ORIGINAL PAPER



Minimally invasive versus open chevron osteotomy for hallux valgus correction: a randomized controlled trial

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Received: 18 December 2017 / Accepted: 28 May 2018 / Published online: 4 June 2018 © The Author(s) 2018

Abstract

Purpose The purpose of this study was to compare a minimally invasive chevron osteotomy technique (MIS group) and the well-established open chevron technique (OC group) for correction of hallux valgus deformity.

Methods Patients who were scheduled to undergo a hallux valgus surgery by means of a distal chevron osteotomy were randomly assigned to one of the two groups. Pre-operatively, six weeks, 12 weeks, and nine months post-operatively the following outcome parameters were determined: Visual Analog Scores (VAS) of pain, the American Orthopedic Foot and Ankle Society (AOFAS) forefoot score, radiographic outcome measures, range of motion (ROM), and patient satisfaction.

Results Forty-seven cases were analyzed (25 MIS group; 22 OC group). Both operative techniques achieved significant correction of the hallux deformity. The intermetatarsal angle (IMA) improved from 15.1° to 5.8° in the OC and from 14° to 6.8° in the MIS group, whereas the hallux valgus angle (HVA) improved from 28.3° to 8.5° in the OC versus 26.4° to 6.9° in the MIS group. No significant differences were observed between the groups by any of the determined outcome parameters. Regarding patient satisfaction, statistically significant differences were found between MIS and open surgery 12 weeks post-operatively in favour of the MIS group (p = 0.022).

Conclusion With the minimally invasive chevron osteotomy, radiological and clinical outcome is comparable to the open technique.

Keywords Bunion · Chevron · Hallux valgus · Minimally invasive · Percutaneous

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Introduction

Multiple different surgical techniques have been established for hallux valgus surgery so far, with each technique having its unique advantages and limitations. The distal chevron method was first described in 1981 [1] and is widely accepted as a method for correcting mild to moderate hallux valgus deformities [2]. Numerous publications presenting the radiological outcome of this surgical technique [3, 4] and the clinical outcome by means of well-established score systems [5, 6] have been published and make this technique, representing distal procedures, today's benchmark.

As a consequence of the open approach to the joint, scarring and decreased range of motion can occur [7]. Patients' demands and soft tissue scarification with open surgery have caused several minimally invasive techniques to be published in the last few years [8–11]. Given to the current literature, these minimally invasive techniques claim, as one of their advantages compared to open techniques, to have minor soft



tissue damage, reduced surgical time, and faster recovery, while same stability of correction and clinical outcome [8–11]. Therefore it seems that minimally invasive techniques are on the rise but the efficiency and stability of correction, as well as the clinical outcome of these minimally invasive techniques, have been discussed controversially. Whereas most studies presented good clinical and radiological results [10, 12], the group Romero et al. found insufficient radiographic HVA correction. However they stated low post-operative pain levels and high patient satisfaction [13]. The study by Lee was prospective, but compared the minimally invasive chevron and the scarf technique [10], which is a distinct issue. In a systematic review of minimally invasive hallux surgery, a clear recommendation could not be made although early results of multiple studies are encouraging [14]. To date and due to the previously published literature, prospectivecomparative studies comparing open versus MIS chevron techniques have not been published and the difference in any of the outcome parameters between the two surgical procedures (open vs. minimal invasive) is not clear given in the current literature.

Given the above-mentioned lack of evidence, it was the aim of the study to prospectively compare conventional open chevron osteotomy and MIS chevron osteotomy. It was hypothesized that the two techniques would show significant differences with regard to clinical score outcome (hypothesis 1), radiographic outcome (hypothesis 2), range of motion (ROM) of the first metatarsophalangeal joint (hypothesis 3), and patient satisfaction (hypothesis 4).

Material and methods

Study design

Patients on the waiting list for a distal chevron osteotomy for hallux valgus were considered for participation. Exclusion criteria were (1) age < 16 years; (2) previous first metatarsal osteotomy or soft tissue intervention for hallux valgus deformity; (3) instability of the first tarsometatarsal joint, defined as abnormal painful motion in this joint; (4) osteoarthritis of the first metatarsophalangeal joint, (5) preoperative hallux valgus angle (HVA) of less than 20° [15, 16]; or (6) an intermetatarsal angle (IMA) of less than 10° [17].

Eligible patients were then invited to participate and provided written informed consent. All patients failed conservative treatment. Patients were randomized to either the OC group or the MIS group. Assignment to these groups was done by lot. After calculation of sample size, we gathered the same amount of envelopes as feet had to be included to the study in a black box. One half was marked with the letters "OC," the other with "MIS." The day before surgery, for every patient, an envelope was drawn and administered to the surgical group

marked on the envelope. Since several patients were treated bilaterally, the cohort is of different sizes.

Surgical technique

In the OC group, a 4-cm-long dorsomedial skin incision was made to perform a V-shaped osteotomy. The apex of the osteotomy was centered 1–2 mm superior to the centre of the metatarsal head, the angle of the chevron was 60° to 90°. To avoid shortening, the osteotomy was directed toward the centre of the third metatarsal head with a slight plantarization of 5° to 10°. The metatarsal head was shifted laterally, sesamoid position was controlled by sight. Fixation was achieved by one cannulated screw (3.0 or 2.5 mm—FRS—Screw [Fusion and Reconstruction System], DePuy-Synthes, Saint Priest, France). Prominent bone ridges were resected with a saw. The distal soft tissue procedure was performed through the same dorsomedial incision. The adductor hallucis tendon was detached from its insertion at the phalangeal bone and from the lateral border of the fibular sesamoid. The transverse intermetatarsal ligament was released and a T-shaped capsulotomy was performed to allow reposition of the sesamoids. Closing of the medial capsula was performed with sutures of coated number 1 polyglactin 910 (Vicryl, Ethicon, Johnson & Johnson). The skin was closed with nylon sutures 4-0.

In the MIS group, the osteotomy was performed percutaneously through a dorsomedial incision of 3–5 mm. An electric motor-driven machine with variable speed was used for the osteotomy (NSK Prado Surgic Pro non-optic, NSK, IL, USA). Speed was limited to 8000 rpm. To prevent overheating, the reamer was frequently rinsed with sterile saline. The medial eminence was excised with a straight reamer of 2.1 or 3.1 mm (Newdeal SAS, Saint Priest, France). With the 2.1-mm reamer, the V-shaped osteotomy was performed (Fig. 1). The apex of the osteotomy was identified by fluoroscopy and centered 1–2 mm superior to the centre of the metatarsal head. The tip of the reamer was directed toward the centre of the head of the third metatarsal with a plantarisation

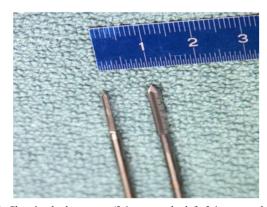


Fig. 1 Showing both reamers (2.1 mm on the left, 3.1 mm on the right side)



of 5° to 10°. After reaming, bone debris was washed out with sterile saline. A lateral soft tissue release was undertaken through a separate lateral incision of 3–5 mm as well (Fig. 2 A + B). The distal fragment was shifted laterally with a slight varus rotation to re-align the joint line. Fixation was achieved with a 1.2-mm K wire that was inserted through the same incision as used for the osteotomy. The K wire was bent at the bone insertion point and cut with an overhang of 3–5 mm. Residual bone ridges were reamed and bone debris washed out. Position of the metatarsal head and the K wire was controlled by fluoroscopy. The skin was closed with a nylon suture 4–0 (Fig. 3).

Post-operative treatment was standardized over both groups. Soft dressings were applied and patients were allowed to weight bear immediately in a custom-made hallux valgus shoe (Ofa GmbH, Bamberg, Germany). After suture removal at two weeks, post-operative wound cover was reduced to allow full movement. The surgical shoe was discarded at six weeks.

Outcome parameters

All outcome parameters were determined pre-operatively and six weeks, 12 weeks, and nine months post-operatively. We did not lose patients for follow-up. Weight-bearing radiographs (Fig. 4) were analyzed in digital manner using the Icoview software (syngo.share, ITH icoserve healthcare GmbH, Siemens, Innsbruck, Austria). The radiographic outcome measures used in this study were as follows: the HVA, which is the angle between the midshaft axes of the first metatarsal bone and the proximal phalangeal bone [18]; the intermetatarsal angle (IMA) which is the angle between the midshaft longitudinal axis of the first and second metatarsal bone. In the post-operative radiographs, it was measured as the angle subtended by the lines from center head to centre base of the first and second metatarsal [18–20].

Clinical outcome was measured in terms of passive range of motion (ROM) of the first metatarsophalangeal joint.

Fig. 2 Showing the incisions before (**a**) and after opening the hydraulic blood barrier (**b**) in a case with combination of chevron and akin osteotomy







Fig. 3 Skin sutures on day 8 after surgery (chevron and akin osteotomy)

Evaluation was performed using a goniometer in maximal plantarflexion as well as maximal dorsiflexion. ROM outcome was categorized in three groups (R1 (ROM < 30°); R2 (ROM 31°–75°); R3 (ROM > 75°); pain was assessed using the Visual Analog Scale (VAS 10). In addition, the outcome was rated with the AOFAS forefoot score (6) (American Orthopedic Foot & Ankle Society Score) and with the subjective patient satisfaction (7). For the latter, patients chose one of four answers for following question: "Are you satisfied with the result of the surgery?" (1. very satisfied, 2. satisfied, 3. don't know, 4. not satisfied).

Data analysis

Statistical data analysis was performed with SPSS (International Business Machines Corporation, Armonk, NY, USA). With a case number < 30 per group, the data were regarded as not normally distributed. Accordingly, medians and interquartile ranges (IQR) were determined as descriptives. To test for differences between the two surgical groups,





Fig. 4 Standing anteroposteriorradiograph (left foot) showing result after a minimally invasive chevron and akin osteotomy

U tests were performed (VAS, AOFAS, radiographic measurements). For categorized data (ordinal data sets), U tests were applied as well (e.g., categorized ROM, patient satisfaction). Alpha level was defined as 0.05.

A sample size calculation was done before study onset on the basis of the AOFAS score. Chan et al. stated a minimal clinically important difference of 19 points on average [21]. Other input parameter of the sample size calculation was a standard deviation of 15 points, an alpha of 0.05, and a beta of 0.1. This resulted in an estimated sample size of 15 per group (30 overall). Sample size calculation was performed with the software G*power 3.1.2 [22]. To account for dropouts, we chose to include at least 20 patients per group.

Results

There were 22 cases in the OC group (19 female, 3 male) and 25 in the MIS group (21 female, 4 male). The groups showed no significant differences with regard to the distribution of sexes (p = 0.820). Age at time of surgery was 44 (IQR 25) years and 52 (IQR 25) years in the OC and MIS groups, respectively (p = 0.117) (Table 1). Baseline clinical data were well comparable between the two groups and are summarized in Tables 2 and 3. Hallux valgus deformity may be accompanied by additional deformity of the proximal phalangeal bone. To address this deformity, a phalangeal akin osteotomy was performed in eight cases in the OC and 13 cases in the MIS group.

None of the post-operative test occasions (6 weeks, 12 weeks, 9 months) showed significant differences between the two groups with regard to VAS or AOFAS score (hypothesis 1, Table 4). Similarly, there were no differences



	OC	MIS
Number of feet	22	25
Right feet	14	13
Left feet	8	12
Bilateral	3	2
Age at time of surgery (years)	44 ± 12	52 ± 7
Female	19	21
Male	3	4

MIS, minimally invasive surgery; OC, open chevron

between groups in any of the radiographic outcome parameters (hypothesis 2, Table 4). Post-operative HVA and IMA were similar in both groups and showed significant improvement at all times of post-operative evaluation. No significant differences between the two groups with regard to ROM were observed at any of the post-operative test occasions (6 weeks, 12 weeks, 9 months) (hypothesis 3, Table 5). Regarding patient satisfaction, statistically significant differences were found between MIS and OC surgery 12 weeks post-operatively in favour of the MIS cohort (p = 0.022). Six weeks and nine months post-operative, there were no such significant differences in patient satisfaction (hypothesis 4, Table 6). In our study sample, only two patients reported poor satisfaction, one in each group.

Besides the hypotheses-based analyses, the following findings are reported: Soft tissue irritation caused by the K wire, which had to be removed in 12 feet. In contrast, in the OC cohort, the screw had to be removed in two feet (2 out of 22). Hardware removal did not affect satisfaction ratings. All cases (12 out of 25 feet) with K wire irritation led to local pain and made K wire removal necessary. In the OC group, hard ware had to be removed in one case due to local pain and soft tissue irritation as well. In the second case, removal was mandatory due to patient wish. In consideration of bone healing, we would not recommend removal of hard ware before week 12 after surgery. Moreover, removal in asymptomatic cases without soft tissue irritation screw or K wire removal is not necessary. There were no cases of fractures, loss of fixation, or avascular necrosis. One case of a hallux varus was identified in the OC group; it remained subclinical and did not need revision surgery. Three feet in the OC and one in the MIS group presented radiologically a mild recurrence of deformity without the need for revision so far. Clinically, all these patients reported a "good" outcome in terms of VAS, AOFAS, and outcome score.

Discussion

It is regarded as the most important finding of this study, that the results of the minimally invasive chevron technique were



Table 2 Pre-operative clinical data for conventional open and MIS chevron osteotomy

	Conventional distal open chevron (OC)		MIS chevron			
	Md	IQR	Md	IQR	p value	
VAS pre-operative	6	4	5	4	0.829	
AOFAS pre-operative	66.5	32	65	18	0.932	
IMA pre-operative	15.15	5.3	14	3.8	0.201	
HVA pre-operative	28.25	8.3	26.4	12	0.957	

ROM, range of motion; MIS, minimally invasive surgery; OC, open chevron

comparable with those of the classical open technique. We identified no difference between groups with regard to (1) clinical outcome score (VAS, AOFAS) (hypothesis 1 rejected), (2) radiographic outcome parameters (HVA, IMA) (hypothesis 2 rejected), or (3) range of motion of the first metatarsophalangeal joint (hypothesis 3 rejected). However, patient satisfaction was significantly superior in the MIS group 12 weeks post-operatively, but not at six weeks or nine months (hypothesis 4 accepted). Both groups experienced a significant pre-to-post-operative correction in terms of the IMA. Over the first nine post-operative months, no significant loss of correction was detected, neither in the minimally invasive nor in the open group.

In an attempt to compare our findings with the reports made by previous researchers, we found 14 retrospective studies outlining results of minimally invasive hallux valgus surgery. Most of them presented good clinical outcome in midterm [12, 23] as well as in long-term follow-up after ten years [24]. Radiological outcome also revealed good results [9, 12]. However, no prospective randomized studies comparing a minimally invasive and an open chevron technique have been published so far. This problem was also issued by Trnka et al. who could not make a definite recommendation in their systematic review analysis [25], since the majority of the studies appeared to be level IV studies performed at centers offering primarily minimally invasive hallux valgus surgery and prospective randomized studies are lacking to date. What is more, the majority of the published studies presented outcome after linear distal metatarsal osteotomies [14], which might have an impact on the joint line. Iyer et al. found higher recurrence rates in individuals with pathological joint lines, whereas another study demonstrated a probability for a metatarsal

 Table 3
 Pre-operative range of motion data (categorized)

Open chevron	MIS	p value
4.5%	4%	
50%	40%	0.491
45.5%	56%	
	4.5%	4.5% 4% 50% 40%

ROM, range of motion; MIS, minimally invasive surgery; OC, open chevron

misload after Reverdin-Isham procedures [26, 27]. Other authors showed already good clinical and radiological results with a minimally invasive chevron technique, but our analysis is the first to compare a minimally invasive chevron type and an open chevron osteotomy in a randomized manner [9, 10, 12]. We consider the chevron type superior to linear osteotomy techniques, since maintenance of the joint line and physiological positioning of the metatarsal head in the coronal plain are more likely with this technique.

Average IMA correction was a marker to determine the corrective potential of a metatarsal osteotomy. Complete reduction of the IMA should result in complete repositioning of the sesamoids as well. Recent literature reports a correlation between IMA and sesamoid position in hallux deformity as well as after hallux valgus correction [28, 29]. We believe that the IMA expresses whether the shift of the metatarsal head is sufficient. In our study, we found good correction of the IMA in both groups (Table 4), comparable to published results in

Table 4 Post-operative clinical data for OC and MIS chevron

	OC		MIS			
	Md	IQR	Md	IQR	p value	
VAS 6 weeks	1	2	1	3	0.956	
VAS 12 weeks	1	2	0	2	0.144	
VAS 9 months	0	3	1	2	0.744	
AOFAS 6 weeks	72	9	77	17	0.157	
AOFAS 12 weeks	83.5	14	85	14	0.237	
AOFAS 9 months	90	14	85	15	0.943	
AOFAS difference pre-12 weeks	13.5	29	22	22	0.285	
AOFAS difference pre-9 months	20	29	23	25	0.781	
IMA 6 weeks	7.7	3.2	7.3	1.9	0.565	
IMA 12 weeks	7.45	3.1	7.7	2.8	0.898	
IMA 9 months	5.85	3.7	6.8	3.0	0.502	
HVA 6 weeks	9.9	7.9	10.5	8.7	0.983	
HVA 12 weeks	9.15	9.7	8.8	8	0.873	
HVA 9 months	8.5	8.8	6.9	7.6	0.839	

Md, median; IQR, interquartile range; VAS, Visual Analog Scale; AOFAS, American Orthopedic Foot and Ankle Society–Score; IMA, intermetatarsal angle; HVA, hallux valgus angle; MIS, minimally invasive surgery; OC, open chevron



 Table 5
 Post-operative range of motion in comparison between surgical groups

ROM (R1 (<3 (31°-75°), R3	**	OC	MIS	p value
6 weeks	R1 R2	31.8% 63.6%	8% 88%	0.075
	R3	4.5%	4%	
12 weeks	R1 R2	9.1% 72.7%	0% 84.0%	0.653
	R3	18.2%	16%	
9 months	R1 R2 R3	5% 60% 35%	0% 66.7% 33.3%	0.910

MIS, minimally invasive surgery; OC, open chevron; ROM, range of motion

the literature [8, 23, 24]. Like the IMA, analysis of the HVA also showed a comparable correction with negligible loss of correction in both groups. A phalangeal osteotomy as well as the quality of the lateral release have a direct impact, the metatarsal osteotomy an indirect impact on this angle. Correction of the HVA in our study (compare Table 2 and Table 4) is in good accordance with the published results in the literature [12].

It is well known that after hallux surgery, clinical scores show significant improvement [30, 31]. In retrospective studies of MIS hallux valgus procedures, this finding was also detected [23, 24]. Our data support this finding as well. With the AOFAS forefoot score, we detected a significant improvement in both cohorts without differences between the two groups (Table 4). This was observed after six and 12 weeks as well as at follow-up after nine months post-operative. A second functional outcome parameter was determined by analyzing the ROM of the greater toe joint. In young athletes, maintenance of the pre-operative levels was recently described as a predictive marker for excellent outcome [32]. Our data show similar ROM values in both cohorts at all

points of follow-up with maintenance of the pre-operative ROM (Table 5).

Evaluation of VAS showed significant improvement at all time points in both groups to a comparable extent (Table 4). Residual pain six months after open hallux surgery with improvement over the following 18 months has been reported [33]. In our study, we could not detect a significant difference between the two groups at any of the time points of our survey, although minor soft tissue irritation was expected with the minimally invasive technique. In contrast to comparable pain levels, we found a statistical difference in patient satisfaction after three months in favor of the MIS cohort, but not at the other time points. This finding may be attributed to minor soft tissue irritation from this technique. Individuals were assigned to the two groups by lot. However, age at time of surgery was higher in the OC group. Since we found no differences in radiological outcome and comparable clinical results between the two groups, we think that this finding can be left out of consideration.

Another remarkable soft tissue irritation event was determined with our investigation. K wire fixation of the metatarsal head in the MIS group led to an extraordinarily high need for hardware removal (12 out of 25 feet). In contrast, in the OC group, only two Fusion and Reconstruction System (FRS) screws had to be removed, which was credited to the oblique insertion of the plain headed screw in one case and to a patient wish in the other case. As a result of this finding, we have meanwhile changed the fixation method in MIS chevron osteotomy. Since we commenced implementation of a cannulated oblique headed compression screw (3.0 mm-MICA-Chevron-Screw, Wright-Medical), no further hardware removal has been necessary (Fig. 5). Therefore, we cannot recommend K wires for routine MIS cases, whereas the use of the oblique headed compression screw seems to result in the lowest rate of soft tissue irritation.

Table 6 Patient satisfaction in comparison between surgical groups

		OC	MIS	p value
Patient satisfaction 6 weeks	Very satisfied Satisfied	54.5% 22.7%	80% 12%	0.061
	Don't know	13.6%	4%	
	Not satisfied	4.5%	4%	
Patient satisfaction 12 weeks	Very satisfied Satisfied	63.6% 13.6%	92% 4%	0.022*
	Don't know	18.2%	0%	
	Not satisfied	4.5%	4%	
Patient satisfaction 9 months	Very satisfied Satisfied	70% 15%	62.5% 25%	0.736
	Don't know	0%	4.2%	
	Not satisfied	15%	8.3%	

MIS, minimally invasive surgery; OC, open chevron





Fig. 5 Cannulated oblique headed 3.0 mm compression screw

The following limitations are acknowledged. First, the investigation was performed at one single centre and is based on cases performed by a single senior surgeon with specialization in foot and ankle surgery (GK). Second, our follow-up of nine months is regarded as a preliminary result. The two cohorts will continue to be followed.

The strength of the study is that it is to our best knowledge the first study to apply a randomized controlled prospective design. Therefore, this study is regarded as relevantly contributing new scientific knowledge.

The findings are also regarded as clinically relevant. This is because there is an increasing demand for minimally invasive surgical techniques in hallux valgus surgery. This demand is driven by patients and by surgeons attempting to meet the patients' needs. This is also seen in the increasing number of publications dealing with minimally invasive hallux valgus surgery. However, until now, no study has been published with a prospective randomized design.

From our findings, we conclude that the outcome of MIS chevron osteotomy is comparable to that of the conventional open technique. We could not identify a difference with regard to clinical outcome score (VAS, AOFAS), radiographic outcome parameters, or range of motion. However, patient satisfaction was significantly superior in the MIS group 12 weeks post-operative.

Acknowledgments Professional language editing of the manuscript by Mary Margreiter is acknowledged.

Funding Information Open access funding provided by University of Innsbruck and Medical University of Innsbruck.

Compliance with ethical standards

Research involving human participants 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.

Ethical approval The Ethics Committee of the medical university approved the study protocol.

Informed consent Informed consent was obtained from all individual participants included in the study.

Conflict of interest The authors declare that they have no conflict of interest.

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