Research Article

Predictors of Diabetic Foot Reulceration beneath the Hallux

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Aims. To evaluate the factors that predict reulceration beneath the hallux in people with a history of diabetic foot ulceration. Methods. A prospective study conducted between January 2012 and December 2014 was performed in a diabetic foot unit to assess the risk factors associated with hallux reulceration. Sixty patients with diabetic neuropathy and a history of previous ulcer were consecutively included. Sociodemographic factors and comorbidities plus the biomechanical and radiographic factors were obtained. Follow-up on participants was conducted every month, and they wore offloading therapeutic footwear and custom-made insoles. Hallux reulceration during the follow-up period was assessed as the main outcome measure in the study. Results. Patients were followed up during 29 (14.2-64.4) months. Twenty-nine patients (52%) developed a new ulceration: 9 patients (31%) in the hallux and 20 (69%) in other locations. Functional hallux limitus ($p=0.005$, 95% CI (2.097–73.128), HR 12.384) and increased body mass index ($p=0.044$, 95% CI (1.003-1.272), HR 1.129) were associated with the hallux ulceration-free survival time in the multivariate Cox model. Conclusions. Obesity and the presence of functional hallux limitus increase the probability of developing hallux reulceration in patients with diabetic neuropathy and a history of ulcers.

1. Introduction

The lifetime incidence of foot ulcers in people with diabetes has been recently estimated to be between 19% and 34% [1]. At least 85% of lower-extremity amputations are preceded by a diabetic foot ulcer (DFU), which severely increases the economic costs of health care and decreases life expectancy [2]. The primary prevention of DFU becomes critical, since death in the first year following diagnosis of the first DFU has been reported in as many as 12% of patients [3].

Diabetic neuropathy, preulcerative lesion, peripheral arterial disease, foot deformity, and increased plantar pressure have been identified as the main risk factors for DFU [4, 5]. Forty percent of patients will have a recurrence within 1 year following healing of the ulcer. The precipitating factors that initially led to the ulcer are generally not resolved after healing [1].

The forefoot is the area of higher prevalence of DFU. In particular, the hallux constitutes one-third of all areas affected by DFUs [6]. Hallux reulceration can lead to hallux amputation which has devastating effects on foot biomechanics and increases the risk of new ulcers and lower-extremity amputation [7]. Research on the risk factors of hallux reulceration can lead to the reduction of the incidence of ulcers and help avoid hallux amputation and its devastating consequences.

The normal range of motion (ROM) of the first metatarsophalangeal joint (first MTPJ) is defined as being more than 65° dorsiﬂexion, while hallux rigidus is the severe limitation of hallux dorsiflexion (<30°) [8, 9]. The ROM of the first MTPJ is routinely evaluated in a non-weight-bearing position in patients with diabetes at high risk of ulcer [10, 11]. However, mobility of the first MTPJ in the resting position has proven to be a poor predictor of abnormal first ray function during gait [12]. Foot examination...
in a non-weight-bearing position can be normal, but dorsiflexion of the first MTPJ can be blocked during gait, which means that this risk factor may remain unrecognized. This condition whereby range of motion is reduced when the forefoot is loaded is referred to as functional hallux limitus [13].

Recurrent studies conducted by Nubé et al. [14] and Cowley et al. [15] evaluated the limited mobility of the first MTPJ in relation to ulcer location on the hallux, but they were unable to find an association. ElMakki et al. [16] reported a relationship between hallux ulceration and a group of deformities which included limited mobility of the first MTPJ. These research studies [14–16] evaluated the factors associated with hallux ulcers in patients with diabetes and measured the range of motion of the first MTPJ in the resting position.

Boffeli et al. [9] explored the ROM of the first MTPJ both in the resting position and in the weight-bearing position in a group of patients with hallux ulceration. These authors [9] demonstrated that almost all patients exhibited limited first MTPJ mobility, and almost half of the patients presented functional hallux limitus.

A higher prevalence of limited mobility of the first MTPJ has been reported in patients with previous hallux ulceration [9]; however, to date, the association between hallux reulceration and limited mobility of the first MTPJ has not yet been explored by means of a prospective follow-up study.

Our study aim was to evaluate the factors that predict reulceration beneath the hallux in people with a history of diabetic foot ulceration.

2. Materials and Methods

A prospective study was conducted between January 2012 and December 2014 involving patients admitted to a diabetic foot unit at the Complutense University of Madrid in Spain, which is an outpatient center. Sixty individuals were consecutively evaluated according to the following criteria: aged over 18 years, diagnosed with type 1 or type 2 diabetes mellitus (DM) according to the criteria of the American Diabetes Association, presence of peripheral neuropathy, presence of a previous ulcer, and location of the ulcer on the forefoot. Baseline clinical data are shown in Table 1.

Patients who met the following criteria were excluded: those with peripheral arterial disease (PAD), active ulcer, diabetic neuropathic osteoarthropathy, history of previous amputation, ulcer caused by trauma, history of rheumatoid arthritis, or disease causing peripheral neuropathy other than DM.

PAD was considered when both distal pulses were absent and/or the ankle brachial index (ABI) was <0.9. In patients whose ABI was >1.4 or in those with diagnostic uncertainty, a toe pressure of <55 mmHg or a toe brachial index of <0.7 was used to diagnose PAD [17].

The local ethics committee approved this study, and all patients signed their informed written consent, in accordance with the principles of the Declaration of Helsinki.

| Table 1: Demographic data of the sample. |
|------------------------------------------|-----------------|
| (N = 60)                                | Patients        |
| Male/female                             | 51 (85)/9 (15)  |
| Mean age (years)                        | 62 ± 8.3        |
| Diabetes mellitus (years)               | 17 ± 13         |
| Diabetes mellitus type 1/type 2         | 14 (23)/46 (77) |
| Ankle brachial index                    | 1.18 ± 0.28     |
| Retinopathy                             | 31 (52%)        |
| Nephropathy                             | 15 (25%)        |
| Body mass index (kg/m²)                 | 29.4 (±5.4)     |
| Hallux                                  | 21 (35)         |
| Location of the previous ulceration     | 2.2. Biomechanical Assessment. Foot type was classified using the validated protocol of the Foot Posture Index (FPI-6) [19]. A total FPI-6 score between 0 and +5 indicates a neutral foot, a score of above +6 indicates a pronated or highly pronated foot, and a score between -1 and -12 indicates a supinated or highly supinated foot.

Hallux deformities were considered when the hallux presented one of the following: hallux valgus, bony prominence of the first metatarsal head, or hallux hammertoe deformity [7, 9, 20].

The range of mobility of the following joints was measured by using a two-armed goniometer: the ankle joint, the subtalar joint, and the first MTPJ.

Ankle dorsiflexion was examined with the patient in the supine position, keeping the subtalar joint position neutral while forcefully dorsiflexing at the ankle joint and measuring the angle formed between the bisections of the fibula and lateral foot, which had been previously marked on the patient’s skin [20].

The ROM of the subtalar joint was examined with the patient in the prone position and holding the calcaneus with one hand and the talus head/neck with the thumb and index finger of the other hand. The adduction (inversion) and abduction (eversion) ROM were assessed with the hand on the calcaneus [20].

Finally, the degree of dorsiflexion of the first MTPJ was recorded with the patient sitting in the resting position (first MTPJ ROM) and with the patient standing in a weight-bearing position (first MTPJ ROMw-b) (Figure 1). The center of the goniometer was placed on the center of the metatarsal head. The proximal arm was placed parallel to the floor, and the foot on the ground was held...
steady with one hand. The distal or mobile arm was placed parallel to the bisection of the proximal phalanx to avoid the influence of the interphalangeal joint ROM and held against the toe with the other hand. The maximum range of passive dorsiflexion was recorded [21].

Hallux rigidus was defined as a hallux dorsiflexion of less than 30° in a non-weight-bearing position. Functional hallux limitus was defined as the limitation of ROMw-b (<30°) of the first MTPJ in the absence of limitation of ROM (>40°) of the first MTPJ [8, 22].

2.3. X-Ray Goniometry. A weight-bearing lateromedial radiographic plane was obtained by using a standardized radiographic technique carried out by the same radiologist belonging to our department. Kodak Quality Control software POC 360 (Eastman Kodak Company, Rochester, NY) was used to calculate measurement angles.

A clinician different from the one who performed the clinical evaluation calculated the first metatarsal declination angle and was later blinded to any of the participants’ clinical and personal data to avoid bias in the interpretation of the radiographic measurements. The following angles in the sagittal plane related to the ankle and the first MTPJ were calculated: calcaneal inclination angle, talar declination angle, talocalcaneal angle, tibiocalcaneal angle, tibiotalar angle, and first metatarsal declination angle [23].

2.4. Offloading Regime. All patients were off-loaded with therapeutic footwear and custom-made insoles and received an in-depth education on how to prevent ulceration. Therapeutic footwear consisting of off-the-shelf shoes with the following characteristics was prescribed: extra-depth toe box, wide heel, laces or Velcro fasteners, seam-free inner lining, folds, and hollows. A list of therapeutic shoes was given to the patients in order to acquire them, and an experienced podiatrist evaluated fit and suitable characteristics. Custom insole was made from a positive plaster cast obtained from a neutral position of the foot. A metatarsal pad of Plastazote was placed behind the metatarsal heads. A cut-out refilled of Poron® was used to relieve local pressure, and a 45-degree shore hardness base of ethyl vinyl acetate (EVA) was used as the main structure of the insole. The dorsal cover was made of 25-degree shore hardness EVA. The total contact built into a multilayer construction with medial arch support replaced the removable standard insole of the shoe.

2.5. Follow-Up Protocol. All patients were followed-up prospectively until the development of an ulcer or until completion of the study (December 2017). Patients were evaluated every month according to the recommendations of the International Working Group on the Diabetic Foot [24]. At each monthly follow-up appointment, each patient was inspected for a new foot ulceration (according to the definition further below); if any callus/preulcerative lesions were observed, they were debrided, therapeutic footwear was monitored for effectiveness, and they were educated on the adherence to care.

Location of ulcer, foot type, presence of deformities, and joint mobility were recorded by the same clinician (RJMB).

2.6. Main Outcome Measure. Hallux reulceration during the follow-up period was assessed as the main outcome measure in the study. Hallux reulceration was defined as a new full-thickness lesion of the skin in the hallux, which is a new wound penetrating through the dermis of the hallux, without reference to time factors [25]. Potential factors responsible for casual pathogenesis of the new ulcers were evaluated according to the patient information and clinical findings.

2.7. Statistical Analysis. The assumption of normality of all continuous variables was verified using the Kolmogorov-Smirnov test. Normally distributed variables (Kolmogorov-Smirnov test with $p \geq 0.05$) were reported as mean and standard deviations, and nonnormally distributed variables (Kolmogorov-Smirnov test with $p < 0.05$) were reported as medians and interquartile ranges.

Statistical differences in risk factors between patients who suffered hallux reulceration and those who did not were evaluated using the chi-square test for categorical variables, and the Student $t$-test was used for normally distributed quantitative variables.

The strength of difference in the effect size was calculated by the phi coefficient for the chi-square test and the $r$ coefficient for the nonparametric test considering the values $>0.01$ as a small effect, $>0.30$ as a medium effect, and $>0.50$ as a large effect. Cohen’s $d$ was calculated as the effect size for the parametric test using an effect size calculator (http://www.uccs.edu/~lbecker/) and considering the values $>0.2$, $>0.5$, and $>0.8$ as small, moderate, and large effects, respectively. SPSS version 20.0 (SPSS, Chicago, IL, USA) was used for the other analyses.

The log-rank test was applied to determine the differences between risk factors in the time to hallux ulceration.

Continuous and categorical variables with $p < 0.10$ were selected as covariates in the univariate analysis to develop a Cox survival model of proportional hazards to determine the time to hallux ulceration and were expressed as a hazard ratio by using a forward stepwise selection method. $p < 0.05$ was accepted as statistically significant with a confidence interval of 95%. The Cox model excluded from the analysis those participants who had a shorter follow-up period than that conducted in the first event of hallux reulceration.
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Table 2: Differences between the risk factors for hallux ulceration.

|                      | Hallux ulceration (n = 9) | Nonhallux ulceration (n = 47) | p value | Effect size |
|----------------------|---------------------------|-------------------------------|---------|-------------|
| Male/female          | 8 (88.9)/1 (11)           | 39 (83)/8 (17)                | 0.658   | -0.059 b    |
| Mean age (years)     | 63 ± 9.7                  | 62 ± 7.7                     | 0.712   | 0.057 b     |
| Duration of DM (years) | 9 ± 10.8                 | 17 ± 13.3                    | 0.072   | -0.314 b    |
| Type 1/type 2 DM     | 0 (0)/9 (100)             | 13 (28)/34 (72)              | 0.072   | 0.241 a     |
| Body mass index (kg/m²) | 33.6 ± 5.0               | 29.1 ± 5.2                   | 0.030*  | 0.404 b     |
| Nephropathy          | 3 (33)                    | 12 (25)                      | 0.628   | 0.065 a     |
| Retinopathy          | 4 (44)                    | 25 (53)                      | 0.630   | -0.064 a    |
| HbA1c (mmol/mol) (%) | 54 ± 6 (7.1 ± 0.8)        | 57 ± 8 (7.3 ± 1.1)           | 0.599   | -0.207 b    |
| Ankle brachial index | 1.43 ± 0.46               | 1.24 ± 0.28                  | 0.254   | 0.242 b     |
| Hallux deformity     | 3 (33)                    | 11 (23)                      | 0.529   | 0.084 a     |
| Hallux valgus        | 1                         | 10                            |         |             |
| Metatarsal prominence | 0                        | 8                            |         |             |
| Hallux hammertoe     | 2                         | 2                            |         |             |
|                      |                           |                               |         |             |
| First MTPJ ROM (degrees) | 47.33 ± 19.36         | 49.79 ± 19.15                | 0.734   | -0.064 b    |
| First MTPJ ROM(0,15) (degrees) | 21.11 ± 7.15           | 33.26 ± 16.99                | 0.041*  | -0.422 b    |
| Functional hallux limitus | 7 (78)                | 11 (23)                      | 0.001*  | 0.428 a     |
| Hallux rigidus       | 4 (44)                    | 18 (38)                      | 0.729   | 0.046 a     |
| Ankle ROM (degrees)  | 90.89 ± 5.49             | 87.60 ± 5.83                 | 0.129   | 0.279 b     |
| Inversion ROM (degrees) | 17.36 ± 4.87            | 16.85 ± 4.64                 | 0.697   | 0.074 a     |
| Eversion ROM (degrees) | 9.33 ± 1.41             | 9.74 ± 3.16                  | 0.411   | -0.083 b    |
| FPI                  | 2.33 ± 2.50              | 0.66 ± 4.30                  | 0.125   | 0.230 b     |
| Tibiotalar angle X-ray | 112.01 ± 5.11         | 111.77 ± 6.21                | 0.902   | 0.021 b     |
| Tibiocalcaneal angle X-ray | 67.96 ± 6.94         | 65.88 ± 7.44                 | 0.433   | 0.143 b     |
| Talocalcaneal angle X-ray | 45.11 ± 3.25         | 46.31 ± 6.13                 | 0.403   | -0.121 b    |
| Talar declination angle X-ray | 26.64 ± 4.02         | 25.35 ± 4.16                 | 0.397   | 0.156 b     |
| Calcanal inclination angle X-ray | 17.91 ± 4.51         | 20.91 ± 6.70                 | 0.114   | -0.254 b    |
| First metatarsal declination angle X-ray | 22.81 ± 2.14         | 24.98 ± 3.50                 | 0.024*  | -0.350 b    |

Abbreviations: HbA1c: glycated hemoglobin; ROM: range of motion; first MTPJ ROM: range of dorsiflexion of the first metatarsophalangeal joint in the resting position; first MTPJ ROM(0,15): range of dorsiflexion of the first metatarsophalangeal joints in the weight-bearing position; FPI: Foot Posture Index. The “nonhallux ulceration” group of patients included other locations of ulceration and patients who did not develop a new ulcer during the follow-up. a n (%) for categorical variables; the phi coefficient was used for the chi-square test: representing effect size values of 0.01 as small effect, 0.30 as medium effect, and 0.50 as large effect. b Mean ± SD for normally distributed variables; for independent samples, Student t-test; effect size as the Cohen’s d: representing effect size values > 0.2 as small effect, > 0.5 as moderate effect, and > 0.8 as large effect; d is positive if the mean difference is in the predicted direction.

3. Results

Four patients dropped out of the study because of their refusal to wear protective therapeutic footwear and/or custom insole and were thus removed from the study. They presented no ulcers at the time of abandoning the study. Finally, 56 patients were included in the analysis. Patients were followed up prospectively for a median time period of 29 months (interquartile ranges (IR) 14.2-64.4 months).

Twenty-nine patients (52%) developed reulceration during follow-up and showed a median time to ulceration of 19 months (IR, 6.0-30.0 months). Different locations of reulceration were as follows: 9 patients (31%) developed a new ulcer in the hallux, 6 (21%) in the minor toes, 13 (45%) beneath the metatarsals, and 1 patient (3%) in the heel. All ulcers were classified as neuropathic reulceration, and there were no ulcers caused by trauma.

Several variables related to demographic and foot characteristics revealed differences between patients who developed hallux ulceration and patients who developed ulceration in other locations or without reulceration (Table 2). Participants who developed hallux ulceration were more likely to present a higher BMI (p = 0.030, Cohen’s d = 0.404), a reduced dorsiflexion of the first MTPJ in a weight-bearing position (p = 0.041, Cohen’s d = -0.422), and functional hallux limitus (p = 0.001, r = 0.428) as baseline characteristics. Hallux rigidus showed no association with hallux ulceration (p = 0.729, r = -0.046).

In the radiographic analysis (Table 2), lower first metatarsal inclination was associated with hallux ulceration (p = 0.024, Cohen’s d = -0.350).
patients with previous hallux ulceration. However, they did not consider a control group, and therefore, statistical association was not investigated. Furthermore, a prospective follow-up of patients with previous hallux ulceration was not carried out.

Biomechanical theories about functional hallux limitus claim that both the elevation of the head of the first metatarsal and the increase in tension in the plantar aponeurosis may alter the joint dynamics in the first MTPJ. Owing to the ground reaction forces that are exerted on the first metatarsal head during midstance and propulsion phases, dorsiflexion of the first ray occurs in some patients, which can lead to blockage of the first MTPJ [13]. In our radiological univariate analysis, an increased dorsiflexion of first metatarsal declination was observed in patients who had previously developed hallux reulceration, which should support this theory. The evaluation of hallux dorsiflexion in a non-weight-bearing position does not regenerate the evidence about the weight borne by the head of the first metatarsal, and the result is normal, despite the fact that the joint dorsiflexion is blocked.

According to our results, we recommend to evaluate dorsiflexion of the hallux in the weight-bearing position and to then identify functional hallux limitus in the biomechanical screening of patients at high risk of foot reulceration.

Factors that contribute to a dorsiflexed first ray remain unknown [26]. An increased BMI value (p = 0.044; HR 1.129) may contribute, yet this is uncertain. A greater body weight has been previously found to be an independent risk factor for foot ulceration in a prospective study on a wide sample of patients with DM [27]. The presence of a high BMI and functional hallux limitus could predict the reulceration beneath the hallux, and we suggest that preventive strategies should be focused on the control of these variables.

Even though some authors have evaluated risk factors for hallux ulceration, this is the first study that prospectively explores functional hallux limitus as a risk factor for great toe reulceration, which represents the main strength of the study. However, our results should be interpreted with caution due to a number of limitations. Cumulative tissue stress has been shown to affect recurrence rates on the foot, and it is the result of the combination of plantar pressure and ambulatory activity [28]. However, neither of the variables was evaluated in this study. Secondly, the level of compliance of the patients was not evaluated in this study; the authors encouraged patients in each monthly visit to use therapeutic footwear, and all participants claimed to use preventive strategies. However, objective methods to evaluate compliance should be used in further studies [29].

5. Conclusions

In conclusion, the present study underscores the importance of identifying functional hallux limitus and demonstrates that patients with a history of DFU with functional hallux limitus and increased BMI have a higher probability of developing reulceration of the hallux.
Data Availability
The datasets used to support this study are not freely available in view of participants’ privacy protection.

Conflicts of Interest
The authors declare that they have no conflicts of interest.

Authors’ Contributions
RJMB and JLLM contributed to the conception and design of the study. RJMB and FJAA performed the analysis and interpretation of the results. RJMB wrote the manuscript. All authors were responsible for the critical revision of the manuscript for important intellectual content. All authors approved the final version of the article. RJMB is the guarantor of the work.

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References
[1] D. G. Armstrong, A. J. M. Boulton, and S. A. Bus, “Diabetic foot ulcers and their recurrence,” The New England Journal of Medicine, vol. 376, no. 24, pp. 2367–2375, 2017.
[2] A. J. Boulton, L. Vileikyte, G. Ragnarson-Tennvall, and J. Apelqvist, “The global burden of diabetic foot disease,” The Lancet, vol. 366, no. 9498, pp. 1719–1724, 2005.
[3] J. W. Walsh, O. J. Hofstad, M. O. Sullivan, and D. J. Margolis, “Association of diabetic foot ulcer and death in a population-based cohort from the United Kingdom,” Diabetic Medicine, vol. 33, no. 11, pp. 1493–1498, 2016.
[4] C. A. Abbott, A. L. Carrington, H. Ashe et al., “The North-West Diabetes Foot Care Study: incidence of, and risk factors for, new diabetic foot ulceration in a community-based patient cohort,” Diabetic Medicine, vol. 19, no. 5, pp. 377–384, 2002.
[5] R. Waaijman, M. de Haart, M. L. J. Arts et al., “Risk factors for plantar foot ulcer recurrence in neuropathic diabetic patients,” Diabetes Care, vol. 37, no. 6, pp. 1697–1705, 2014.
[6] D. G. Armstrong, L. A. Lavery, and L. B. Harkless, “Validation of a diabetic wound classification system. The contribution of depth, infection, and ischemia to risk of amputation,” Diabetes Care, vol. 21, no. 5, pp. 855–859, 1998.
[7] R. J. Molines-Barroso, J. L. Lazaro-Martinez, F. J. Alvaro-Afonso, I. Sanz-Corbalan, J. L. Garcia-Klepzig, and J. Aragon-Sanchez, “Validation of an algorithm to predict ulceration in amputation patients with diabetes,” International Wound Journal, vol. 14, no. 3, pp. 523–528, 2017.
[8] C. Formosa, A. Gatt, and N. Chockalingam, “The importance of clinical biomechanical assessment of foot deformity and joint mobility in people living with type-2 diabetes within a primary care setting,” Primary Care Diabetes, vol. 7, no. 1, pp. 45–50, 2013.
[9] T. J. Moffeli, J. K. Bean, and J. R. Natwick, “Biomechanical abnormalities and ulcers of the great toe in patients with diabetes,” The Journal of Foot and Ankle Surgery, vol. 41, no. 6, pp. 359–364, 2002.
[24] K. Bakker, J. Apelqvist, N. C. Schaper, and International Working Group on the Diabetic Foot Editorial Board, "Practical guidelines on the management and prevention of the diabetic foot 2011," *Diabetes/Metabolism Research and Reviews*, vol. 28, Supplement 1, pp. 225S–231S, 2012.

[25] R. J. Molines-Barroso, J. L. Lazaro-Martinez, J. Aragon-Sanchez, E. Garcia-Morales, J. V. Beneit-Montesinos, and F. J. Alvaro-Afonso, "Analysis of transfer lesions in patients who underwent surgery for diabetic foot ulcers located on the plantar aspect of the metatarsal heads," *Diabetic Medicine*, vol. 30, no. 8, pp. 973–976, 2013.

[26] B. Durrant and N. Chockalingam, "Functional hallux limitus: a review," *Journal of the American Podiatric Medical Association*, vol. 99, no. 3, pp. 236–243, 2009.

[27] E. J. Boyko, J. H. Ahroni, V. Stensel, R. C. Forsberg, D. R. Davignon, and D. G. Smith, "A prospective study of risk factors for diabetic foot ulcer. The Seattle Diabetic Foot Study," *Diabetes Care*, vol. 22, no. 7, pp. 1036–1042, 1999.

[28] J. J. Van Netten, J. G. van Baal, A. Bril, M. Wissink, and S. A. Bus, "An exploratory study on differences in cumulative plantar tissue stress between healing and non-healing plantar neuropathic diabetic foot ulcers," *Clinical Biomechanics*, vol. 53, pp. 86–92, 2018.

[29] R. Waaijman, R. Keukenkamp, M. de Haart, W. P. Polomski, F. Nollet, and S. A. Bus, "Adherence to wearing prescription custom-made footwear in patients with diabetes at high risk for plantar foot ulceration," *Diabetes Care*, vol. 36, no. 6, pp. 1613–1618, 2013.