Does the intra-operatively measured leg length correction compare to the post-operative radiograph in total hip replacement surgery?

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Abstract

Background: This study aims to analyse the accuracy of the Vertical Measurement System™ (VMS) in assessing the leg length correction (LLC) during total hip arthroplasty (THA) by comparing the intra-operative measurements to the radiographic measurements obtained six weeks post-operatively.

Patients and methods: A prospective cohort study was conducted in which patients undergoing primary THA were enrolled at two centres in Cape Town, over a period of 19 weeks. THAs were performed by four surgeons. Pre-operative leg length discrepancy (LLD) measurements were obtained in 92 patients. The VMS was used to predict intra-operative LLC, and this measurement was compared to the post-operative LLC measured on the six-week follow-up X-ray. These measurements were statistically compared using the Mann–Whitney U test.

Results: The difference between the intra-operative VMS calculation and the six-week radiological measurement was not significant (p>0.05), with the difference in their mean values being 0.1±3.3 mm. In the cohort, 82% of the patients (n=75) were within 5 mm of the target LLC, and 96% of patients (n=88) were within 10 mm of the target LLC. The mean absolute residual LLD at six weeks was 3.2±3.1 mm.

Conclusion: The intra-operative LLC measurement obtained using the VMS accurately predicts the six-week post-operative radiographic LLC measurement.

Level of evidence: Level 4

Keywords: total hip replacement, leg length discrepancy, leg length correction, vertical measurement system, comparative study, longitudinal study


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Introduction

Total hip arthroplasty (THA) is one of the most successful orthopaedic operations, with high patient satisfaction and low revision rates. Accurate leg length correction (LLC) in THA is imperative for a good clinical outcome. Therefore, equalisation of leg length remains one of the primary objectives of THA. Nevertheless, leg length inequality remains a recognised complication of the procedure. Leg length discrepancy (LLD) accounts for 5% of all medical errors, as per the Joint Commission on Accreditation of Healthcare Organizations (JCAHO), and remains one of the leading causes of litigation against orthopaedic surgeons in the USA. The complications of LLD after THA include sciatic, femoral and peroneal nerve palsies, hip or low back pain, abnormal gait and posture, and aseptic loosening.

The incidence of LLD after THA has been reported to range from 1% to 27%, with some studies reporting values of LLD from 3 mm to 70 mm (mean 3–17 mm). Small discrepancies may be a source of dissatisfaction for some patients; however, several studies have shown that up to 10 mm of LLD may be well tolerated by most patients. Leaving the operated leg short seems to be more acceptable to patients than lengthening the operated leg, since patients can detect relatively small increases in length, and are particularly unhappy if they have to wear a shoe raise on the contralateral, unoperated side.

The importance of attempting to equalise leg length is recognised among all orthopaedic surgeons in all sub-specialties, not just arthroplasty surgeons. This is attested to by the large amount of literature on LLD in THA. In order to mitigate the occurrence of LLD after THA, various methods have been used. These include pre-operative templating, a wide range of intra-operative techniques, such as measurements from a fixed point on the pelvis using a suture or ruler, to drilling Steinman pins or K-wires into a point in the pelvis. More recently, computer navigation has been used. In order to achieve consistent LLC, the surgeon needs to be familiar with the various surgical techniques and the accuracy of these in the clinical or operative setting.

The objective of this study was to assess the accuracy of a method we use to quantify the LLC intra-operatively, namely the Vertical Measurement System (VMS), and compare it to six-week post-operative X-rays. The basic principle of this system is that the difference in vertical height between the excised femoral head and neck, and the combined vertical height of the implants, determines the change in leg length. Our hypothesis was that the LLC measured using the intra-operative VMS method would equal the post-operative radiological measurement.

Patients and methods

A prospective cohort study was conducted at two hospitals in the Western Cape, South Africa. Patients who were booked for THA were invited to participate, after careful explanation of the study design and methods. Informed consent for the study was obtained from all patients. Inclusion criteria were all patients undergoing primary THA, as per standard protocols utilised in the arthroplasty units at the two hospitals. Exclusion criteria were THAs performed for trauma (fractures of the femoral neck or pelvis) and revision THA. Patients were recruited between May and October 2019, over a period of 19 weeks.

Pre-operative assessment

Prior to surgery (at the routine pre-operative clinic visit), clinical assessment of the true and apparent leg length was performed to exclude other causes of LLD such as hip adduction, abduction or flexion contractures, or knee flexion contractures. Digital X-rays were obtained using the Philips IntelliSpace PACS Enterprise system. A standard AP pelvis standing X-ray, scaled using a radiological sphere marker at the level of the greater trochanter, was used for planning. OrthoView Digital Planning software was used for pre-operative templating, sizing and positioning of implants and calculation of the pre-operative radiological LLD. The method described by Woolson, using the distance measured between a line drawn at the inferior aspect of each acetabular teardrop (the reference line) and the medial vertex of each lesser trochanter, was used to measure LLD (Figure 1). The three possible pelvic reference points include the inferior aspect of the obturator foramen, the ischial tuberosities, and the acetabular teardrop. The teardrop is the most reproducible and accurate when calculating limb length discrepancy. This measurement, in combination with the clinical assessment of LLD, was used to inform the intra-operative LLC to be achieved.

Intra-operative measurement and calculation

The THAs were performed by four surgeons at two hospitals. Each THA proceeded in the routine manner, utilising the modified Hardinge or direct anterior approach. Implanted components

![Image 1](https://via.placeholder.com/150)

**Figure 1.** The method used by Woolson to measure LLD. A reference inter-teardrop line is drawn between the most inferior aspect of each teardrop. The distance to the medial vertex of each lesser trochanter is measured (WoA and WoN).

![Image 2](https://via.placeholder.com/150)

**Figure 2.** The measurement jig utilised by the Vertical Measurement System (VMS) to measure the vertical height of the excised femoral head and neck.
were mostly Triloc, Summit and C-stem stems with Pinnacle cups (De Puy Synthes, Warsaw, IN, USA), while a small proportion were Accolade stems and Tritanium cups (Stryker, Kalamazoo, MI, USA). After the femoral neck osteotomy, the vertical height (VH) of the excised bone (resection measurement) was measured using the VMS or Vertical Measurement System™ (Peninsula Orthopaedics, Cape Town, South Africa) jig (Figure 2). Acetabular and femoral preparation, trial implantation and reduction were performed, and the hip tested for range of movement (ROM), stability and tension. The resection measurement and implanted component data were then utilised by the available application (VMS), an online calculator with a database of implant sizes and measurements that obviates the need to use multiple charts, to calculate the LLC. The difference between what is resected, i.e. the height of the excised femoral head and neck (VH), and the height of the implanted components (IC) (Figure 3) determines the LLC.14

At this point, if it was found that the LLC achieved (using the VMS system) did not match what was planned (as per the pre-operative X-ray measurement), intra-operative adjustments were made to further correct the leg length, until the objective was achieved. The surgery was concluded in the normal manner.

Post-operative

Standard rehabilitation protocols were followed, and the patients were followed up at six weeks. Standardised, calibrated X-rays and templating software were again utilised to measure the radiological LLC achieved. This radiological LLC was compared to the intra-operative LLC measurement provided by the VMS.

Statistics

All data analyses were performed using IBM SPSS ver. 25 (Armonk, New York, USA) and G*Power ver. 3.1.9 (open source).15,16 The distribution of VMS and X-ray measurement data were analysed using the Shapiro-Wilk test for normality. The two sets of measurements were compared using the Mann–Whitney U test for statistical significance. The cut-off for type I error (α) was set at 0.05.

Results

For this study, 98 patients were enrolled over the period of 19 weeks. Prior to the six-week follow-up, one patient died from an unrelated cause. A further four patients were later excluded from the final analysis due to incomplete data, and one patient failed to return for their six-week follow-up. This left 92 patients who completed the six-week follow-up and whose data was complete for analysis. Baseline characteristics of the study group are listed in Table I.

The difference between the means of the VMS measurements and the X-ray measurements was –0.1±3.3 mm (Figure 4). The mean absolute measurement difference between the two sets of values was 2.4±2.2 mm. The difference of each patient’s values (VMS and X-ray) was plotted against their mean (Figure 5). The mean difference of all these values was very close to zero, which was ideal, and most measured differences were found to lie within the 95% confidence interval.

When compared to the target LLC decided on pre-operatively, the mean absolute residual LLD post-op was 3.2±3.1 mm. Of the 92 patients, 82% (n=75) had a residual post-operative LLD of ≤5 mm, while 96% patients (n=88) had an LLD of ≤10 mm.

Discussion

The primary goals of THA include pain relief and the restoration of normal hip biomechanics, gait and function. However, restoring or maintaining equal leg lengths is critical for patient satisfaction and return to function. The orthopaedic literature is replete with articles on LLD, the effects thereof, and methods to achieve adequate LLC during THA. Nevertheless, the amount of LLD at which it becomes...
clinically significant, or that leads to symptoms, is still debated. Generally, an LLD of less than 10 mm is widely accepted.\textsuperscript{11} Beard et al. found patients had worse Oxford Hip scores at three years if LLD was greater than 10 mm.\textsuperscript{17} Our clinical aim was to achieve equal leg lengths since even small discrepancies are associated with functional impairment and pain.\textsuperscript{18,19}

In our study, the desired LLC was decided on pre-operatively, using a combination of measuring the LLD on a templating pelvic X-ray and clinical measurement. We then aimed to achieve this LLC intra-operatively, by using the VMS. Intra-operative adjustments were therefore possible (in component sizing and positioning), allowing restoration of leg length to near equal.

When comparing the intra-operative VMS measurements to the six-week post-operative radiographic measurement, there was no statistically significant difference (p>0.05) between the two sets of values. The mean absolute difference of 2.4±2.2 mm is very similar to the values quoted in other studies, where an intra-operative method was compared to the post-operative radiograph. Barbier et al.\textsuperscript{20} utilised a mechanical measurement device (LOOD – length and offset optimisation device) fixed to the pelvis to correct LLD, and the mean deviation from target length was 2.3 mm (range 0.04–10.6 mm). Other studies have reported post-operative radiographic LLD of between 1.8 mm and 3.5 mm.\textsuperscript{21,22}

Using intra-operative fluoroscopy is an available option, particularly in the anterior approach where supine positioning is conducive to imaging, as discussed by Austin et al., who compared two different techniques of LLC.\textsuperscript{23} Using a radiographic overlay technique, the LLD was 4.8 mm, and their transverse rod method yielded a LLD of 4.4 mm. However, this involved increased surgical time, radiation exposure and increased surgical cost.

More invasive measures have been utilised, which involve fixing a reference device into the pelvis and obtaining measurements to the greater trochanter or other reference point on the femur. The reference can be iliac fixation pins, intra-operative callipers, infracotyloid pins, and fixed suture lengths. In order for these devices to work properly, the operating table must be level with the floor and the position of the hip must be reproduced precisely in all planes before and after reconstruction is performed.\textsuperscript{24}

Ranawat et al. used a Steinman pin fixed to the ischium in the posterior acetabulum and achieved LLD<6 mm in 87% of their cases.\textsuperscript{25} Shiramizu et al. compared a series of patients operated on with or without the use of a calliper fixed to the anterior superior iliac crest, and found a mean post-op LLD of 2.1 mm using the calliper versus 8.2 mm without.\textsuperscript{26} A plethora of other examples of similar techniques have been reported. However, due to these techniques having their own problems – inconsistent leg positions during measurement, extra skin incisions, additional invasiveness of inserting devices into the pelvis, reference pins or devices loosening during surgery, greater surgical time and greater cost – most of them are not widely used.

More recently Tagomori et al. proposed a simpler intra-operative technique of LLC. They utilised a reference mark cut into the posterior acetabular wall with a saw and referenced this off a marking on the greater trochanter. Their measurement error, as calculated by intra-operative measurement versus post-operative CT LLD measurement, was 1.9±1.4 mm.\textsuperscript{27}

Modern advancements in arthroplasty include the use of computer-assisted navigation to enhance the accuracy of implant placement. This method of computer-assisted surgery (CAS) uses two different techniques, i.e. imageless and image-based (using CT, MRI or intra-operative fluoroscopy). Imageless systems use a generic simulated model, whereas CT-based systems allow
visualisation of a patient-specific model.²⁹ CAS systems require the registration of landmarks on the pelvis and femur. This requires placement of a reference frame on the pelvis, commonly involving placement of Steinman pins or similar into the iliac crest, and other landmarks on the pubis sometimes requiring mini incisions to accurately locate them. Femur landmarks are registered using a dynamic sensor array, which the surgeon controls. This intra-operative method can lead to complications during surgery, including failure to calibrate the CAS station and fracture of the iliac crest, greater trochanter and distal femur when inserting the pins for the sensor arrays.

In a study by Brown et al., where CAS was compared to conventional freehand technique, no difference was found in component positioning, leg length and Harris Hip Scores (HHS) in their series. They reported an increased operative time of 18 minutes in the CAS group, increased blood loss (69 ml), and a higher cost of surgery, with no additional benefit over freehand THA.²⁸ In contrast, Elapparajda et al. used navigation in a series of 152 THAs, and produced very good results, with 96% of THAs restoring the leg length to within 6 mm of the contralateral side. They also reported minimal extra surgical time or surgical cost required in the navigated THAs.²⁹ Similarly, Renkawitz et al. compared the intra-operative values provided by the CAS system they used to the post-operative LLC measured on radiographs, and found a high degree of correlation between the two measurement methods, and recommended CAS as a good intra-operative tool.³⁰

According to Rajapaul and Rasool, CAS enables the surgeon to more accurately and reproducibly correct leg length, with fewer outliers and no major complications. However, the improved accuracy does not translate into better outcome scores, and the technique is associated with complications including fractures, pin-site infections and pain.³¹ Longer-term studies are required to assess the effect of CAS on implant longevity and revision rates.

All the methods discussed here have their drawbacks. Some intra-operative tools are invasive, cumbersome or expensive; many are not user-friendly or accurate enough; more modern tools have steep learning curves, are very costly to acquire and have potential complications with their use. A simple, accurate and reliable method that is easy to use, and that gives live feedback or results, allowing intra-operative adjustments to be made in order to accurately achieve the desired LLC, would be the panacea of LLC tools. However, all tools have their drawbacks, and none are perfect. Even with the most accurate method, there is always the possibility of human error, which could skew the effectiveness of whatever method is used to correct LLC. In this study, we found that the VMS method offers the surgeon a reliable, accurate, simple and inexpensive method of quantifying LLC intra-operatively, where adjustments can be made to fine-tune the outcome. Provided that the surgeon pays careful attention while templating and with intra-operative measurements, the VMS can accurately predict the post-operative radiographic LLC.

Conclusion

In this study, we found that the VMS method offers the surgeon a reliable, accurate, simple and inexpensive method of quantifying LLC intra-operatively, where adjustments can be made to fine-tune the outcome. Provided that the surgeon pays careful attention while templating and with intra-operative measurements, the VMS can accurately predict the post-operative radiographic LLC.

Ethics statement

Prior to commencement of the study, ethical approval was obtained from the following ethical review boards: University of Cape Town, Faculty of Health Sciences, Human Research Ethics Committee, HREC REF: 117/2019; Institutional Review Board (IRB) Number: IRB00001839. Informed written consent was obtained from all patients prior to being included in the study. The authors declare that this submission is in accordance with the principles laid down by the Responsible Research Publication Position Statements as developed at the 2nd World Conference on Research Integrity in Singapore, 2010. All procedures were in accordance with the ethical standards of the responsible committee on human experimentation (institutional and national) and with the Helsinki Declaration of 1975, as revised in 2008.

Declarations

The authors declare authorship of this article and that they have followed sound scientific research practice. This research is original and does not transgress plagiarism policies.

Author contributions

ZM contributed to the conceptualisation, study design and data collection, performed surgeries and prepared the manuscript. MBN contributed to conceptualisation and design, performed surgeries, supervised the study and reviewed the manuscript. RD contributed to data and statistical analysis, as well preparation of graphs and manuscript review.

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References


