preoperative functioning increases with 5 points (on a 0–100 scale). Taken together, this results in a 1.2 points higher postoperative outcome. The second scenario is that a surgeon thinks a patient is too young to perform a THA. If a patient
receives this THA 10 years later, and during this 10 years the patient also gains weight due to an inactive lifestyle (e.g. 10 points of BMI) and the functioning also reduces with 10 points (on a 0–100 scale), his/her postoperative functioning will be 9 points lower compared to the situation if she/he had received THA surgery 10 years earlier. The effect of these scenarios on HRQoL and pain are also shown in Table 3. Overall effects vary between 1.2 and 6.5 points better postoperative outcomes for scenario 1 and between 1.6 and 9 points worse postoperative outcomes for scenario 2.

Discussion

The present pooled analysis of 1783 knee and 2400 hip OA patients shows that patients with a higher preoperative quality of life or functioning and less pain also have better postoperative outcomes but that they improve less. Furthermore, women and patients with a higher BMI had more postoperative pain and less improvement after both TKA and THA. Higher age and higher BMI was associated with lower postoperative HRQoL and functioning and more pain after a THA. However, preoperative quality of life, functioning and pain seem to be most consistently associated with outcomes after both TKA and THA.

Our results regarding the effect of preoperative status on outcomes are consistent with other studies that also found that patients with worse preoperative functioning had greater improvements [18–21], but did not achieve the postoperative level of those with higher preoperative functioning [22–28]. Contrary, other studies showed opposite results regarding the direction and size of the effect of age, gender and BMI. Santaguida et al. [29] performed a systematic review about patient characteristics affecting the prognosis after TKA/THA and concluded that an older age is
related to worse functioning, but that age and sex do not influence postoperative pain level. We found that women had more pain after a TKA (4 points on a 100 point scale) and THA (2 points on a 100 point scale), even though this may not be a clinically relevant difference [30]. For TKA no association with age or gender and functioning was found. In addition, a previous review about prognostic determinants in THA reported that preoperative functioning was most consistently associated with better outcomes [13]. In addition, another systematic review on preoperative predictors on outcomes in THA [31] concluded that only patients’ poor preoperative functioning affects the outcome after THA. This was also found for patients with a TKA [32, 33]. Consistent with our finding, Lingard et al. [33] found that patients with severe pain had worse outcomes after a TKA. Other studies also identified other determinants, such as radiological scores, severity of inflammations or comorbidities. A disadvantage of using multiple cohorts with different protocols for data acquisition was that we could not include these determinants. The linear mixed model had to be the equal for each cohort, so that regression coefficients in each cohort have the same meaning. Thus the prognostic determinants found in this present study are not exhaustive; there may be other determinants that have an additional effect on the outcome.

The effect of different preoperative determinants on the postoperative outcomes after TKA and THA may seem to be small on itself, but if taken together they may add up to a clinically relevant effect. However, the scenarios should be interpreted with care, because these are hypothetical examples based on observational data and cannot be interpreted causally. The overall effects of the virtual scenarios which were calculated as examples vary between 1.2 and 6.5 points better postoperative
outcomes and between 1.6 and 9 points worse postoperative outcomes. These scenarios provide more insights how small differences may add up or cancel each other out. This probably explains why most effects do not reach a clinically significant difference. Usually a 10% difference (i.e. 10 points on a 0–100 scale [30]) is considered as clinically relevant, but is a 10% difference the right criterion? Postoperative TKA/THA scores increases on average by 20–40 points on a 0–100 scale (results not shown) compared to preoperative scores regardless of the preoperative status. Thus is it realistic to use a difference of 10 points to define whether it is clinically relevant to operate now or wait, based on differences in preoperative determinants?

It is important to realize that the effects found in our study are not only the effect of the surgery, but also the effect of regression to the mean (RTM). RTM occurs because values are observed with random error, such as random fluctuations in a subject [34]. This means that patients with low preoperative scores are more likely to have higher scores during the next measurement and that patients with high preoperative scores are more likely to have lower scores during the next measurement, even without surgery. This results on average in a larger “improvement” for patients with lower preoperative scores compared to patients with higher baseline scores. Although different methods have been proposed to estimate the size of the RTM effect, but no solution is available to distinguish the real change due to surgery from the change due to RTM. Furthermore, we had to standardize different questionnaires measuring the same domain. Ideally, a minimal dataset should be composed, so that is more easily comparable without the need of standardization since standardized regression coefficients are more difficult to interpret [35]. A strength of our study is that we pooled existing cohort studies. Most of these studies collected a comprehensive set of questionnaires. Although national arthroplasty registries are established, these registries differ from clinical studies. Most registries focus on long-term data collection and therefore focus on minimal data sets and collect patient and operative information, but not all registries collect patient-reported outcomes [36]. If registries collect patient-reported outcomes such as HRQoL, function or pain most often short questionnaires are used, with only 1 or 2 questions covering the domain e.g. VAS-scale for pain or the EQ-5D to measure HRQoL. Most of the in our study included cohort studies used more comprehensive questionnaires with the ability to measure each outcome more reliable. On the other hand using questionnaires with composite scores has some weaknesses. Different patients may have very different domain scores but these may still result in the same composite score. In our study we therefore used domain scores of different questionnaires (functioning and pain) besides the overall HRQoL composite score, which may reduce this problem. Another potential problem is that there may be cultural differences between countries in how questionnaires are answered, but this would only influence our results if these cultural differences would affect e.g. elderly patients differently than younger patients thereby resulting in a different relationship of age with the outcomes.

**Conclusion**

The information regarding the combined effects of preoperative determinants on postoperative outcomes will support orthopaedic surgeons to estimate differences in outcome after a joint replacement for specific patient groups, i.e. poorer outcomes for patients with a worse preoperative status, but with greater
postoperative improvement compared to patients with higher preoperative scores. In addition, preoperative status may decline during a long surgical delay period and thereby lead to worse postoperative outcomes if no other non-surgical treatments are started. On the other hand, it may sometimes be better to first optimize the patient’s preoperative condition or to reduce for example their BMI. The present study may support orthopaedic surgeons in their decision making by giving an estimate of the magnitude of the effect for different scenarios. Future studies should combine the results of our study with observational cohort studies among OA patients who did not have surgery yet, specific survival data from medical literature and the effects on survival of the artificial joint to assess optimal timing of surgery. This is needed to assess the long-term impact for the patient of the decision to perform surgery at a certain preoperative state of specific patient groups.

**Abbreviations**

ADL: Activities of Daily Living; BMI: Body Mass Index; CI: Confidence interval; EQ-5D: EuroQoL 5 Dimensions; HOOS: Hip disability and Osteoarthritis Outcome Score; HRQoL: Health Related Quality of Life; KOOS: Knee disability and Osteoarthritis Outcome Score; LMM: Linear Mixed Models; MoM: Metal-on-metal; OA: Osteoarthritis; OHS: Oxford Hip Score; OKS: Oxford Knee Score; PF: Physical Function; SF-12: 12-item short form health survey; SF-36: 36-item short form health survey; THA: Total Hip Arthroplasty; TKA: Total Knee Arthroplasty; VAS: Visual Analogue Scale; WOMAC: Western Ontario & McMaster Universities Osteoarthritis Index

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**Table 3 Hypothetical examples of combined data within scenarios**

| Arthroplasty | Assessed outcome | Effect of age | Effect of BMI | Effect of preoperative status | Total effect on postoperative outcome (points)* |
|-------------|-----------------|--------------|--------------|-------------------------------|----------------------------------|
| TKA         | HRQoL           | 0            | 0            | 5*−0.51                       | 2.6                              |
|             | Functioning     | 0            | 0            | 5*−0.31                       | 1.6                              |
|             | Pain            | 0            | 5*−0.47      | 5*−0.18                       | 3.3                              |
| THA         | HRQoL           | 5*−0.08      | 5*−0.23      | 5*−0.37                       | 2.6                              |
|             | Functioning     | 5*−0.33      | 5*−0.35      | 5*−0.22                       | 1.2                              |
|             | Pain            | 5*−0.18      | 5*−0.36      | 5*−0.15                       | 1.7                              |

**Scenario 1:** A patient loses weight (X points) and increases preoperative status by Y points, this takes Z years

\[ X, Y, Z = 5 \text{ (e.g. in 5 years BMI decreases from 30 to 25, KOOS HRQoL/ functioning/ pain increases from 35 to 40)} \]

TKA HRQoL 0 0 5*−0.51 2.6
Functioning 0 0 5*−0.31 1.6
Pain 0 5*−0.47 5*−0.18 3.3

THA HRQoL 5*−0.08 5*−0.23 5*−0.37 2.6
Functioning 5*−0.33 5*−0.35 5*−0.22 1.2
Pain 5*−0.18 5*−0.36 5*−0.15 1.7

**Scenario 2:** A patient gains weight (X points) and decreases preoperative status by Y points, this takes Z years

\[ X, Y, Z = 5 \text{ (e.g. in 5 years BMI increases from 25 to 30, KOOS HRQoL/ functioning/ pain decreases from 40 to 35)} \]

TKA HRQoL 0 0 5*−0.51 −2.6
Functioning 0 0 5*−0.31 −1.6
Pain 0 5*−0.47 5*−0.18 −3.3

THA HRQoL 5*−0.08 5*−0.23 5*−0.37 −3.4
Functioning 5*−0.33 5*−0.35 5*−0.22 −4.5
Pain 5*−0.18 5*−0.36 5*−0.15 −3.5

**Scenario 2:** A patient gains weight (X points) and increases preoperative status by Y points, this takes Z years

\[ X, Y, Z = 10 \text{ (e.g. in 10 years BMI decreases from 35 to 25, KOOS HRQoL/ functioning/ pain increases from 35 to 40)} \]

TKA HRQoL 0 0 10*−0.51 5.1
Functioning 0 0 10*−0.31 3.1
Pain 0 10*−0.47 10*−0.18 6.5

THA HRQoL 10*−0.08 10*−0.23 10*−0.37 5.2
Functioning 10*−0.33 10*−0.35 10*−0.22 2.4
Pain 10*−0.18 10*−0.36 10*−0.15 3.3

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*On a 0–100 scale*
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Availability of data and materials
The data that support the findings of this study are available from third parties (see acknowledgements) but restrictions apply to the availability of these data, which were used under license for the current study, and so are not publicly available. Data are however available from the authors upon reasonable request and with permission of the third parties (see acknowledgements).

Authors’ contributions
SH takes responsibility for the integrity of the data and the accuracy of the data analysis. Study conception and design: SH, MG, RN, PM. Acquisition of data: SH, MG, RN, PM. Cleaning of the datasets: SH and MG. Analysis: SH performed the analyses and TS gave statistical advices. Interpretation of data: SH, MG, TS, RN, PM. SH wrote the article and PM supervised writing of the manuscript. All authors were involved in drafting the article or revising it critically for important intellectual content, and all authors approved the final version.

Ethics approval and consent to participate
The Medical Ethical Committee of the Leiden University Medical Center (CMIE P15.043/SH/sh) confirmed on February 13 2015 that ethical approval for this type of study is not required under the Dutch Medical Research (Human Subjects) Act. The hospitals that supplied anonymous data obtained written informed consent from the study participants.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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References
1. OECD. Hip and knee replacement. In: Health at a glance: Europe 2012; 2012.
2. Otten R, van Roermund PM, Picavet HS. Trends in the number of knee and hip arthroplasties: considerably more knee and hip prostheses due to osteoarthritis in 2030. Ned Tijdschr Geneesk. 2010;154:A1534.
3. van Steenbergen LN, Denissen GA, Spooren A, van Rooden SM, van Oosterhout FJ, Morrenhof JW, Nelissen RG. More than 95% completeness of reported procedures in the population-based Dutch arthroplasty register. Acta Orthop. 2015;86(4):498–505.
4. Spooren A, van Steenbergen LN, Denissen GAW, van der Togt CR, van Rooden SM, LROI Report 2013 Insight into quality & safety. Den Bosch: Dutch Arthroplasty Register (LROI); 2013.
5. Labek G, Thaler M, Janda W, Agreiter M, Stockl B. Revision rates after total hip and knee replacement: cumulative results from worldwide joint register datasets. J Bone Joint Surg Br. 2011;93(3):293–300.
6. Bennell KL, Hunter DJ, Hinnan RS. Management of osteoarthritis of the knee. BMJ. 2012;345:e4934.
7. Dieppe P, Judge A, Williams S, Ikvueke I, Guenther KP, Floreen M, Huber J, Ingvanson T, Lernam Hy, Lohmander LS, et al. Variations in the pre-operative status of patients coming to primary hip replacement for osteoarthritis in European orthopaedic centres. BMC Musculoskel Disord. 2009;10:19.
8. Ackerman IN, Diepepe PA, March LM, Roos EM, Nilsdotter AK, Brown GC, Sloan KE, Osborne RH. Variation in age and physical status prior to total knee and hip replacement surgery: a comparison of centers in Australia and Europe. Arthritis Rheum. 2009;61(2):166–73.
9. Nilsdotter AK, Toksvig-Larsen S, Roos EM. Knee arthroplasty: are patients’ expectations fulfilled? A prospective study of pain and function in 102 patients with 5-year follow-up. Acta Orthop. 2009;80(1):55–61.
10. Anderson KG, Watson RJ, Tsai D, Stulberg SD, Chang RW. Functional outcome and patient satisfaction in total knee total hip patients over the age of 75. J Arthroplast. 1996;11(7):831–40.
11. Baker PN, van der Meulen JH, Lewsey J, Gregg PJ. The role of pain and function in determining patient satisfaction after total knee replacement. Data from the National Joint Registry for England and Wales. J Bone Joint Surg Br. 2007;89(7):789–900.
12. Gandhi R, Davey JR, Mohomed NN. Predicting patient dissatisfaction following joint replacement surgery. J Rheumatol. 2008;35(12):2415–8.
13. Hofstede SN, Gademam MG, Vliet Vlieland TP, Nelissen RG, Marang-van de Meehen PJ. Preoperative predictors for outcomes after total hip replacement in patients with osteoarthritis: a systematic review. BMC Musculoskelet Disord. 2016;17(1):212.
14. Yang W, Xie YM. How to read and understand registries for evaluating patient outcomes: a User’s guide (2nd edition). China J Chin Materia Medica. 2013;38(18):2958–62.
15. Harris KK, Price AJ, Beard DJ, Fitzpatrick R, Jenkinson C, Dawson J. Can pain and function be distinguished in the Oxford hip score in a meaningful way?: An exploratory and confirmatory factor analysis. Bone Joint Res. 2014;3(11):305–9.
16. Harris K, Dawson J, Dell H, Field RE, Murray DW, Fitzpatrick R, Jenkinson C, Price AJ, Beard DJ. Can pain and function be distinguished in the Oxford knee score in a meaningful way? An exploratory and confirmatory factor analysis. Qual Life Res. 2013;22(9):2561–8.
17. Higgins JP, Thompson SG, Deeks JJ, Altman DG. Measuring inconsistency in meta-analyses. BMJ. 2003;327(7414):557–60.
18. Clement ND, MacDonald D, Howie CR, Biant LC. The outcome of primary total hip and knee arthroplasty in patients aged 80 years or more. J Bone Joint Surg Br. 2011;93(9):1265–70.
19. Cuschniriahn J, Coggon D, Reading I, Croft P, Byng P, Cox K, Dieppe P, Cooper C. Long-term outcome following total hip arthroplasty: a controlled longitudinal study. Arthritis Rheum. 2007;57(8):1375–80.
20. Judge A, Jawad MK, Arden NK, Cuschniriahn J, Reading I, Croft P, Dieppe P, Cooper C. Clinical tool to identify patients who are most likely to achieve long-term improvement in physical function after total hip arthroplasty. Arthritis Care Res (Hoboken ). 2012;64(6):881–9.
21. Judge A, Cooper C, Arden NK, Williams S, Hobbs N, Dixon D, Gunther KP, Dreinhofer K, Dieppe PA. Pre-operative expectation predicts 12-month post-operative outcome among patients undergoing primary
total hip replacement in European orthopaedic centres. Osteoarthr Cartil. 2011;19(6):659–67.

22. Fortin PR, Penrod JR, Clarke AE, St-Pierre Y, Joseph L, Belisle P, Liang MH, Ferland D, Phillips CB, Mahomed N, et al. Timing of total joint replacement affects clinical outcomes among patients with osteoarthritis of the hip or knee. Arthritis Rheum. 2002;46(12):3327–30.

23. Nilsdotter AK, Lohmander LS. Age and waiting time as predictors of outcome after total hip replacement for osteoarthritis. Rheumatology (Oxford). 2002;41(11):1261–7.

24. Johansson HR, Bergschmidt P, Skripitz R, Finze S, Bader R, Mittelmeier W. Impact of preoperative function on early postoperative outcome after total hip arthroplasty. J Orthop Surg (Hong Kong). 2010;18(1):6–10.

25. Heiberg KE, Bieland A, Bruun-Olsen V, Mengshoel AM. Recovery and prediction of physical functioning outcomes during the first year after total hip arthroplasty. Arch Phys Med Rehabil. 2013;94(7):1352–9.

26. Kennedy DM, Stratford PW, Robarts S, Gillish JD. Using outcome measure results to facilitate clinical decisions the first year after total hip arthroplasty. J Orthop Sports Phys Ther. 2011;41(4):232–9.

27. Roder C, Staub L, Eggli S, Dietrich D, Busato A, Muller U. Influence of preoperative functional status on outcome after total hip arthroplasty. J Bone Joint Surg Am. 2007;89(1):11–7.

28. Judge A, Arden NK, Batra RN, Thomas G, Beard D, Javaid MK, Cooper C, Murray D, group EPOSE. The association of patient characteristics and surgical variables on symptoms of pain and function over 5 years following primary hip-replacement surgery: a prospective cohort study. BMI Open. 2013;3(3):2013.

29. Santaguida PL, Hawker GA, Hudak PL, Glazier R, Mahomed NN, Keder HJ, Coyte PC, Wright JG. Patient characteristics affecting the prognosis of total hip and knee joint arthroplasty: a systematic review. Can J Surg. 2008;51(6):428–36.

30. Pijls BG, Dekkers OM, Middeldorp S, Valstar ER, van der Heide HJ, Van der Linden-Van der Zwaag HM, Nelissen RG. AQUILA: assessment of quality in lower limb arthroplasty. An expert Delphi consensus for total knee and total hip arthroplasty. BMC Musculoskeletal Disord. 2011;12:173.

31. Montin L, Leino-Kilpi H, Suominen T, Lepisto J. A systematic review of empirical studies between 1966 and 2005 of patient outcomes of total hip arthroplasty and related factors. J Clin Nurs. 2008;17(1):40–5.

32. Fortin PR, Penrod JR, Clarke AE, Joseph L, Liang MH, Tanzer M, Ferland D, Phillips CB, Partridge AJ, Belisle P, Fossel AH, et al. Outcomes of total hip and knee replacement: preoperative functional status predicts outcomes at six months after surgery. Arthritis Rheum. 1999;42(8):1722–8.

33. Lingard EA, Katz RN, Wright EA, Sledge CB. Predicting the outcome of total knee arthroplasty. J Bone Joint Surg Am. 2004;86-A(10):2179–86.

34. Barnett AG, van der Pols JC, Dobson AJ. Regression to the mean: what it is and how to deal with it. Int J Epidemiol. 2005;34(1):215–20.

35. Greenland S, Schlesselman JJ, Criqui MH. The fallacy of employing standardized regression coefficients and correlations as measures of effect. Am J Epidemiol. 1986;123(2):203–8.

36. Kolling C, Simmen BR, Labek L, Goldhahn J. Key factors for a successful National Arthroplasty Register. J Bone Joint Surg Br Vol. 2007;89(12):1567–73.