



Endovascular treatments of the extended iliac vein occlusion: what are the limits?

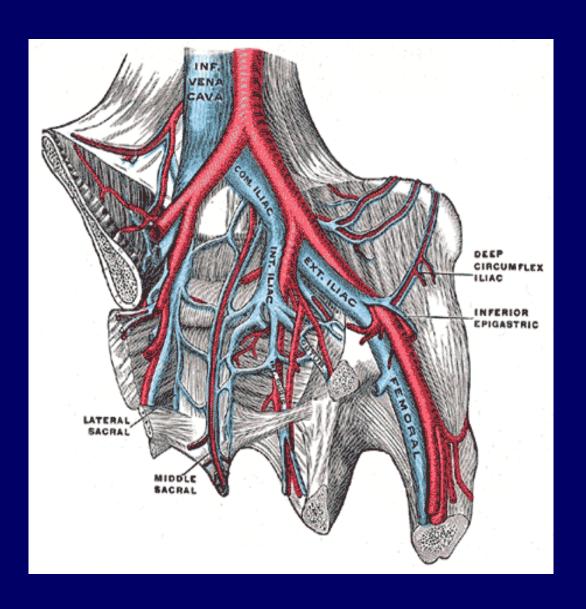
Professor Alun H Davies

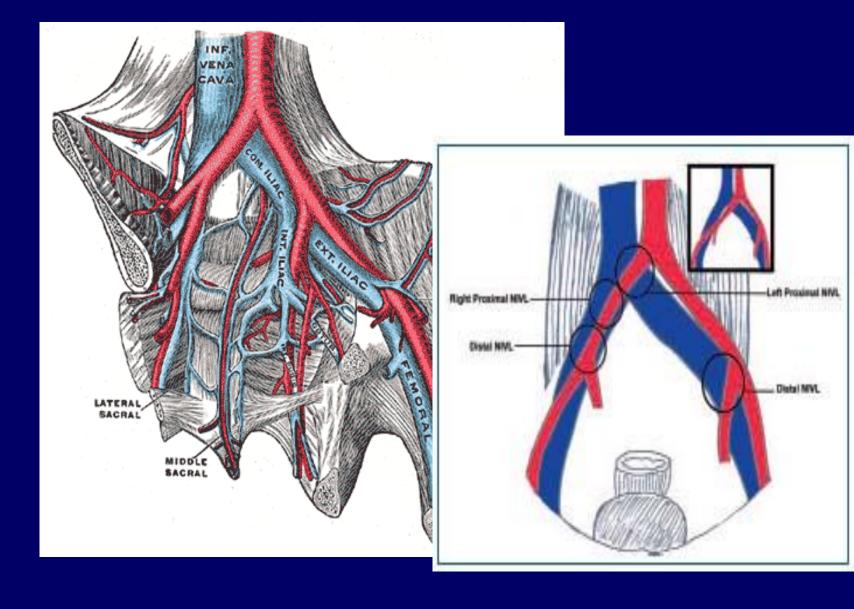
Section of Vascular Surgery

Charing Cross & St Mary's Hospitals

London



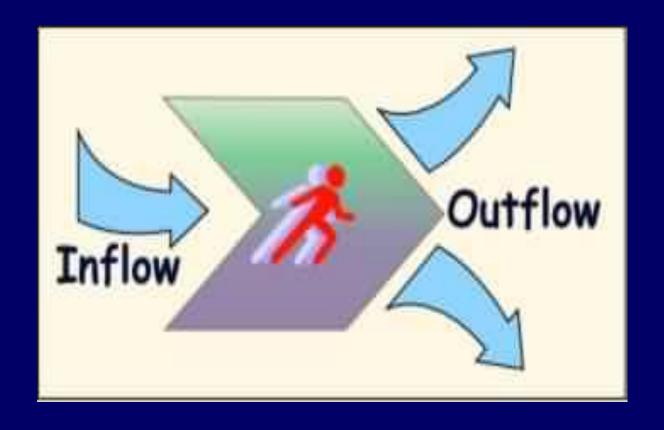




Iliac Vein Stenting for Chronic Venous Insufficiency

Chronic venous insufficiency has devastating sequelae in terms of patients' lifestyles and negative economic impact on society. Traditional surgical procedures have yielded variable patency results, and follow-up has not always been reported. This review summarizes the current applications, patency rates, stent selection, and complications of balloon angioplasty and stenting in the treatment of chronic venous outflow obstruction in the lower extremity. We conclude that endovenous stenting is the current method of choice in the treatment of chronic venous obstruction. (**Tex Heart Inst J 2007;34:60-6**)

Firas F. Mussa, MD Eric K. Peden, MD Wei Zhou, MD Peter H. Lin, MD Alan B. Lumsden, MD Ruth L. Bush, MD, MPH



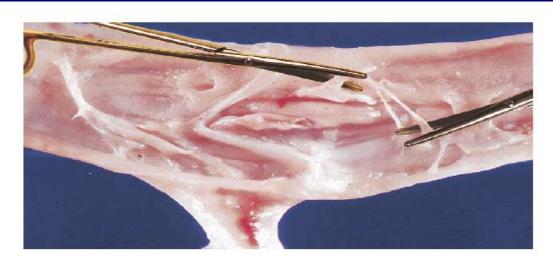
Best management options for chronic iliac vein stenosis and occlusion

Seshadri Raju, MD, FACS, Jackson, Miss

First author, year	Comments
Halliday,17 1985	Palma bypass; 47 PTS limbs. Venographic cumulative patency of 75% at 5 years
Jost,18 2001	18 Palma bypass • 17 prosthetic bypass
	6 spiral vein bypass
	• 1 vein patch
	62% overall cumulative patency at 3 years and 83% for Palma bypass at 4 years
Puggioni, 19 2004	Technique of endophlebectomy
Garg, ²⁰ 2011 Neglen, ²¹⁻²⁵	Hybrid and open techniques of correction
Neglen, ²¹⁻²⁵ 2000-2010	Technique of iliac vein stenting; need for large stents and extension into inferior vena cava. Stent extension across inguina ligament. Bilateral iliac vein stents. Stenting across stenotic/occluded IVC filters. Incidence of ISR after iliac vein stent
Raju, ²⁶ _2009	CTO recanalization technique
Kolbel, ²⁷ 2009	Jugular approach for CTO recanalization; ancillary techniques
Raju, ²⁸ 2009	Different types of stent malfunction and reinterventional techniques to correct them

Surgical disobliteration of postthrombotic deep veins—endophlebectomy—is feasible

Alessandra Puggioni, MD,^{a,c} Robert L. Kistner, MD,^{a,b} Bo Eklof, MD, PhD,^{a,b,c} and Fedor Lurie, MD, PhD,^{a,b,c} Honolulu, Hawaii



On opening the vein the synechiae attached to the intimal layer are placed on tension and carefully removed with scissors by snipping their attachments to the intimal surface of the vein.

Surgical disobliteration of postthrombotic deep veins—endophlebectomy—is feasible

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Results: In 10 patients (77%) the treated segments remained primarily patent at median follow-up of 8 months (range, 1-28 months). Early thrombosis near the endophlebectomy site occurred in 3 patients, at 2, 5, and 12 days, respectively, after surgery. In 2 patients with early thrombosis further interventions were carried out with success. In a third patient with early postoperative thrombosis the final outcome was recanalization and reflux. These results yielded an overall secondary patency rate of 93%. No perioperative pulmonary embolism was observed.

their attachments to the intimal surface of the veni.

Surgical disobliteration of postthrombotic deep veins—endophlebectomy—is feasible

Alessandra Puggioni, MD, a,c Robert L. Kistner, MD, a,b Bo Eklof, MD, PhD, a,b,c and Fedor Lurie, MD, PhD, a,b,c Honolulu, Hawaii

Conclusion: This series demonstrates that surgical disobliteration of postthrombotic deep veins is technically feasible, and led to patency of the segments for the duration of follow-up for up to 28 months (mean, 10.8 ± 8.2 months). We used this technique with the objective of disobstructing postthrombotic veins, to increase flow through a previously narrowed lumen. Postoperative thrombosis at the site of endophlebectomy occurred in 23% of patients. Although this early experience is encouraging, further studies and longer follow-up are necessary to assess the durability of the procedure. (J Vasc Surg 2004;39:1048-52.)

their attachments to the mumai surface of the veni.

Indications for endophlebectomy and/or arteriovenous fistula after stenting

M A F de Wolf*†, C W K P Arnoldussen† and C H A Wittens*†§

*Department of General Surgery Maastricht University Medical Center (MUMC), Maastricht, the Netherlands;
†Cardiovascular Research Institute (CARIM), Maastricht, the Netherlands;
†Department of Radiology MUMC+, Maastricht, The Netherlands;
§Department of Vascular Surgery University Hospital Aachen, Aachen, Germany



Use additional imaging to quantify inflow.

If extensive disease do AVF as well.



Venous stenting across the inguinal ligament

Peter Neglén, MD, PhD, T. Paul Tackett Jr, BS, and Seshadri Raju, MD, Flowood, Miss (J Vasc Surg 2008;48:1255-61.)

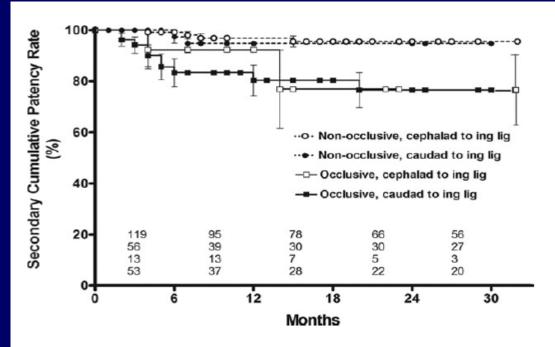
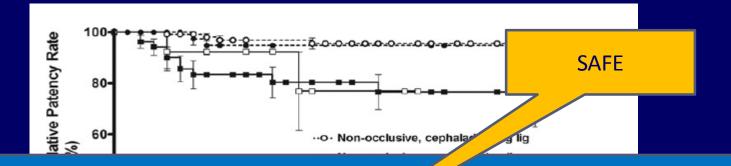


Fig 6. Secondary patency rates of stents placed in thrombotic limbs for non-occlusive obstruction and occlusion with stenting terminating cephalad and caudad to the inguinal ligament (ing lig). The lower numbers represent total limbs at risk for each time interval (± SEM).

Venous stenting across the inguinal ligament

Peter Neglén, MD, PhD, T. Paul Tackett Jr, BS, and Seshadri Raju, MD, Flowood, Miss (J Vasc Surg 2008;48:1255-61.)



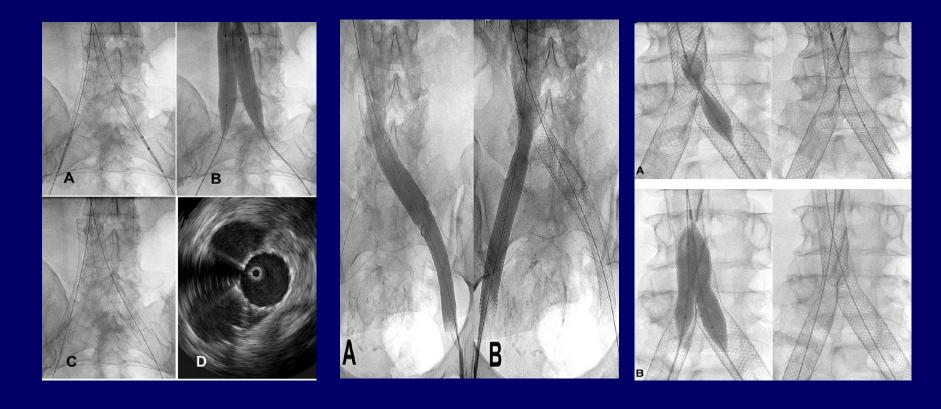
Conclusion: Contrary to arterial stenting, braided stainless steps can be safely placed in the venous system across the inguinal crease with no risk of stent fractures, narrowing due to external compression, focal development of severe in-stent restenosis, and no effect on long-term patency. The patency rate is not related to the length of stented area or the placement of the stent across the inguinal ligament, but is dependent upon the etiology and whether the treated postthrombotic obstruction is occlusive or non-occlusive. (J Vasc Surg 2008;48:1255-61.)

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The lower numbers represent total limbs at risk for each time interval (\pm SEM).

Bilateral stenting at the iliocaval confluence

Peter Neglén, MD, PhD, a Rikki Darcey, BS, a Jake Olivier, PhD, and Seshadri Raju, MD, a Flowood, Miss; and Sydney, New South Wales, Australia (J Vasc Surg 2010;51:1457-66.)



Bilateral stenting at the iliocaval confluence

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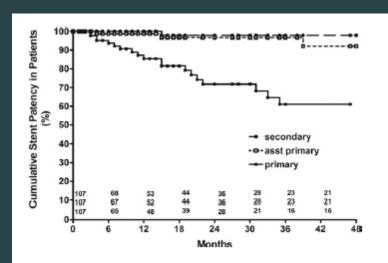


Fig 7. Cumulative primary, assisted primary, and secondary patency rates are shown for 107 patients with a bilateral stent system. The lower numbers represent limbs at risk at each time interval (all standard of means <10%).

Fenestrated stents do worse

Double barrel preferred

CLINICAL INVESTIGATION

Iliocaval Confluence Stenting for Chronic Venous Obstructions

Rick de Graaf • Mark de Wolf • Anna M. Sailer • Jorinde van Laanen • Cees Wittens • Houman Jalaie

Results Recanalization was achieved for all patients. In 15 (38 %) patients, a hybrid procedure with endophlebectomy and arteriovenous fistula creation needed to be performed because of significant involvement of inflow vessels below the inguinal ligament. Mean follow-up was 443 ± 438 days (range 7–1683 days). For all patients, primary, assisted-primary, and secondary patency rate of 27, 73, and 78 %, respectively. Twelve-month patency rates in the SECS group were 85, 85, and 95 % for primary, assisted-primary, and secondary patency. In the BECS group, primary patency was 100 % during a mean follow-up period of 134 ± 118 (range 29–337) days.

Conclusion Stenting of chronic bi-iliocaval obstruction shows relatively high patency rates at medium follow-up. Short-term patency seems to favor confluence stenting with balloon-expandable stents. Good secondary patency

Open and hybrid deep vein reconstructions: to do or not to do?

P Gloviczki*, M Kalra*, A A Duncan*, G S Oderich*, T J Vrtiska† and T C Bower*
*Division of Vascular and Endovascular Surgery; †Department of Radiology, Mayo Clinic, Rochester, MN, USA

Hybrid reconstructions



When to do it?

(1) The main indication for hybrid procedures, that we first reported on in 2001, 1 is extensive thrombosis of the common femoral vein that prevents primary iliofemoral venous stenting. If these patients have good inflow at least through the profunda femoris vein, a groin incision is performed, the femoral veins and saphenous vein are exposed and endophle-bectomy of the chronically occluded common femoral vein is done. The procedure is finished with saphenous vein or bovine pericardial patch angioplasty. Once inflow is assured, iliofemoral stenting is done with standard techniques.

When not to do it?

- (1) Poor inflow even to the profunda femoris vein;
- (2) If there are other contraindications for stenting and indications for open repair as listed above.

But...

REVIEW

Editor's Choice — A Systematic Review of Endovenous Stenting in Chronic Venous Disease Secondary to Iliac Vein Obstruction

M.J. Seager, A. Busuttil, B. Dharmarajah, A.H. Davies

Department of Surgery and Cancer, Imperial College London, Charing Cross Hospital, London, UK

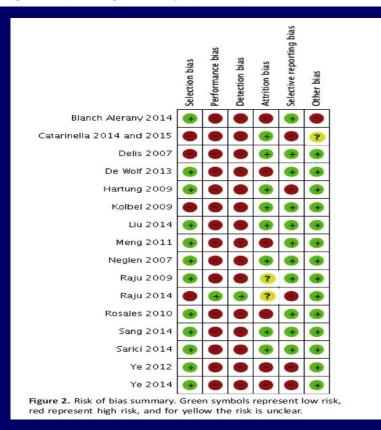


Table 2. GRADE assessment of the evidence.

Outcomes	Relative effect (95% CI)	No of events (studies)	Quality of the evidence (GRADE)
Severity of chronic venous disease			
Validated measures of CVD severity (CEAP, Villalta if post-thrombotic, etc.) Follow-up: 6—38 months	Not estimable	295 patients stented (5 studies ^{24,26,30,31,39})	$\oplus\ominus\ominus\ominus$ very low ^{a,b}
Other signs/symptoms of CVD			
See Table 3 Follow-up: 6—48 months	Not estimable	1984 patients stented (14 studies ^{24–26,29–36,38–40})	⊕⊖⊖⊖ very low ^c
Ulcer healing			
Complete healing (re- epithelialization) Follow-up: 0—46 months	Not estimable	400 ulcers (12 studies ^{24–26,30,31,33–36,38–40})	$\oplus \oplus \ominus \ominus$ low ^{d,e}
Quality of life			
SF-36, CIVIQ-2, CIVIQ-20, VEINES-QOL and VEINES-Sym Follow-up: 6—48 months	Not estimable	340 patients or limbs stented (3 studies ^{24,27,36})	$\oplus\ominus\ominus\ominus$ very low ^f
Stent patency			
% patent Follow-up: 6—48 months	Not estimable	2410 limbs stented (15 studies ^{24-27,29,31-40})	$\oplus\ominus\ominus\ominus$ very low ^g
Complications	N	500 11 11	
Incidence (%) Follow-up: 6—48 months	Not estimable	599 complications (15 studies ^{24–26,29–40})	$\oplus\ominus\ominus\ominus$ very low ^{h,i}

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

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Quality of life			
SF-36, CIVIQ-2, CIVIQ-20, VEINES-QOL and VEINES-Sym Follow-up: 6—48 months	n = 3	atients or limbs stented dies ^{24,27,36})	$\oplus\ominus\ominus\ominus$ very low ^f
Stent patency	\sim 3 sQoL + 1 s	gQoL	
% patent Follow-up: 6—48 months	Not estimable	2410 limbs stented (15 studies ^{24–27,29,31–40})	$\oplus\ominus\ominus\ominus$ very low ^g
Complications			
Incidence (%) Follow-up: 6—48 months	Not estimable	n=1	15
GRADE Working Group grades of evider High quality: Further research is very u		nce in the estimate of effect. Prim	ary

Moderate quality: Further research is likely to have an important impact on our confidence in the estimated patency

Low quality: Further research is very likely to have an important impact on our confidence in the estimate 32 - 99%the estimate.

Very low quality: We are very uncertain about the estimate.

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Eur J Vasc Endovasc Surg (2016) 51, 100-120

REVIEW

Editor's Choice — A Systematic Review of Endovenous Stenting in Chronic Venous Disease Secondary to Iliac Vein Obstruction

M.J. Seager, A. Busuttil, B. Dharmarajah, A.H. Davies

Department of Surgery and Cancer, Imperial College London, Charing Cross Hospital, London, UK

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WHAT THIS PAPER ADDS

This review demonstrates that quality of evidence behind the use of deep venous stenting to treat obstructive chronic venous disease is weak. However, the consistent effects and marked changes to disease course mean that it should be considered as an acceptable treatment. This review intended to influence clinical practice so vascular teams are aware of this, and it will serve to guide the future research that is needed.



Figure 2. Risk of bias summary. Green symbols represent low risk, red represent high risk, and for yellow the risk is unclear.



Deep vein reconstruction

There is no level 1 evidence

Axillary vein transfer in trabeculated postthrombotic veins

Seshadri Raju, MD, Petae Noodén MD Loffery Doolittle BS and Edward E Sesnadri Raju, NID, Peter Nachan Neovalve construction in postthromhotic Meydrech, PhD, Jackson Neovalve construction in postthromhotic

Purpose: This study assessed in trabeculated veins and w Methods: A total of 102 as Oscar Maleti, MD, and Marzia Lugli, with trabeculated veins. Me second axillary valve in 19

superficial and deep femora leaky axillary valves was po

from recurrent ulceration obtained in a matched grou preoperative ambulatory ve seconds), which was presen

of patients with severe pain Although these preliminary results are enc.

Contclusion: Axillary vein trechnique. (J Vasc Surg 2006;43:794-9.)

Technique indicated is eafer effe severe forms of postthromby.

severe forms of postthromby or other types of surgery fall. (1 Yasc Surg 1999), 2 yr. t. servative therapy or other types.

syndrome

Objective: The purpose of this study wa syndrome. The surgical procedure is descr Methods: From December 2000 to June (8 male and 8 female; median age, 55.5 ye mattus or utceration was pt (8 maic and 8 temale; median age, 55.5 yet a successful valve reconstruct postthrombotic syndrome. Surgical treatment of the postthrombotic syndrome. appearance, classification class C(0), Preoperative duple, routinely performed, Valvular cusps were seeming.

Results: In 16 lower extremities (89%), the Occurred. Neovake competence was confir showed a significant improvement in hem muscle pump function. In 17 limbs (95%) plethysmography) improved muscle pump function. In 17 limbs (95%) months. Early thrombosis below the neoral months of the plant of the provided in 55% of particular and provided in the provi ative swelling, the swelling occurred in three patients (17%), sens occurred in three patients (17%). Sensor patients in the patients of the swelling occurred in three patients (17%). 37 patients), it became wors Conclusions. Neovaire construction seems of Patients with severe pain of Patients with severe pain transfer actions a subsection for the construction of Vise Surv 2006;43:794-9.)

Deep Venous Reconstruction Primary or Post-throm

A. Lehtola, A. Oinonen, 12

¹Department of Vascular Surgery, Hels. ⁵²Department of Vascular and Apple

Objectives. To evaluate long-term durability and c consecutives. To estimate tong-term aurantizy and the instafficency as a part of rotatine workload at a unit Study design. Retrospective analysis of cluttal set PA Hents. Between 1996 and 2000, 43 patients under era menus. Democen 1290 una 2001, 93 puntents unaco Of these, 29 had secondary valve disease. Median fol

or nation ticers.

Methods. Socrety of the disease was assessed by the disease was asset which the disease was asset which was asset when the disease was asset which was a second was as a second was as a second was a second was a second was a s are around the control of the alse are the duplex evaluation. Certain for christal success use to aupoca communion. Orierts 30° carried success were constructions included 12 internal and 5 Surgical reconstructions included 12 internal and ourgem reconstructions anchured is internal and transferance of the 38 patients, six method at least transferance of the 38 patients, six method at least transferance of the six method at least transferance transpearmentons. Of the 35 persons, 8x needed at least months, and thereafter articular Two independs see montans, and increases arranged a results. The cornell cumulative distal success and Results. The oremal cumulative distant success risk were the most durable techniques with a cumulative were the most durable techniques with a cumulative were the most aurune techniques until a cumulative units 45%, and for transplantations 15%, O/13 real urus 45%, and for trianspartiations 16%. Cy 13 rec Conclusions. Overall results were unsatts actory. were acceptable.
© 2007 European Society for Vascular Surgery. Pub Keywords: Chronic remus insufficiency. Deep venen

Effect of external valvuloplasty of the deep vein in the treatment of chronic venous insufficiency of

the lower extremity

Shen-Ming Wang, MD, PhD, Zuo-Jun Hu and Cai-Sheng Ye, MD, PhD, Guangabou,

Objective: This study was conducted to verify the primary chronic venous insufficiency (PCVI). Methods: Forty patients with PCVI of the bilaters 80 limbs were classified as CEAP C2 to C4 with randomized into one of two groups according to canuormazeu 11500 one of two groups according to A, regardless of whether it was the right or left cs, regardness of whemer it was the right or let valvuloplasty of the femoral vein was combined w superficial venous system alone was performed compared by color duplex scanning, a color Dop score at 1 month, 1 year, and 3 years postoperar Results Within each group of limbs, no significant of limbs. The viricose veins resolved, there were a all cases. Leg heaviness was relieved completely in AND VEROUS VAIVE COMPREHENCE WAS achieved in 100%. venous varve competence was achieved in 100%, poetroperatively, respectively. The amount of veno, have postoperatively, respectively. The amount of relatively different between the two groups preoperatively unretent ocerreen the two groups preoperature) scores, and mulcle pumping indices improved mi Scores, and answer pringing indices did not improve (P < 01); musck pumping indices did not improve (1° < .01); muscle pumping indices did not improve significant differences in the amount of venous reagmucant curreptuces in the amount of venous reliable and 1 month and 1 year postoperatively (P) impos at 1 month and 1 year postoperatively (1) between group A and B limbo 3 years postoperation netween group A and 8 limbs 3 years portoperate.

Canclustions: External valvidoplasty of the femoral CONCESSIONS EXTERNAL VARIAGED AS OF the femore is improved the hemodynamic status of the lower is improved the nemodynamic status of the lower outcomes than surgical repair of the superficial scattering of the superficial scattering of the superficial scattering outcomes than surgical repair outcomes that such a surgical repair outcomes than surgical repair outcomes that such a surgical repair outcomes th

External Venous Valve Plasty (EVVP) in Patients with Primary Chronic Venous Insufficiency (PCVI)

A. Rosales,1* C.E. Slagsvold, ² A.J. Kroese, ¹ E. Stranden, ² Ø. Risum¹ and J.J. Jorgensen¹

¹Oslo Centre for Vascular Surgery, and ²Department of Vascular Diagnosis and Research,

Objectives. To evuluate the patency of EVVP and its effect in symptom relief, uteer heating and uter-free period in

Methods. Between 1993 and 2004, 1800 patients with CVI were evaluated and seventeen with PCVI were selected for Methods. Between 1893 and 2004, and analysis are recovered evacuate measurement (AVPI, colour durier ultresund Methods. Between 1993 and 2004, 1800 patients with CVI were evaluated and screnters with EVVI were selected for EVVP. They there all investigated with ambulatory tenous pressure measurement (AVP), colour dupler ultrasumd.

EVVP. They there all investigated with ambulatory tenous pressure. The CEAP classification was used to rouse the suices.

(CDLI), ascend into tenous modes and docconding video renew random. The CEAP classification was used to rouse the suices. EVVP. They were all investigated with ambidulary benous pressure measurement (AVP), colour duplex ultrasund.

(COU), ascending senggraphy and descending video verify raphy. The CEAP classification was used to group the patients.

See partieurs where CA four CS and consum CS. All had down rether and high brooks of AVP.

Six patients were C4, four C5 and seven C6. All had deep reflux and high levels of AVP.

Results. All procedures were technically successful. The ulcer healing rate ms 47 (57%) within 3 months. All C4 (mage 60 and 15 an Results. All procedures were technically successful. The ulcor healing rate was 47 (57%) within 3 months. All C4
patients experienced symptom improvement postopenitody and had a modian recurrence free period of 22 (range 6).

122) months. The C5 orough had an median where free period of 61 months (12—72) and the C8 of median 48 (12—72). patients experienced symptom improvement postoperatively and had a median recurrence free period of 72 (range 69-122) months. The CS group had an median user free period of 61 months (12-72), 12 multiple tube plastics into the CS group had an median user free period of 48 months (12-72), 12 multiple tube plastics into the CS group had an median competence period of 48 months (12-72), 2 multiple tube plastics into the plastics (4) roached a median competence period of 48 months (12-72). 122) months. The C5 group had at median tuker free period of 61 months (12-72) and the 65 of median 48 (12-72)
months. Single tube plasties (4) reached a median competence period of 45 months (12-72), 12 multiple value
at the same bruel sharot a modden 78 months (63-122) and 10 multilevel repairs median S4 months (12-96). Multiple value
at the same bruel sharot a modden 78 months (63-122) and 10 multilevel repairs median S4 months (12-96). months. Single take plasties (4) rowhed a median competence period of 48 months (12—72), 12 multiple rates plasties at the same level show a median 78 months (63—122) and 10 multilevel spairs median 54 months (22—96). Multiple rates at the same level show a median 78 months (63—122) and 10 multilevel spairs median 54 months (12—96). Multiple rates at the same level (multi-station plastics) performed on the C4 group seemed to yield the longest durability rate and the same level (multi-station plastics) performed on the C4 group seemed to yield the longest durability rates. at the same level shows a median 78 months (63–122) and 10 multilevel spairs median 54 months (12–96). Multiple value plastics at the same level (multi-station plastics) performed on the C4 group seemed to yield the longest durability ruth a median of 103 months (84–122).

a mediant of 103 months (84-122).

Conclusion. EVP with an ulcor healing rate of 57% and satisfactory symptom improposes to be related to clinical second surgicial treatment for selected partients with PCVI. The durability of this technique second to be related to clinical second and the multiplicity of repairs.

Keynourds: Vensus insufficiency, Valve plasty. Clinical and functional outcome, Vensus pressure measurements; Vensus refuse.

RCT vs Registry

RCTs

- Randomized
- Controlled selection criteria
- Study specific interventions
- · Hypothesis driven
- (Relatively) small population
- · Fixed timeframe
- Powered

Real-world studies

- Observational
- No or broad selection criteria
- No study specific interventions
- Hypothesis generating
- Large population
- · Long timeframe
- Flexible

RCT vs Registry

RCTs

Real-world studies





Powered

Flexible

