





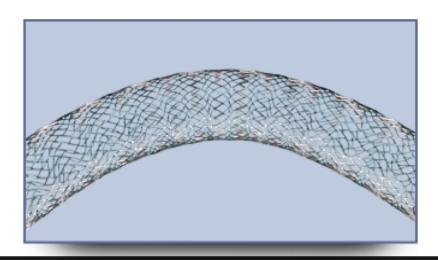
12 M Outcomes of the Veniti Vici Venous Stent System for iliofemoral deep vein thrombosis

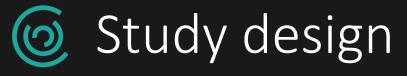
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- Self-expanding stent
- Nitinol alloy
- Over the wire system
- High radial force and flexibility
- Harmonious alignment
- Closed-cell geometry
- Compatible with .035 inch (.89mm) guidewire
- 9 French introducer sheath.

VEN12060	12mm x 60mm Stent System
VEN12090	12mm x 90mm Stent System
VEN12120	12mm x 120mm Stent System
VEN14060	14mm x 60mm Stent System
VEN14090	14mm x 90mm Stent System
VEN14120	14mm x 120mm Stent System
VEN16060	16mm x 60mm Stent System
VEN16090	16mm x 90mm Stent System
VEN16120	16mm x 120mm Stent System





- Retrospective analysis
- Acute and chronic obstructive deep venous thrombosis (DVT) of the pelvic vessels
- December 2015 February 2017
- Two centers
- 26 patients



Endpoints:

Primary Endpoint

- patency rate of the stent

Secondary Endpoint

- postoperative clinical improvement

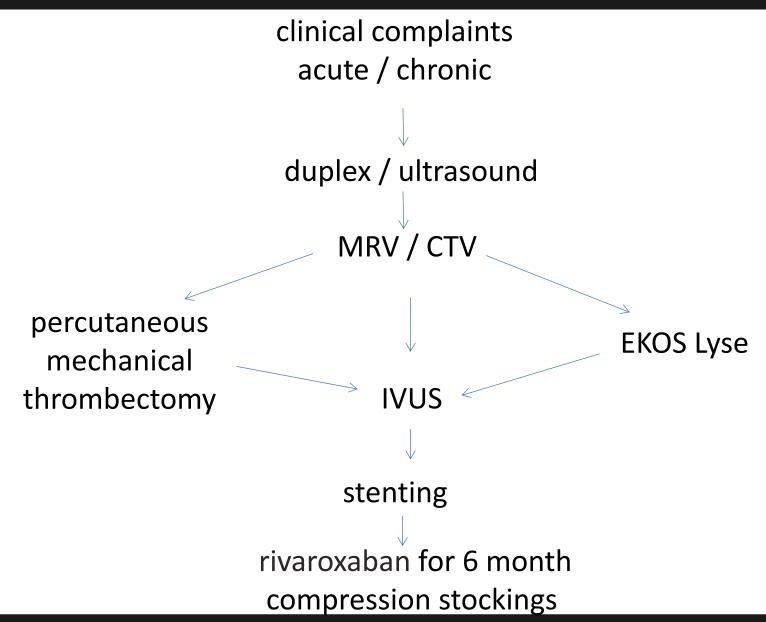
(CEAP - classification system)

(VCSS – venous clinical severity score)





Prior after





Baseline characteristics

Age	18-81 (Ø52)	
Female gender	21 (80%)	
Current smokers	6 (23%)	
Pulmonary embolism	1 (4%)	
Previous pulmonary embolism	7 (27%)	
Previous deep vein thrombosis	10 (38%)	
Venous claudication	15 (58%)	
Clotting disorders	1 (4%)	
Neoplastic disease	3 (12%)	
Diabetes	3 (12%)	
Hypertension	7 (27%)	

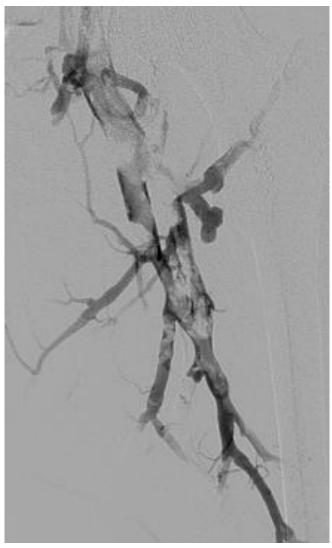


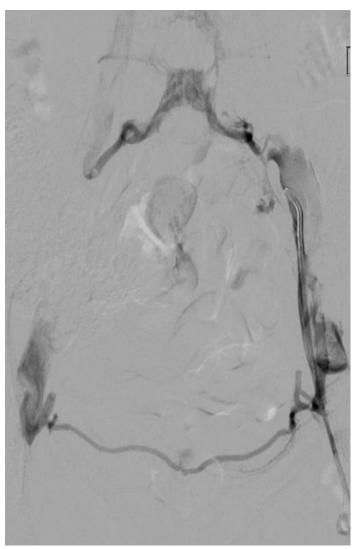
Baseline characteristics

characteristics	number	
Acute < 30 days	13 (50%)	
Chronic > 30 days	13 (50%)	
Occlusion	21 (81%)	
may thurner syndrom	15 (58%)	
left ilofemoral vein	18 (69%)	
EKOSE (ultrasound enhanced thrombectomy>)	4 (15%)	
percutaneous mechanical thrombectomy	7 (27%)	



© Case presentation





DVT of the left pelvic veins



Final result after stent deployment



Early and long term Outcomes

Endpoint	N	(%)
Technical success	26	100%
Major bleeding	0	0%

Endpoint	@ 6 Monts	@ 12 Months
primary patency	92%	85%
secondary patency	100 %	100%
CEAP improvement		76 %
venous clinical severity score VCSS improvement		69%

- Early Occlusion (< 30 days):
- 1 patient (4%)
- Secondary interventions @12 Months
- 4 patients (15%)
 - thrombectomy with Stent-in-Stent n-3 (12%)
 - endovascular phlebectomy n-1 (4%)

- Dedicated stent design
- High Technical success
- Acceptable clinical outcomes
- CEAP / VCSS improvement @ 12 months
- Need for re-interventions

