# ORIGINAL ARTICLE

# Correlation of post-operative anterior knee pain in two different total knee arthroplasty techniques: the conventional instrumentation (CI) and the patient specific instrumentation (PSI).

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# ABSTRACT

**Purpose:** Evaluation of the clinical outcome, as assessed by the Visual Analogue Scale score and clinical questionnaires, in total knee arthroplasty undertaken with patient specific or conventional instrumentation. **Methods:** This is a prospective comparative clinical study of 115 consecutive patients who underwent total knee arthroplasty in two different orthopedic centers. Patients were assessed using the 0-10 Visual Analog Scale in different activities, the Tegner -Lysholm Functional Score, the Knee Society Scores (KSS), measured preoperatively and at 3-6-12 months postoperatively.

**Results:** Both groups improved significantly over time on all score clinical outcomes. Statistically significant differences were observed between the two groups in the KSS knee score postoperatively at 6 and 12 months (p=0.007 and p=0.004 respectively), and in the Tegner- Lysholm score only at 6 months postoperatively (p=0.001), both in favour of patient specific intrumentation group. The mean KSS was 91.32±4.29 and 93.15±4.72 for the conventional and patient specific groups respectively at 12 months follow up (p=0.063). No statistically significant difference was found at 12 months between two groups when the visual analog scale was measured related to the activity of standing up from the sitting position (conventional group: 1.15±0.36 versus patient specific: 1.10±0.30, p=0.380).

Conclusion: We did not find major significant differences for pain scores, functional scores and clinical out-

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KEYWORDS: total knee arthroplasty, anterior knee pain, patient specific instrumentation, visual analog scale, Tegner - Lysholm Score, Knee Society Score

### Introduction

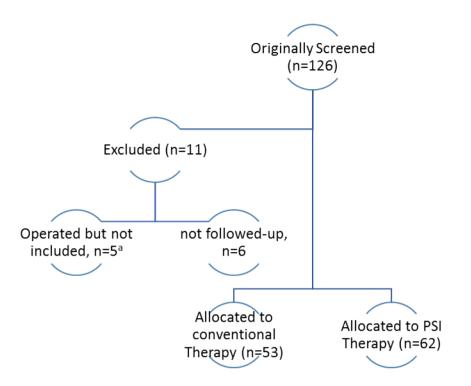
Patient specific instrumentation (PSI) is a modern technique in total knee arthroplasty (TKA), aiming to facilitate the implant of the prosthesis. PSIs were introduced to increase the accuracy of the surgical technique and avoid issues related to the complexity of the navigation system, such as procedural costs, surgical time, and learning curve. PSI is expected to improve component alignment and positioning, postoperative functional recovery, and patient satisfaction (16,17). The customized cutting blocks of the PSI are generated from pre-operative computer-aided three-dimensional (3D) reconstruction, 3D printing from a disposable template, using computed tomography (CT) or magnetic resonance imaging (MRI), aiming at an accurate intraoperative placement of the cutting blocks and an accurate osteotomy (16,17). A correct surgical plan is mandatory for a good surgical implant. (18,19,23) The PSI guide takes into account any slight deformities or osteophytes and applies preoperative planning for bone resection, using the pre-determined implant size, position, and rotation. PSI is hypothesized to have advantages with respect to improving component alignment, shortening the surgical time and length of hospital stay, and decreasing perioperative blood loss. Many manufacturers have invested in PSIs (19,21,26). Large debates have taken place about this topic during the last years and, now, there is no consensus in the.. current . literature regarding the accuracy and reliability of PSI, as many studies have shown controversial and inconsistent results. (18,20,21,22,24,25,26,27) In a recent comprehensive systematic review and meta-analysis, Gong et al (30) concluded that PSI has advantages for axial alignment of the femoral component, operative time, and perioperative blood loss compared to conventional instrumentation (CI) total knee arthroplasty. However, no significant differences were observed between PSI and CI with respect to the alignment of the remaining components, number of outliers, and length of hospital stay.

As far as we know there is no clinical study comparing especially the postoperative anterior knee pain (AKP) after primary TKA between CI and PSI of the exact same knee prosthesis. Background of the hypothesized less AKP after TKA with patient specific instrumentation were the causal relationship between malalignment and malrotation component mistakes after TKA and more severe amount of AKP on the one hand (41-49), and the already published superiority of the PSI TKA in the coronal and sagittal alignment and component rotation in comparison to the CI TKA on the other hand (50-55).

## Materials and Method

They were 53 patients who took part in the study in the first Orthopaedic center between 2015-2016 (group 1) and 62 patients in the second Orthopaedic center between 2017-2018 (group 2) (Figure 1, Table 1). The inclusion criteria were correct prosthetic components alignment (34), complete one-year follow-up scores, X-rays, and surgery performed by the same OP team for each Orthopaedic center, supervised by an Orthopaedic fellow-trained in Joint Replacement Surgery. Exclusion criteria were major postoperative complications, inflammatory systemic disease, impaired cognitive status and inability

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*Figure 1* Flow chart of patients to be considered in this study (a=Not included due to post-operative complications, Haematoma, Fracture, Infection, Iatrogenic Intra-OP mistakes)

TABLE 1.						
Demographic characteristics of patients (SD= standard deviation).						
DEMOGRAPHICS						
FACTOR	Conventional	PSI				
NUMBER OF PATIENTS	53	62				
MEN/WOMEN RATION	14/39	12/50				
AGE (YEARS) MEAN (SD)	70,11 (3,55)	70,35 (3,84)				
BMI MEAN (SD)	27,41 (3,10)	25,91 (4,5)				
ASA GRADE, NUMBER OF PATIENT						
1	7	3				
2	40	50				
3	6	9				
4	0	0				

for follow up. Finally, 11 patients were excluded from the originally 126 registered patients. Written consent was obtained from each patient of each group, after detailed information about the study, and ethical approval was obtained from each respective local ethical committee about this clinical study.

All patients of the study received no patellar pros-

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Comparison of Anterior Knee Pain (AKP) at different activities between groups during the observation period

period								
АКР	GROUP	3 MONTHS	6 MONTHS	12 MONTHS	P-VALUE WITH IN GROUP	TIME INDEPENDENT\$ MEAN (95%CI)		
	CI TKA	2.74±0.74	1.64±0.74 ª	1.15±0.36 <sup>a,b</sup>	<0.001	1.84 (1.71-1.98)		
RAISE FROM	PSI TKA	2.53±0.59	1.55±0.56 ª	1.10±0.30 <sup>a,b</sup>	<0.001	1.73 (1.60-1.85)		
A CHAIR	p-value between groups	0.104	0.444	0.380		0.206		
	CI TKA	2.45±0.72	1.51±0.75 °	1.19±0.39 <sup>a,b</sup>	<0.001	1.72(1.58-1.85)		
GOING	PSI TKA	2.39±0.55	1.34±0.54 ª	1.10±0.30 <sup>a,b</sup>	<0.001	1.61 (1.49-1.73)		
UPSTAIRS	p-value between groups	0.582	0.160	0.158		0.229		
	CI TKA	3,70±0.89	2,36±0.83 ª	1.32±0.58 <sup>a,b</sup>	<0.001	2,46( 2,28-2,64)		
GOING	PSI TKA	3,60±0.76	2,39±0.71 ª	1.32±0.50 <sup>a,b</sup>	<0.001	2,43 (2,27-2,60)		
DOWNSTAIRS	p-value between groups	0.511	0.843	0.986		0.846		
DURING WALKING	CI TKA	2,62±0.81	1,58±0.69 ª	1.11±0.32 <sup>a,b</sup>	<0.001	1,77(1,64-1,91)		
	PSI TKA	2,55±0.59	1,58±0.59 ª	1.05±0.22 a,b	<0.001	1,73 (1,60-1,85)		
	p-value between groups	0.573	0.972	0.200		0.601		

All values are presented as mean ±SD

\$ Results based on Two-Way ANOVA model using as factors the intervention and time

<sup>a</sup> P< 0.005 vs 3 months, <sup>b</sup> P< 0.005 vs 6 months

CI TKA : Conventional Instrumentation Total Knee Arthroplasty

PSI TKA : Patient Specific Instrumentation Total Knee Arthroplasty

thesis, but routine intraoperative denervation of the patella with electro cautery and osteophytes removal. All patients received two doses of first-generation cephalosporin preoperatively and 8 h postoperatively. A standard medial parapatellar approach was used for the CI TKA group, where a subvastus approach for the PSI TKA group.

For the control group of the CI, standard intramedullary instrumentation was used for the femoral component. The femoral rotational axis was defined using Whiteside's line, the epicondyle axis, and posterior condylar axis. The tibial component was placed according to the mechanical axis using extramedullary instrumentation. For the study group, the preoperative planning from 3D reconstructed MRI images was performed using planning software (Materialise NV, Leuven, Belgium). The femoral component was set at 3 degrees of flexion. The surgical epi- condylar axis obtained from 3D MRI reconstructed images was used to set femoral rotational reference. The tibial component was planned according to the ideal mechanical axis and with 3 degrees of posterior slope. Intraoperatively, the patient-specific cutting guides were placed on the femur and tibia guiding the bone resection.

The PSI TKA group received 2 g of tranexamic acid (TXA) perioperatively, a standard analgesic cocktail of periarticular, intermuscular und subcutaneous infiltration consisted of Bupivacaine, Morphine-sulfate, Epinephrine, Methylprednisolone diluted in NaCl 0.,9% perioperatively, and a systemic Patient-controlled analgesia for pain management

# TABLE 3.

Comparison of VAS for pain at different sites of the knee joint, between groups during the observation period

VAS	GROUP	3 MONTHS	6 MONTHS	12 MONTHS	P-VALUE WITH IN GROUP	TIME INDEPENDENT\$ MEAN ( 95%CI)	
POPLITEAL	CI TKA	3,92±0.85	2,49±0.85 ª	1.53±0.64 <sup>a,b</sup>	<0.001	2,65(2,46-2,83)	
	PSI TKA	3,85±0.83	2,47±0.74 ª	1.48±0.62 <sup>a,b</sup>	<0.001	2,60 (2,42-2,77)	
FOSSA	p-value between groups	0.658	0.878	0.706		0.723	
MEDIAL SURFACE OF	CI TKA	3,45±0.85	2,15±0.86 ª	1.28±0.50 <sup>a,b</sup>	<0.001	2,30(2,12-2,47)	
	PSI TKA	3,55±0.74	2,39±0.71 ª	1.29±0.49 <sup>a,b</sup>	<0.001	2,41 (2,25-2,57)	
THE KNEE	p-value between groups	0.541	0.110	0.947		0.347	
LATERAL SURFACE OF	CI TKA	2,72±0.82	1,72±0.66 ª	1.09±0.30 <sup>a,b</sup>	<0.001	1,84(1,70-2,00)	
	PSI TKA	3,08±0.77	2,10±0.69 ª	1.18±0.39 a,b	<0.001	2,11 (1,98-2,25)	
THE KNEE	p-value between groups	0.016	0.013	0.203		0.006	

All values are presented as mean ±SD

\$ Results based on Two-Way ANOVA model using as factors the intervention and time

 $a \ P < 0.005 \ vs \ 3 \ months$  ,  $b \ P < 0.005 \ vs \ 6 \ months$ 

CI TKA : Conventional Instrumentation Total Knee Arthroplasty

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postoperatively. On the other hand, the patients of the CI TKA group received epidural catheter analgesia until the second postoperative day, and then systemic Patient-controlled analgesia for pain management postoperatively. Both groups received low-molecular-weight heparin for DVT prophylaxis.

The same homogenous rehabilitation protocol for each separate center with an active range of motion exercises and continuous passive motion exercises (CPM) started on the first postoperative day and mobilization with a walker started on the second postoperative day for the CI TKA group. Fast track Knee TKA protocol with CPM motion exercises until full range of knee motion starting in the recovery room at the operation day and mobilization with walker started on the first postoperative day for the PSI TKA group.

Patients were followed up 3 months, 6 months and 1 year postoperatively. The knee pain was as-

sessed between the 2 groups at different sites of the knee joint (Table 3), and the AKP when climbing stairs and standing up from the sitting position (Table 2). Knee function was assessed using KSS Knee and Function Score, and Tegner - Lysholm score (56,57) (Table 4).

Data were expressed as mean±SD or mean±SE (for two way ANOVA analysis results) for continuous variables and as percentages for categorical data. The Kolmogorov-Smirnov test was utilized for normality analysis of the parameters. Homogeneity between compared groups was examined using the Independent samples t-test, Chi-square test and Fisher's exact test. Two-way mixed ANOVA model was used to examine the interaction between "intervention" and "time" factors and compare the variables at each time point and time independently between groups. One factor Repeated Measures ANOVA model was used for the comparison of different time measurements of variables for each

TABLE 4.									
Comparison of Evaluation knee scores between groups during the observation period									
SCORES	GROUP BASELINE		3 MONTHS	6 MONTHS	12 MONTHS	P-VALUE WITH IN GROUP	% CHANGE FROM BASELINE TO		
		BASELINE					3 MONTHS	6 MONTHS	12 MONTHS
	CI TKA	8.55±0.70	4.09±0.90 ª	2.58±20.86 <sup>a,b</sup>	1.55±0.75 a,b,c	<0.001	-52.0±10.5	-69.7±10.1	-81.9±8.6
NAC	PSI TKA	8.34±0.68	3.92±0.77 ª	2.52±0.74 <sup>a,b</sup>	1.45±0.56 <sup>a,b,c</sup>	<0.001	-53.0±8.7	-69.9±8.2	-82.7±6.2
VAS	p-value between groups	0.107	0.266	0.647	0.437		0.591	0.878	0.533
	CI TKA	46.42±7.70	65.70±5.61 ª	72.36±5.81 <sup>a,b</sup>	80.91±4.93 <sup>a,b,c</sup>	< 0.001	44.85±24.6	59.60±26.6	78.59±28.88
TEGNER- LYSHOLM	PSI TKA	46.24±4.62	66.82±5.63 ª	76.23±6.50 <sup>a,b</sup>	84.74±5.62 <sup>a,b,c</sup>	< 0.001	45.69±17.1	66.10±18.69	84.85±20.1
	p-value <sub>bg</sub>	0.882	0.287	0.001	< 0.001		0.829	0.130	0.175
KSS KNEE	CI TKA	54.45±4.94	71.81±6.50 ª	78.00±5.08 <sup>a,b</sup>	83.75±4.02 <sup>a,b,c</sup>	<0.001	32.55±13.8	44.11±13.1	54.95±14.72
	PSI TKA	54.60±4.45	74.53±6.80 ª	80.60±4.95 <sup>a,b</sup>	86.03±4.23 <sup>a,b,c</sup>	<0.001	36.99±13.1	48.33±12.4	58.38±12.38
	p-value <sub>bg</sub>	0.870	0.031	0.007	0.004		0.079	0.079	0.178
KSS FUNCTION	CI TKA	64.72±6.00	75.66±3.93 ª	81.98±5.31 <sup>a,b</sup>	91.32±4.29 a,b,c	<0.001	17.83±12.0	27.74±14.7	42.37±15.7
	PSI TKA	66.13±46.55	75.16±4.15 ª	82.26±5.48 <sup>a,b</sup>	93.15±4.72 <sup>a,b,c</sup>	<0.001	14.48±10.2	25.25±11.7	41.95±13.1
	p-value <sub>bg</sub>	0.234	0.511	0.785	0.063		0.110	0.315	0.875

All values are presented as mean  $\pm SD$ 

<sup>*a*</sup> P < 0.005 vs baseline , <sup>*b*</sup> P < 0.005 vs 3 months , <sup>*c*</sup> P < 0.005 vs 6 months

CI TKA : Conventional Instrumentation Total Knee Arthroplasty

PSI TKA : Patient Specific Instrumentation Total Knee Arthroplasty

intervention. Pairwise multiple comparisons were performed using the Bonferroni test. Comparison of percentage change from baseline to other time points of variables between interventions was analyzed using the Independent samples t-test or Mann-Whitney test in case of violation of normality. All tests are two-sided, statistical significance was set at p < 0.05. All analyses were carried out using the statistical package SPSS vr 21.00 (IBM Corporation, Somers, NY, USA).

# Results

There is statistically significant reduction of pain during the observation period for both types of surgical instrumentation for all pain variables (p<0.001). There is no statistically significant difference between the two types of surgical instrumentation in relation to absolute values of anterior knee pain at different activities at 3, 6, 12 months and time independently. (Table 2, 3) There was statistically significant lower pain at the lateral only surface of the knee for the CI group compared with the patient specific instrumentation group at 3 months (p=0.016), 6 months (p=0.013) and time independently (p=0.006)

There is statistically significant increase of functional scores during the observation period for both types of intervention (p<0.001). There is no statistically significant difference between the two types of surgical instrumentation in relation to percentile change from baseline to 3, 6 and 12 months of functional scales.

Statistically significant differences were observed between the 2 groups in the absolute values of the KSS Knee score postoperatively at 6 and 12 months, and in the absolute values of the Tegner - Lysholm

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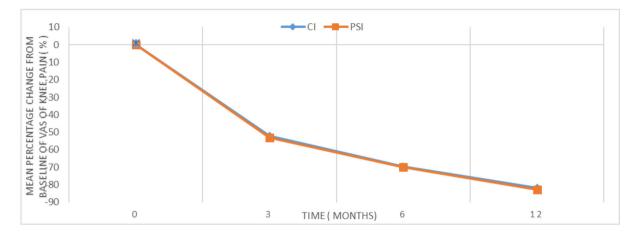


Figure 2 Mean percentage change from baseline of VAS of knee pain (%).

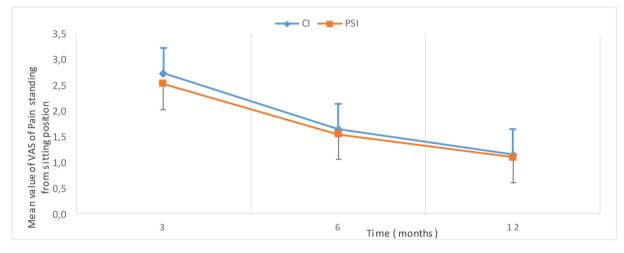


Figure 3 Mean value of VAS of Pain standing from sitting position.

score only at 6 months postoperatively, both in favour of the PSI group. (Table 4)

## Discussion

The reported incidence of anterior knee pain following primary TKA is 8% (39). Several studies have been conducted to determine the cause of anterior knee pain following TKA with variable results (40). Various causes are responsible for anterior knee pain after primary total knee arthroplasty, such as functional problems due to muscle imbalance, iatrogenic mistakes such as patellofemoral compartment overstuffing, patello-femoral instability or maltracking, different prosthetic design and mainly the design of the femoral component or even patella resurfacing or not strategy (1-15,35,36). Malalignment and malrotation mistakes of the femoral and tibial components play a very important role. Component malalignment following primary TKA has a prevalence ranging between 9.4% and 11.8% (41,42). Isolated internal rotation of the femoral component has been described as a potential source of prosthetic dysfunction, anterior knee pain, and potential early failure (43,44,45). Malrotation of the tibial prosthetic component constitutes another potential cause of a suboptimal clinical outcome following primary TKA (41,43,44). A strong correlation is reported between anterior or medial knee pain and isolated excessive tibial rotation (44). Femoral component rotation also plays a key factor in patellar tracking and can contribute to patellofemoral

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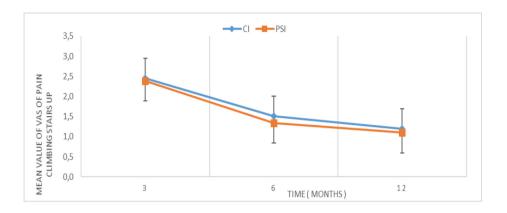


Figure 4 Mean value of VAS of Pain climbing stairs up.

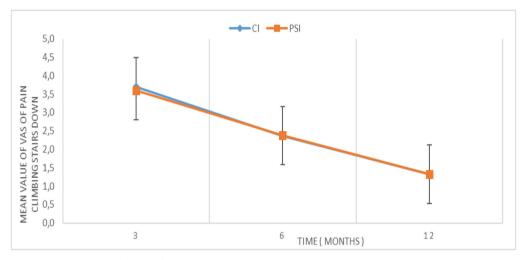


Figure 5 Mean value of VAS of Pain climbing stairs down.

complications following TKA. External rotation of the femoral component relative to the posterior femoral condyles facilitates central patellar tracking by reducing patellofemoral lateral shear forces (46,47,48). External rotation of the femoral component leads to lateral positioning of the sulcus and preserves sulcus height, which facilitates a more anatomic orientation of the trochlear groove (49). Therefore it is crucial for TKA systems to have instrumentation that allows for perfect external rotation of the femoral component in order to reproduce a more natural patellofemoral joint (47,49).

The effectiveness of patient specific instrumentation (PSI) compared to that of standard instrumentation (SI) is not completely clear, and the existing data are conflicting. There are studies showing that PSI and SI exhibited significant difference in the coronal and sagittal alignment of the femoral and tibial component (50, 51, 52). Morover other published studies have showed improvement of femoral and tibial rotation in primary TKA with PSI systems in comparison with conventional instrumentation (CI) systems (CI) (53). Khuangsirikul et al (54) and Silva et al (55) showed a more accurate rotational alignment of femoral and of tibial components between custom cutting block (PSI) and CI technique in total knee arthroplasty.

Taking into consideration the causal relationship between malalignment and malrotation mistakes of the femoral and tibial components by primary Total Knee Arthroplasty and Anterior Knee Pain on the

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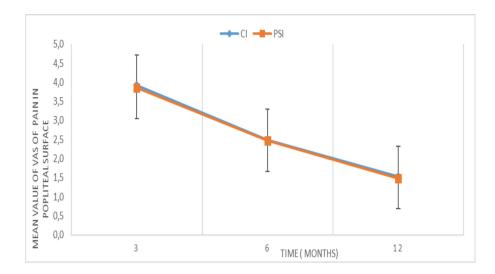


Figure 6 Mean value of VAS of Pain in popliteal surface.

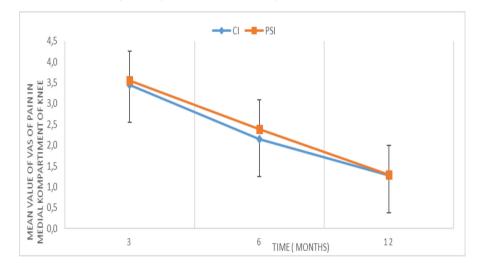


Figure 7 Mean value of VAS of Pain in medial compartment of knee.

one hand (41-49), and the already published superiority of the PSI TKA in the coronal and sagittal alignment and rotation of the femoral and tibial components in comparison to the CI TKA on the other hand by several authors (50-55), we hypothesized that PSI would have a better clinical outcome for the postoperative anterior knee pain and in general for patients' satisfaction after primary TKA than conventional Instrumentation, reflected in VAS scores after specific activities and in standard functional knee scores. Thus we compared, between the 2 centers, the exact same Prothesis Nex-Gen CR-Flex Fixed Bearing (Zimmer – Biomet Inc, Warsaw, IN USA), which is already worldwide often implanted, with only difference the PSI planning and PSI surgical technique for the second center. As far as we know there is no clinical study comparing specifically the postoperative AKP after primary TKA between CI and PSI of the exact same knee prosthesis.

The most important finding of this study was that there was no difference in clinical outcome and especially what concerns anterior knee pain after total knee arthroplasty between patient specific and conventional instrumentation 3-6-12 post-operatively, as was hypothesised. We did not find major signifi-

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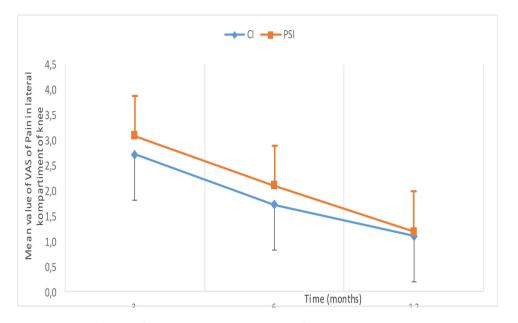


Figure 8 Mean value of VAS of Pain in lateral compartment of knee

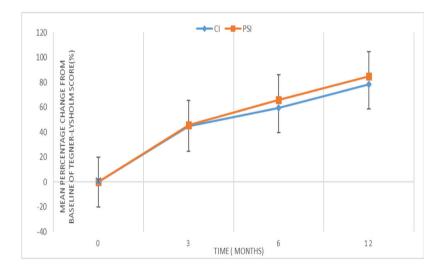


Figure 9 Mean percentage change from baseline of Tegner-Lysholm score (%).

cant differences for pain scores, knee evaluation scores and clinical outcomes between the conventional instrumentation and the use of patient specific instrumentation for total knee replacement. Our observations concerning the clinical outcome are in line with other authors. Yan et al (62) and Abane et al (63) found no difference in clinical outcome on short-term follow-up. Anderl et al (50) and Chen et al (64) also found no difference in clinical outcome two years post-operatively. In the present study, we make the hypotheses that patients who underwent a patient specific instrumentation total knee arthroplasty (PSI TKA) would have statistically fewer symptoms of anterior knee pain than those who underwent conventional instrumentation total knee arthroplasty (CI TKA), and generally better clinical improvement reflected in standard knee scores and better subjective satisfaction reflected in VAS scores. Actually, we found no statistically significant difference in VAS

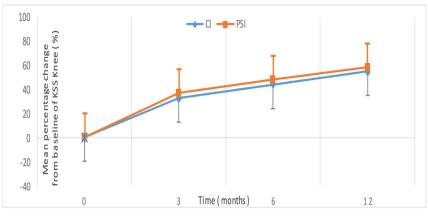


Figure 10 Mean percentage change from baseline of KSS Knee (%).

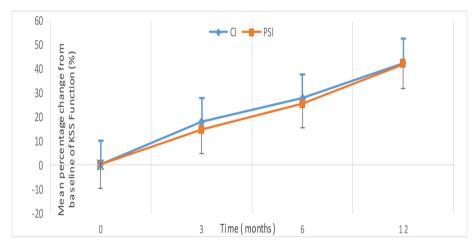


Figure 11 Mean percentage change from baseline of KSS Function (%)

score for knee pain (Table 4), in VAS score for the popliteal knee pain (Table 3), and in VAS score for AKP by standing from the sitting position and by climbing stairs (Table 2) between the CI and the PSI TKA group. All the used scores (Tegner Lysholm, KSS Function und KSS Knee) showed no magor difference between the two groups (Table 4). The statistically significant lower pain at the lateral surface for the CI group at 3 months and 6 months postoperativly, and the statistically significant better clinical scores of the KSS Knee score postoperatively at 6 and 12 months, and of the Tegner - Lysholm score only at 6 months postoperatively (both in favour of the PSI group), could be attributed to the lower preoperative BMI score of the patients of the PSI group, and to the more aggressive rehabilitation of the patients of the PSI group (all the patients rehabilitate

to special Reha-Clinics, with daily Reha-Sport program). Boonen et al. (32) did not show a significant difference between the conventional and patient specific instrumentation operation when using the Oxford Knee score and did not find a difference for the pain scores also. Woolson et al. (33) did not report any significant difference in range of motion between the 2 operation methods, consistent with our findings. Van Leeuwen J. et al. (31) showed that all KOOS sub-scores and the pain scores were similar between groups of conventional and patient specific instrumentation total knee arthroplasty.

There are limitations to our study. First, the total number of included patients was lower than planned, which was mainly due to problems of follow-up. Second, our study did not compare radiological alignment postoperatively, due to inability

for postoperatively long leg view and patella axial in the first Orthopaedic center. Third, we assessed no interobserver agreement between the different clinical scores. Forth, the two different Orthopaedic centers used different surgical approaches and different rehabilitation protocols, but still homogenous for each group. Fifth, the data of this study were based on only one type of patient specific instrumentation of one specific company and the findings should therefore not be generalized for other PSI systems. Sixth, the follow up time of one year was relatively short and as a result, no reliable data could be provided on the survival of the TKA in both groups.

The strengths of our study were the double center head to head study design, with homogeneity of demographic and clinical characteristics between compared groups and the fact that same surgeons team for each center respectively, performed the primary knee arthroplasties, implanting exactly the same knee prosthesis of one specific company. Before the study start the surgeons of the second Orthopaedic center were already familiar with the PSI technique.

No clear advantage of PSI seems to exist over conventional instruments. The cost effectiveness of the PSI technique needs to be considered. Potential cost savings include a shortened operating time (51), reduction in the number of sets of instruments (and additional sterilisation costs in most cases), reduced processing time (65). On the other hand, additional costs include the cost of an MRI or CT scans (hospital specific), costs of the patient specific instrumentation (manufacturer and hospital specific), and time needed for logistical tasks, depending on the available personnel. These requirements include the scanning process, transfer of the images to the manufacturer, monitoring the delivery of PSI to the hospital and approval of the digital plan by the surgeon prior to fabrication of the PSI. This last item is essential when using PSI in order to avoid time-consuming intra-operative changes to the proposed size of the components and the levels of resection (66). Literature does not suggest PSI techniques as a gold standard in TKA, and therefore it cannot be recommended as a standard technique and specifically in order to minimize the anterior knee pain (AKP) after TKA. One could only suggest a positive effect of the PSI instrumentation for the less experienced surgeons and in cases of preoperatively femoral or tibia posttraumatic deformities, by minimizing femoral and tibia cutting, without needing intramedullary orientation and therefore minimizing OP time (22,23).

### Conclusion

Patient specific instrumentation leads to equal clinical outcome in the short term with no major difference in anterior knee pain and other clinical scores when compared with conventional instruments in TKA surgery.

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