

Modification of Primary Avalvular Varicose Anomaly after endovenous radiofrequency ablation

Primary Avalvular Varicose Anomalies before and after RFA

Veränderungen von Primary Avalvular Varicose Anomalies nach endovenöser Radiofrequenzablation.

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ABSTRACT

Primary avalvular varicose anomaly (PAVA) is a new medical concept defined as primary varicose veins resembling neovascularized tissue on ultrasound examination. PAVAs could be misdiagnosed as recurrence at the saphenofemoral or saphenopopliteal junction, but no studies have yet examined their role before and after venous invasive procedure. In this report, we describe a case of PAVA in a 39-year-old man with symptomatic varicose veins and great saphenous vein truncal incompetence. Six months after radiofrequency ablation of the great saphenous vein, duplex ultrasound revealed complete occlusion of great saphenous vein and partial thrombosis of the still incompetent PAVA.

ZUSAMMENFASSUNG

Primär avalvuläre variköse Anomalien (PAVA) sind ein neues medizinisches Konzept, definiert als primäre Varikosis, die duplexsonographisch dem neovaskularisierten Gewebe ähnelt. PAVA können mit Rezidiven im Bereich der Crosse assoziiert sein; keine Studie hat jedoch ihre Rolle vor und nach dem invasiven Eingriff untersucht.

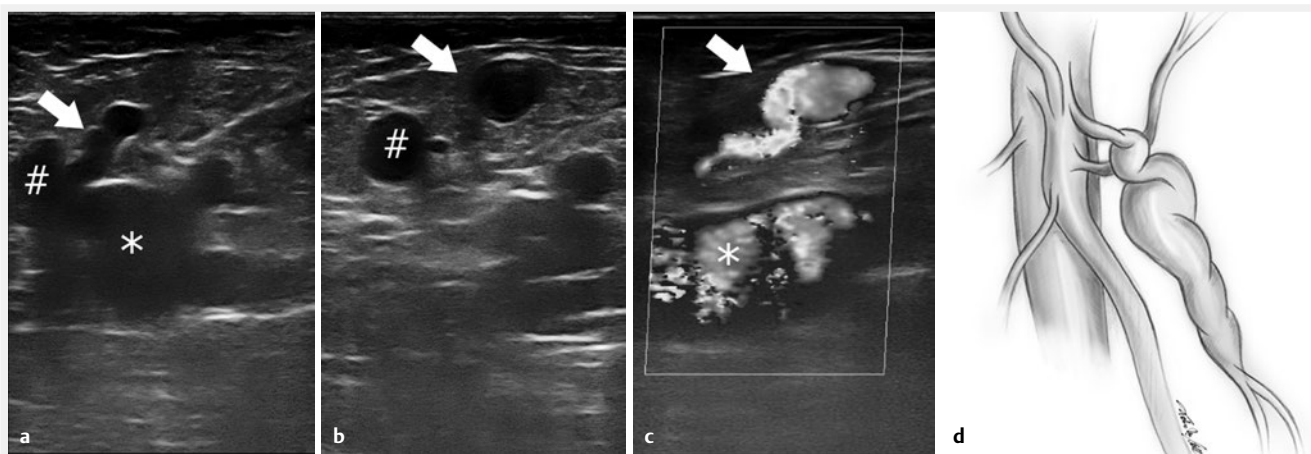
In diesem Report beschreiben wir ein Fall von PAVA in einem 39-jährigen Patienten mit symptomatischen Varizen und Saphena magna-Stamminkompetenz. Sechs Monate nach Radiofrequenzablation der Vena Saphena Magna zeigte die Duplexsonographie einen kompletten Verschluss der Vena Saphena Magna und eine partielle Thrombose der weiterhin inkompetenten PAVA.

Introduction

Primary avalvular varicose anomaly (PAVA) is a new phenomenon defined as tortuous, thin-walled and incompetent veins in patients presenting with primary varicose veins (VVs), resembling neovascularized tissue on ultrasound and surgical examination [1].

The concept of PAVA was introduced to justified saphenofemoral junction (SFJ) recurrence with an aspect of neovascularization, after endovenous or surgical saphenous intervention. In particular, the misdiagnosis of a PAVA – already present before surgery – mistak-

en for neovascularized tissue in recurrent VVs, may lead to underestimating the role of these avalvular residual SFJ collaterals after ablative intervention as a source of recurrence. The recognition of SFJ collaterals' incompetence has a greater relevance in case of endovenous procedure, in which only saphenous axis is treated. We present a case of PAVA, originating from SFJ, in a patient who underwent radiofrequency ablation (RFA) for incompetent great saphenous vein (GSV) and describe Doppler ultrasound scan (CDUS) PAVA modifications during follow-up period.



► **Fig. 1** Preoperative CDUS analysis of saphenofemoral junction with femoral vein (asterisk), great saphenous vein (hashtag) and PAVA (arrow), proximal cross section, A and B. Longitudinal section of incompetent PAVA, C. Graphic overview of preoperative saphenofemoral junction with its collaterals, great saphenous vein and PAVA, D.

The patient provided written consent for the publication of this paper in a medical journal.

Case report

A 39-year-old man presented to our attention with primary varicose veins (C2-S; Ep; As; Pr2,5). He has a past medical history of appendectomy and adenoidectomy during his youth, prior episode of prostatitis and occasional asthma treated with bronchodilator medication (Salmeterol and Fluticasone) when needed. He also reported a familiar asymptomatic Fahr's syndrome, genetically diagnosed with mutation in SLC20A2 gene, with no basal ganglia or electrolyte homeostasis involvement, as demonstrated by head computed tomography scan and blood test performed one month before the procedure. Intolerance to acetylsalicylic-acid was reported.

A preoperative CDUS analysis revealed SFJ reflux with incompetence of both terminal and preterminal valves during Valsalva maneuver. A pathological reflux >0.5 s extending along the course of GSV, up to the knee, was detected. Extra-saphenous varicose vein along the leg were also present. GSV diameter was measured at proximal, middle and distal thigh, being 8.7 mm, 10.7 mm and 9.5 mm, respectively. GSV-skin distances were also measured at same thigh levels: 17 mm, 19 mm and 8 mm, respectively. Neither thigh GSV incompetent collaterals nor anterior accessory saphenous vein (AASV) were detected.

During the CDUS exam, a SFJ collateral, next to the connection with femoral vein, was noted (► Fig. 1). It was a serpiginous, avascular, dilated, refluxing collateral, extending from SFJ to saphenous space, wrapping the GSV in its proximal tract. Its maximum diameter was 9.9 mm proximal to the SFJ. Distally at middle third level of the thigh, the collateral became suprafascial, with a marked decrease in caliber and the abolition of reflux. It was not possible to describe neither where this collateral originated from nor which district it drained. The patient underwent endovenous ablation of GSV using radiofrequency with ClosureFAST™ catheter (Medtronic, San Jose, CA, USA) and concomitant phlebectomies of the leg under local anesthesia (► Fig. 2). Neither intraprocedural nor post-procedural complications occurred. Patient was discharge the same

day of the procedure with compression stockings (20–30 mmHg) for two weeks and prophylactic dose of low molecular weight heparin for twelve days.

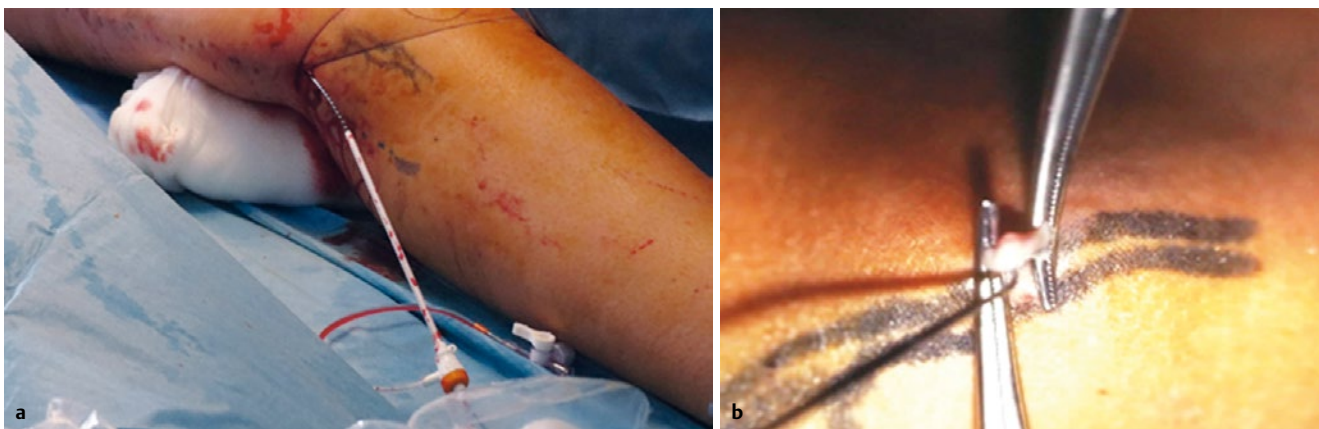
After six months, the patient underwent CDUS control examination that revealed good results of endovenous ablation, with patency of SFJ through the superficial epigastric vein. The tortuous SFJ collateral was still patent, with maximum diameter of 7.8 mm and partial circumferential thrombosis of the distal portion, with a residual lumen of 4.8 mm in diameter (► Fig. 3). Residual reflux was detected in this tributary vein during Valsalva maneuver. No clinical signs of residual or recurrent VVs was present. Reviewing pre-interventional CDUS images and comparing them with the follow-up images, a truncal PAVA was then diagnosed.

Discussion

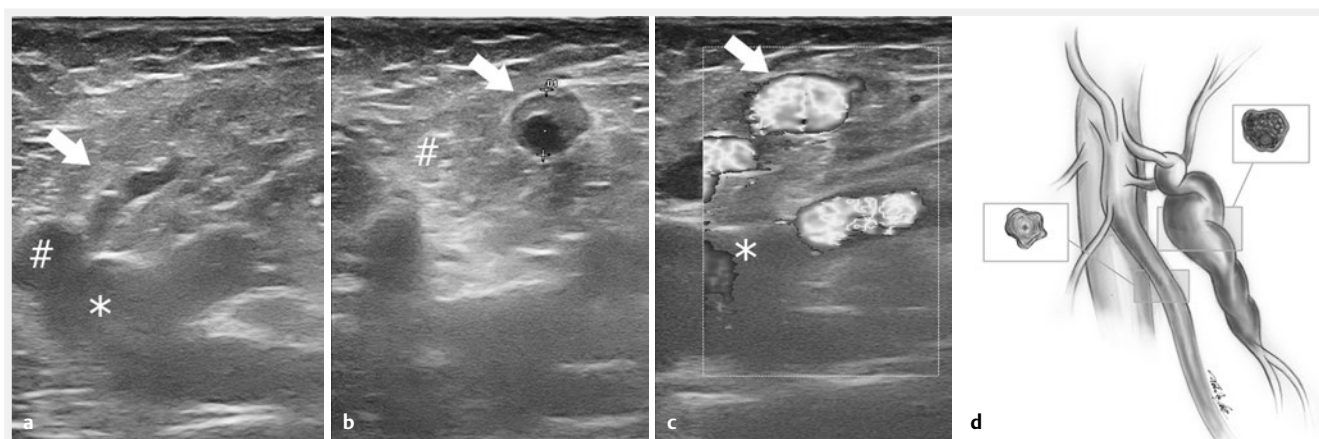
Varicose veins recurrence remains one of the most undisclosed and debated topic in phlebology with a wide incidence range from 7% to 65% [2]. Moreover, the pathogenesis of recurrent SFJ reflux remains incompletely understood, even if several studies have proposed different pathological causes [3–7]. Among these, neovascularization has been reported to account from 8% to 60% of recurrent VVs [2, 8, 9]. Tactical and technical errors were detected in 80% of cases, although incidences remain very heterogeneous among studies [2, 10, 11] with unclear correlation between recurrent VVs defined with CDUS, surgical or histological examination [12].

PAVA is a new anatomical entity consisting of serpiginous, avascular and usually incompetent veins arising from SFJ or directly from femoral vein, detected during preoperative CDUS evaluation [1]. Three patterns of PAVA were described by Ostler and collaborators [1]:

- Lymph node pattern: PAVA emerge directly from a groin lymph nodes
- Peritruncal pattern: PAVA wrap around truncal veins (the GSV, small saphenous vein, or anterior accessory saphenous vein)



► **Fig.2** Endovenous ablation of GSV was performed using radiofrequency ablation technique with ClosureFAST™ catheter (Medtronic, San Jose, CA, USA), A. Concomitant phlebectomies of the leg under local anesthesia were also performed, B



► **Fig.3** Post-RFA six-month CDUS analysis, revealing patency of femoral vein (asterisk), saphenofemoral junction, proximal great saphenous vein (hashtag) and proximal PAVA (arrow), A. Total occlusion of great saphenous vein due to RFA (hashtag), with partial PAVA occlusion (arrow), still incompetent, B and C. Graphic overview of postoperative saphenofemoral junction with its collaterals, great saphenous vein, PAVA and their cross sectional lumen view, D.

- Atypical pattern: PAVA have the same appearance as the other two patterns but lie in various locations and are not connected to lymph nodes or truncal veins.

The incidence of PAVA is around 5% among CDUS examinations, with a female-to-male ratio of about 5:1 and with no significant correlation with CEAP grade [1].

The recognition of PAVA during CDUS examination is challenging because it may be confused with SFJ collateral or with GSV groin tributaries with poor or none clinical significance. The differences have not yet been clearly defined. PAVA differs from AASV for the absence of valves and the serpiginous course, not parallel to GSV.

We report a case of a patient with truncal PAVA undergoing to a RFA procedure.

PAVA was not treated because it was recognized only during retrospective imaging.

At 6-month CDUS follow-up, a good clinical and instrumental result were evident, although PAVA was still incompetent, with distal partial circumferential vein lumen thrombosis (residual lumen of 4.8 mm) and a decrease in maximum diameter (from 9.9 mm to

7.8 mm). The reasons for these two changes remains unknown. Furthermore, we do not know if in future this collateral could become a source of recurrent VVs, but we would like to introduce the hypothesis that some recurrent VVs cases considered as neovascularization, could be a residual PAVA. The misdiagnosis derives from similarity in recurrent VVs and PAVA appearance (avalvular, shaped, next to SFJ, ...). The fact that, 6 month after endovenous procedure, PAVA is still incompetent may justify its role as a future source of reflux in the SFJ thus creating recurrent VVs.

For these reasons, it should be recommended to detect PAVA during preoperative CDUS examination and during follow up controls, in order to clarify:

- if PAVA is primarily observed as a patent and competent/incompetent vessel, and remains or become incompetent after RFA or others venous treatment
- if PAVA modifies its diameter after intervention, producing new varices or dilated veins
- if PAVA represents a potential new source of pathological/symptomatic reflux

- if it is necessary to remove PAVA in case of surgery or sclerotherapy for recurrent VVs.

The answers to these questions might strongly modify CDUS approach and change management and treatment of primary and recurrent VVs.

CONCLUSION

Primary Avalvular Vein Anomalies are not so rare in patients performing a CDUS analysis for varicose veins. It is important to detect PAVA during preoperative examination and report its presence in order to correlate it to a possible recurrence of VVs. The need for new studies that could identify a correlation between recurrent VVs patterns and preoperative PAVA is necessary to confirm the role of these veins in the recurrence of symptomatic chronic venous disease.

Conflict of Interest

The authors declare no conflict of interest.

Acknowledgment

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