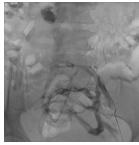


**VEITH SYMPOSIUM**  
Connecting The Vascular Community

## Update Of Post-Thrombotic Syndrome Prevention

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## Disclosure

**AlfaSigma – Speaker bureau, Research grant, Advisory board member**  
**Bayer – Speaker bureau**  
**Pfizer – Speaker bureau**  
**Balton – Research grant**  
**Medi – Speaker bureau**

### PTS risk factors - risk factors related to the patient initial status:

Age*	++
Obesity**	++
Preexisting varicose veins ***	++
Trombophilia presence	-

\*Older age: ↑ PTS risk from 30% to 3 fold  
 \*\* BMI>30: More than 2-fold ↑ PTS risk  
 \*\*\* Preexisting varicose veins – up to 2-fold ↑ risk of PTS

Kahn et al. et al. Determinants and time course of the postthrombotic syndrome after acute deep vein thrombosis. *Ann Intern Med* 2008; 148: 898-907.  
 Klok et al. et al. Prediction of post-thrombotic syndrome in a population with DVT after venous thrombolysis and primary venous occlusion. *J Thromb Haemostasis* 2013; 13: 479-85.  
 Kahn et al. et al. American Heart Association Council on Peripheral Vascular Disease, Council on Clinical Cardiology, and Council on Cardiovascular and Stroke Nursing. The postthrombotic syndrome: evidence-based prevention, diagnosis, and treatment strategies: a scientific statement for health-care professionals. *Circulation* 2014; 129: 2038-50.  
 Kahn et al. et al. Guidelines for prevention and treatment of the post-thrombotic syndrome. *J Thromb Haemostasis* 2014; 14: 144-53.  
 Klok et al. et al. Journal of Thrombosis and Haemostasis. 2013; 13: 240-249.  
 Urbanek et al. Post-thrombotic syndrome: a clinical review. *J Thromb Haemostasis* 2013; 13: 70-80.

### PTS risk factors related to the initial DVT characteristics:

Symptomatic DVT vs Asymptomatic	+/-
Provoked DVT vs Unprovoked	-
DVT location /massive proximal vs. distal/*	++

\*But distal DVT does not exclude PTS occurrence (PTS Rate 14-47%)  
/Poulsen P. (2004), Brown AL (2006), Lindblad M (2006), Klok VE (2006).

Kahn et al. et al. Determinants and time course of the postthrombotic syndrome after acute deep vein thrombosis. *Ann Intern Med* 2008; 148: 898-907.  
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 Ståhl et al. et al. Clinical Presentation and Deep and Long-Term Outcomes in Patients With Isolated Distal Deep Vein Thrombosis in the RETE Registry. *JAMA Cardiol* 2022; 7: 827-835.  
 Galis et al. et al. Long-term risk of postthrombotic syndrome after symptomatic distal deep vein thrombosis: the DVTx-PTS study. *J Thromb Haemostasis* 2020; 18: 827-834.

### Post-thrombotic syndrome prediction models

**SOX – PTS index /range 0-5/**  
 High risk predictors at baseline:

1 point: iliac vein DVT	1
2 points: BMI>35	2
1 point: Villata score 9-14 at baseline	1
1 point: Villata score >14 at baseline	1

**MEAN model** (proximal or distal DVT age >65):

Mean M et al. 2018	score
Age > 75	1
NSAID/antiplatelet	1
Multilevel thrombosis	1
Prior varicose vein surgery	1

**SOX** Several factors studied and proposed but still very difficult to predict on the individual basis which patients will develop and which one will do not develop prothrombotic syndrome ....

**Robins**

1 point: Heparin	1
1 point: Provoked DVT	1
1 point: Iliofemoral DVT	1
1 point: History of DVT	1

**HUANG model** (Chinese cohort, first proximal or distal DVT)

iliac vein compression	Score
Occlusion	2.5
Residual iliac – femoral vein thrombosis	3
Residual femoral – popliteal vein thrombosis	3
Insufficient anticoagulation	6


Kabatavci A, Durrant T, Adair SA. Six-Factor Investigation: Development of a Distal Prediction Model for the Postthrombotic Syndrome in a Cohort of Patients With Proximal Deep Vein Thrombosis. *J Thromb Haemostasis* 2018; 18: 242-250. Urbanek T, Urbanek A, Adair A, Agrebi B, Matuszewska M. Determinants and time course of post-thrombotic syndrome in a population with DVT after venous thrombolysis and primary venous occlusion. *J Thromb Haemostasis* 2013; 13: 479-85. Urbanek T, Urbanek A, Adair A, Agrebi B, Matuszewska M. Determinants and time course of post-thrombotic syndrome in a population with DVT after venous thrombolysis and primary venous occlusion. *J Thromb Haemostasis* 2013; 13: 479-85. Urbanek T, Urbanek A, Adair A, Agrebi B, Matuszewska M. Determinants and time course of post-thrombotic syndrome in a population with DVT after venous thrombolysis and primary venous occlusion. *J Thromb Haemostasis* 2013; 13: 479-85. Urbanek T, Urbanek A, Adair A, Agrebi B, Matuszewska M. Determinants and time course of post-thrombotic syndrome in a population with DVT after venous thrombolysis and primary venous occlusion. *J Thromb Haemostasis* 2013; 13: 479-85. Urbanek T, Urbanek A, Adair A, Agrebi B, Matuszewska M. Determinants and time course of post-thrombotic syndrome in a population with DVT after venous thrombolysis and primary venous occlusion. *J Thromb Haemostasis* 2013; 13: 479-85.

### PTS risk factors related to the treatment phase



Poor INR control in the treatment phase /especially within first 3 months/	++
LMWH vs VKA (in favor of LMWH)	+
„Residual thrombosis“ /non complete recanalisation and thrombus resolution/	+
Incomplete resolution of the symptoms within 1st month of the treatment	+

Kahn et al. et al. Determinants and time course of the postthrombotic syndrome after acute deep vein thrombosis. *Ann Intern Med* 2008; 148: 898-907.  
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**ExACT study** **Anticoagulant therapy duration in PTS prevention ?**  
 Extended AntiCoagulation Treatment for venous thromboembolism  
 281 patients with unprovoked proximal DVT after 3 months of anticoagulation  
 anticoagulation vs anticoagulation continuation  
 Follow up - 24 months  
 Mean Villalta score: 5.09 vs 5.0; p=0.9  
 No difference in the VEINES-QoL and EQ-5D-3L quality of life scores  
**Conclusion: No direct influence of the anticoagulation duration on the PTS occurrence.**  
**VTE recurrence rate 13.54 vs 2.75 events/100 patients years**  
 Bradbury C, Fletzner K, Sun Y, Heneghan C, Gardiner C, Roelle A, et al. A randomised controlled trial of extended anticoagulation treatment versus standard treatment for the prevention of recurrent venous thromboembolism (VTE) and post-thrombotic syndrome in patients being treated for a first episode of unprovoked VTE (the EXACT study). *Br J Haematol.* 2020; 188: 962-975

**DVT recurrence (ipsilateral)**  
 Post – thrombotic syndrome risk 2-6x 

Optimal duration and quality of antithrombotic VTE treatment

Risk of recurrence  Risk of bleeding 

Kahn S, Comella A, Guzman M, Evans N, Gruber J, Gutierrez M, Gupta DK, Prandoni P, Vittinghoff DM, Villalta ME, Wells J, on behalf of the American Heart Association Council on Clinical Cardiology, Council on Clinical Cardiology and Council on Cardiovascular and Stroke Nursing, The post-thrombotic syndrome: evidence-based prevention, diagnosis and treatment strategies: a scientific statement from The American Heart Association. *Circulation.* 2014

**DVT recurrence prevention options:**

- Full anticoagulation  
 VKA, DOACS /high efficacy, continuous risk of bleeding/
- Reduced doses of anticoagulants  
 VKA INR 1.5-2.0,  
 Rivaroxaban 10 mg /EINSTEIN CHOICE/  
 Apixaban 2x 2,5 mg /AMPLIFY EXTENSION/
- Sulodexide /SURVET/  
 Sulodexide 2x500 LSU
- ASA /WARFASA, ASPIRE/

**DOAC's in PTS prevention ?**

- DOACs\* improve the compliance to the effective anticoagulant treatment
  - no need of INR control
  - simplified therapy
  - high clinical efficacy and safety
- DOACs and PTS rate decrease ? Do all DOACs works in the same way?

\*DOACs - Direct Oral Anticoagulants

**Rivaroxaban DVT treatment in PTS rate reduction**

Karathanos C, et al. Efficacy of rivaroxaban in prevention of post-thrombotic syndrome: **A systematic review and meta-analysis.** *J Vasc Surg Venous Lymphat Disord.* 2021; 9: 1568-1576.e1.  
 6 retrospective studies, 2 RCT; 59199 patients

**PTS rate reduction in rivaroxaban group (vs warfarin treated patients)**  
**OR 0.52; 95% CI 0.43-0.63; (p < 0.001)**

Severe PTS reduction: 3.7% (rivaroxaban) vs 6.4% (warfarin); OR = 0.55; 95% CI, 0.36-0.85; p = 0.024

Jeraj L, Jezovnik MK, Poredos PI. Rivaroxaban versus warfarin in the prevention of post-thrombotic syndrome. *Thromb Res.* 2017; 157: 46-48

/prospective RCT/  
 Rivaroxaban (61 pts.) vs Warfarin (39 pts.)  
 Follow up: 23 months (median) after DVT episode

**PTS (Villalta scale): 25% - rivaroxaban vs. 49% - warfarin /p=0.013/**

OR 2.9 (1.2-6.8; p=0.014) for PTS development in warfarin group /compared to rivaroxaban/  
 adjusted OR 3.5 (1.1-11.0; p=0.035).  
 PTS patients: more frequently recurrent DVT 15% vs 3% (no PTS); p = 0.03

**CONCLUSIONS:**  
 Treatment of DVT with rivaroxaban might be associated with a lower risk for PTS development. A larger randomized trial would be needed for stronger evidence.

**/prospective RCT/**  
 de Athayde Soares R, Malieo MF, Brochado Neto FC, Nogueira MP, Almeida RD, Sacilotto R  
**Comparison of the recanalization rate and postthrombotic syndrome in patients with deep venous thrombosis treated with rivaroxaban or warfarin.** *Surgery.* 2019;166:1076-1083.

Prospective, consecutive, randomized, blind cohort study - 84 DVT patients  
**Oral anticoagulation for 6 months:**  
**rivaroxaban vs warfarin**  
 /follow up 12 months/

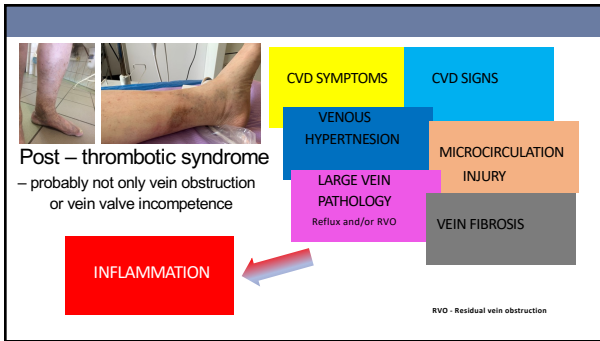
**PTS rate 8.7% vs 28.9% (P < .001; OR 4.278)**

Total venous recanalization @ day 360: 13.2% vs 76.1%; (P < .001).  
 Five patients in the total cohort (6%) showed no venous recanalization - all of them in the warfarin

**PTS rate reduction by DOACs?**  
**Unknown for other DOACs - very limited data, no prospective dedicated studies**

**Dabigatran data (no effect on PTS occurrence proved) – RECOVER study patient follow up**  
 Wik HS, Kahn SR, Eriksson HI, et al. Post-thrombotic syndrome in patients with venous thromboembolism treated with dabigatran or warfarin: a long-term cross-sectional follow-up of RECOVER study patients. *J Thromb Haemost.* 2021; 19: 2495-2503

**Edoxaban data (no effect on PTS occurrence proved) – HOKUSAI – VTE substudy:** 316 patients, mean time since randomisation 7.0 yrs.  
 Bistervels I, et al. Postthrombotic syndrome and quality of life after deep vein thrombosis in patients treated with edoxaban versus warfarin. *Res Pract Thromb Haemost.* 2022; 6: e12748



**PTS and anti-inflammatory treatment ?**

**801 consecutive patients with proximal DVT and no active cancer**  
**Including 82 patients with Chronic Inflammatory Disease (CID)**  
 /Inflammatory bowel disease, systemic rheumatoid disease, gout/

2 year prospective follow-up  
 PTS (Villata score), Residual Venous Obstruction – RVO (US based evaluation)

**PTS rate: CID patients 35.4% vs. No CID 18.9% (p < 0.001) HR 1.72 [1.15–2.58]**

**Residual Venous Obstruction /Less commonly in antiinflammatory treated CID patients/**  
 CID treated 26.7% vs CID with no antiinflammatory treatment 51.6% (p<0.05)

**PTS in CID DVT patients in relationship to the antiinflammatory treatment:**  
**CID treated 26.1% vs CID with no antiinflammatory treatment 47.2% p=0.047**

**Local anti-inflammatory treatment in PTS prevention ?**

**DEXTERITY AFP TRIAL**  
Local perivenous injection of dexamethasone sodium phosphate after successful vein thrombectomy

1st phase trial /21 patients/  
 6 month follow up (20 pts.): **mild PTS 5% /no moderate to severe PTS reported/**  
 Villata score reduction /from baseline to 6 months/ 7.3 ± 0.7 → 1.6 ± 0.4

2nd phase /60 participants/ RCT on the local drug delivery

D. Dexter @VEINS 2023 Las Vegas October 28-30

Category	Venoactive drugs
<b>Plant derived</b>	
	Micronised purified flavonoid fraction
	Diosmin
<b>Gamma-benzopyrones (flavonoids)</b>	Rutin, Rutosides
	O-(beta-hydroxyethyl)-rutosides
	Kaempferol glucoside, Quercetin glucuronide
	Procyanidins
	Anthocyanins
<b>Alpha-benzopyrones</b>	Cumarin
<b>Saponins</b>	Ruscus extract
	Horse chestnut seed extract, Escin
<b>Other plant extracts</b>	Ginkgo Biloba extracts
<b>Chemical synthesis</b>	
<b>Synthetic products</b>	Calcium dobesilate
	Naftazone
	Banzarone
<b>Animal derived</b>	
<b>Glycosaminoglycans</b>	Sulodexide

**VENOACTIVE DRUGS IN PTS PREVENTION**  
?

**Statins in PTS prevention**

RCT: 234 DVT patients

**LMWH vs LMWH + rosuvastatin**

<u>CRP levels after 3 months</u>	22.39 vs 4.17	p=0.018
<u>Villalta score after 3 months</u>	7.79 vs 3.45	p = 0.035
<b>PTS rate</b>	<b>48.5% vs 38.3%</b>	<b>p = 0.019</b>

*Conclusion: adjuvant rosuvastatin treatment in patients diagnosed of DVT improve CRP level and reduce PTS incidence*

San Norberto EM et al. Effects of rosuvastatin as an adjuvant treatment for deep vein thrombosis. Vasa 2016; 45: 133-140

**SAVER Study**

**Statins for Venous Event Prevention**

**SAVER study results**

312 pts – RCT rosuvastatin 20 mg (180 days) vs no rosuvastatin

**Overall PTS rate (Villalta >4): 29.7% (rosuvastatin) vs 25.5% (control arm) p=0.04**

**Severe PTS @ day 180: 2.0% vs 2.7% (control arm) p=1**

Delluc A, Ghanima W, Kovacs M, et al. Prevention of post-thrombotic syndrome with rosuvastatin: A multicenter randomized controlled pilot trial (SAVER). Thromb Res. 2022; 213: 119-124

*/randomisation and beginning of rosuvastatin therapy up to 30 days from DVT diagnosis ?/*

**Statins in PTS prevention – the hypothesis has to be confirm in further trials !**

**Caiano L. Role of statins in the prevention of post-thrombotic syndrome after a deep vein thrombosis event: a systematic review and meta-analysis. J Thromb Haemost. 2023; 21(4): 944-952.**

5 studies (2 retrospective cohorts/3 randomized controlled trials [RCTs])

**The pooled PTS incidence:**  
34.8% per patient-year (95% CI, 9.5-127.4) (statins) vs 41.6% per patient-year (95% CI, 13.2-132) (control)

**[22% PTS rate reduction with statins in systemic review]**

**Meta-analysis of 2 retrospective cohorts: significant reduction in the risk of developing PTS (IRR, 0.68; 95% CI, 0.51-0.91)**

**Meta-analysis of 3 RCTs: no reduction in PTS occurrence (IRR, 0.92; 95% CI, 0.68-1.25)**

*Conclusions: Although this systematic review suggests that statins may reduce PTS incidence by 22% after deep vein thrombosis, meta-analysis of RCTs showed no risk reduction. Confirmation of the efficacy of statins on the prevention of PTS should be assessed in larger RCTs.*

**Prospective RCT**

**The Effects of 3-Month Rosuvastatin Adjuvant Therapy on Post Thrombotic Syndrome following Deep Vein Thrombosis; a Randomized Clinical Trial**

3 months follow up (PTS rate – Brandjes criteria)  
182 patients: overall PTS rate 17%

	severe PTS	Moderate PTS	Mild PTS
Warfarin	8%	12%	12%
Warfarin + Rosuvastatin	0	4.0%	10.2%
Rivaroxaban	4.5%	6.8%	9.1%
Rivaroxaban + Rosuvastatin	0	2.5%	10%

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**Prospective RCT**

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3 months follow up (PTS rate – Brandjes criteria)  
182 patients: overall PTS rate 17%

	severe PTS	Moderate PTS	Mild PTS
Warfarin	8%	12%	12%
<b>Warfarin + Rosuvastatin</b>	<b>0</b>	<b>4.0%</b>	<b>10.2%</b>
Rivaroxaban	4.5%	6.8%	9.1%
<b>Rivaroxaban + Rosuvastatin</b>	<b>0</b>	<b>2.5%</b>	<b>10%</b>

**Conclusion: Rosuvastatin administration can significantly reduce the incidence of PTS and cause a difference in the size of the lower limbs within 3 month. Rosuvastatin administration in combination with rivaroxaban or warfarin significantly reduces the level of inflammatory factors including CRP and D-dimer, compared to patients receiving anticoagulants alone.**

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**Early thrombus removal in PTS prevention ?**

- Surgical thrombectomy
- Endovascular /CDT, pharmacomechanical, mechanical/ methods

**„Open vein“ concept**

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Javed A. et al. Meta-analysis of lytic catheter-based intervention for acute proximal deep vein thrombosis in the reduction of post-thrombotic syndrome. J Vasc Surg Venous Lymphat Disord 2023;11:866-75

**CAVENT ATTRACT CAVA**

**A**

Study or Subgroup	LCBI Events	Total Events	Total	Weight	M-H, Forest, 95% CI	Year
Wang 2016	17	87	63	89	0.40 [0.45, 0.79]	2016
Vendramin 2017	17	316	171	155	0.87 [0.81, 1.16]	2017
Nelson 2021	29	62	40	58	0.88 [0.49, 0.93]	2021
<b>Total (95% CI)</b>			<b>485</b>	<b>302</b>	<b>0.84 [0.74, 0.95]</b>	
<b>Total events:</b>	<b>223</b>	<b>274</b>				
<b>Heterogeneity:</b>	<b>Chi<sup>2</sup> = 10.55, I<sup>2</sup> = 2.00, P = 0.001, I<sup>2</sup> = 83%</b>					
<b>Test for overall effect:</b>	<b>Z = 2.73, P = 0.006</b>					

**B**

Study or Subgroup	LCBI Events	Total Events	Total	Weight	M-H, Forest, 95% CI	Year
Wang 2016	6	87	84	12	0.41 [0.13, 1.00]	2016
Vendramin 2017	49	316	84	155	0.75 [0.56, 1.01]	2017
Nelson 2021	17	62	58	58	0.90 [0.56, 1.76]	2021
<b>Total (95% CI)</b>			<b>485</b>	<b>302</b>	<b>0.73 [0.58, 0.97]</b>	
<b>Total events:</b>	<b>82</b>	<b>314</b>				
<b>Heterogeneity:</b>	<b>Chi<sup>2</sup> = 2.24, I<sup>2</sup> = 2.00, P = 0.331, I<sup>2</sup> = 11%</b>					
<b>Test for overall effect:</b>	<b>Z = 5.02, P = 0.001</b>					

**A/ PTS rate reduction: RR 0.84; 95% CI 0.74-0.95; P = 0.006**  
**B/ Severe and moderate PTS rate reduction: RR 0.75; 95% CI 0.58-0.97; P = 0.03**  
**LCBI – Lytic catheter based intervention**

Conclusions: Pooling of current best evidence suggests that LCBI in acute proximal DVT decreases the rate of PTS and moderate to severe PTS with a number needed to treat of 12 and 18, respectively. However, this is complicated by a significantly higher rate of major bleeding with a number needed to treat of 37. This evidence supports the use of LCBI in selected patients, including those who are at low risk of major bleeding.

The CLOUT (ClotTriever Outcomes) registry /ClotTriever for DVT: CLOUT Registry/  
 Prospective, multicenter study evaluating patient outcomes for proximal lower extremity deep vein thrombosis (DVT) treated with mechanical thrombectomy (MT) using the ClotTriever system

**Mechanical methods of thrombus removal**

**PTS occurrence (overall): 19.3%**  
**Moderate-to-severe post-thrombotic syndrome (PTS; Villalta score ≥ 10): 8.8%**  
 Venous patency /presence of flow with normal or partial compressibility on duplex US/ - 94.2%

D. Dexter @ VIVAOctober 28-30 2023, Las Vegas, Nevada.

Published: Bishara MB et al. One-Year Clinical Outcomes Following Mechanical Thrombectomy for Deep Vein Thrombosis: A CLOUT Registry Analysis J Soc Cardiovasc Angiogr Interv. 2024 Feb 15;33(Part A):101307.

**RCT: ClotTriever vs anticoagulation /DEFIANCE TRIAL/**

**MEDICAL COMPRESSION in PTS PREVENTION**

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**Compression stocking use after DVT episode for post-thrombotic syndrome prevention**

**Proximal DVT**  
**Knee-length elastic CS /30 – 40mmHg/ - 2 yrs.**  
**Follow up - 5 yrs**

**PTS (after 49 months (mean)): CS - 26% vs Control - 49% NNT 5 (3-11)**

Prandoni P et al. Below – knee elastic compression stocking to prevent the post thrombotic syndrome. Ann Int Med. 2004; 141: 4: 249-256

Class II ECS vs „Placebo stockings“  
 Post-thrombotic syndrome /Ginsberg Criteria/

**SOX TRIAL**

Randomization: ECS - 410 pts vs Placebo ECS

Primary outcome - the cumulative incidence of PTS from 0 to 24 months follow-up

**Cumulative incidence of PTS**  
**14.2% active ECS/ vs. 12.7% /placebo ECS/ (HR 1. 13; 95% CI 0.73-1.76; p=0.58)**

Most proximal extent of DVT	Active ECS	Placebo ECS	Active ECS:
Iliac	10.8%	12.4%	Knee – length
Common femoral	26.7%	27.2%	30-40 mm Hg compression.
Femoral	31.3%	31.2%	
Popliteal	31.3%	29.2%	

Kahn S. et al. Compression stockings to prevent post-thrombotic syndrome: a randomised placebo controlled trial. Lancet 2014; 383:880-88

Meng J et al. It is necessary to wear compression stocking and how long they should be worn for prevention of post thrombotic syndrome? A meta –analysis of randomized controlled trials. Thrombosis Research. 2023; 225: 79-86

9 RCT (including 2 studies concerning treatment duration)  
 Elastic Compression Stocking in acute DVT patients vs no compression or placebo

Study or Subgroup	Compression stockings Events	Total	None stockings or placebo Events	Total	Weight	M-H, Random, 95% CI	Risk Ratio	M-H, Random, 95% CI
Ashworth 2008	11	66	17	66	100%	0.68 [0.31, 1.52]		
Brandes 1997	30	56	69	56	17.2%	0.44 [0.22, 0.93]		
Chen 2011	0	24	1	23	0.0%	0.02 [0.01, 0.46]		
Jiang 2015	29	38	23	35	18.2%	1.00 [0.46, 1.52]		
Kahn 2014	136	428	168	368	20.0%	1.05 [0.86, 1.28]		
Parvizi 2004	23	90	44	80	19.0%	0.52 [0.30, 0.79]		
Yang 2015	42	113	89	110	19.7%	0.75 [0.56, 1.02]		
<b>Total (95% CI)</b>		<b>882</b>	<b>842</b>	<b>788</b>	<b>100.0%</b>	<b>0.73 [0.63, 1.08]</b>		
<b>Total events:</b>	<b>311</b>		<b>301</b>					
<b>Heterogeneity:</b>	<b>I<sup>2</sup> = 0.00, Chi<sup>2</sup> = 0.00, I<sup>2</sup> = 0.00, P = 0.99, I<sup>2</sup> = 0.0%</b>							
<b>Test for overall effect:</b>	<b>Z = 1.89, P = 0.06</b>							

**Post thrombotic syndrome (overall) rate reduction (RR 0.73, 95% CI 0.53 to 1.00; p = 0.05)**  
**But...no significant difference in severe post thrombotic syndrome rate.**

**DVT patient**

compression or no compression (in 2024)  
in DVT patients in acute DVT treatment and PTS prevention?

Be carefull with final conclusion!

**MEDICAL COMPRESSION in PTS PREVENTION**

or rather ...

**Immediate MEDICAL COMPRESSION and early MOBILISATION ...**

**DVT treatment – early mobilisation and ambulation decrease the Postthrombotic Syndrome occurrence**

- Immobilisation (9 days) without compression stocking - 17
- Unna boots + early mobilisation - 18
- Elastic compression + early mobilisation - 18

**Follow up 28,9 (+/- 4,9) months**

Assesment of PTS severity

Villalta scale: < 5 points (**no PTS**)

- Walking patients - 46%
- Immobilisation - 18%

Patsch H et al. Immediate mobilisation in acute vein thrombosis reduces post thrombotic syndrome. Int Angiol 2004; 23: 206-12

Reduced incidence of vein occlusion and postthrombotic syndrome after **immediate compression for deep vein thrombosis**

**No compression vs acute compression within 24 hours of diagnosis of DVT with either multilayer bandaging or compression hosiery (PROGRESS)**

Number: 30, 46, 41, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 100

**4 RCT studies (687 patients) which met inclusion criteria for early initiation and consistency over of ECS in patients with an asymptomatic DVT.**

ECS significantly reduced the incidence of mid-moderate PTS (OR: 0.46; 95% CI: 0.36-0.63)

ECS significantly reduced severe PTS (OR: 0.44; 95% CI: 0.28-0.58)

ECS significantly reduced the incidence of post-thrombotic syndrome rate (after 24 months)

No Residual Venous Obstruction	46.0%
Residual venous obstruction	54.0%

OR, 0.65; 95% confidence interval, 0.49-0.92; P=0.013

**Compression in PTS prevention**

**Consensus Statement for compression stockings in venous & lymphatic disorders**  
International Consensus Group: E. Falck, H. Parfitt, J. Heller, G. Lattimer, G. Moss, B. Neumann, T. Ustianski, H. Weibner, S. Gaillard, P. Carpentier. Phlebology 2018

**Recommendation 18**

**Use of medical compression stocking as early as possible after diagnosis of DVT in order to prevent PTS**

/1B\*

\*downgraded from 1A in the 2008 consensus document

**CHAPS**  
**Compression Hosiery to Avoid Post-Thrombotic Syndrome**

**Objective**  
To measure the difference in incidence of post-thrombotic syndrome at a median of 18 months follow up after first, acute DVT between standard clinical care (anticoagulation) and the intervention arm (a graduated compression stocking and the standard clinical care (anticoagulation)).

**Inclusion Criteria**  
• Symptomatic presentation of first deep vein thrombosis, <2 weeks from diagnosis  
• Imaging confirmed, lower limb deep vein thrombosis (popliteal, femoral, iliac or combination)

**Primary endpoint:** incidence of Post Thrombotic Syndrome (PTS) using the validated Villalta criteria over a median 18 month follow up (range 6 to 30 months).

**NIHR** National Institute for Health Research  
**CHAPS** Compression Hosiery to Avoid Post-Thrombotic Syndrome (CHAPS) Trial

European Venous Forum in Cracov/Poland (26-28.06.2025)



**25<sup>th</sup>** Annual Meeting of the European Venous Forum

26-28 June 2025  
Krakow, Poland

**Save the Date**

For more information, visit: [europeanvenousforum.org](http://europeanvenousforum.org)

**Call for abstract – deadline 14th February 2025**