# A propensity score-matched analysis on the impact of patient and surgical factors on early periprosthetic joint infection in minimally invasive anterolateral and transgluteal total hip arthroplasty 

Matthias Luger ${ }^{1,2(1)} \cdot$ Marcel de Vries $^{1,2}$. Sandra Feldler ${ }^{1,2}$. Günter Hipmair ${ }^{1,2} \cdot$ Tobias Gotterbarm $^{1,2}$ Antonio Klasan ${ }^{2}$

Received: 17 March 2022 / Accepted: 29 December 2022 / Published online: 11 January 2023
© The Author(s) 2023


#### Abstract

Introduction Increased risk of periprosthetic joint infection (PJI) in minimally invasive (MIS) total hip arthroplasty (THA) is still debated. This study aimed to identify differences in surgical and patient-related risk factors for PJI between an MIS anterolateral approach and transgluteal-modified Hardinge approach. Methods A retrospective cohort of 5315 THAs performed between 2006 and 2019 at a single institution was screened. Short stem THAs performed via an MIS anterolateral approach in the supine position and standard straight stem THAs performed via a transgluteal modified Hardinge approach were included. Propensity score matching was performed to control for selection bias. After matching, 1405 (34.3\%) short stem THAs implanted via MIS anterolateral approach and 2687 ( $65.7 \%$ ) straight stem THAs implanted via a transgluteal modified Hardinge approach were included. The risk of PJI due to patient-specific and surgical factors was retrospectively analyzed using chi-square test and multivariate regression analysis. Results PJI occurred in $1.1 \%$ in both MIS anterolateral and transgluteal approach ( $p=0.823$ ). Multivariate regression showed an increased infection risk for patients with a BMI between 35 and $39.99 \mathrm{~kg} / \mathrm{m}^{2}$ (OR 6.696; CI 1.799-24.923; $p=0.005$ ), which could not be demonstrated for transgluteal approach (OR 0.900 ; CI $0.900-4.144 ; p=0.93$ ). A BMI $\geq 40 \mathrm{~kg} / \mathrm{m}^{2}$ (OR 14.150; CI 2.416-82.879; $p=0.003$ ) was detected as a risk factor for PJI only in anterolateral approach. Increased operation time $\geq 121$ min showed a significantly increased risk for PJI in the general cohort (OR 6.989; CI1.286-37.972; $p=0.024$ ). Conclusion Minimally invasive anterolateral and transgluteal THA show a comparable rate of early PJI within the first year of index surgery. A BMI of $\geq 35 \mathrm{~kg} / \mathrm{m}^{2}$ was detected as a clear risk factor for infection in the anterolateral approach. Prolonged operation time $\geq 121 \mathrm{~min}$ increases the risk of PJI regardless of approach.


Keywords Minimally invasive • Total hip arthroplasty • Cementless • Anterolateral approach • Transgluteal approach • Periprosthetic joint infection

## Abbreviations

| ASA Score | American Society of Anesthesiologists <br> Score |
| :--- | :--- |
| BMI | Body Mass Index $\left(\mathrm{kg} / \mathrm{m}^{2}\right)$ |

[^0]| CI | Confidence interval |
| :--- | :--- |
| DAA | Direct anterior approach |
| DLA | Direct lateral approach |
| MIS approach | Minimally invasive surgical approach |

Antonio Klasan
Klasan.antonio@me.com
1 Department for Orthopedics and Traumatology, Kepler University Hospital GmbH, Krankenhausstrasse 9, 4020 Linz, Austria

2 Johannes Kepler University Linz, Altenberger Strasse 69, 4040 Linz, Austria

| OR | Odds Ratio |
| :--- | :--- |
| PA | Posterior approach |
| PJI | Periprosthetic joint infection |
| THA | Total hip arthroplasty |
| TJA | Total joint arthroplasty |

## Introduction

Total hip arthroplasty (THA) is one of the most successful surgeries in orthopedics, providing pain reduction, good functional outcomes, and improvement in quality of life [1, 2]. Although complication rates in THA are relatively low, periprosthetic joint infection (PJI) is a devastating complication, that can lead to revision surgery with increased morbidity and mortality [3-5].

Several patient-specific factors such as obesity [6-12], diabetes [6, 12-14], rheumatoid arthritis [8, 12], alcohol abuse [12] and smoking status [15] are considered as potential risk factors for postoperative wound complications and PJI after total joint arthroplasty (TJA). Apart from patientspecific aspects, various surgical factors seem to be related to an increased risk of PJI [8].

In recent years, minimally invasive surgical (MIS) ante-rior-based approaches have gained popularity because they are associated with faster postoperative rehabilitation, less pain, and better functional outcomes than conventional surgical approaches [2, 16]. One of these MIS approaches is the MIS anterolateral approach. The risk of complications in the MIS anterolateral approach and in particular of the risk for PJI increases significantly in severely and morbidly obese patients [7]. In a big registry study, Smith et al. [8] found an increased PJI revision rate by about 1.6 -fold when compared to the posterior approach. In contrast, Sheth et al. [17], do not report a significantly increased risk for surgical complications and especially septic revision in the anterolateral approach compared to the posterior approach. The rate of septic revision was reported of being two times higher in the direct lateral approach (DLA) $(0.5 \%$ vs. $1.1 \%)$ with a hazard ratio of 2.15 for DLA compared to 0.98 for the anterolateral approach, however without statistical significance [17]. In a recent meta-analysis by Acuña et al. [18] did not find a significantly increased risk for PJI in the anterolateral approach when compared to the direct anterior approach (DAA). However, data about the incidence of PJI and potential risk factors for infection after THA via MIS anterolateral approach compared to conventional standard approaches are inconclusive. Therefore, the aim of this study was to identify risk factors and differences in risk for periprosthetic joint infection (PJI) in primary THA using a minimally invasive anterolateral cementless short stem THA and transgluteal cementless straight stem THA within 12 months after index surgery.

## Patients and methods

The institutional electronic database was used to obtain information on patients who underwent THA between 2006 and 2019. In total, 5315 THAs in 5205 patients have been performed in this period. Inclusion criteria were defined as cementless short stem THA via a mini-mally-invasive anterolateral approach in supine positioning [19] or cementless straight stem THA via a modified Hardinge approach [20]. Diagnosis for inclusion was primary osteoarthritis, avascular necrosis of the head and hip dysplasia. All forms of secondary osteoarthritis due to posttraumatic deformities or rheumatoid arthritis and all cases with previous surgeries on the affected side were excluded. Additionally, all forms of other approaches were excluded. Cemented THA, the use of deviating implants such as revision cups or stems were excluded. We retrospectively screened every case in this time period that was revised for any reason within the first year. In a second step, all revisions were screened if they met criteria for a periprosthetic joint infection (PJI). A PJI was defined according to the new scoring system from 2018 by Parvizi et al. [21]. As the relevant parameters for the minor criteria were not available in all cases, only PJIs could be included, that met one of the major criteria. Alpha-Defensin test was performed in selected cases. The cases, in which the alpha-defensin test was performed, also fulfilled the major criteria and therefore overruled the minor criteria.

Transgluteal approach was performed as the standard approach between the years 2006 and 2011 at the institution. In 2011 anterolateral approach was introduced at the institution. Between the years 2011 and 2015 MIS anterolateral approach was performed in parallel with the transgluteal approach. From the beginning of the year 2016 until the end of 2019 MIS anterolateral approach was performed as the standard approach. From 2016 transgluteal approach was only performed in selected cases by the preference of the performing surgeon. With transitioning from transgluteal to anterolateral approach as the standard approach for primary THA at our institution, also residents were primarily trained in the anterolateral approach from 2016 and onwards.

The study was approved by the institutional review board (EK-No.: 1194/2021). Because of the retrospective anonymized evaluation of pre-existing medical records, an informed consent was not required. All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.

## Surgical technique

In total 28 surgeons performed the surgeries. The surgeries were performed by 8 consultants and 10 residents. 10 surgeons performed the surgeries as residents and consultants. All surgeons performed the surgeries in a standardized manner and were partly or fully trained at the authors' institution. The institutional transition from transgluteal straight stem THA to MIS anterolateral short stem THA was introduced by two experienced consultants. After gaining enough experience, the transition was then extended to further surgeons in the team under the supervision of these two consultants. Operation time was defined as the time in minutes from skin incision to skin closure.

All surgeries were performed under laminar airflow. Extremity preparation was performed with threefold antiseptic scrub with alcohol disinfectant in all cases. Routinely draping with sterile adhesive surgical iodine film was used only by a certain number of surgeons. The standardized periand postoperative protocol was identical in all cases, including single-shot antibiotics (Cefuroxime 1.5 g i.v. directly pre-operatively), Indomethacin 75 mg twice daily for the prevention of heterotopic ossification on day one to four post-operatively, and 40 mg low-molecular-weight heparin or Rivaroxaban 10 mg for 28 days post-operatively as venous thromboembolic event prophylaxis.

Minimally invasive anterolateral approach was performed in supine positioning. A skin incision was centered over the greater trochanter. An incision at the border between the tensor fasciae latae and the tractus iliotibilias was performed. Then, the Watson-Jones interval between tensor fasciae latae and gluteus medius was bluntly dissected. A capsulectomy was performed in each case. Full weight-bearing was allowed immediately on the day of surgery. Drainage was used until the end of 2017 in every case in an anterolateral approach. In 2018 drainage was used only to the surgeon's preference. From 2019 drainage was not used routinely in the anterolateral approach.

The direct lateral approach (DLA) by Hardinge was first described in 1982 [22]. The modified Hardinge approach has previously been described by Frndak et al. [20]. The modified Hardinge approach was performed with the positioning the patient in supine positioning. A lateral skin incision was used centered over the greater trochanter. Access to the hip joint was gained through an abductor muscle split approach. The fibers of the gluteus medius were split longitudinally at the junction of the anterior third to posterior two-thirds of the muscle belly. The gluteus minimus and capsule were then divided vertically along the same incision parallel to the gluteus medius split. Then a capsulectomy of the anterior capsule was performed. Full weight-bearing was allowed on day one after surgery. Drainage was used in every case in transgluteal approach.

Suturing was done either by skin clamping or intracutaneous suturing. Intracutaneous suturing was the standard wound closure until 2019. Skin clamping was the standard wound closure from the beginning of 2019. The sutures were removed by the family practitioner or by the rehabilitation staff or in certain cases at the outpatient department of the institution after 12-14 days of surgery. Patients were informed at dismissal by the medical report and orally to readmit at the institution in case of any signs PJI. Follow-up was scheduled at 3 months and 1 year postoperatively.

## Implants

In a minimally invasive anterolateral approach, a cementless, curved short stem (Fitmore ${ }^{\circledR}$ stem, Zimmer Biomet, Warsaw, IN, USA) was digitally templated using mediCAD ${ }^{\circledR}$ version 5.1 (Hectec GmbH, Altdorf, Germany). Fitmore ${ }^{\circledR}$ hip stem is a titanium alloy stem ( $\mathrm{Ti} \mathrm{Al6V} 4$ ) that has a porolock TiVPS coating in the proximal part to enhance bone ingrowth and is available in four different neck angle options ( $127^{\circ}$, $129^{\circ}, 137^{\circ}, 140^{\circ}$ ). A cementless titanium press-fit cup with or without screws (Allofit ${ }^{\circledR} /-\mathrm{S}, \mathrm{Zimmer}$ Biomet, Warsaw, IN, USA) or two types of cementless threaded cups (Alloclassic $\mathrm{CSF}^{\circledR}$ / Alloclassic Variall ${ }^{\circledR}$, both Zimmer Biomet, Warsaw, IN, USA) were used. In the transgluteal approach a cementless Zweymüller straight stem in two variations was used (Alloclassic SL/SLO; Alloclassic SLV; both Zimmer Biomet, Warsaw, IN, USA). In the transgluteal approach, a cementless titanium press-fit cup with or without screws (Allofit ${ }^{\circledR} /$-S/IT, Zimmer Biomet, Warsaw, IN, USA) or two types of cementless threaded cups (Alloclassic $\mathrm{CSF}^{\circledR} /$ Alloclassic Variall ${ }^{\circledR}$, both Zimmer Biomet, Warsaw, IN, USA) were used. Independent from the approach, highly cross-linked polyethylene liners (Alpha Durasul ${ }^{\circledR}$, Gamma Durasu ${ }^{\circledR}$, Alloclassic CSF Durasul ${ }^{\circledR}$, Longevity IT Liner ${ }^{\circledR}$, all Zimmer Biomet, Warsaw, IN, USA) were used in every case. As femoral heads two types of ceramic heads were used (BIOLOX forte, CeramTec GmbH, DE; Sulox, Zimmer Biomet, Warsaw, IN, USA) as well as Cobalt-chrome (CoCr) metal heads (Durasul CoCr, Zimmer Biomet, Warsaw, IN, USA).

## Statistics

Descriptive analysis was performed for patient demographics. A Shapiro-Wilk test for normality was performed to determine whether continuous data were normally distributed. As the variables were normally distributed, a Pearson's chi-square test was performed for categorical variables and a student's $t$ test was performed for continuous variables. Because of statistically significant differences in the patient demographics a propensity score matching was performed using the caliper technique. The caliper was set at 0.2 . The
propensity score matching was performed for patient age at operation, Body Mass Index (BMI; $\mathrm{kg} / \mathrm{m}^{2}$ ) ASA Score (American Society of Anesthesiologists Score), gender, diagnosis, operation side, smoking status, alcohol consumption, diabetes and the surgical factors approach, operation time and the surgeon's experience. The risk of PJI was calculated for all patient and surgical factors that were included in the propensity score matching. A post hoc power analysis was performed. With the total sample size of 4092 patients, an alpha of 0.05 and an omega $(\omega)$ of 0.003 , a power (beta) of 0.54 was calculated. The rates of revision due to PJI were recorded for all patients and divided by approach. A multivariate regression model was calculated for all patient and surgical factors on the risk of PJI for the general cohort. Additionally, a multivariate regression model was calculated and divided by approach. All significant factors of the univariate analysis were then used for multivariate regression analysis. Data were analyzed using SPSS version 28 (IBM SPSS statistics, Chicago, IL, USA).

## Results

## Propensity score matching

In total, 4511 THAs have met the inclusion criteria included in this study. A total of 806 THAs did not meet the inclusion criteria, Fig. 1. Of these 806 patients, 80 patients have been lost to follow-up. Both groups differed significantly in the patient age at operation $(p=0.011)$, experience of the surgeon $(<0.001)$, operation time $(<0.001)$, ASA Score ( $p<0.001$ ) and smoking status ( 0.046 ), Table 1. By propensity score matching 409 patients were excluded. Therefore, 4092 THAs were included after the propensity score matching in the final analysis. The patient demographics did not differ between both approaches after matching in
all categories, Table 1. In 1405 cases, short stem THA ( $34.3 \%$ ) was performed via a minimally-invasive anterolateral approach and 2687 straight stem THAs ( $65.7 \%$ ) were implanted via a transgluteal modified Hardinge approach.

## Comparison between both approaches

In total, 45 PJIs (1.1\%) were detected within 12 months of index surgery. All cases met the major criteria by Parvizi et al. [21]. In all cases, either two positive cultures of the same organism or a sinus tract with evidence of communication to the joint or visualization of the prosthesis or both were documented. Rate of PJI was $1.1 \%$ in the anterolateral approach compared to $1.1 \%$ in the transgluteal approach ( $p=0.862$ ), Table 2. The number of infections and the testing for the occurrence of PJI in the general cohort are shown in Table 2. Increased BMI was statistically significant in the general cohort ( $p=0.022$ ), Table 2. In the anterolateral approach, the number of infections were significantly higher in patients with increased BMI ( $p<0.001$ ), Table 3. In the transgluteal approach the number of infections were significantly higher in patients with increased ASA Score ( $p=0.034$ ) and in diabetic patients $(p=0.044)$, Table 4.

## Regression analysis

The multivariate regression analysis for the general cohort and separated by approach is shown in detail in Table 5. Multivariate regression analysis showed a significantly increased odds ratio (OR) for PJI in the total study group in patients between 70 and 79 years at operation (OR 4.687; CI 1.629-14.536), Table 5. The OR was also increased in patients 80 years of age at operation or older (OR 3.723; CI 0.955-14.522) but without statistical significance ( $p=0.059$ ), Table 5. The OR for increasing operation time increased throughout all groups but only showed a

Fig. 1 Consort Diagram for inclusion and exclusion of patients


Table 1 Patient demographics

|  | Pre-matched cohort |  | $P$ value | Post-matched cohort |  | $P$ value |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Anterolateral <br> Mean ( $\pm$ SD) | Lateral <br> Mean ( $\pm$ SD) |  | Anterolateral <br> Mean ( $\pm$ SD) | Lateral <br> Mean ( $\pm$ SD) |  |
| Number of patients | 1410 (31.3\%) | 3101 (68.7\%) |  | 1405 (34,3\%) | 2687 (65,7\%) |  |
| Age at operation | 67.18 ( $\pm 11.55)$ | 66.29 ( $\pm 12.07)$ | 0.011 | 67.15 ( $\pm 11.56)$ | 66.81 ( $\pm 11.96)$ | 0.383 |
| BMI | 27.9 ( $\pm 4.8)$ | 28 ( $\pm 4.74)$ | 0.527 | $27.92( \pm 4.8)$ | $27.77( \pm 4.5)$ | 0.318 |
| Sex |  |  | 0.343 |  |  | 0.606 |
| Female | 783 (55.5\%) | 1675 (54\%) |  | 780 (55.5\%) | 1469 (54.7\%) |  |
| Male | 627 (44.5\%) | 1426 (46\%) |  | 625 (44.5\%) | 1218 (45.3\%) |  |
| Diagnosis |  |  | 0.077 |  |  | 0.168 |
| Primary OA | 1177 (83.5\%) | 2501 (80.7\%) |  | 1172 (83.4\%) | 2177 (81.1\%) |  |
| AVN | 146 (10.3\%) | 276 (12.1\%) |  | 146 (10.4\%) | 321 (11.9\%) |  |
| Hip Dyplasia | 87 (6.2\%) | 224 (7.2\%) |  | 87 (6.2\%) | 189 (7\%) |  |
| Surgeon's experience |  |  | <0.001 |  |  | 0.060 |
| Consultant | 1128 (80\%) | 2155 (69.5\%) |  | 1123 (79.9\%) | 2079 (77.4\%) |  |
| Resident | 282 (20\%) | 946 (30.5\%) |  | 282 (20.1\%) | 608 (22.6\%) |  |
| Operation time | 80.29 ( $\pm 23.32)$ | 84.73 ( $\pm 22.76)$ | <0.001 | 80.38 ( $\pm 23.3)$ | 81.75 ( $\pm 20.25)$ | 0.052 |
| Side |  |  | 0.646 |  |  | 0.522 |
| Left | 658 (46.7\%) | 1470 (47.4\%) |  | 654 (46.5\%) | 1279 (47.6\%) |  |
| Right | 752 (53.3\%) | 1631 (52.6\%) |  | 751 (53.5\%) | 1408 (52.4\%) |  |
| ASA |  |  | <0.001 |  |  | 0.146 |
| 1 | 276 (19.6\%) | 494 (15.9\%) |  | 272 (19.4\%) | 469 (17.5\%) |  |
| 2 | 823 (58.4\%) | 2011 (64.9\%) |  | 823 (58.6\%) | 1673 (62.3\%) |  |
| 3 | 305 (21.6\%) | 581 (18.7\%) |  | 304 (21.6\%) | 533 (19.8\%) |  |
| 4 | 6 (0.4\%) | 15 (0.5\%) |  | 6 (0.4\%) | 12 (0.4\%) |  |
| Diabetes | 176 (12.5\%) | 385 (12.4\%) | 0.950 | 176 (12.5\%) | 323 (12\%) | 0.639 |
| Smoking | 240 (17\%) | 456 (14.7\%) | 0.046 | 235 (16.7\%) | 410 (15.3\%) | 0.221 |
| Alcohol | 300 (21.3\%) | 645 (20.8\%) | 0.715 | 299 (21.3\%) | 557 (20.7\%) | 0.680 |

Bold letters indicate significant values
$B M I$ Body Mass Index $\mathrm{kg} / \mathrm{m}^{2}$, primary $O A$ primary osteoarthritis, $A V N$ avascular necrosis of the femoral head, $A S A$ American Society of Anesthesiologists
statistically increased risk in THAs with $\geq 121 \mathrm{~min}$ of operation time (OR 6.989; CI 1.286-37.972), Table 5.

The multivariate analysis separated by approach showed a significantly increased risk for PJI in the anterolateral approach for patients with a BMI $\geq 35-39.99 \mathrm{~kg} / \mathrm{m}^{2}$ (OR 6.696; CI 1.799-24.923) and BMI $\geq 40 \mathrm{~kg} / \mathrm{m}^{2}$ (OR 14.150 ; CI 2.416-82.879), Table 5. In the transgluteal approach, a patient aged between 70 and 79 years at operation (OR 4.404; CI 1.206-16.085) and smoking (OR 3.023; CI 1.126-8.119) were identified as independent risk factors for PJI.

## Discussion

In the current study, we retrospectively analyzed the rates of periprosthetic joint infection within 12 months from index surgery in propensity-score matched cohorts including
minimally invasive anterolateral short stem THA and transgluteal straight stem THA and evaluated potential risk factors for infection. We did not find a statistically significant difference in the rates of PJI within the first year of index surgery between an anterolateral MIS approach and a transgluteal Hardinge approach, while the risk for occurrence of a PJI was significantly higher in severely obese patients in an anterolateral approach and increased surgical time longer than 120 min was a significantly increased risk factor in both approaches.

The incidence of PJI is reported within a range of $0.3-3 \%$ [9, 23]. We found comparable PJI rates of $1.1 \%$ in both groups ( $p=0.862$ ). Ilchmann et al. [24] found a rate of PJI of $1.7 \%$ for DLA. Shohat et al. [25] reported a rate of $1.3 \%$ in DLA. Some authors suggest higher rates of PJI and numbers of revision surgeries due to PJI with MIS approaches [8, 26]. Smith et al. [8] reported THAs implanted via an anterolateral approach at a higher risk of revision for postoperative

Table 2 Chi-Square test for PJI and patient or surgical factors for the general cohort

|  | Total (n) | Infection ( $n$ ) | Infection (\%) | $P$ value |
| :---: | :---: | :---: | :---: | :---: |
| Approach |  |  |  | 0.862 |
| Anterolateral | 1405 | 16 | 1.1 |  |
| Transgluteal | 2687 | 29 | 1.1 |  |
| BMI ( $\mathrm{kg} / \mathrm{m}^{2}$ ) |  |  |  | 0.022 |
| <35 | 3792 | 37 | 1.0 |  |
| 35-40 | 240 | 6 | 2.5 |  |
| $>40$ | 60 | 2 | 3.3 |  |
| ASA |  |  |  | 0.087 |
| 1 | 741 | 6 | 0.8 |  |
| 2 | 2496 | 23 | 0.9 |  |
| 3 | 837 | 16 | 1.9 |  |
| 4 | 18 | 0 | 0.0 |  |
| Gender |  |  |  | 0.084 |
| Female | 2249 | 19 | 0.8 |  |
| Male | 1843 | 26 | 1.4 |  |
| Age (years) |  |  |  | 0.059 |
| <60 | 1112 | 6 | 0.5 |  |
| 60-69 | 1138 | 11 | 1.0 |  |
| 70-79 | 1318 | 22 | 1.7 |  |
| $\geq 80$ | 512 | 6 | 1.2 |  |
| Diagnosis |  |  |  | 0.222 |
| Primary OA | 3349 | 36 | 1.1 |  |
| AVN | 467 | 8 | 1.7 |  |
| Hip dysplasia | 1 | 1 | 0.4 |  |
| Side |  |  |  | 0.706 |
| Left | 1933 | 20 | 1.0 |  |
| Right | 2159 | 25 | 1.2 |  |
| Smoking |  |  |  | 0.709 |
| Yes | 645 | 8 | 1.2 |  |
| No | 3447 | 37 | 1.1 |  |
| Alcohol |  |  |  | 0.559 |
| Yes | 856 | 11 | 1.3 |  |
| No | 3236 | 34 | 1.1 |  |
| Diabetes |  |  |  | 0.108 |
| Yes | 499 | 9 | 1.8 |  |
| No | 3593 | 36 | 1.0 |  |
| Operation time (min) |  |  |  | 0.095 |
| $\leq 60$ | 472 | 2 | 0.4 |  |
| 61-90 | 2550 | 26 | 1.0 |  |
| 91-120 | 870 | 12 | 1.4 |  |
| $\geq 121$ | 200 | 5 | 2.5 |  |
| Experience |  |  |  | 0.775 |
| Consultant | 3202 | 36 | 1.1 |  |
| Resident | 890 | 9 | 1.0 |  |

Bold letters indicate significant values
BMI Body Mass Index $\mathrm{kg} / \mathrm{m}^{2}$, primary $O A$ primary osteoarthritis, $A V N$ avascular necrosis of the femoral head, ASA American Society of Anesthesiologists

Table 3 Chi-Square test for PJI and patient or surgical factors separated for anterolateral approach

|  | Total ( $n$ ) | Infection (n) | Infection (\%) | $P$ value |
| :---: | :---: | :---: | :---: | :---: |
| BMI ( $\mathrm{kg} / \mathrm{m}^{2}$ ) |  |  |  | <0.001 |
| $<35$ | 1282 | 10 | 0.8 |  |
| 35-40 | 99 | 4 | 4.0 |  |
| $>40$ | 24 | 2 | 8.3 |  |
| ASA |  |  |  | 0.899 |
| 1 | 272 | 2 | 0.7 |  |
| 2 | 823 | 10 | 1.2 |  |
| 3 | 304 | 4 | 1.3 |  |
| 4 | 6 | 0 | 0.0 |  |
| Gender |  |  |  | 0.655 |
| Female | 780 | 8 | 1.0 |  |
| Male | 625 | 8 | 1.3 |  |
| Age (years) |  |  |  | 0.421 |
| <60 | 208 | 1 | 0.5 |  |
| 60-69 | 864 | 9 | 1.0 |  |
| 70-79 | 258 | 4 | 1.6 |  |
| $\geq 80$ | 75 | 2 | 2.7 |  |
| Diagnosis |  |  |  | 0.350 |
| Primary OA | 1172 | 13 | 1.1 |  |
| AVN | 146 | 3 | 2.1 |  |
| Hip dysplasia | 87 | 0 | 0.0 |  |
| Side |  |  |  | 0.821 |
| Left | 654 | 7 | 1.1 |  |
| Right | 751 | 9 | 1.2 |  |
| Smoking |  |  |  | 0.259 |
| Yes | 1170 | 15 | 1.3 |  |
| No | 235 | 1 | 0.4 |  |
| Alcohol |  |  |  | 0.388 |
| Yes | 299 | 2 | 0.7 |  |
| No | 1106 | 14 | 1.3 |  |
| Diabetes |  |  |  | 0.997 |
| Yes | 176 | 2 | 1.1 |  |
| No | 1229 | 14 | 1.1 |  |
| Operation time (min) |  |  |  | 0.319 |
| $\leq 60$ | 373 | 1 | 0.3 |  |
| 61-90 | 401 | 6 | 1.5 |  |
| 91-120 | 467 | 7 | 1.5 |  |
| $\geq 121$ | 164 | 2 | 1.2 |  |
| Experience |  |  |  | 0.262 |
| Consultant | 1123 | 11 | 1.1 |  |
| Resident | 282 | 5 | 1.8 |  |

Bold letters indicate significant values
BMI Body Mass Index $\mathrm{kg} / \mathrm{m}^{2}$, primary $O A$ primary osteoarthritis, $A V N$ avascular necrosis of the femoral head, ASA American Society of Anesthesiologists

Table 4 Chi-Square test for PJI and patient or surgical factors separated for transgluteal approach

|  | Total (n) | Infection (n) | Infection (\%) | $P$ value |
| :---: | :---: | :---: | :---: | :---: |
| BMI ( $\mathrm{kg} / \mathrm{m}^{2}$ ) |  |  |  | 0.761 |
| <35 | 2510 | 27 | 1.1 |  |
| 35-40 | 141 | 2 | 1.4 |  |
| > 40 | 36 | 0 | 0.0 |  |
| ASA |  |  |  | 0.034 |
| 1 | 469 | 4 | 0.9 |  |
| 2 | 1673 | 13 | 0.8 |  |
| 3 | 533 | 12 | 2.3 |  |
| 4 | 12 | 0 | 0.0 |  |
| Gender |  |  |  | 0.069 |
| Female | 1469 | 11 | 0.7 |  |
| Male | 1218 | 18 | 1.5 |  |
| Age (years) |  |  |  | 0.110 |
| <60 | 749 | 5 | 0.7 |  |
| 60-69 | 737 | 5 | 0.7 |  |
| 70-79 | 851 | 15 | 1.8 |  |
| $\geq 80$ | 250 | 4 | 1.1 |  |
| Diagnosis |  |  |  | 0.539 |
| Primary OA | 2177 | 23 | 1.1 |  |
| AVN | 321 | 5 | 1.6 |  |
| Hip dysplasia | 189 | 1 | 0.5 |  |
| Side |  |  |  | 0.764 |
| Left | 1279 | 13 | 1.0 |  |
| Right | 1408 | 16 | 1.1 |  |
| Smoking |  |  |  | 0.181 |
| Yes | 410 | 7 | 1.7 |  |
| No | 2277 | 22 | 1.0 |  |
| Alcohol |  |  |  | 0.169 |
| Yes | 2130 | 9 | 0.9 |  |
| No | 557 | 20 | 1.6 |  |
| Diabetes |  |  |  | 0.044 |
| Yes | 323 | 7 | 2.2 |  |
| No | 2364 | 22 | 0.9 |  |
| Operation time (min) |  |  |  | 0.304 |
| $\leq 60$ | 264 | 1 | 0.4 |  |
| 61-90 | 1686 | 17 | 1.0 |  |
| 91-120 | 612 | 8 | 1.3 |  |
| $\geq 121$ | 125 | 3 | 2.4 |  |
| Experience |  |  |  | 0.253 |
| Consultant | 2079 | 25 | 1.2 |  |
| Resident | 608 | 4 | 0.7 |  |

Bold letters indicate significant values
BMI Body Mass Index $\mathrm{kg} / \mathrm{m}^{2}$, primary $O A$ primary osteoarthritis, $A V N$ avascular necrosis of the femoral head, ASA American Society of Anesthesiologists
infection compared to the posterior approach (PA) (OR 1.61; CI 1.16-2.23; $p=0.005$ ). Other studies based on nationwide registries could not find a negative influence of MIS approaches on the risk of revision due to infection [27-29]. Sheth et al. [17] could not report a statistically significant increased risk for septic revision in an anterolateral approach, with a reduced rate of early dislocation, concluding to be the main advantage of an anterolateral approach. We also report comparable rates of PJI in anterolateral and transgluteal approach. Additionally, the low rate of dislocation of an MIS anterolateral approach and a cementless short stem has been previously [7]. Therefore, an anterolateral MIS approach might be favorable due to the reduced rate of early complications without leading to an increased rate of PJI compared to a standard transgluteal approach.

Although the overall infection rate was equivalent in both cohorts, the anterolateral approach was associated with higher infection rates as BMI increased. Severely ( $\mathrm{BMI} \geq 35 \mathrm{~kg} / \mathrm{m}^{2}$ ) and morbidly obese patients (BMI $\geq 40 \mathrm{~kg} / \mathrm{m}^{2}$ ) receiving THA via anterolateral approach were at a higher risk of developing PJI than obese patients in the transgluteal approach group. Obesity has previously been demonstrated to increase the risk of postoperative wound complication, deep infection, and revision surgery due to infection in THA via MIS approaches [7, 8, 30, 31]. A recent systematic review by Shah et al. [32] did not find any significantly increased risk for PJI in the anterolateral approach. However, the used cut-off was a BMI of $30 \mathrm{~kg} / \mathrm{m}^{2}$. In the present study, a BMI above $35 \mathrm{~kg} / \mathrm{m}^{2}$ did not have a statistically significant impact on PJI rates in the transgluteal approach group. Therefore, the transgluteal approach might be favorable compared to the MIS anterolateral approach regarding the risk of early PJI in obese patients.

Prolonged surgical duration has previously been shown to increase the risk of surgical site infection in total joint arthroplasty [12, 33]. Every 20-min increase in operation time is related to an almost $25 \%$ higher risk of PJI in primary TJA [33]. In the current study, an operation time $\geq 121$ min was identified as an independent risk factor for PJI. However, this increased risk was only significant in the general cohort. When separated by approach, the OR increased with longer operation times, but without statistical significance.

Diabetes is a well-known risk factor for the risk PJI in THA [8]. Jämsen et al. [6] report a more than twofold increase in PJI risk for patients diagnosed with diabetes (OR 2.31, CI 1.12-4.72), independent of BMI. Iorio et al. [34] found a four times higher risk of infection in patients with diabetes undergoing total hip or knee arthroplasty. In the present study, testing for significance revealed a significantly higher number of PJIs in patients diagnosed with diabetes in the transgluteal approach cohort. However, multivariate analysis did not show a significant influence of diabetes on the occurrence of PJI. Some studies suggest that the higher incidence of surgical

Table 5 Multivariate analysis for the risk of PJI for patient or surgical factors for all patients and divided by approach

|  | All <br> OR (CI) | $P$ value | Anterolateral OR (CI) | $P$ value | Transgluteal OR (CI) | $P$ value |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| BMI (kg/m ${ }^{2}$ ) |  |  |  |  |  |  |
| <35 | 1.000 | - | 1.000 | - | 1.000 | - |
| 35-40 | 2.486 (0.993-6.223) | 0.052 | 6.696 (1.799-24.923) | 0.005 | 0.900 (0.900-4.144) | 0.93 |
| >40 | 2.851 (0.621-13.089) | 0.178 | 14.150 (2.416-82.879) | 0.003 | NV | - |
| ASA |  |  |  |  |  |  |
| 1 | 1.000 | - | 1.000 | - | 1.000 | - |
| 2 | 0.758 (0.268-2.144) | 0.602 | 0.754 (0.145-3.931) | 0.738 | 0.614 (0.173-2.178) | 0.451 |
| 3 | 1.056 (0.331-3.368) | 0.926 | 0.650 (0.090-4.485) | 0.650 | 1.345 (0.333-5.430) | 0.678 |
| 4 | NV | - | NV | - | NV | - |
| Gender |  |  |  |  |  |  |
| Female | 1.000 | - | 1.000 | - | 1.000 | - |
| Male | 1.694 (0.893-3.215) | 0.106 | 1.624 (0.559-4.179) | 0.373 | 1.787 (0.781-4.089) | 0.169 |
| Age (years) |  |  |  |  |  |  |
| <60 | 1.000 | - | 1.000 | - | 1.000 | - |
| 60-69 | 2.291 (0.765-6.964) | 0.139 | 5.822 (0.622-54.467) | - | 1.462 (0.365-5.856) | 0.592 |
| 70-79 | 4.687 (1.629-14.536) | 0.005 | 8.851 (0.867-90.327) | 0.066 | 4.404 (1.206-16.085) | 0.025 |
| $\geq 80$ | 3.723 (0.955-14.522) | 0.058 | 8.484 (0.556-129.393) | 0.124 | 3.240 (0.626-16.755) | 0.161 |
| Diagnosis |  |  |  |  |  |  |
| Primary OA | 1.000 | - | 1.000 | - | 1.000 | - |
| AVN | 1.490 (0.670-3.313) | 0.328 | 1.915 (0.495-7.411) | 0.347 | 1.239 (0.446-3.441) | 0.763 |
| Hip dysplasia | 0.741 (0.087-5.672) | 0.741 | NV | - | 0.829 (0.093-7.405) | 0.867 |
| Side |  |  |  |  |  |  |
| Left | 1.000 | - | 1.000 | - | 1.000 | - |
| Right | 1.165 (0.640-2.119) | 0.617 | 1.181 (0.423-3.299) | 0.751 | 1.141 (0.538-2.418) | 0.731 |
| Smoking |  |  |  |  |  |  |
| No | 1.000 | - | 1.000 | - | 1.000 | - |
| Yes | 1.869 (0.803-4.350) | 0.147 | 0.526 (0.062-4.466) | 0.556 | 3.023 (1.126-8.119) | 0.028 |
| Alcohol |  |  |  |  |  |  |
| No | 1.000 | - | 1.000 | - | 1.000 | - |
| Yes | 1.054 (0.509-2.183) | 0.887 | 0.505 (0.062-4.466) | 0.526 | 1.433 (0.579-3.442) | 0.388 |
| Diabetes |  |  |  |  |  |  |
| No | 1.000 | - | 1.000 | - | 1.000 | - |
| Yes | 1.295 (0.590-2.841) | 0.519 | 0.530 (0.107-2.634) | 0.438 | 1.897 (0.748-4.816) | 0.178 |
| Operation time (min) |  |  |  |  |  |  |
| $\leq 60$ | 1.000 | - | 1.000 | - | 1.000 | - |
| 61-90 | 2.581 (0.607-10.980) | 0.199 | 2.232 (0.272-18.299) | 0.454 | 3.049 (0.400-23.224) | 0.272 |
| 91-120 | 3.663 (0.794-16.909) | 0.096 | 2.414 (0.240-24.236) | 0.454 | 4.304 (0.522-35.466) | 0.175 |
| $\geq 121$ | 6.989 (1.286-37.972) | 0.024 | 6.856 (0.517-90.862) | 0.144 | 9.318 (0.906-95.817) | 0.061 |
| Experience |  |  |  |  |  |  |
| Consultant | 1.000 | - | 1.000 | - | 1.000 | - |
| Resident | 0.719 (0.339-1.528) | 0.392 | 2.068 (0.663-6.449) | 0.210 | 0.389 (0.132-1.150) | 0.088 |

Bold letters indicate significant values
BMI Body Mass Index $\mathrm{kg} / \mathrm{m}^{2}$, primary OA primary osteoarthritis, $A V N$ avascular necrosis of the femoral head, ASA American Society of Anesthesiologists
site infections in patients diagnosed with diabetes might be limited to those with poorly controlled disease [35, 36]. Most THAs implanted via transgluteal approach were performed at
the beginning of the study period before the transition from transgluteal to anterolateral as the standard approach at our institution. Possibly, antidiabetic treatment and, therefore,
glycemic control in patients diagnosed with diabetes have improved over the study period. However, consistent data on preoperative glucose level and glycated hemoglobin were not available in the present retrospective study.

Limitations of this study mainly conclude the retrospective study design. Therefore, baseline differences could be found for age, experience of the surgeon, ASA score and smoking status between both study groups. To control for selection bias and to eliminate possible confounders, propensity score matching incorporating patient demographics, comorbidities and surgery-related variables was performed. However, the anterolateral and transgluteal approach were not performed concurrently over the study period as the standard approach transitioned from the transgluteal to MIS anterolateral approach. Additionally, the follow-up period was defined as 12 months after index surgery. However, the retrospective data analysis of our institutional electronic data does not provide reliable data for a longer follow-up period because of increasing patients lost-to-follow-up after 12 months. Because of the very long time period of included patients, we cannot provide full information and data to fulfill the minor criteria for the new scoring system from 2018 by Parvizi et al. [21]. Therefore, only PJIs that fulfilled the major criteria could be included, leading to the possibility of overseeing low-grade PJIs. Due to the retrospective study design, data on preoperative glucose levels or glycated hemoglobin cannot be presented consistently. Additionally, preoperative risk factors were handled individually by the operating surgeon. Therefore, we cannot give conclusive information on the different preoperative thresholds for operating diabetic patients or patients with elevation of inflammatory markers such as C-reactive protein as it was handled individually. Intraoperative differences between surgeons were also not recorded consistently such as the use of iodine film. However, as a strength, apart from the differences between surgeons, we report a very standardized study collective. Furthermore, we
report a very large study cohort with clear inclusion and exclusion criteria. The number of patients is unequally distributed with fewer cases in the short-stem group. However, we controlled for this unequal distribution by performing the propensity score matching to reduce the risk for bias due to unequal group sizes. A further limitation of the study is the presentation of only one complication. As there may be an increased risk for PJI in certain groups, a minimally invasive approach might have the potential to lead to a decrease of other complications such as deep vein thrombosis or other medical complications because a faster mobilization might be feasible. Another limitation of the study is the high number of different surgeons and also the inclusion of training operations of residents. However, all surgeons performed the surgeries in a standardized manner and were partly or fully trained at the authors' institution. The institutional transition from transgluteal straight stem THA to MIS anterolateral short stem THA was introduced by two experienced consultants. After gaining enough experience, the transition was then extended to further surgeons in the team under the supervision of these two consultants.

## Conclusion

Minimally invasive anterolateral and transgluteal THA show a comparable rate of early PJI within the first year of index surgery. A BMI of $\geq 35 \mathrm{~kg} / \mathrm{m}^{2}$ was detected as a clear risk factor for infection in the anterolateral approach. Prolonged operation time $\geq 121 \mathrm{~min}$ increases the risk of PJI regardless of approach.

## Appendix

See Table 6

Table 6 Distribution of PJIs by the year of operation for all patients and separated by approach

| Year | All |  | Anterolateral |  | Transgluteal |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Total | Infections, $n(\%)$ | Total | Infections, $n(\%)$ | Total | Infections, $n(\%)$ |
| 2006 | 234 | 3 (1.3) | 0 | 0 (0.0) | 234 | 3 (1.3) |
| 2007 | 261 | 1 (0.4) | 0 | 0 (0.0) | 261 | 1 (0.4) |
| 2008 | 252 | 0 (0.0) | 0 | 0 (0.0) | 252 | 0 (0.0) |
| 2009 | 251 | 3 (1.2) | 0 | 0 (0.0) | 251 | 3 (1.2) |
| 2010 | 313 | 3 (1.0) | 0 | 0 (0.0) | 313 | 3 (1.0) |
| 2011 | 303 | 5 (1.7) | 4 | 0 (0.0) | 299 | 5 (1.7) |
| 2012 | 302 | 1 (0.3) | 10 | 0 (0.0) | 292 | 1 (0.3) |
| 2013 | 293 | 3 (1.0) | 37 | 0 (0.0) | 256 | 3 (1.2) |
| 2014 | 307 | 2 (0.7) | 117 | 0 (0.0) | 190 | 2 (1.1) |
| 2015 | 294 | 4 (1.4) | 156 | 2 (1.3) | 138 | 2 (1.4) |
| 2016 | 309 | 3 (1.0) | 216 | 2 (0.9) | 93 | 1 (1.1) |
| 2017 | 320 | 6 (1.9) | 271 | 3 (1.1) | 49 | 3 (6.1) |
| 2018 | 327 | 9 (2.8) | 295 | 8 (2.7) | 32 | 1 (3.2) |
| 2019 | 326 | 2 (0.6) | 298 | 1 (0.3) | 27 | 1 (3.7) |

Acknowledgements The statistical analysis was supported by the Johannes Kepler University Linz, Center for Clinical Studies (CCS Linz) at the Center for Clinical Research, Altenberger Strasse 69, 4040 Linz and Krankenhausstraße 5, 4020 Linz, Austria.

Author contributions ML: Wrote the manuscript, performed the statistical analysis, designed the study, acquisition of data, interpretation of the data. MV: Involved in the acquisition of data. SF: Co-wrote the manuscript. GH: Revised the manuscript, interpretation of the data. TG: Revised the manuscript. AK: Jointly conceived the study, edited the manuscript, interpretation of the data.

Funding Open access funding provided by Johannes Kepler University Linz. The study was conducted without any funding or benefits from a commercial party. Two co-authors have received or will receive benefits for personal or professional use from a commercial party outside the conduction of this study.

Availability of data and materials Data and materials are available on request.

## Declarations

Conflict of interest One co-author (G.H.) has received consultant honoraria of Zimmer Biomet, Europe, outside the submitted work. We report personal fees paid to one co-author (T.G.) during the conduct of the study from Zimmer Biomet, Europe and from Depuy Synthes Orthopädie Gmbh, Peter Brehm GmbH, ImplanTec GmbH outside the submitted work. We report research grants paid to our institution during the conduct of the study from Zimmer Biomet, Europe, Mathys AG Switzerland, Anika Therapeutics outside the submitted work.

Ethical approval This study received ethical approval from the local institutional review board (EK-No.: 1194/2021) of the "Ethikkommission OÖ" of the Johannes Kepler University Linz (JKU Linz).

Informed consent All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. Informed consent was not required because of the retrospective study design.

Consent to participate The study was approved by the institutional review board (EK-No.: 1239/2019) in accordance with the World Medical Association Declaration of Helsinki. Because of the retrospective evaluation of pre-existing medical records informed consent was not required.

Consent to publish All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. Informed consent for publishment was not required because of the retrospective study design.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not
permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by/4.0/.

## References

1. Learmonth ID, Young C, Rorabeck C (2007) The operation of the century: total hip replacement. Lancet 370(9597):1508-1519
2. Gkagkalis G, Goetti P, Mai S, Meinecke I, Helmy N, Bosson D et al (2019) Cementless short-stem total hip arthroplasty in the elderly patient - is it a safe option? A prospective multicentre observational study. BMC Geriatr 19(1):112
3. Barton CB, Wang DL, An Q, Brown TS, Callaghan JJ, Otero JE (2020) Two-stage exchange arthroplasty for periprosthetic joint infection following total hip or knee arthroplasty is associated with high attrition rate and mortality. J Arthroplasty 35(5):1384-1389
4. Aggarwal VK, Elbuluk A, Dundon J, Herrero C, Hernandez C, Vigdorchik JM et al (2019) Surgical approach significantly affects the complication rates associated with total hip arthroplasty. Bone Joint J 101-B(6):646-51
5. Osmon DR, Berbari EF, Berendt AR, Lew D, Zimmerli W, Steckelberg JM et al (2012) Diagnosis and management of prosthetic joint infection: clinical practice guidelines by the infectious diseases society of America. Clin Infect Dis 56(1):e1-e25
6. Jamsen E, Nevalainen P, Eskelinen A, Huotari K, Kalliovalkama J, Moilanen T (2012) Obesity, diabetes, and preoperative hyperglycemia as predictors of periprosthetic joint infection: a singlecenter analysis of 7181 primary hip and knee replacements for osteoarthritis. J Bone Joint Surg Am 94(14):e101
7. Luger M, Hochgatterer R, Schopper C, Pisecky L, Allerstorfer J, Klasan A et al (2021) Obesity in short stem total hip arthroplasty using a minimally invasive supine anterolateral approach-a risk factor for short-term complications? Int Orthop 45(11):2833-2841
8. Smith JO, Frampton CMA, Hooper GJ, Young SW (2018) The impact of patient and surgical factors on the rate of postoperative infection after total hip arthroplasty-a New Zealand joint registry study. J Arthroplast 33(6):1884-1890
9. Pulido L, Ghanem E, Joshi A, Purtill JJ, Parvizi J (2008) Periprosthetic joint infection: the incidence, timing, and predisposing factors. Clin Orthop Relat Res 466(7):1710-1715
10. Antoniadis A, Dimitriou D, Flury A, Wiedmer G, Hasler J, Helmy N (2018) Is direct anterior approach a credible option for severely obese patients undergoing total hip arthroplasty? A matched-control, retrospective, clinical study. J Arthroplast 33(8):2535-2540
11. Namba RS, Inacio MC, Paxton EW (2012) Risk factors associated with surgical site infection in 30,491 primary total hip replacements. J Bone Joint Surg Br 94(10):1330-1338
12. Kong L, Cao J, Zhang Y, Ding W, Shen Y (2017) Risk factors for periprosthetic joint infection following primary total hip or knee arthroplasty: a meta-analysis. Int Wound J 14(3):529-536
13. Malinzak RA, Ritter MA, Berend ME, Meding JB, Olberding EM, Davis KE (2009) Morbidly obese, diabetic, younger, and unilateral joint arthroplasty patients have elevated total joint arthroplasty infection rates. J Arthroplast 24(6 Suppl):84-88
14. Tsang ST, Gaston $P$ (2013) Adverse peri-operative outcomes following elective total hip replacement in diabetes mellitus: a systematic review and meta-analysis of cohort studies. Bone Joint J 95-b(11):1474-9
15. Bedard NA, DeMik DE, Owens JM, Glass NA, DeBerg J, Callaghan JJ (2019) Tobacco use and risk of wound complications and periprosthetic joint infection: a systematic review and
meta-analysis of primary total joint arthroplasty procedures. J Arthroplast 34(2):385-96.e4
16 Gustke K (2012) Short stems for total hip arthroplasty: initial experience with the fitmore stem. J Bone Joint Surg Br 94(11 Suppl A):47-51
16. Sheth D, Cafri G, Inacio MC, Paxton EW, Namba RS (2015) Anterior and anterolateral approaches for THA are associated with lower dislocation risk without higher revision risk. Clin Orthop Relat Res 473(11):3401-3408
17. Acuna AJ, Do MT, Samuel LT, Grits D, Otero JE, Kamath AF (2022) Periprosthetic joint infection rates across primary total hip arthroplasty surgical approaches: a systematic review and meta-analysis of 653,633 procedures. Arch Orthop Trauma Surg 142(10):2965-2977
19 Pfeil J (2010) Minimally invasive surgery in total hip arthroplasty. Springer, Berlin. https://doi.org/10.1007/978-3-642-00897-9
18. Frndak PA, Mallory TH, Lombardi AV Jr (1993) Translateral surgical approach to the hip. The abductor muscle "split." Clin Orthop Relat Res 295:135-41
21 Parvizi J, Tan TL, Goswami K, Higuera C, Della Valle C, Chen AF et al (2018) The 2018 Definition of Periprosthetic Hip and Knee Infection: An Evidence-Based and Validated Criteria. J Arthroplast 33(5):1309-14
19. Hardinge $K$ (1982) The direct lateral approach to the hip. J Bone Joint Surg Br 64(1):17-19
20. Dale H, Skramm I, Lower HL, Eriksen HM, Espehaug B, Furnes O et al (2011) Infection after primary hip arthroplasty: a comparison of 3 Norwegian health registers. Acta Orthop 82(6):646-654
21. Ilchmann T, Zimmerli W, Bolliger L, Graber P, Clauss M (2016) Risk of infection in primary, elective total hip arthroplasty with direct anterior approach or lateral transgluteal approach: a prospective cohort study of 1104 hips. BMC Musculoskelet Disord 17(1):471
22. Shohat N, Goswami K, Clarkson S, Chisari E, Breckenridge L, Gursay D et al (2021) Direct anterior approach to the hip does not increase the risk for subsequent periprosthetic joint infection. J Arthroplast 36(6):2038-2043
23. Aggarwal VK, Weintraub S, Klock J, Stachel A, Phillips M, Schwarzkopf R et al (2019) 2019 Frank Stinchfield Award: A comparison of prosthetic joint infection rates between direct anterior and non-anterior approach total hip arthroplasty. Bone Joint J. 101-B(6_Supple_B):2-8
24. Hoskins W, Bingham R, Lorimer M, Hatton A, de Steiger RN (2020) Early rate of revision of total hip arthroplasty related to surgical approach: an analysis of 122,345 primary total hip arthroplasties. J Bone Joint Surg Am 102(21):1874-1882
25. Lindgren V, Garellick G, Karrholm J, Wretenberg P (2012) The type of surgical approach influences the risk of revision in total hip arthroplasty: a study from the Swedish hip arthroplasty register of 90,662 total hipreplacements with 3 different cemented prostheses. Acta Orthop 83(6):559-565
26. Mjaaland KE, Svenningsen S, Fenstad AM, Havelin LI, Furnes O, Nordsletten L (2017) Implant survival after minimally invasive anterior or anterolateral Vs. conventional posterior or direct lateral approach: an analysis of 21,860 total hip arthroplasties from the Norwegian arthroplasty register (2008-2013). J Bone Joint Surg Am 99(10):840-7
27. Purcell RL, Parks NL, Gargiulo JM, Hamilton WG (2016) Severely obese patients have a higher risk of infection after direct anterior approach total hip arthroplasty. J Arthroplast 31(9 Suppl):162-165
28. Hartford JM, Graw BP, Frosch DL (2020) Perioperative complications stratified by body mass index for the direct anterior approach to total hip arthroplasty. J Arthroplast 35(9):2652-2657
29. Shah NV, Huddleston HP, Wolff DT, Newman JM, Pivec R, Naziri Q et al (2022) Does surgical approach for total hip arthroplasty impact infection risk in the obese patient? A systematic review. Orthopedics 45(2):e67-e72
30. Wang Q, Goswami K, Shohat N, Aalirezaie A, Manrique J, Parvizi J (2019) Longer operative time results in a higher rate of subsequent periprosthetic joint infection in patients undergoing primary joint arthroplasty. J Arthroplast 34(5):947-953
31. Iorio R, Williams KM, Marcantonio AJ, Specht LM, Tilzey JF, Healy WL (2012) Diabetes mellitus, hemoglobin A1C, and the incidence of total joint arthroplasty infection. J Arthroplast 27(5):726-9.el
32. Marchant MH Jr, Viens NA, Cook C, Vail TP, Bolognesi MP (2009) The impact of glycemic control and diabetes mellitus on perioperative outcomes after total joint arthroplasty. J Bone Joint Surg Am 91(7):1621-1629
33. Pedersen AB, Mehnert F, Johnsen SP, Sorensen HT (2010) Risk of revision of a total hip replacement in patients with diabetes mellitus: a population-based follow up study. J Bone Jt Surg Br 92(7):929-934

Publisher's Note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.


[^0]:    Matthias Luger
    m.n.luger@gmail.com

    Marcel de Vries
    Devries.marcel@icloud.com
    Sandra Feldler
    Sandra_feldler@aon.at
    Günter Hipmair
    Guenter.hipmair@kepleruniklinikum.at
    Tobias Gotterbarm
    Tobias.gotterbarm@kepleruniklinikum.at

