The Effect of Perioperative Vitamin D Levels on the Functional, Patient-Related Outcome Measures and the Risk of Infection Following Hip and Knee Arthroplasty: A Systematic Review

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Introduction: The aim of this study was to evaluate the effect of perioperative vitamin D levels in terms of functional results, patient-related outcome measures (PROMs) and infection risk after hip or knee replacement.

Materials and Methods: A systematic search in PubMed, Cochrane library, ScienceDirect and ClinicalTrials.gov was conducted according to the PRISMA guidelines from inception to January 2020.

Results: Eighteen studies with more than 8000 knee and 1500 hip joint arthroplasties were included. The mean follow-up ranged from 6 weeks to 1 year and mean patients’ age from 59.4 to 76 years. Hypovitaminosis was diagnosed in 26.7% of cases. Most studies did not find significant differences in pre- and postoperative functional results, PROMs and length of hospital stay between hypovitaminosis and euvitaminosis groups. Deficient patients may be at higher risk of postoperative joint stiffness. Patients suffering from hip and knee periprosthetic joint infection seem to have lower vitamin D levels compared to those with aseptic loosening of implants.

Conclusion: The necessity of pre-operative correction of vitamin D levels to achieve better functional results and minimize the risk of infection following hip and knee arthroplasty remains inconclusive. Extend of exposure to low vitamin D levels and comparison between outliers needs further evaluation.

Keywords: vitamin D, hip arthroplasty, THA, knee arthroplasty, TKA

Introduction

Vitamin D deficiency is the most common nutritional deficiency worldwide. The incidence of hypovitaminosis D is rising globally, affecting approximately one billion people. Staying indoors for prolonged periods, wearing clothes that cover the skin, and using sunscreen are some of the primary causes of this increasing epidemic, the repercussions of which remain obscure.

The metabolism of vitamin D is well described. Vitamin’s D main role is the regulation of calcium metabolism by controlling its absorption by the small intestine. It promotes bone resorption by stimulating the activity of osteoclasts. Vitamin D plays a determining role in muscle function through regulating the deposition of phosphorus in myocytes, in the metabolism of creatine phosphate as well as in muscle cells’ response to insulin and muscle action. As a result, the role of vitamin D in bone resistance is crucial, promoting the stimulation of bone turnover and muscle function.
Inclusion and Exclusion Criteria

The search was limited to articles that were published in the English language. The inclusion criteria included randomised controlled trials, retrospective and prospective cohort studies as well as case-control studies that a) reported mean pre- and postoperative vitamin D values in patients undergoing knee or hip arthroplasty b) reported pre- and postoperative functional scores and PROMs for these patients and c) compared functional outcomes and PROMs between vitamin D deficient and sufficient patient groups. Case reports, reviews or conference presentations were excluded from the study.

Outcome Measures

The primary outcomes were the effect of vitamin D levels in the functional outcomes and PROMs following knee or hip arthroplasty. Secondary outcomes were comparison of a) survivorship b) length of stay (LOS) and c) complications, namely infection, between vitamin D sufficient and deficient patients undergoing knee or hip arthroplasty.

Study Selection and Data Extraction

Two authors (PK and DK) independently searched the electronic databases for potentially relevant studies. After removing duplicates, the titles and abstracts of the retrieved records were screened for eligibility. In case of disagreement, a third investigator (ET) independently evaluated the study. Consequently, any discrepancies about the study selection process were solved by discussion and eligibility was defined by agreement with all authors. The various reasons for ineligible studies are presented in the flowchart of Figure 1.

Data extraction was independently performed by two investigators (PK and EK) in duplicate for each eligible study. Data collected included the year of publication, study design, patients’ demographics, methodological features of the trial, PROMs, functional results, LOS and infection rate. Any potential disagreement was resolved by discussion between the reviewers. Where necessary, a third investigator (ET) was contacted, and independently extracted data before consensus was reached.

Data Analysis and Methodological Quality Assessment

We performed a narrative synthesis of the literature to evaluate the included data. The quality of the studies was assessed using the Newcastle – Ottawa Scale (NOS). The NOS assesses the selection and comparability of study
groups and ascertainment of the exposure-outcome using a “star system” where studies can be granted up to nine stars.

Results
Search Results
The initial search of the literature retrieved 3828 available articles. After removing duplicates and irrelevant studies by screening the title and abstract, 34 studies remained for review. Sixteen records were excluded because they were either incomplete studies, review studies or considered irrelevant after reading the full text. In total, 18 studies were included in the systematic review. Details of the study screening and selection are displayed in the flowchart of Figure 1.

Methodological Quality Assessment
Five studies were of high methodological quality, 11,14,15,18,21 12 were of moderate quality 12,13,16,17,19,20,22-25,27,28 and 1 of low quality.26

The results of the quality assessment are demonstrated in Supplementary Table 2.

Study Characteristics
The included studies were published between 2008 and 2018. There were ten retrospective case series, six prospective cohort studies, one retrospective and one prospective case-control study. The sample size ranged from 42 to 6593, with a mean follow-up from six weeks to one year. We extracted data from more than 8000 knee and 1500 hip
replacements. The mean reported age of the patients ranged from 59.4 to 76 years. Seventeen studies reported on the prevalence of hypovitaminosis D; 11–23,25–28 2598 out of 9720 (26.7%) patients were diagnosed with low vitamin D levels. The detailed descriptive characteristics of the included studies are presented in Table 1.

Table 1 Descriptive Characteristics of the Included Studies

| Author (Year) | Design | Type | No of Patients | Male (%) | Mean Age of Involved Groups | Follow-Up | Type of Surgery | Vitamin D Sufficiency Cut Off Level | Patients with Hypovitaminosis (%) |
|---------------|--------|------|----------------|----------|----------------------------|-----------|----------------|------------------------------------|-----------------------------------|
| Hedge et al17 | R      | CS   | 6593           | 1599 (24.2) | I: 65 S:70                  | 12 m      | KA             | ≥20 ng/mL                          | 868 (13.2)                        |
| Janssen et al15 | R   | CS   | 138            | 57 (41.3)   | I: 69.9 S: 72.8             | 6m        | KA             | >40 ng/mL                          | 33 (23.9)                         |
| Allain et al16 | P    | C    | 92             | 38 (41.3)   | W: 75.1 M:73.6              | 6 m       | KA (9 RKA)     | ≥50 ng/mL                          | 33 (36)                           |
| Maniar et al12 | R    | CS   | 120            | 23 (19.1)   | I: 67 S:69                  | 3m        | KA             | ≥30 ng/mL                          | 64 (53.3)                         |
| Lee et al13    | P     | C    | 191            | 38 (19.8)   | I:68 s:67                   | 3m        | KA             | >50 nmol/L                         | 84 (44)                           |
| Jansen et al22 | R    | CS   | 139            | 58 (42)     | 71.4                        | 6m        | KA             | >40 ng/mL                          | 23 (22.5)                         |
| Shin et al11   | P     | C    | 87             | NA (10.3)   | I:70.7 S:72.4               | 3m        | KA             | ≥12 ng/mL                          | 44 (50.5)                         |
| Maier et al23  | R     | CS   | 1083           | 516 (47.5)  | 76                          | NA        | HA, KA         | ≥30 ng/mL                          | 681 (62.8)                        |
| Traven et al27 | R    | CC   | 126            | 56 (44.4)   | 65.4                        | 3m        | 64 RKA, 62 RHA | ≥30 ng/mL                          | 69 (54.7)                         |
| Maier et al24  | R     | CS   | 190            | 84 (44.2)   | PA:65, Pji:68, AL:68,4      | NA        | HA, KA         | ≥30 ng/mL                          | 129 (68)                          |
| Zajonz et al16 | P    | CC   | 240            | 113 (47)    | PA:71, Pji:74, AL:70        | NA        | HA, KA         | >30 ng/mL                          | 203 (84.5)                        |
| Visser et al18 | P    | C    | 87             | 21 (24.1)   | 74                          | 6w        | HA             | ≥75ng/ml                           | 55 (63.2)                         |
| Signori et al26 | R   | CS   | 42             | 9 (21.4)    | Pji:63.5, AL:59.4           | NA        | 21 RHA, 21 RKA | >20mg/dl                           | NA                               |
| De Cuhna et al28 | P  | C    | 93             | 48 (51.6)   | 59.7                        | 3m        | HA             | NA                                 | 77 (82.8)                         |
| Lavernia et al19 | R | CS   | 60             | 12 (20)     | 70                          | 11m       | HA             | 20 or 30 ng/mL                      | 39 (65)                           |
| Unnanuntana et al21 | P | C    | 219            | 93 (42.5)   | 1.65 s:69.3                 | 6w        | HA             | >30 ng/mL                          | 102 (46.6)                        |
| Unnanuntana et al22 | R | CS   | 200            | 88 (44)     | 66.5                        | NA        | HA             | 32 ng/mL                           | 79 (39.5)                         |
| Naswab et al20 | R    | CS   | 62             | 23 (37)     | 71                          | 6m        | HA             | 40nmol/l                           | 15 (24.2)                         |

Notes: *Values are given as raw numbers with percentages in parentheses.

Abbreviations: R, retrospective; P, prospective; HA, hip arthroplasty; KA, knee arthroplasty; SA, shoulder arthroplasty; RHA, revision hip arthroplasty; RKA, revision knee arthroplasty; NA, non available; C, cohort; CC, case-control; CS, case series; m, months; w, weeks; S, sufficiency group; I, insufficiency group; Pji, periprosthetic joint infection; AL, aseptic loosening; PA, primary arthroplasty; W, women; M, men.
Functional Outcomes in Knee Replacement

Seven studies reported functional outcomes and PROMs according to vitamin D status on 7360 knee arthroplasties (Table 2). The most commonly used PROMs were Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC) and Knee Society Score (KSS). The majority of studies showed no significant differences in preoperative WOMAC score between deficient and sufficient patients. Only one study demonstrated a higher preoperative WOMAC score in vitamin D deficient patients. Mean preoperative WOMAC score, as evaluated in four studies, ranging from 41.2 to 79.6 in hypovitaminosis group and 42.3 to 75.8 in euvitaminosis group.

Postoperatively, three studies reported insignificant differences in overall WOMAC score between vitamin D deficient and sufficient patients. However, one study found that WOMAC stiffness subset had significantly less postoperative improvement in deficient patients. The series with the longest follow up, revealed significantly worse functional results in deficient patients. Mean postoperative WOMAC score ranged from 17.6 to 62.2 in the hypovitaminosis group and 15.8 to 52.5 in the euvitaminosis group.

KSS was evaluated in three studies. Mean preoperative KSS score ranged from 31.5 to 56.2 in vitamin D deficient and 37.1 to 58.9 in vitamin D sufficient patients. Two studies reported comparable preoperative outcomes in both deficient

Table 2 Effect of Vitamin D in Functional Outcomes in Patients Undergoing Knee Arthroplasty

| Author          | Mean Vitamin D Level (ng/mL) | Scores Evaluated | Groups Compared | Main Clinical Findings                                                                 |
|-----------------|------------------------------|------------------|-----------------|----------------------------------------------------------------------------------------|
| Hedge et al17   | NA                           | NA               | D <20 ng/mL     | Deficient patients: higher incidence of postoperative stiffness requiring manipulation under anesthesia (OR, 1.69; 95% CI, 1.39–2.04) |
| Janssen et al15 | D:32, S:65                   | WOMAC            | D <40 ng/mL     | Deficient patients: worse functional outcome after eight years (p<0.01)                |
| Allain et al14  | NA                           | WOMAC, ABC       | D <30 ng/mL     | No differences in preoperative and postoperative total scores between groups           |
|                 |                              |                  | I: 30–49 ng/mL  | Deficient patients: significantly less postoperative improvement in WOMAC stiffness score |
|                 |                              |                  | S ≥50 ng/mL     |                                                                                         |
| Maniar et al12  | NA                           | SF-12, KSS, WOMAC| D <30 ng/mL     | No differences in preoperative total scores except WOMAC                               |
|                 |                              |                  | S ≥30 ng/mL     | (p = 0.04)                                                                              |
|                 |                              |                  |                 | No differences in postoperative total scores between groups                             |
| Lee et al13     | NA                           | WOMAC, QoR, EQ-5D| I< 50 nmol/L (Severe <12.5, Moderate 12.5–29, Mild 30–49) S > 50 nmol/L | Moderate-to-severe hypovitaminosis D group: lower preoperative WOMAC function score     |
|                 |                              |                  |                 | No differences in postoperative total scores between groups                             |
| Jansen et al25  | 58.9                         | KSS              | D <40 ng/mL     | Deficient patients: significantly lower mean preoperative KSS                          |
|                 |                              |                  | S >40 ng/mL     | Deficient patients: not statistically significant lower mean postoperative KSS at six months |
| Shin et al11    | D:9.1, S:18.3                | KSS, AST, SMT, STS, TUGT | D <12 ng/mL | No differences in preoperative total scores between groups                               |
|                 |                              |                  | S ≥12 ng/mL     | No differences in postoperative mean time of STS and TUGT                              |
|                 |                              |                  |                 | Deficient patients: significantly lower post-operative functional KSS (p=0.045)         |
|                 |                              |                  |                 | Deficient patients: significantly longer mean time taken for post-operative AST (p = 0.033) and SMT (p = 0.012) |

Notes: *Mean vitamin D level is given for the whole group or for separate groups accordingly.
Abbreviations: NA, non available; D, deficient; I, insufficient; S, sufficient; OR, odds ratio; CI, 95% confidence interval; WOMAC, Western Ontario and McMaster Universities Osteoarthritis Index; SF-12, 12-item Short Form Survey; KSS, Knee Society Score; AST, alternative step test; TUGT, timed up and go test; SMT, six meter walk test; STS, sit to stand test; ABC, Activities-specific Balance Confidence Scale; QoR, Quality of Recovery; EQ-5D, European Quality of Life-5 Dimensions; 25-(OH)D, 25-hydroxyvitamin D.
and sufficient groups.\textsuperscript{11,12} One study showed significantly lower KSS score in vitamin D deficient patients.\textsuperscript{25}

Postoperatively, there was no significant difference in KSS between groups in two studies.\textsuperscript{12,25} One study found that functional KSS was significantly lesser in the vitamin D deficient group than in the vitamin D non-deficient group postoperatively; clinical KSS score remained comparable in both groups.\textsuperscript{11} Mean postoperative KSS score ranged from 74.6 to 91.9 in the hypovitaminosis group and 80.4 to 91.2 in the euvitaminosis group.\textsuperscript{11,12,25}

Among other functional scores, 12-Item Short Form Survey mental component summary (SFMCs), 12-Item Short Form Survey physical component summary (SFPCs), alternative step test (AST), timed up and go test (TUGT), six-meter walk test (SMT), sit to stand test (STS) and Activities-specific Balance Confidence Scale (ABC) did not differ significantly preoperatively between patients with low and normal vitamin D levels. Postoperatively, SFPCs, SFMCS, ABC, STS, TUGT remained comparable between the two groups. However, the mean time taken for postoperative AST and SMT (8.8 vs 7.7 s, \(p = 0.012\)) was significantly longer in the vitamin D-deficient group than in the vitamin D non-deficient group.\textsuperscript{11,12,14} Mean pre- and postoperative scores are demonstrated in detail in Supplementary Table 3.

**Functional Outcomes in Hip Replacement**

Six studies reported functional outcomes and PROMs according to vitamin D status on 721 hip replacements (Table 3).\textsuperscript{18–22,28} Preoperative WOMAC score showed no significant difference between sufficient and deficient patients.\textsuperscript{18,19,21} Mean preoperative WOMAC score ranged from 40.4 to 59 in the deficient and from 43.5 to 54 in the sufficient group. Postoperatively, the mean WOMAC score remained comparable between the two groups.\textsuperscript{19,21} One study found that an increase in 25-hydroxyvitamin D\(_3\) (25(OH)D\(_3\)) concentration six weeks postoperatively was correlated with improved hip function.\textsuperscript{18}

Harris Hip Score (HHS) was evaluated in two studies.\textsuperscript{19,20} One study found significantly higher preoperative HHS in vitamin D sufficient group.\textsuperscript{20} The result of the second study was dependent on the vitamin D cut off point. When the threshold of hypovitaminosis was set in 20 ng/mL, the authors found comparable pre- and postoperative HHS independent of vitamin D levels. However, using the 30 ng/mL as a cut-off point, deficient patients found to have significantly lower pre- and postoperative HHS than sufficient patients.\textsuperscript{19}

Among the other functional outcomes evaluated, there was no association between serum vitamin D levels and Chair Stand Test (CST), 10 Meter Walking Test (10MWT), in-hospital functional milestones and 36-Item Short Form Survey (SF-36).\textsuperscript{18,21,22} TUGT was recorded in two studies.\textsuperscript{18,21} One study showed a significant improvement in TUGT over time but independent of vitamin D status.\textsuperscript{18} Contrary to these findings, a significant improvement in TUGT only in the low vitamin D group was revealed in another study.\textsuperscript{21} Mean pre- and postoperative scores are demonstrated in detail in Supplementary Table 4.

**LOS**

Five studies reported on the duration of hospitalization according to preoperative vitamin D status.\textsuperscript{13,15,22,26,27} Mean LOS was longer in the hypovitaminosis D group (3.8 to 15.6 days) compared to the euvitaminosis D group (3.6 to 11.3 days). The difference in LOS found to be statistically significant in two studies, with a mean difference between 1 to 4.3 days.\textsuperscript{15,25} (Table 4)

**Survival Rate of the Prosthesis**

Survivorship was recorded in one study with the largest sample.\textsuperscript{17} Out of 6593 total knee arthroplasties (TKA), 868 suffered from hypovitaminosis D in this retrospective cohort study. At first-year follow-up, the incidence of revision arthroplasty for prosthesis explantation was found to be higher among the vitamin D deficient patients (OR, 2.97; 95% CI, 2.04–4.31; \(p=0.001\)). However, this was mainly attributed to periprosthetic joint infection (PJI) as the rates of non-infectious component revision were comparable between the two groups.

**Complications - Risk of Infection**

Six studies evaluated complication rate between vitamin D deficient and sufficient patients.\textsuperscript{16,17,22,24,26,27} They comprised of 7385 joint replacements; 6635 primary knee arthroplasties, 238 primary total hip arthroplasties (THA), 186 revision TKA, 186 revision THA, 109 unclarified primary, and 31 unclarified revision joint arthroplasties. Among the complications reported, PJI was the best described. Two studies reported that patients with low vitamin D levels are more likely to suffer from PJI.\textsuperscript{17,27} One study found no significant differences in complication rates between sufficient and deficient groups.\textsuperscript{22}

Other failures reported were postoperative deep venous thrombosis, myocardial infarction, cerebrovascular accident and postoperative stiffness requiring manipulation under anesthesia, all of which were found to be higher in vitamin D deficient patients as documented by one study only.\textsuperscript{17}
Table 3 Effect of Vitamin D in Functional Outcomes in Patients Undergoing Hip Arthroplasty

| Author            | Mean Vitamin D Level* | Scores                          | Groups Compared | Main Clinical Findings                                                                 |
|-------------------|-----------------------|---------------------------------|-----------------|----------------------------------------------------------------------------------------|
| Visser et al18     | 68.1 nmol/l           | WOMAC, HGS, CST, TUGT, 10-MWT, LAPAQ | D <50 ng/mL I: 50–75 ng/mL S ≥75 ng/mL | No differences in preoperative total scores between groups No differences in postoperative HGS, CST, 10-MWT, WOMAC Insufficient patients: higher postoperative LAPAQ score |
| Cuhna et al28      | 13.8 ng/mL            | Gait Analysis                    | NA              | Baseline Vitamin D levels: weak correlation with change in peak extension (p = 0.017) and peak power generation (p = 0.04) Vitamin D levels: no effect in change of gait pattern |
| Lavernia et al19   | NA                    | WOMAC, HHS, PMA, QWB-7, SF-36   | I <20 or 30 ng/mL S >20 or 30 ng/mL | No differences in preoperative and postoperative total scores between groups when the threshold was set in 20 ng/mL No differences in preoperative and postoperative total scores of WOMAC, QWB-7 and SF-36 between groups when the threshold was set in 30 ng/mL Significant differences in preoperative and postoperative total scores of HHS and Merle d’Aubigne'-Postel score between groups when the threshold was set in 30 ng/mL favoring sufficient group |
| Unmanuntana et al22| D:15.7 ng/mL l: 26.7 ng/mL s:43.8 ng/mL | Transferring in and out of bed, ability to walk with assisting devices and ability to ascend/descent stairs | I <32 ng/mL S>32 ng/mL | No association between vitamin D level and the attainment of hospital functional milestones |
| Nawabi et al20     | 58 nmol/l             | HHS                             | S >40nmol/l D <40nmol/l | Sufficient patients: significantly higher preoperative HHS (p = 0.018) Sufficient patients: non-significantly higher postoperative HHS at 6 months (p = 0.067) Deficient patients: significantly less likely to attain an excellent outcome (p = 0.038) |
| Unmanuntana et al21| NA                    | WOMAC, SF-36, TUGT, Two-min walk test | S>30 ng/mL l<30 ng/mL | No differences in preoperative scores between groups No differences in postoperative WOMAC and SF-36 between groups Deficient patients: significantly improved TUGT postoperatively |

Notes: *Mean vitamin D level is given for the whole group or for separate groups accordingly.

Abbreviations: CST, Chair Stand Test; TUGT, timed up and go test; 10-MWT, 10-meter walking test; NA, non available; WOMAC, Western Ontario and McMaster Universities Osteoarthritis Index; HHS, Harris Hip Score; SF-36, 36-Item Short Form Survey; HGS, hand grip strength; LAPAQ, LASA Physical Activity Questionnaire; QWB-7, Quality of Well-being Scale; 25-(OH)D3, 25-hydroxyvitamin D3; D, deficient; S, sufficient; I, insufficient; PMA, Merle d’Aubigne'-Postel.

Vitamin D Levels in Patients with PJI vs Aseptic Loosening

Four studies compared vitamin D levels of patients following revision joint replacement due to infection or aseptic loosening with conflicting outcomes.16,24,26,27 Two of them concluded that patients undergoing revision for PJI were more likely to have low preoperative vitamin D levels as compared to those for aseptic loosening.26,27 One study revealed lower vitamin D levels in patients with aseptic loosening compared to those undergoing revision for PJI.24 Another study found no significant differences as to vitamin D status in the 25(OH)D3 levels between these two groups.16

Vitamin D Levels in Patients with PJI vs Primary Arthroplasty

Two studies compared vitamin D status in patients scheduled for primary or revision joint arthroplasty due to PJI.16,26 One of them reported non-significant differences (p=0.846), while the other one found a significantly lower vitamin D status in the PJI group (p<0.001).16,26
Table 4 Length of Hospital Stay Based on the Vitamin D Level of Groups of Patients

| Author              | Deficient (Range) | Insufficient | Sufficient (Range) | p    |
|---------------------|-------------------|--------------|-------------------|------|
| Maier et al.        | 15.6 (7.2)        | NA           | 11.3 (7.9)        | 0.014|
| Trenen et al.       | 5                 | NA           | 3.6               | 0.097|
| Lee et al.          | 10 (8–15)         | 9            | 10                | 0.76 |
| Janssen et al.      | 7 (2–28)          | NA           | 7 (2–19)          | 0.03 |
| Unnanuntana et al.  | 4                 | 3.8          | 3.7               | 0.724|

Notes: *Values are given as mean with the standard deviation in parentheses.

Vitamin D Levels in Patients with Aseptic Loosening vs Primary Arthroplasty
Two studies reported no significant differences in vitamin D status between patients undergoing primary or revision joint replacement due to aseptic loosening.16,26 (Table 5)

Discussion
In this systematic review, we aimed to configure the potential role of vitamin D levels in lower limb arthroplasty in terms of PROMs, implant survivorship, LOS, and complications. The majority of studies did not report any differences in postoperative functional scores based on vitamin D status.13,14,21,22,25,28 It seems that patients with lower Vitamin D levels may be at higher risk of joint stiffness postoperatively.17 Vitamin D may also be a protective factor against PJI.26

The existing literature shows a high prevalence of vitamin D deficiency in patients with osteoarthritis undergoing arthroplasty.25 The incidence of hypovitaminosis D in patients undergoing TKA ranges from 36.4% up to 76.2%.12–14,21,22,25,28 Similar rates were reported in patients scheduled for THA, ranging from 22% up to 82.8%.19,22,28,35 In our systematic review, we found 26.7% of patients undergoing joint replacements suffering from low vitamin D levels. This, however, might be an incidental finding attributed to factors, such as advanced age, osteoporosis and sun restriction.

The role of vitamin D in the pathogenesis of osteoarthritis could be explored further through analysis of preoperative data of patients undergoing TJA. Most studies reported no connection between hypovitaminosis D and preoperative physical function of the knee or hip joint.11,13–15,18,19 These findings contrast with other studies reporting that deficient patients scheduled for TKA had worse preoperative WOMAC score.12 The similar finding was reported in patients who had undergone THA as preoperative HHS was higher in sufficient patients scheduled for THA.20 However, these are only point measurements and potentially, underestimate the disease course and the effect of hypovitaminosis D on the disease progress.

On the other hand, the postoperative progress of patients undergoing THA and TKA is dependent on a number of factors, including the patients’ age, the types of prostheses, and the disease duration.36 The potential for a positive correlation between vitamin D levels and improved postoperative results is open to research and might, in fact, actually be given the possible positive effect of vitamin D on the function of muscular system and integration of prostheses.

Postoperatively, the outcomes regarding the knee function after TKA remain inconclusive. More specifically, several studies showed comparable postoperative clinical results between vitamin D sufficient and deficient groups.12,14 Contrary to this finding, many studies showed an association between low vitamin D levels and worse postoperative clinical outcomes after TKA, such as KSS & AST as well as stiffness subcategory of WOMAC score.11,17 The short follow-up period and the inability to measure any potential vitamin D intake are the main drawbacks of these studies. Moreover, the absence of

Table 5 Comparison of Vitamin D Levels Between Groups of Patients Undergoing Revision for Periprosthetic Joint Infection or Aseptic Loosening or Primary Arthroplasty

| Author       | Number of Patients | Vitamin D Level (ng/mL)* | p-value |
|--------------|--------------------|--------------------------|---------|
|              | PJI                | PA          | AL       | PJI       | PA         | AL       | PJI vs AL | PJI vs PA | PA vs AL |
| Zajonc et al | 80                 | 80          | 80       | 17.9 (8.9)| 16.8 (6.9)| 19.7 (7.9)| 0.771     | <0.001    | NA       |
| Maier et al  | 50                 | 109         | 31       | 13.29 (6.5)| 19.46 (9.4)| 20.5 (9.1)| <0.001    | <0.001    | 0.58     |
| Signori et al| 23                 | NA          | 19       | 18.5 (6.5)| NA        | 13.6 (9.4)| <0.05     | NA        | NA       |

Note: *Values are given as mean with standard deviation in parentheses.

Abbreviations: PJI, periprosthetic joint infection; NA, non available; PA, primary arthroplasty; AL, aseptic loosening.
comparability as regards the preoperative levels of vitamin D, the disease duration, the type of prosthesis, and the surgical methods all reduce the reliability of the results.

As for the postoperative function of the hip joint, the results are equivocal. There seems to be a positive association between vitamin D levels and improved postoperative PROMs. 18,20 A weak association has also been found between vitamin D deficiency and a negative change in peak extension and peak power generation when analyzing the gait patterns after THA.28 Once again, the exclusion of the most severe cases, the short follow-up period and the absence of comparability in regard to prostheses type and surgical methods all affect the reliability of the studies. So further research is needed to determine the long-term effects of vitamin D deficiency in patients undergoing hip arthroplasty.

Unfortunately, there is limited literature evidence on the effect of vitamin D on implant survival and the ways in which the vitamin can prolong prostheses survivorship.17 It might be the case that it is associated with higher osseointegration yet, this primarily concerns uncemented prostheses. A sole study demonstrated that vitamin D deficiency was an independent risk factor for prosthesis explantation following knee arthroplasty.17 However, this was mainly attributed to infectious reasons as the non-infectious component revision rate was found to be comparable between deficient and sufficient groups.17

The role of vitamin D in the prevention of septic conditions is well-documented.37-40 It prevents the overexpression of inflammatory cytokines, such as interleukin-6 and C-reactive protein and is a crucial mediator in the aggregation of leukocytes, local inflammation, and anti-bacterial responses in innate immunity.41 The levels of vitamin D have also been inversely associated with the risk of infectious complications after surgery.42,43 In orthopaedic procedures, the evidence is relatively scarce.16,24,26,27 Several studies found lower vitamin D levels in patients with PJI compared to those with aseptic loosening.26,27 Interestingly, there was one study that reported higher vitamin D levels in patients with PJI.24 However, the absence of measurements of serum vitamin D in the postoperative follow-up negatively impacts upon the quality of these findings.

The main limitation of this systematic review is the different vitamin D reference range reported in the studies. The classification of vitamin D status based on serum 25-hydroxyvitamin has not been unanimously accepted, which complicates the interpretation of the data.44 What is thus, needed is the standardization of the definition of vitamin D insufficiency. Another concern that could be raised is the retrospective design in the majority of the studies. High-quality randomized control trials are necessary to prove the effectiveness of vitamin D on PROMs following lower limb arthroplasty. A third objection is the comparability of the groups. Functional results following TJA are multifactorial. There was no clear indication in the studies that vitamin is connected to some extent to the outcomes.

Conclusions
In conclusion, our study identified a high prevalence of hypovitaminosis in osteoarthritis patients undergoing joint arthroplasty. Although a positive correlation of vitamin D status and postoperative functional outcomes is reported in some studies, the necessity of preoperative correction of vitamin D remains an open question. Moreover, despite the acknowledged role of vitamin D in infectious diseases, it is yet to be clarified whether supplementation of vitamin D has a protective effect in PJI.

Abbreviations
TJA, total joint arthroplasty; PROMs, patient-related outcome measures; PRISMA, Preferred Reporting Items for Systematic Reviews and Meta-Analyses; LOS, length of stay; NOS: Newcastle – Ottawa Scale; WOMAC, Western Ontario and McMaster Universities Osteoarthritis Index; KSS, Knee Society Score; SFPCS, 12-Item Short Form Survey mental component summary; SFMCS, 12-Item Short Form Survey physical component summary; AST, alternative step test; TUGT, timed up and go test; SMT, six-meter walk test; STS, sit to stand test; ABC, Activities-specific Balance Confidence Scale; 25(OH)D3, 25-hydroxyvitamin D3; HHS, Harris Hip Score; CST, Chair Stand Test; 10MWT, 10 Meter Walking Test; SF36, 36-Item Short Form Survey; TKA, total knee arthroplasty; THA, total hip arthroplasty; PJI, periprosthetic joint infection.

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**References**

1. Mithal A, Wahl DA, Bonjour J-P, et al. Global vitamin D status and determinants of hypovitaminosis D. *Osteoporos Int*. 2009;20(11):1807–1820. doi:10.1007/s00198-009-0954-6
2. Neal S, Sykes J, Rigby M, Hess B. A review and clinical summary of vitamin D in regard to bone health and athletic performance. *Phys Sportsmed*. 2015;43(2):161–168. doi:10.1080/00913647.2015.1020248
3. Bikle D. Vitamin D and bone. *Curr Osteoporos Rep*. 2012;10(2):151–159. doi:10.11191/012-0098-z
4. Dawson-Hughes B. Vitamin D and muscle function. *J Steroid Biochem Mol Biol*. 2017;173:313–316. doi:10.1016/j.jsbmb.2017.03.018
5. Tomlinson PB, Joseph C, Angiò M. Effects of vitamin D supplementation on upper and lower body muscle strength levels in healthy individuals. A systematic review with meta-analysis. *J Sci Med Sport*. 2015;18(5):575–580. doi:10.1016/j.jsams.2014.07.022
6. Dhaliwal R, Aloia JF. Effect of vitamin D on falls and physical performance. *Endocrinol Metab Clin North Am*. 2017;46(4):919–933. doi:10.1016/j.ecl.2017.07.004
7. Wicherts IS, van Schoor NM, Boeke AJP, et al. Vitamin D status predicts physical performance and its decline in older persons. *J Clin Endocrinol Metab*. 2007;92(6):2058–2065. doi:10.1210/jc.2006-1525
8. Cao Y, Winzenberg T, Nguo K, Lin J, Jones G, Ding C. Association between serum levels of 25-hydroxyvitamin D and osteoarthritis: a systematic review. *Rheumatology (Oxford)*. 2013;52(7):1323–1334. doi:10.1093/rheumatology/ket132
9. Sanghi D, Mishra A, Sharma AC, et al. Does vitamin D improve osteoarthritis of the knee: a randomized controlled pilot trial. *Clin Orthop Relat Res*. 2013;471(11):3556–3562. doi:10.1007/s11999-013-3201-6
10. Manoy P, Yuktanandana P, Tanavaale A, et al. Vitamin D supplementation improves quality of life and physical performance in osteoarthritis patients. *Nutrients*. 2017;9(8):799. doi:10.3390/nut9080799
11. Shin K-Y, Park KK, Moon S-H, Yang IH, Choi H-J, Lee W-S. Vitamin D deficiency adversely affects early post-operative functional outcomes after total knee arthroplasty. *Knee Surg Sports Traumatol Arthrosoc*. 2017;25(11):3424–3430. doi:10.1007/s00167-016-4209-8
12. Maniar RN, Patil AM, Maniar AR, Gangaraju B, Singh J. Effect of preoperative vitamin D levels on functional performance after total knee arthroplasty. *Clin Orthop Surg*. 2016;8(2):153–156. doi:10.4055/cios.2016.8.2.153
13. Lee A, Chan SKC, Samy W, Chiu CH, Gin T. Effect of hypovitaminosis D on postoperative pain outcomes and short-term health-related quality of life after knee arthroplasty: a cohort study. *Medicine*. 2015;94(42):e1812. doi:10.1097/MD.0000000000001812
14. Allain TJ, Berosford PA, Newman JH, Swinkels A. Vitamin D levels in patients undergoing knee arthroplasty: does vitamin D status affect postoperative outcomes? *E Spor Eur J Clin Nutr Metab*. 2008;3(1):e17–e21. doi:10.1016/j.echmn.2007.10.002
15. Jansen J, Tahmassabi J, Haddad FS. Vitamin D deficiency is associated with longer hospital stay and lower functional outcome after total knee arthroplasty. *Acta Orthop Belg*. 2017;83(4):664–670.
16. Zajonz D, Prager FL, Edel M, et al. The significance of the vitamin D metabolism in the development of periprosthetic infections after THA and TKA: a prospective matched-pair analysis of 240 patients. *Clin Interv Aging*. 2018;13:1429–1435. doi:10.2147/CIA.S171307
17. Hegde V, Arshi A, Wang C, et al. Preoperative vitamin D deficiency is associated with higher postoperative complication rates in total knee arthroplasty. *Orthopedics*. 2018;41(4):e489–e495. doi:10.3928/01474472-20180424-04
18. Visser E, de Roos NM, Oosting E, Endenburg SC, Dronkers JJ. Association between preoperative vitamin D status and short-term physical performance after total hip arthroplasty: a prospective study. *Ann Nutr Metab*. 2018;73(3):252–260. doi:10.1159/000492938
19. Lavermia CJ, Villa JM, Iacobelli DA, Rossi MD. Vitamin D insufficiency in patients with THA: prevalence and effects on outcome. *Clin Orthop Relat Res*. 2014;472(2):681–686. doi:10.1007/s11999-013-3172-7
20. Nawabi DH, ChinKF, Keen RW, Haddad FS. Vitamin D deficiency in patients with osteoarthritis undergoing total hip replacement: a cause for concern? *J Bone Joint Surg Br*. 2010;92(4):496–499. doi:10.1002/jb/jvs.23353
21. Unnamantana A, Saleh A, Nguyen JT, et al. Low vitamin D status does not adversely affect short-term functional outcome after total hip arthroplasty. *J Arthroplasty*. 2013;28(2):315–322.e2. doi:10.1016/j.arth.2012.04.027
22. Unnamantana A, Rebollode BJ, Gladnick BP, et al. Does vitamin D status affect the attainment of in-hospital functional milestones after total hip arthroplasty? *J Arthroplasty*. 2012;27(3):482–489. doi:10.1016/j.arth.2011.05.023
23. Maier GS, Maus U, Lazovic D, Horas K, Roth KE, Kurfth A. Is there an association between low serum 25-OH-D levels and the length of hospital stay in orthopaedic patients after arthroplasty? *J Orthop Traumatol*. 2016;17(4):297–302. doi:10.1007/s11991-016-0414-y
24. Signori V, Romano CL, De Vecchi E, Mattina R, Drago L. May osteoarticular infections be influenced by vitamin D status? An observational study on selected patients. *BMC Musculoskelet Disord*. 2015;16:183. doi:10.1186/s12891-015-0648-5
25. Jansen JA, Haddad FS. High prevalence of vitamin D deficiency in elderly patients with advanced osteoarthrosis scheduled for total knee replacement associated with poorer preoperative functional state. *Ann R Coll Surg Engl*. 2013;95(8):569–572. doi:10.1308/003066913x13429796592
26. Maier GS, Horas K, Seeger JB, Roth KE, Kurfth A, Maus U. Is there an association between periprosthetic joint infection and low vitamin D levels? *Int Orthop*. 2014;38(7):1499–1504. doi:10.1007/s00464-014-3338-6
27. Traven SA, Chiaramonti AM, Barfield WR, et al. Fewer complications following revision hip and knee arthroplasty in patients with normal vitamin D levels. *J Arthroplasty*. 2017;32(9S):S193–S196. doi:10.1016/j.arth.2017.02.038
28. Cunha BM, da Cunha BM, Gava AD, de Oliveira SB, de David AC, Dos Santos-Neto LL. Vitamin D is related to gait recovery after total hip arthroplasty: a prospective analysis. *Gait Posture*. 2016;50:96–101. doi:10.1016/j.gaitpost.2016.08.014
29. Garfinkel RJ, Dilisio MF, Agrawal DK. Vitamin D and its effects on articular cartilage and osteoarthritis. *Orthop J Sports Med*. 2017;5(6):2325967117711376.
30. Pizzuti NS, George J, Kholapas A, et al. High prevalence and seasonal variation of hypovitaminosis D in patients scheduled for lower extremity total joint arthroplasty. *Ann Transl Med*. 2018;6(16):321. doi:10.21037/atm.2018.08.21
31. Liberati A, Altman DG, Tetzlaff J, et al. The PRISMA statement for reporting systematic reviews and meta-analyses of studies that evaluate healthcare interventions: explanation and elaboration. *BMJ*. 2009;339:b2700. doi:10.1136/bmj.b2700
32. Wells GA, Shea B, O’Connell D, et al. The Newcastle-Ottawa Scale (NOS) for assessing the quality of nonrandomised studies in meta-analyses; 2016. Available from http://www.ohri.ca/programs/clinical_epidemiology/oxford.asp. Accessed December 2016.

33. Reid D, Toole BJ, Knox S, et al. The relation between acute changes in the systemic inflammatory response and plasma 25-hydroxyvitamin D concentrations after elective knee arthroplasty. Am J Clin Nutr. 2011;93(5):1006–1011. doi:10.3945/ajcn.111.008490

34. Kelly MA, Campbell J, Sheahan J, Murphy P. Vitamin D insufficiency in patients undergoing total knee arthroplasty in Ireland. Ir Med J. 2017;110(10):649.

35. Glowacki J, Hurwitz S, Thornhill TS, Kelly M, LeBoff MS. Osteoporosis and vitamin-D deficieny among postmenopausal women with osteoarthritis undergoing total hip arthroplasty. J Bone Joint Surg Am. 2003;85-A:2371–2377. doi:10.2106/0004623-200312000-00015

36. Dennis DA, Komistek RD, Stehl JB, Walker SA, Dennis KN. Range of motion after total knee arthroplasty: the effect of implant design and weight-bearing conditions. J Arthroplasty. 1998;13(7):748-752. doi:10.1016/s0883-5403(98)90025-0

37. Torres C, Sánchez de la Torre M, García-Moruja C, et al. Immuno-phenotype of vitamin D receptor polymorphism associated to risk of HIV-1 infection and rate of disease progression. Curr HIV Res. 2010;8(6):487–492.3. doi:10.2174/157016210793499330

38. Abhimanyu A, Coussens AK. The role of UV radiation and vitamin D in the seasonality and outcomes of infectious disease. Photochem Photobiol Sci. 2017;16(3):314–338. doi:10.1039/C6PP00355A

39. Trongtrakul K, Feemuchang C. Prevalence and association of vitamin D deficiency and mortality in patients with severe sepsis. Int J Gen Med. 2017;10:415–421. doi:10.2147/IJGM.S147561

40. Tiwari S, Pratyush DD, Gupta SK, Singh SK. Vitamin D deficiency is associated with inflammatory cytokine concentrations in patients with diabetic foot infection. Br J Nutr. 2014;112(12):1938–1943. doi:10.1017/S0007114514003018

41. Shojaii M, Sabzeghabaei A, Helia Valaei Barhagh H, Soheil Soltani S. The correlation between serum level of vitamin D and outcome of sepsis patients; a cross-sectional study. Arch Acad Emerg Med. 2019;7(1):e1.

42. Braun A, Chang D, Mahadevappa K, et al. Association of low serum 25-hydroxyvitamin D levels and mortality in the critically ill. Crit Care Med. 2011;39(4):671–677. doi:10.1097/CCM.0b013e318206edcf

43. Matthews LR, Ahmed Y, Wilson KL, Griggs DD, Danner OK. Worsening severity of vitamin D deficiency is associated with increased length of stay, surgical intensive care unit cost, and mortality rate in surgical intensive care unit patients. Am J Surg. 2012;204(1):37–43. doi:10.1016/j.amjsurg.2011.07.021

44. Thacher TD, Clarke BL. Vitamin D insufficiency. Mayo Clin Proc. 2011;86(1):50-60. doi:10.4065/mcp.2010.0567