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SVT Medical Management In 2024: Is There A Role For Surgical Intervention And When?



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Disclosures

I have the following potential conflicts of interest:

- Consulting / Conferences / Honoraria:
 - Medtronic
 - Cook Medical
 - Boston Scientific
 - BD
 - Gore Medical
- Research funding:
 - Medtronic
 - Gore Medical
 - Laboratoires Urgo

Introduction



Common problem
Important recent advances in understanding
Poorly covered topic in venous guidelines

Superficial vein thrombosis – POST study

Annals of Internal Medicine

Superficial Venous Thrombosis and Venous Thromboembolism A Large, Prospective Epidemiologic Study

844 patients with SVT in primary and secondary care units in France
210 (24.9%) had DVT or symptomatic PE at presentation

Superficial vein thrombosis – POST study

Thromboembolic Event (n = 586)	Incidence (95% CI), n (%) ^a
Any	58 (10.2 [7.7–12.7])
Asymptomatic	40 (6.7 [5.0–8.6])
PE or DVT	18 (3.1 [1.8–4.8])
DVT	15 (2.8 [1.4–4.2])
Proximal	7
Distal	8
SVT	3 (0.5 [0–1.2])
Recurrence	10 (1.9 [0.7–3.0])
Extension	18 (3.1 [1.8–4.8])
Asymptomatic	12 (2.1 [0.9–3.2])

Patients with isolated SVT given anticoagulation for median 11 days
Follow-up to 3 months
10.2% risk of VTE

INSIGHTS SVT study

Prospective observational study including 872 patients in Germany

197 (22.5%) underwent invasive procedure
817/872 (93.4%) received anticoagulation

INSIGHTS SVT study

Only 5 patients underwent acute junction ligation

Majority of procedures were superficial venous interventions

136/197 (64.5%) of interventions were after 3 months

Other studies

REVIEW: VENOUS DISEASE
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Current evidence on the effectiveness and safety of oral anticoagulants in superficial venous thrombosis: a systematic review and meta-analysis
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PDF | Supplementary Materials

INTRODUCTION: Previous studies suggest fondaparinux as an effective regimen for superficial venous thrombosis (SVT), but the incorporation of prolonged parenteral injection, has prompted investigations into oral anticoagulants (OACs). This study aims to evaluate the current evidence on the effectiveness and safety of OACs in the treatment of SVT.

EVIDENCE ACQUISITION: Following the PRISMA 2020 guidelines, we conducted a systematic review and meta-analysis registered in PROSPERO CRD420230262424. A comprehensive literature search was performed across multiple databases up to April 2024. Studies were included if they involved adult patients diagnosed with SVT, treated with OACs, and reported relevant efficacy and safety outcomes. Both randomized controlled trials (RCTs) and observational studies were considered. Data extraction and risk of bias assessments were independently performed by two reviewers.

EVIDENCE SYNTHESIS: The analysis identified 1231 results, with six studies (n=167) being three prospective cohort studies meeting inclusion criteria. Meta-analysis for Rivaroxaban-treated group showed DVT occurrence was 1.20% (95% CI: 0.1-2.3%), SVT recurrence was 0.20% (95% CI: 0.00-1.0%), SVT recurrence was 0.79% (95% CI: 0.0-2.30%), clinically relevant non-major bleeding was 1.01% (95% CI: 0.46-1.71%), minor bleeding was 5.04% (95% CI: 3.2-8.01%). These estimates were for patients treated with rivaroxaban 15-20 mg once daily over 45 days to 37 months. No major bleeding was reported with rivaroxaban 10 mg once daily.

CONCLUSIONS: This systematic review and meta-analysis demonstrate that OACs, especially rivaroxaban, are effective and safe for the treatment of SVT. They offer a convenient alternative to parenteral anticoagulants, generally improving patient compliance and outcomes. However, further large-scale studies are warranted to confirm these findings.

Published guidelines

Clinical Practice Guidelines for Vascular Surgery (ESVS) 2021 Clinical Practice Guidelines on the Management of Venous Thrombosis¹²

Specific and extensive chapter on management of SVT

A few recommendations, focused on anticoagulation

Antithrombotic Therapy for VTE Disease Second Update of the CHEST Guideline and Expert Panel Report

Other important considerations

- Most studies have not included SVT close to the saphenofemoral or saphenopopliteal junctions
- The follow-up period is usually short (long-term outcomes are unclear)
- Vast majority of cases occur in patients with varicose veins
- Consider thrombophilia / malignancy / other diagnosis (such as Burger's disease) if no varicose veins

Superficial vein thrombosis

- SVT is not a benign pathology
- Significant risk of VTE at presentation
- Ongoing risk of VTE
- Several types:
 - In varicosities / short segment of truncal vein
 - In truncal vein (>5cm), but away from junction
 - In truncal vein <3cm from junction
 - Extending into deep vein

ESVS Venous Thrombosis guidelines 2021

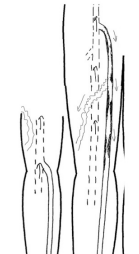
Exclude concomitant DVT and evaluate anatomical position / extent of SVT

High risk: treatment dose for 3 months

Intermediate risk: low dose prophylaxis 6 weeks

Ablate reflux once acute phase settled

Superficial venous ablation





Very common to see recanalized, scarred, incompetent saphenous vein after SVT

High risk of recurrent VTE

Reasonable to ablate superficial venous reflux

May be challenging and remember VTE prophylaxis

Conclusions



Superficial vein thrombosis is not a benign entity and more aggressive diagnosis and treatment is needed

Acute surgical intervention is very rarely needed (anticoagulation is very safe and effective)

Vast majority of surgical interventions are for residual superficial reflux and probably best performed out of the acute phase