Direct anterior versus posterior approach for total hip arthroplasty: a multicentre, prospective, randomized clinical trial

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Background: The ideal approach for a total hip arthroplasty (THA) would be kind to soft tissues, have the lowest complication rates and be easily reproducible. Although there have been several attempts to find the best approach for THA in the last decade, a definitive answer has not been found. We performed a prospective study to compare the direct anterior and posterior approaches for THA in terms of hospital length of stay, functional outcome, pain, implant position, complications and surgical time.

Methods: A prospective, randomized, multicentre clinical study was conducted between February 2011 and July 2013, with an average follow-up of 55 months. Patients undergoing the direct anterior or posterior approach for THA were enrolled. Hospital length of stay, surgical time and complications were documented. The Harris Hip Score and visual analogue scale were used to monitor functional outcome and pain until 5 years postoperatively. Radiologic analysis was used to assess implant position.

Results: Fifty-five patients (28 undergoing the direct anterior approach, 27 undergoing the posterior approach) were enrolled in this study. Length of stay, functional outcome, pain, implant position and complications were similar for the 2 approaches. There was a trend toward a better functional outcome for patients who underwent the direct anterior approach in the first 3 months postoperatively, with a peak at 4 weeks (Harris Hip Score 76.7 v. 68.7; p = 0.08). Average surgical time for the direct anterior approach was significantly longer (69.9 v. 45.7 min; p = 0.002).

Conclusion: The direct anterior approach for THA appears to be a safe and effective option. However, there is no significant difference in hospital length of stay or post-operative recovery between the 2 approaches.

Clinical trial registration: Clinicaltrials.gov, no. NCT03673514

Contexte : L'approche idéale pour l'arthroplastie totale de la hanche (ATH) serait douce pour les tissus mous, aurait le taux de complications le plus bas et serait facilement reproductible. Dans les 10 dernières années, on a tenté à de nombreuses reprises de déterminer quelle est la meilleure approche, sans obtenir de réponse concluante. Nous avons mené une étude prospective visant à comparer la durée du séjour à l'hôpital, les résultats fonctionnels, la douleur, la position de l'implant, les complications et le temps de chirurgie associés aux approches antérieure directe et postérieure pour l'ATH.

Méthodes : Un essai clinique randomisé prospectif multicentrique a été mené auprès de patients ayant subi une ATH par voie antérieure directe ou postérieure entre février 2011 et juillet 2013; le suivi moyen était de 55 mois. La durée du séjour à l'hôpital, le temps de chirurgie et les complications ont été notés. Le score de Harris pour la hanche et l'échelle analogique visuelle ont servi au suivi des résultats fonctionnels et de la douleur dans les 5 ans suivant l'opération. Des clichés radiologiques ont été analysés pour évaluer la position de l'implant.

Résultats : Au total, 55 patients ont été recrutés (28 ayant subi une ATH par voie antérieure directe, et 27, une ATH par voie postérieure). La durée du séjour, les résultats fonctionnels, la douleur, la position de l'implant et les complications étaient sensiblement les mêmes, quelle que soit l'approche utilisée. Dans les 3 premiers mois suivant l'opération, les patients ayant subi une ATH par voie antérieure directe avaient tendance à présenter de meilleurs résultats fonctionnels que les autres, en particulier à la quatrième semaine postopératoire (score de Harris pour la hanche : 76,7 c. 68,7; p = 0,08). Le temps de chirurgie moyen pour l'approche antérieure directe était significativement plus long (69,9 c. 45,7 min; p = 0,002).

Conclusion : La voie antérieure directe semble être une approche efficace et sûre. Aucune différence significative n'a toutefois été observée entre les 2 approches quant à la durée du séjour à l'hôpital ou au rétablissement postopératoire.

Enregistrement de l'essai : Clinical Trials.gov, no NCT03673514

he best approach for a total hip arthroplasty (THA) is still a matter of debate. The ideal approach would be kind to soft tissues, have the lowest complication rates and be easily reproducible. Although there have been several attempts to resolve this issue in the last decade, controversy remains.

There are 3 classic approaches for THA, each with disadvantages: the direct anterior approach, the posterior approach and the lateral approach. The main drawback of the posterior approach (PA), the most commonly used approach in the United States and probably worldwide,^{1,2} is the disinsertion of the external rotators. Posterior dislocation is a potential complication of this approach, although recent studies have showed no increased risk when adequate capsulorrhaphy or enhanced posterior soft tissue repair was performed.^{3–6} Relying on the Hueter interval, the direct anterior approach (DAA) has a steep learning curve^{7–9} but is by its very nature truly intermuscular and internervous.¹⁰ Two studies have found less iatrogenic muscular damage, as seen on magnetic resonance imaging, with the DAA than with the lateral or posterior approaches.^{11,12}

Consequently, proponents of the DAA infer that the associated postoperative recuperation will be easier and faster and could lead to a better functional outcome. Although recent literature has showed promising short-term results in some respects,^{7,13-16} other studies have failed to demonstrate long-term advantages^{13,14,17} or any advantage^{9,18-21} of the DAA over other surgical approaches to the hip. Furthermore, reported complication rates associated with the DAA have been relatively high in some series.^{21,22} Therefore, we performed a prospective study to compare the DAA and the PA in terms of hospital stay as the primary outcome and functional recovery, pain, implant position, complications and surgical time as secondary outcomes.

METHODS

Patients

This was a multicentre, prospective, randomized clinical trial that enrolled patients who underwent total hip arthroplasties between February 2011 and July 2013. The trial was registered retrospectively with ClinicalTrials.gov (NCT-03673514). Institutional review board approval was granted for this study by the Comité d'éthique de la recherche of the CIUSSS NIM Hôpital du Sacré-Coeur de Montréal (CER 2010-415). After providing written informed consent to participate in this study, patients meeting the inclusion criteria underwent surgical treatment using the DAA or PA, according to a randomization process. This process used random blocks of 2 and 4, ensuring that group allocation was equal throughout the recruitment period. Group allocation was made immediately before surgery by an independent research coworker using sequentially numbered, sealed envelopes containing the designated surgical approach.

Inclusion criteria were as follows: primary total hip replacement because of osteoarthrosis or osteonecrosis, and patients older than 50 years. Patients were excluded if they suffered from inflammatory arthritis, had previously undergone any ipsilateral hip surgery, had a proximal femoral deformity, had a body mass index over 40, had an active infection, had severe contralateral hip disease (Tönnis grade 3 or any dysplasia), had neuromuscular pathology or required structural bone grafts. Patients were enrolled by the 2 surgeons involved in the study (H.G, B.B.).

Clinical and radiologic assessment

Clinical follow-up was scheduled at 2 weeks, 4 weeks, 3 months, 6 months, 1 year, 2 years and 5 years postoperatively. The Harris Hip Score (HHS) and visual analogue scale (VAS) were used to monitor pain and functional outcome at these visits. Data were collected by an independent research assistant at the outpatient clinic of both surgeons.

Radiologic analysis assessed implant position, limb lengthening, and potential implant-related complications. Radiography was performed at subsequent follow-up visits and consisted of an anteroposterior (AP) pelvic radiograph and a lateral projection of the hip.23 Two independent observers (K.M., P.D.), not involved with patients or surgeries, reviewed all postoperative radiographs independently. Cup version was measured with the Lewinnek method. Radiographic analysis of radiolucency and sclerotic changes around the implants was performed using the DeLee zones for the acetabulum and the Gruen zones for the femoral stem. An orthopedic templating software (Orthoview) was used for surgical planning. Preoperatively, an AP scaled radiograph of the pelvis was obtained on which the desired limb length and offset were planned, according to the contralateral side. During surgery, abductor tension (shuck test and hip pistoning) and the kick test were performed by the surgeon to confirm the accuracy of preoperative planning and hip stability.

Surgical procedure

Two fellowship-trained surgeons in 2 separate hospitals performed all procedures. Before the initiation of this study, both surgeons had each performed more than 100 cases with each approach.

Patients underwent either the PA or DAA. The PA to the hip, which is considered to be a minimally invasive approach because of the smaller operative scar (under 10 cm), has been described by many authors^{24–27} and could potentially yield better results.²⁸ The modified Hueter approach, based on the Smith-Peterson approach, was performed for the direct anterior minimally invasive surgery.^{29,30} A traction table was used for DAA as the surgeons were trained to use this method. No intraoperative fluoroscopy was used for implant confirmation. The 2 groups of patients received the same implants (Quadra-H stem and Versacup hip system, Medacta), with metal on polyethylene bearing. All implants were non-cemented. Surgical approaches were compared in terms of surgical time (skin to skin), length of hospital stay and complications.

Study blinding

Because of the nature of the intervention, it was impossible for the investigator and the patients to be blinded. However, the decision to discharge patients was made by physiotherapists who were blinded to treatment group assignment, and they made the decision on the basis of objective criteria. These criteria were as follows: (a) the patient had to be able to autonomously transfer from bed/chair to the upright position, (b) the patient had to be able to walk with a walking aid, (c) the patient had to be able to climb stairs in a safe way and (d) the patient's pain had to be controlled by painkillers. No restrictions were recommended for either group. The same rehabilitation and pain protocols were used for the 2 groups. In addition, statistical analyses were performed by an independent consultant who remained blinded to treatment group assignment.

Statistical analysis

The sample size was determined by power analysis (statistical power of 0.80, α of 0.05) with a large effect size (0.80) and an allocation ratio of 1 using G*Power,³¹ yielding 26 patients per group for a total of 52 patients. Primary and secondary outcome continuous variables were analyzed using the Student *t* test. The Mann–Whitney *U* test was used when data were not equally distributed. Nominal variables were analyzed using χ^2 tests. All *p* values were 2-tailed, and the significance level was set at 0.05. All analyses were performed using SPSS version 25.0 software.

RESULTS

Fifty-five total hip athroplasties were performed on 50 patients between February 2011 and July 2013. Demographic data did not differ in terms of age, American Society of Anesthesiologists (ASA) class and body mass index. Only sex distribution differed significantly between the 2 groups (Table 1). One patient in the PA group was lost to follow-up after 6 months; 4 patients (2 in the PA group, 2 in the DAA group) were lost to follow-up after 1 year. All 50 other patients completed the 5-year follow-up. In our cohort of 55 patients, the percentage of hip dysplasia was similar in the 2 groups (7.1% in the DAA group v. 7.4% in the PA group). Only dysplasias of Hartofilakidis type A were present; there were no low or high hip dislocations.

The mean length of hospital stay was similar in the 2 groups: 3.5 days (range 2–10 d) for PA and 3.8 days

Table 1. Characteristics of patients who underwent the direct anterior or posterior approach for total hip arthroplasty

	Approach		
Characteristic	PA n = 27	DAA n = 28	p value
Age, yr, mean ± SD	68.9 ± 8.8	70.4 ± 9.1	0.52
ASA class, mean ± SD	2.0 ± 0.8	1.8 ± 0.7	0.38
Body mass index, mean ± SD	26.5 ± 4.3	27.6 ± 4.4	0.36
Sex, no.			0.042
Men	18	11	
Women	9	17	
ASA = American Society of Anesthesia	ologists; DAA = dire	ct anterior approac	h; PA =

ASA = American Society of Anesthesiologists; DAA = direct anterior approach; PA = posterior approach; SD = standard deviation.

Table 2. Length of stay, surgical time, and pain and functional outcomes in patients who underwent the direct anterior or posterior approach for total hip arthroplasty

	Appr	Approach	
Variable	PA n = 27	DAA n = 28	<i>p</i> value
Length of stay, d, mean ± SD	3.5 ± 2.2	3.8 ± 1.8	0.53
Surgical time, min, mean ± SD	45.7 ± 17.9	59.9 ± 12.7	0.002
VAS score, mean ± SD			
Preoperative	6.9 ± 2.1	5.0 ± 2.4	0.029
Postoperative			
2 wk	2.1 ± 2.0	2.0 ± 2.0	0.79
4 wk	1.6 ± 1.9	1.4 ± 2.0	0.63
3 mo	1.1 ± 1.9	1.0 ± 1.7	0.66
6 mo	0.4 ± 1.0	0.4 ± 0.8	0.61
1 yr	0.6 ± 1.2	0.3 ± 0.5	0.38
2 yr	1.0 ± 1.9	0.5 ± 0.8	1.00
HHS score, mean \pm SD			
Preoperative	48.2 ± 10.1	52.1 ± 19.7	0.66
Postoperative			
2 wk	60.0 ± 15.1	66.9 ± 17.1	0.12
4 wk	68.7 ± 16.8	76.7 ± 16.4	0.08
3 mo	83.3 ± 15.1	88.4 ± 11.8	0.18
6 mo	90.3 ± 12.3	90.1 ± 11.3	1.00
1 yr	91.4 ± 13.0	94.4 ± 8.0	0.72
2 yr	88.7 ± 20.0	89.4 ± 11.9	0.58
5 yr	80.0 ± 20.4	82.0 ± 19.8	0.72
DAA = direct anterior approach; visual analogue scale; SD = stan		e; PA = posterior appr	oach; VAS =

(range 2–7 d) for DAA (Table 2). Functional preoperative HHS scores were similar for the 2 groups and HHS scores indicated a trend toward faster functional recovery with the DAA until 4 weeks postoperatively (Mann–Whitney U test, 76.7 v. 68.7; p = 0.08). However, no statistical difference was found in terms of overall functional recovery, with similar VAS and HHS scores at each visit for both groups. Figure 1 shows the HHS scores over time. None of the scores for any of the subcategories of the HHS were significantly different at any postoperative time (Table 2).

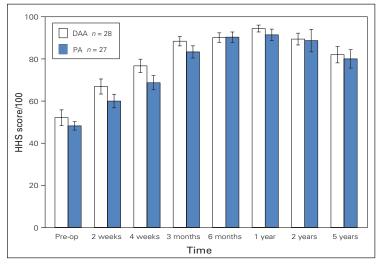


Fig. 1. Comparison of Harris Hip Score (HHS) values preoperatively and at different time points postoperatively for patients who underwent the direct anterior approach (DAA) and posterior approach (PA) for total hip arthroplasty. The maximum score possible is 100. Preop = preoperative.

Preoperative VAS pain scores were significantly different between the groups (Mann–Whitney *U* test, p = 0.029), but postoperative scores were similar at each visit for the 2 groups (Table 2).

Radiologic analysis confirmed that the implants were well positioned in terms of version (Student *t* test, DAA group 26.9°, range 11° to 45°; PA group 21.9°, range 4° to 41°; *p* = 0.103) and abduction (Student *t* test, DAA group 43.3°, range 20° to 56°; PA group 39.8°, range 24° to 49°; *p* = 0.064), with both surgical approaches (Table 3). The data suggest more anteversion and abduction with the DAA, but none of these values were significant. Cup abduction was statistically different between the 2 surgeons, with a mean difference of 4.8° (Student *t* test, *p* = 0.013). With regard to the mean planned leg length discrepancy, the reconstructed hip was 2.2 mm (range –14 to 9 mm) shorter than the ideal planned length for DAA, and a 1.0 mm (range –9 to 10 mm)

Table 3. Radiologic assessment of postoperative pelvic

	Appr		
Variable	PA n = 27	DAA n = 28	<i>p</i> value
Implant			
Cup anteversion, degrees, mean ± SD	21.9 ± 13.4	26.9 ± 8.6	0.10
Cup abduction, degrees, mean ± SD	39.8 ± 5.4	43.3 ± 8.4	0.06
Preoperative LLI, mm, mean ± SD	-2.4 ± 4.3	-2.2 ± 4.8	0.87
Planned leg length discrepancy, mm, mean ± SD	1.0 ± 5.6	-2.2 ± 6.8	0.06

leg lengthening was reported, compared with the ideal planned length, for PA (Student *t* test, p = 0.061) (Table 3). Acetabular and femoral implants showed no migration or progressive radiolucency at the latest follow-up.

Few complications were documented for the 2 groups. Two intraoperative periprosthetic fractures occurred in the PA group and were found to be stable (not requiring fixation) according to surgeon assessment: 1 occurred on the acetabular side and the other on the femoral side. These 2 patients were advised to refrain from weight-bearing activities for a period of 6 weeks; both patients had good outcomes and did not require revision surgery. One patient in the DAA group experienced an early prosthetic joint infection with *Enterococcus faecalis* and was successfully treated with débridement, lavage and intravenous antibiotics.

Surgical time was significantly longer in the DAA group (59.9 min, range 24 to 85 min) than in the PA group (45.7 min; range 26 to 85 min), for a mean difference of 14 minutes (Mann–Whitney U test, p = 0.002). Surgical duration was significantly different between the 2 surgeons (Student *t* test, p < 0.001); however, each surgeon was significantly faster using the PA (Table 2). The distribution of patients who underwent the DAA and PA was similar for the 2 surgeons.

DISCUSSION

The DAA approach has been widely promoted in recent years as the only truly minimally invasive hip approach, using the intermuscular and internervous interval. The major benefit of the DAA is thought to be a shorter length of stay, accelerated rehabilitation and less postoperative pain. To our knowledge only a few randomized controlled trials have compared clinical outcomes between the DAA and PA,13,32-34 and there is a need for larger multicentre, prospective, randomized controlled trials, as these are sparse.³⁵ Higgins and colleagues concluded, following a meta-analysis, that current evidence comparing outcomes for the DAA versus the PA failed to demonstrate clear superiority of either approach.³⁶ Our findings suggest that the choice of surgical approach should be based on patient characteristics and surgeon experience and preference. As patient satisfaction and clinical follow-up were similar with the 2 approaches and surgical duration was shorter with the PA, both surgeons favour the PA.

Length of stay was similar for the 2 approaches in our study (mean 3.5 d for PA and 3.8 d for DAA). The length of stay in our study may appear to be overly long, but it could have been because of the exacting criteria for hospital discharge, which were applied for both approaches. There was a recent decrease in length of stay for our THA patients resulting from better patient education, which also increased the number of ambulatory THA.

The literature suggests that functional recovery is accelerated in patients who undergo the DAA. However, most of these studies have been retrospective and have had small sample sizes. Barrett and colleagues assessed this issue in a randomized controlled trial comparing 43 patients who underwent the DAA with 44 patients who underwent the PA and showed that DAA patients performed better in the immediate postoperative period.¹³ More DAA patients were climbing stairs normally and walking freely at 6 weeks. There were no significant differences at later time points. Studies by Reininga and colleagues and Ward and colleagues found improved gait after surgery for both approaches without any significant difference between the 2.^{32,33} In the present study, functional outcome seemed to improve in the early postoperative period (first 3 months), with an HHS score that almost reached statistical significance at 1-month follow-up. However, this was not borne out in the long-term follow-up. Miller and colleagues reviewed 13 prospective studies comparing DAA and PA in the first 90 days postoperatively and found better hip function, less pain and lower narcotic consumption in the DAA group.¹⁶

Findings similar to those of our study have recently been published.^{32–34} A prospective clinical trial of 54 patients, randomly assigned to undergo the DAA or PA, showed a faster voluntary cessation of use of all walking aids in the DAA group (22 v. 28 d). At 3 weeks, Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC) function scores and SF-12 mental scores favoured the PA group. There were no differences in HHS, SF-12 or WOMAC scores at long-term follow-up.³² Another study comparing the DAA and mini-posterior approaches found no difference in return to activities of daily living, length of stay, complication rate and use of pain medication. However, the DAA group had a higher HHS score at 8 weeks, but these patients also had a higher VAS score for pain in the hospital, and fewer patients returning to work.³³ In 2017, Cheng and colleagues also reported a randomized trial involving 72 patients who underwent either the DAA or the PA approach for primary hip arthroplasty.³⁷ They found no difference between the groups with regard to total scores for primary outcomes with the WOMAC, EuroQol and 10 m walk test. Functional outcomes may be improved in the early postoperative period when the DAA is performed; however, this advantage seems to be short lived with negligible longterm benefits. Furthermore, postoperative pain was comparable in both groups at each recorded time.

Implant positioning showed that cup anteversion seemed to be greater with the DAA, but this difference was not significant. Conversely, Barrett and colleagues found significantly more inclination and less anteversion with the DAA, suggesting that it is probably a surgeon-dependent factor.¹³ Surgeons using an anterior approach will usually avoid too much anteversion to prevent anterior dislocation. All cups were placed within the Lewinnek safe zone with no outliers. There was no statistically significant difference in planned leg lengthening between the cohorts. Previous reports have demonstrated increased risks of postoperative complications and/or poor outcomes during the surgeon's learning curve period when adopting the DAA.^{21,35} Others have indicated a theoretical advantage for the DAA in terms of dislocation risk over the PA. In the present study, a low complication rate was seen with both approaches with no dislocations. Only 1 patient was reoperated for infection and then for a periprosthetic fracture of the femur after sustaining a fall.

Surgical time was significantly longer with the DAA than with the PA. Two meta-analyses concluded that the mean operative time was significantly longer with the DAA and many also reported a steep learning curve.^{35–38} The only study that found a significantly longer surgical time with the PA³⁹ compared 1 surgeon who performed the DAA surgeon with 2 surgeons who performed the PA. Therefore, these results could be attributed to the surgeon.

The strengths of our study include the random assignment of patients, with few lost to follow-up, and a complete clinical and radiologic follow-up at different periods for a mean of 55 months. Patients were evaluated for discharge by a physiotherapist who was blinded to study group allocation, according to established criteria, unlike a previous publication by Barrett and colleagues.¹³

Limitations

Limitations of the study include the relatively small number of patients. However, most studies analyzing prospective outcomes following orthopedic surgeries have had a similar sample size. Patients were not equally distributed in the 2 groups in terms of sex, and preoperative VAS score was higher in the PA group, which could have affected the results.

CONCLUSION

This study could not demonstrate significant advantages for the PA or the DAA in the early nor in the late postoperative recovery. Length of hospital stay was similar in the 2 groups, with a low complication rate for both approaches. Surgical time was greater in the DAA group.

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Competing interests: G. Laflamme and S. Leduc have received consulting fees from Stryker, outside the submitted work. B. Benoit has received consulting fees from Medacta, Stryker and Bioventus, outside the submitted work. The Hôpital Sacré-Coeur de Montréal has received funding for research and educational purposes from Arthrex, ConMed, Depuy, Linvatec, Medacta, Smith & Nephew, Stryker, Synthes, Tornier, Wright and Zimmer Biomet. No other competing interests were declared.

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Data sharing: The data sets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Contributors: K. Moerenhout designed the study. P. Derome, S. Leduc, H. Gaspard and B. Benoit acquired the data, which K. Moerenhout, P. Derome, Y. Laflamme analyzed. K. Moerenhout, P. Derome, Y. Laflamme wrote the manuscript, which Y. Laflamme, S. Leduc, H. Gaspard and B. Benoit critically revised. All authors gave final approval of the article to be published.

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