Locations and etiologies of extracranial venous lesions in MS

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What is the background rationale that supports an association between MS and venous obstruction?



The Association of Chronic Cerebro-Spinal Venous Insufficiency (CCSVI) and Multiple Sclerosis Background and Hypothesis

• MS plaques venocentric

- Lesions extend counter-current to normal venous flow direction
- Distribution of lesions often peri-ventricular where higher vein density
- Peri-venous cuffs similar to appearance noted in chronic venous disease

BBB breakdown

- Vessel wall breakdown which leads to micro-bleeds
- Iron acts as an inflammatory agent (histo and MR SWI show increased iron content in plaques developing in pattern identical to venous counter-current
- Ischemic areas associated with shunting of blood volume and vessel atrophy

Extracranial venous obstruction

- Lesion site is non-specific (dural sinus, jugular, brachiocephalic, azygous veins alone or in combination)
- Lesion etiology is non-specific (congenital/hereditary, osseous impingement, arterial compression, post-inflammatory, arachnoid granulation, etc., alone or in combination)

Is there evidence that similar narrowing of other venous territories causes symptoms?





MAY-THURNER ANATOMY

LEFT COMMON ILIAC VEIN COMPRESSION BY RIGHT COMMON ILIAC ARTERY

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44 year old male 4 months post lumbar spine surgery with leg pain, swelling and prominent venous collaterals





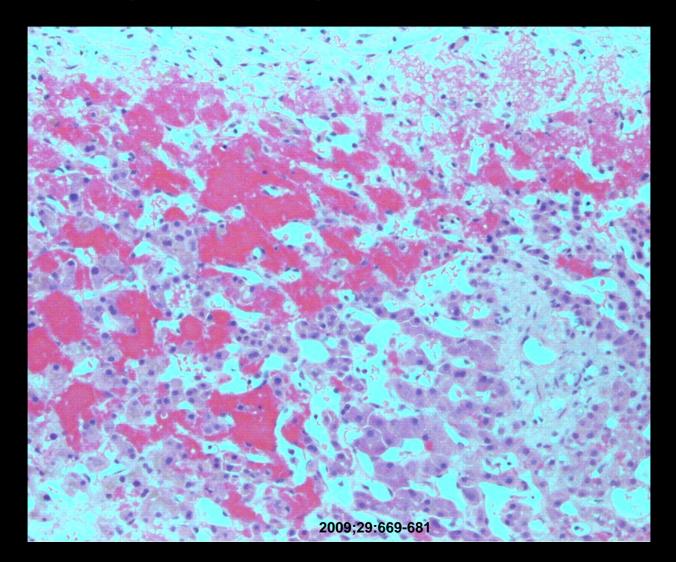
Pathent Radio horacic Surgery, Stanford University School of Medicine

Cavogram in a 24-year-old woman with Budd-Chiari syndrome and a history of oral contraceptive use depicts a narrowed intrahepatic IVC stenosis (*) and collateral vessels (arrows)



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Photomicrograph (original magnification, x40; hematoxylin-eosin stain) of a transjugular liver biopsy specimen from a 27-year-old woman with acute Budd-Chiari syndrome shows sinusoidal dilatation and clustering of red blood cells in the sinusoidal spaces around the hepatic vein, findings indicative of congestion in the perivenular zone





Why are stents used in certain situations; what is their role?



Based on prior experience with endovascular management of venous obstruction

- Balloon angioplasty of veins is usually ineffective
- Stents in veins counteract recoil
- Both PTA and stents may be associated with venous re-narrowing; this is especially common in dialysis patients



When POBA?

- Always first choice -- initial therapy
- Valve issues: stuck/stiff leaflet(s)
- Membranous lesions
- Annular ostial/junctional lesions
- Dural sinus lesions



Why Stents Instead of PTA Only?

- Residual stenosis
- Residual pressure gradient
- Residual collateral network flow
- Residual symptoms



Are any stents FDA-approved specifically for treatment of patients with venous obstruction?



Brief Reports

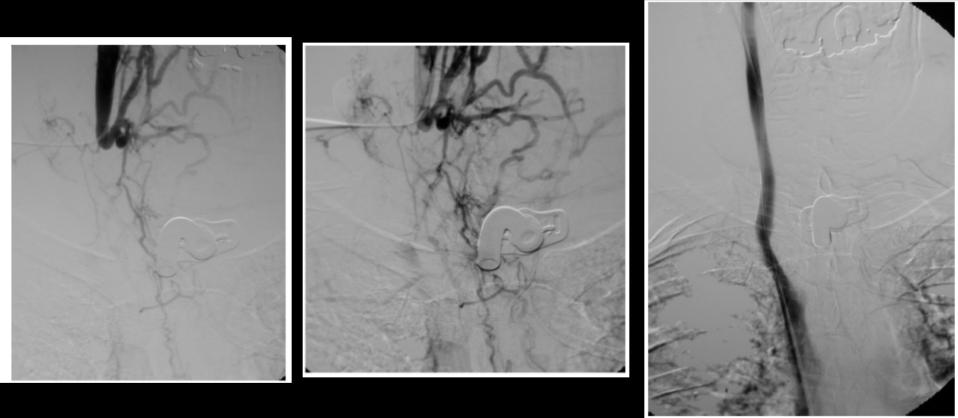
SVC Syndrome with a Patent SVC: Treatment of Internal Jugular Venous Occlusion after Surgical and Radiation Therapy of Esophageal Cancer

Justin C. Riutta, MD, Andrea L. Cheville, MD, and Scott O. Trerotola, MD

A patient is described in whom recanalization and stent placement in an occluded internal jugular vein was performed for the treatment of refractory facial edema initially thought to be lymphedema. The authors describe the combination of venous obstructions leading to this clinical presentation, which they term "SVC syndrome with a patent SVC."

J Vasc Interv Radiol 2005; 16:727-731

Abbreviations: IJV - internal jugular vein, SVC - superior vena cava





18-year-old obtunded s/p assault; normal CT of brain, carotids, but bilateral IJV thrombosis and ICP of 33 mmHg



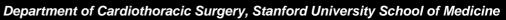


Neurosurgery 1997; 41:680-683

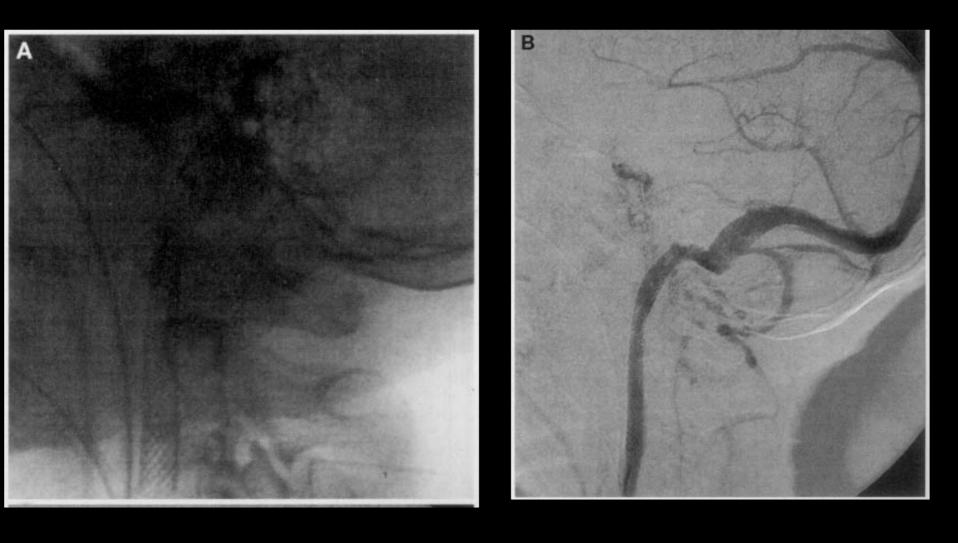
Left IJV occluded at C1 level With 8 mmHg pressure gradient; catheter-directed thrombolytic infusion ineffective







S/P stent placement with immediate physiological and clinical improvement; Coumadin administered for 8 weeks post procedure; stent widely patent on f/u venography





Neurosurgery 1997; 41:680-683

Where does the extra-cranial venous obstruction occur?



High Jugular Lesions





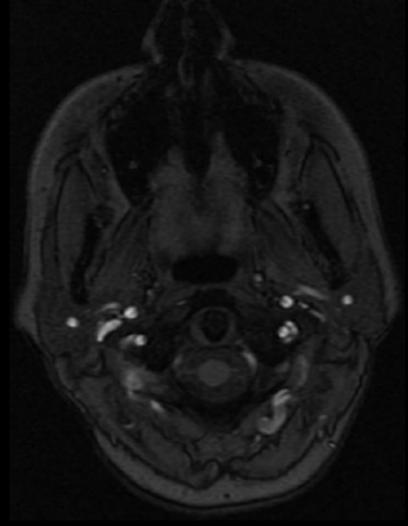


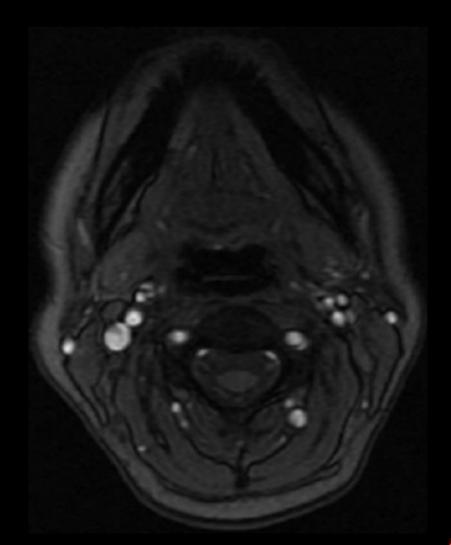


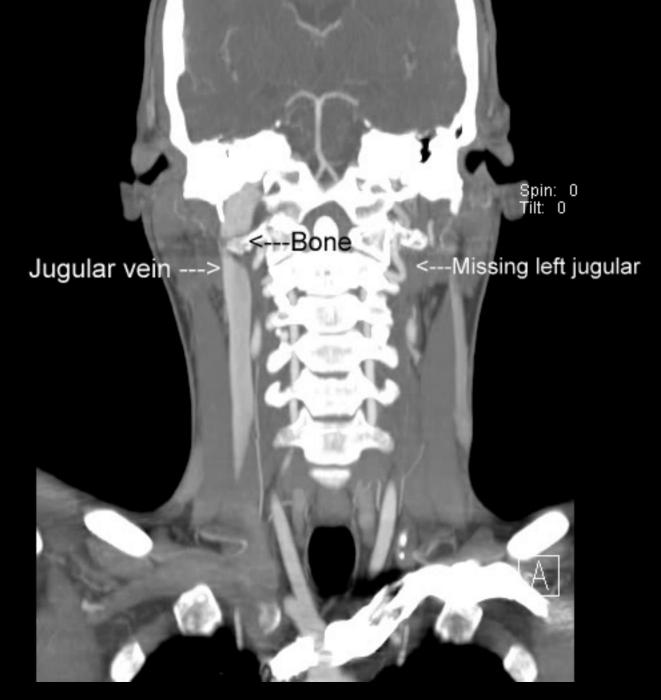




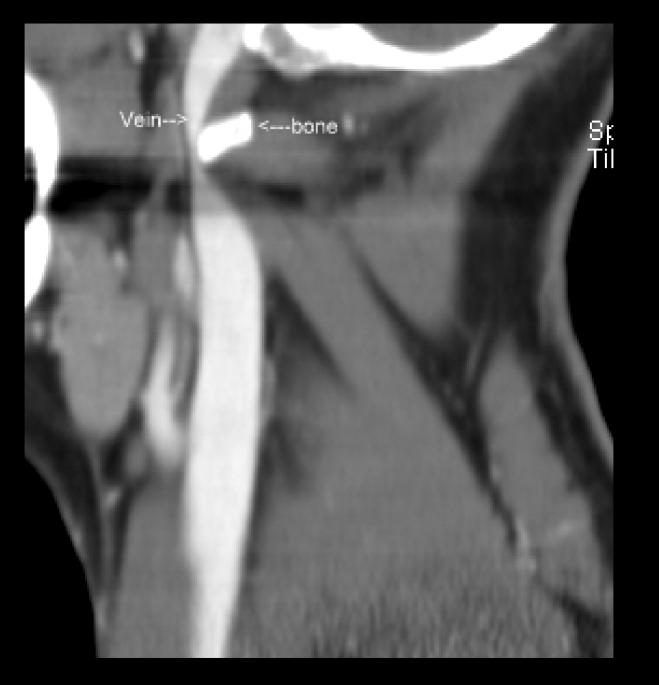
Asymmetric compression of nondominant left IJ by C1 lateral mass



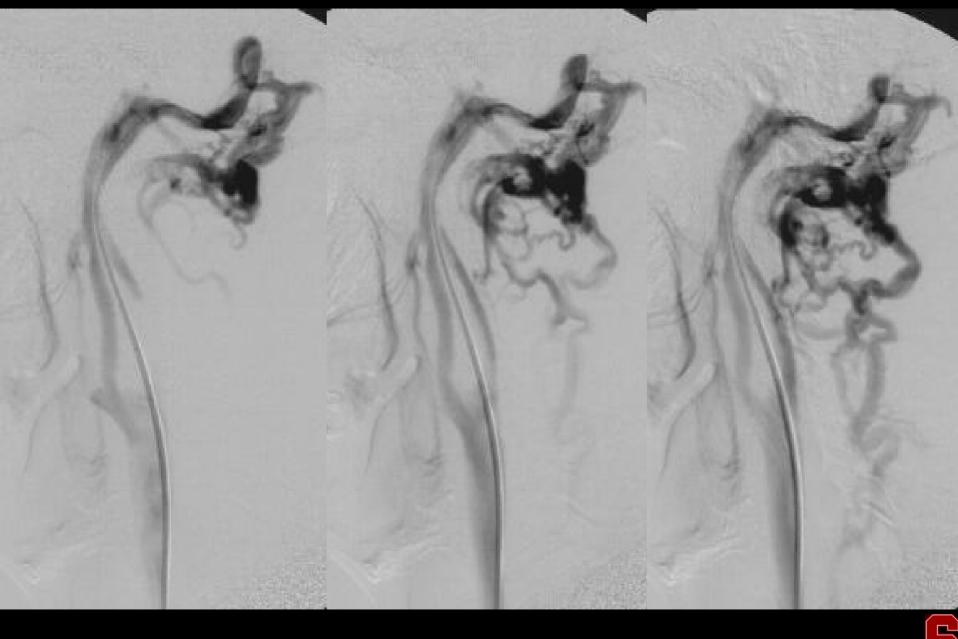




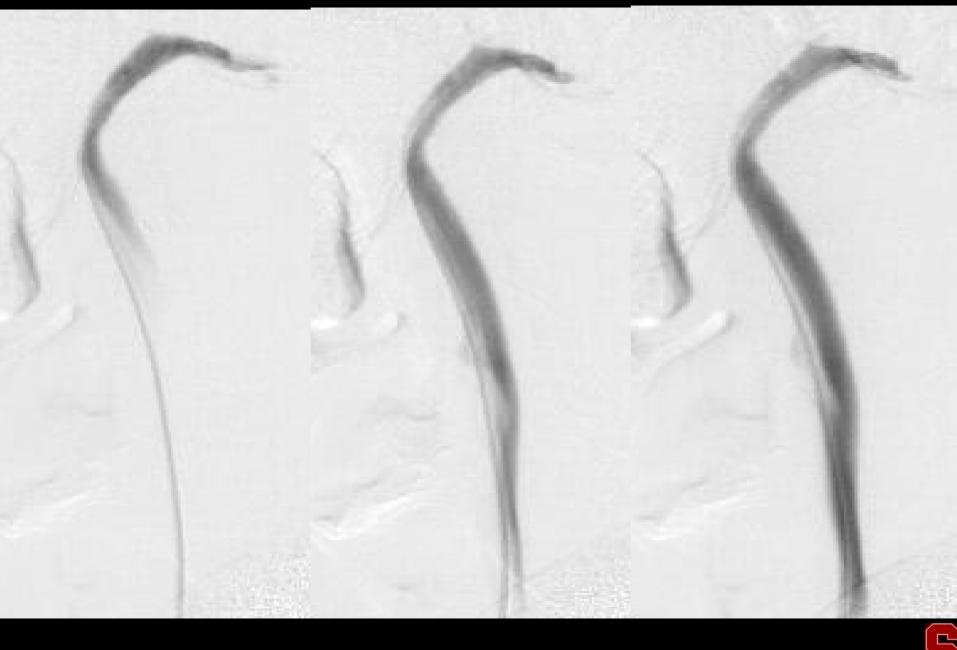
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Collaterals Associated with Mid RIJ Stenosis



RIJ Post Stent Placement



Mid Jugular Lesions



Axial MR with mid jugular compression by ectatic carotid

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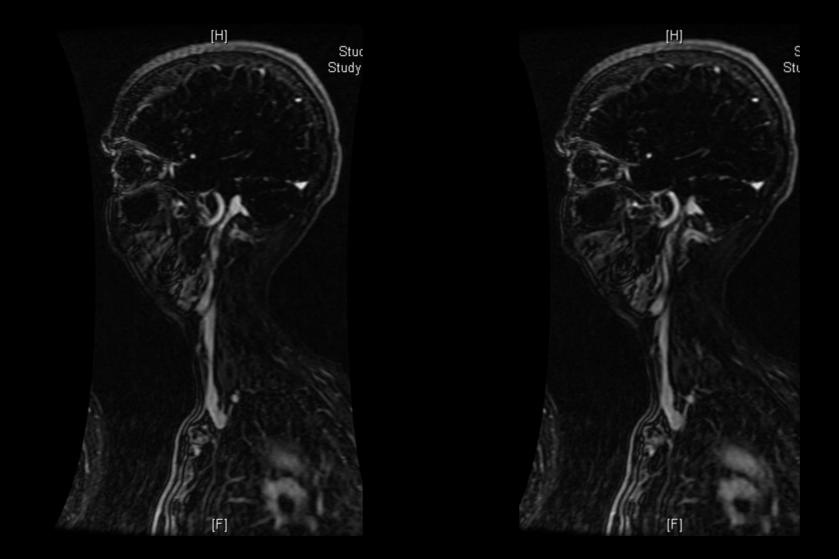
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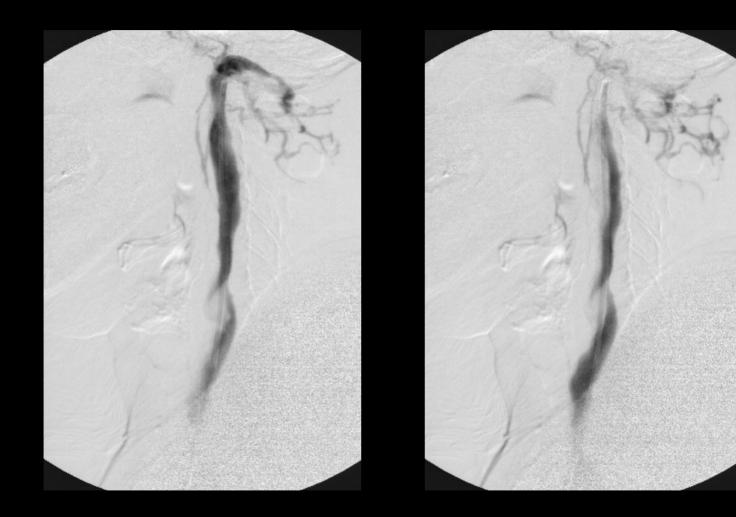
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Carotid impression upon mid-jugular





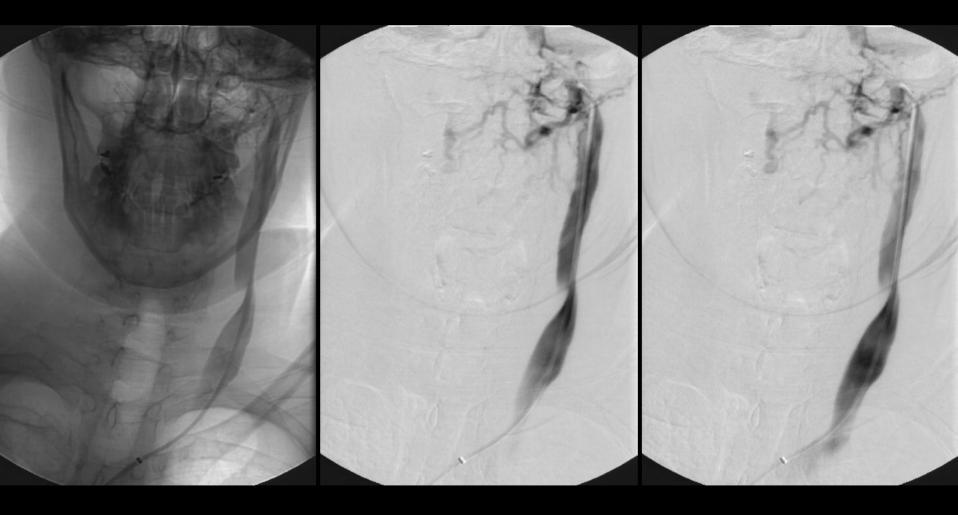
Lateral view of left mid-jugular narrowing





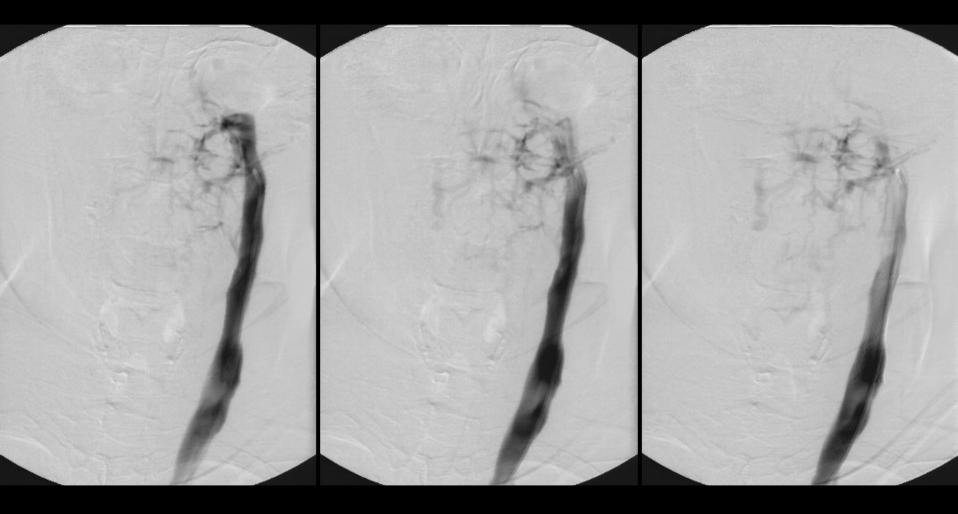


Mid-jugular carotid compression



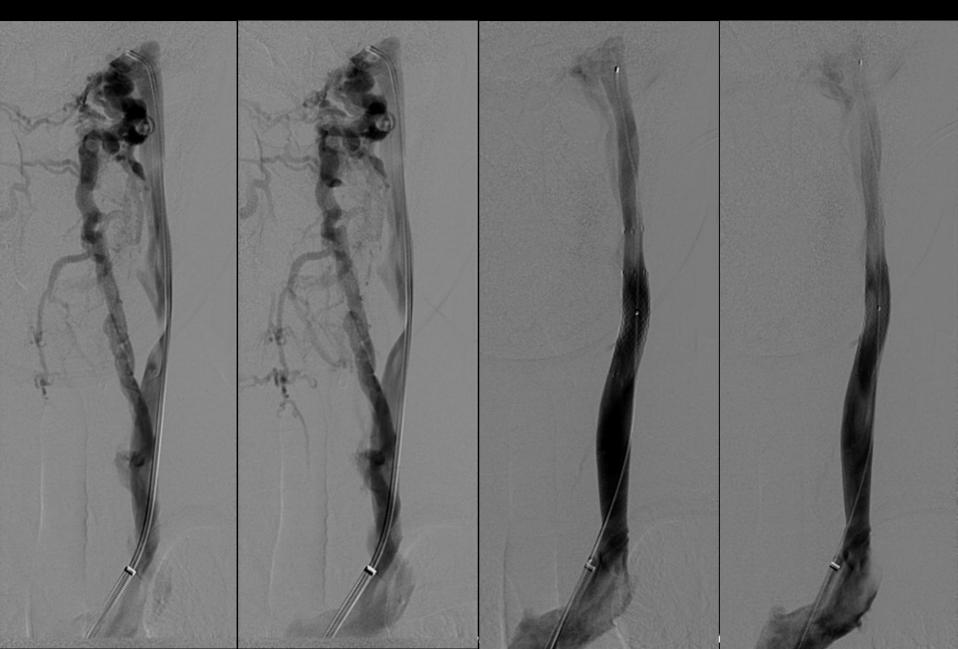


Post-stent placement in mid-jugular



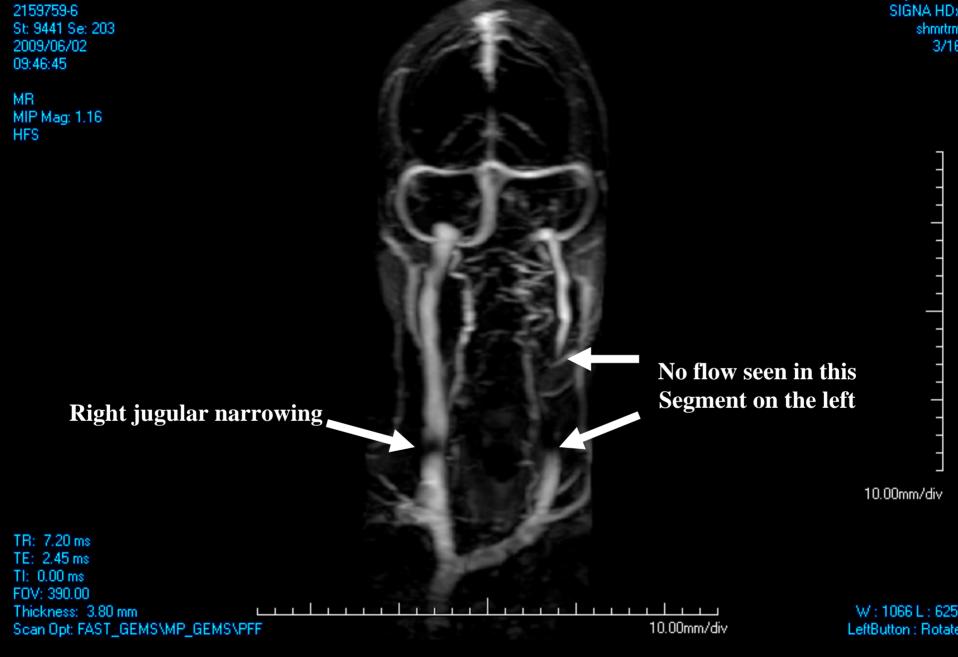


Venography Pre/Post stent



Low Jugular Lesions

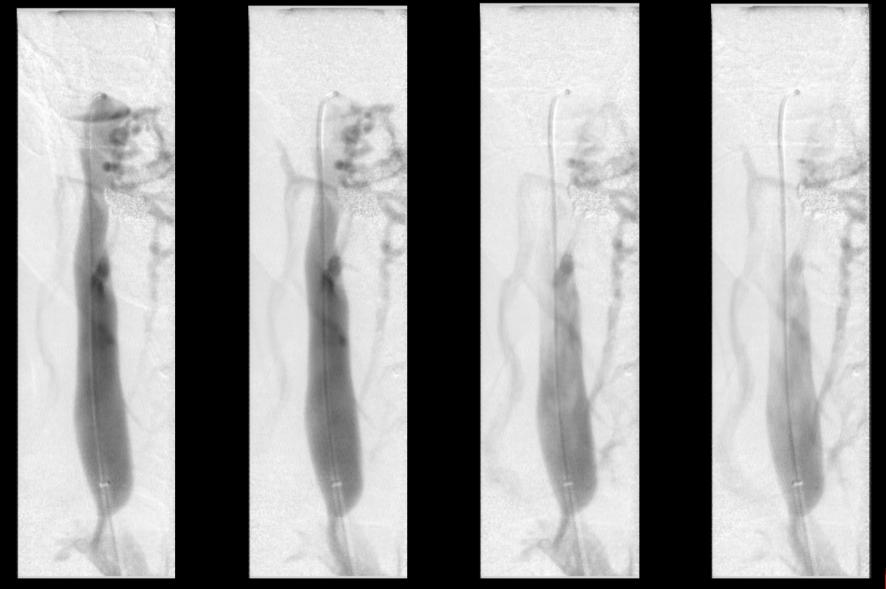




Jugular Valves

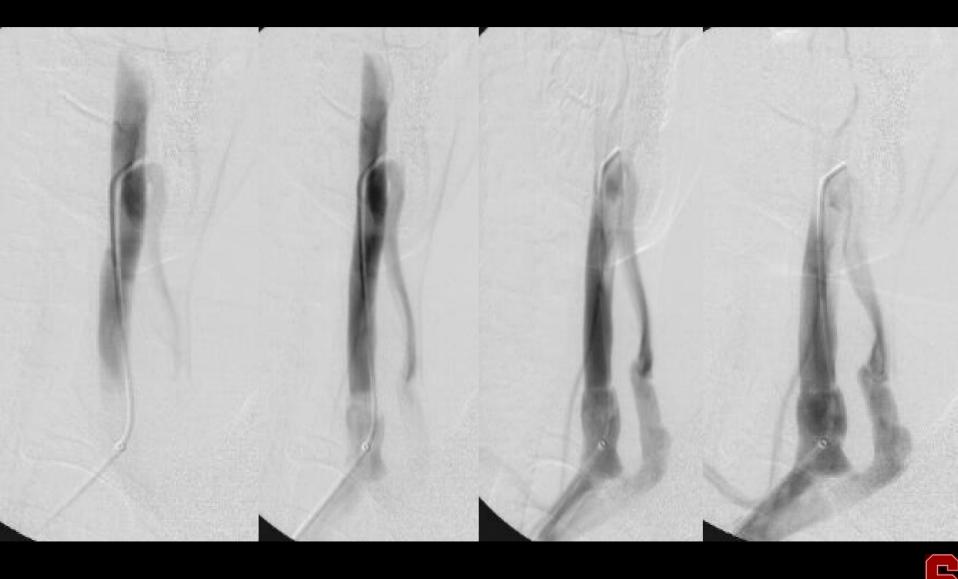


Narrowed valve orifice

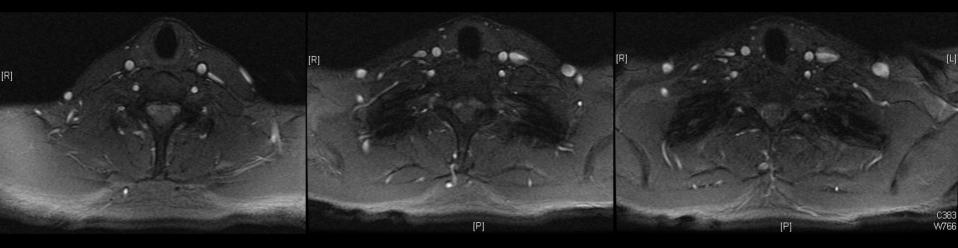


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LIJ Venogram with Large Collateral

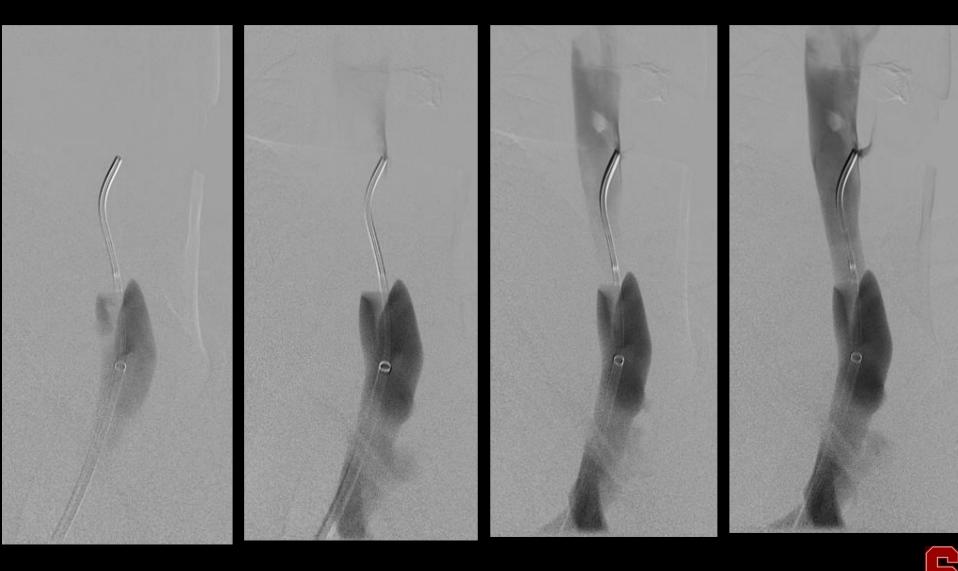


Prominent valve cusps more conspicuous on left





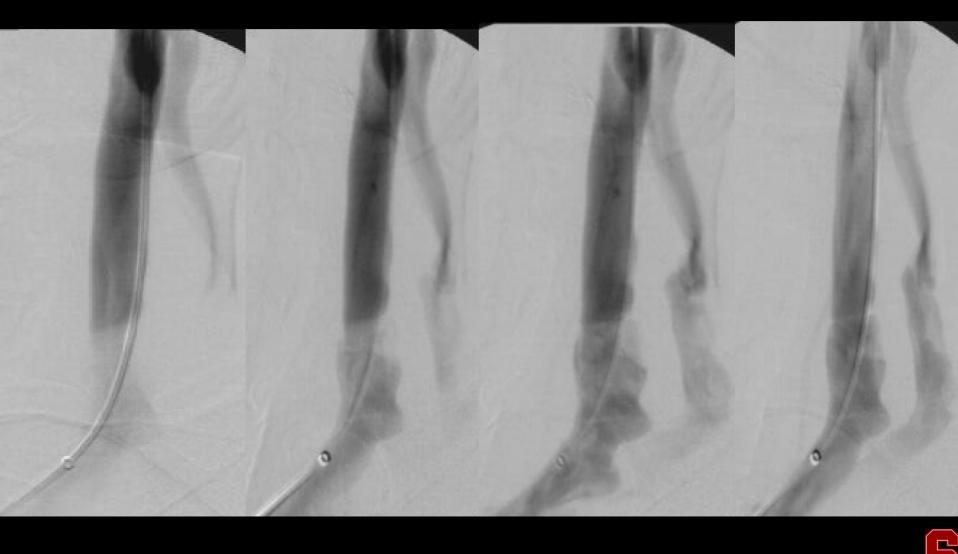
Too Competent...Stuck Leaflet?



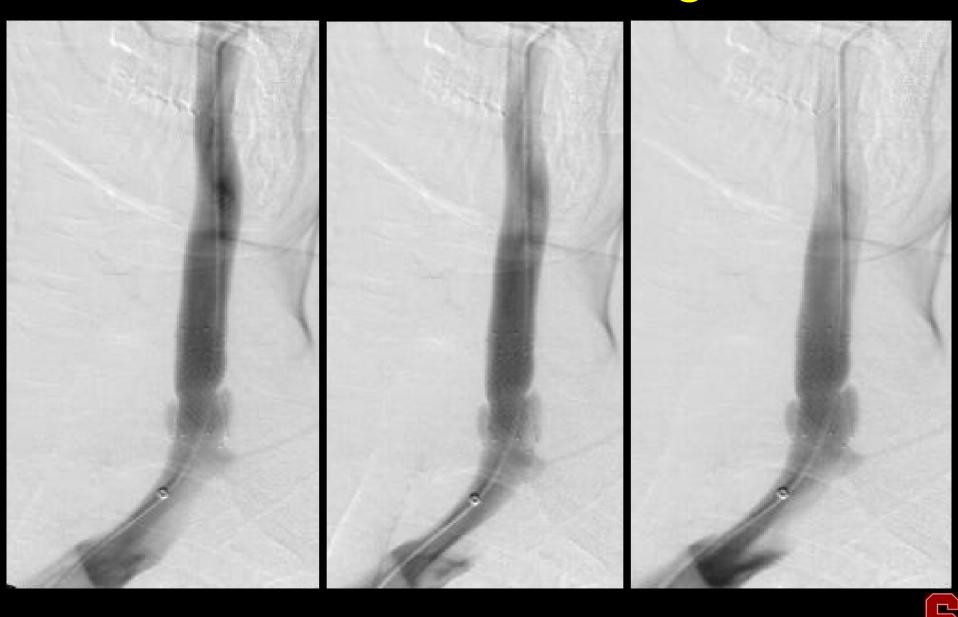
12 mm PTA of LIJ Valve



Post-PTA LIJ Venogram



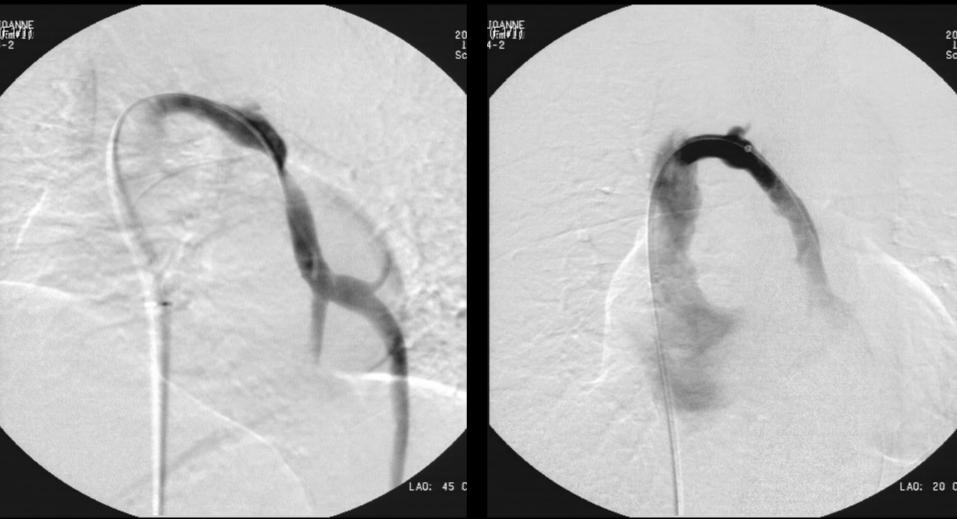
Post-Stent LIJ Venogram



Azygous



Pre/Post stenting of twisted descending azygous vein





What causes the venous narrowing?



Constellations of locations and etiologies of venous obstruction in patients with MS

- High Internal Jugular (C1-3 vertebral level)
 - Often at C1 associated with impingement from anterior aspect of broad-based transverse process
 - May be a dynamic obstruction related to position (vein stretched when supine with neck extended)
 - PTA ineffective; stent commonly improves, but outward force may cause pressure injury to CN XI (accessory nerve) which runs in jugular sheath

• Mid Internal Jugular (middle third between upper and lower segments, C4-6)

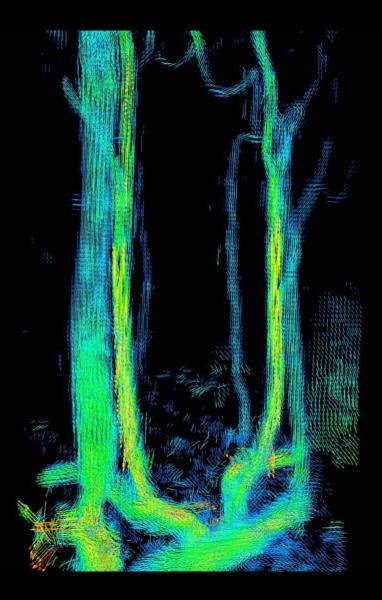
- Usually secondary to arterial compression from ectatic carotid located medial to vein (side by side) as opposed to anterior/posterior relationship
- PTA ineffective; stent placement relieves compression
- Low Internal Jugular (lower third, including valvular segment, C7-T2)
 - Pancake flattening subjacent to sternocleidomastoid muscles has a dynamic character of unknown etilology or physiological import
 - 2nd loci related to abnormal valve function (stuck or inverted leaflets, stiff/stenosed valve, etc.)
 - These sites may exist in combination or alone. Valve abnormalities frequently respond tp PTA, with stent placement reserved for persistent obstruction

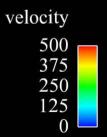
How do we know if the venous lesion is significant?



Collaterals Pressure Gradients Flow Patterns

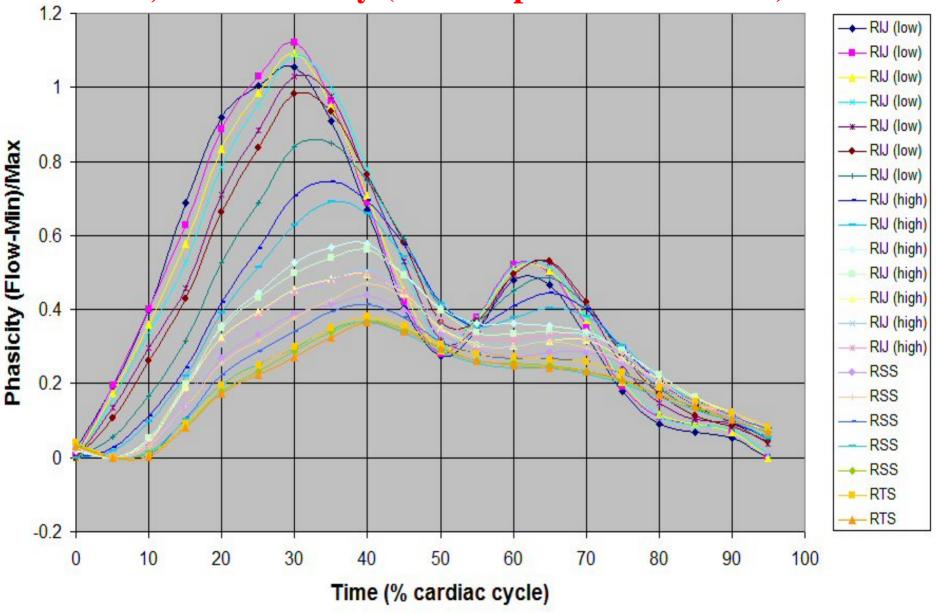






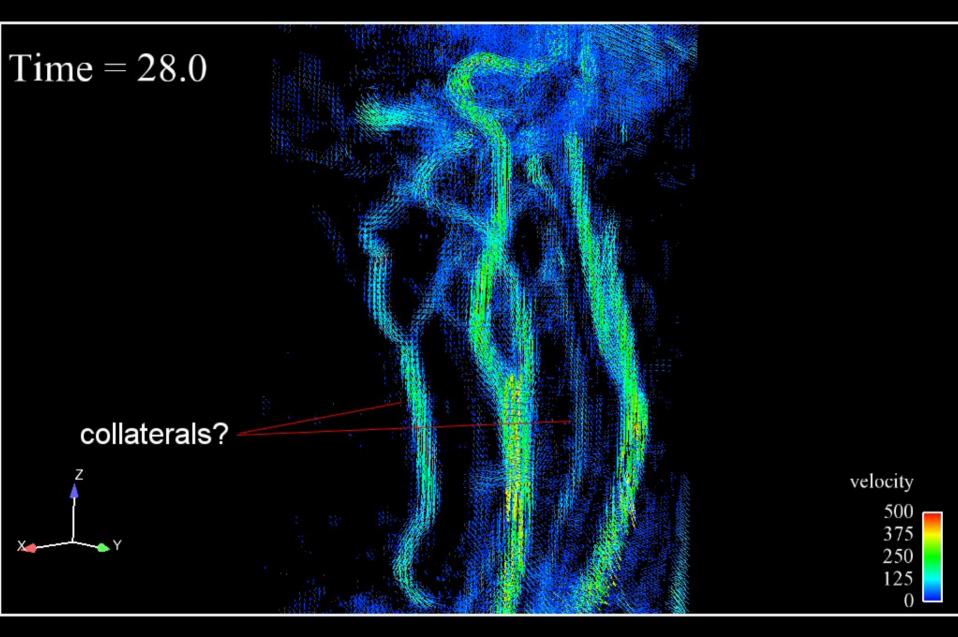


RIJ, RSS Phasicity (cardiac phase re-centered)



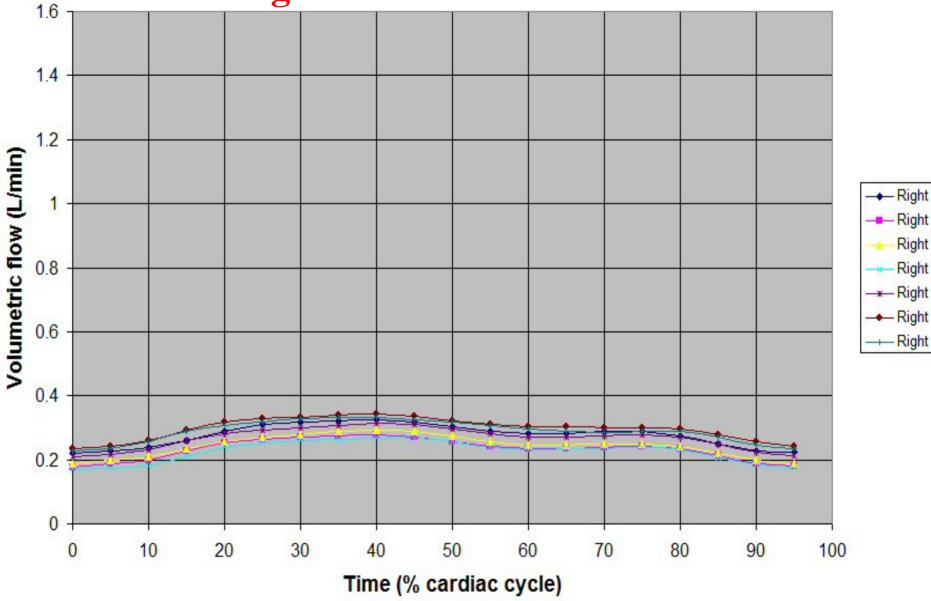
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Right-sided collateral flow



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What are the anticipated short and long-term outcomes of endovascular management of extra-cranial venous insufficiency?



Venous Obstruction (CCSVI) and MS

- Initial Observations Recorded after Endovascular Treatment of Venous Stenotic Lesions
 - Global symptoms attributable to MS, but not referrable to a specific neuro-anatomic loci (ie., fatigue, headache, heat sensitivity, "brain fog", urinary urgency,etc.), show short-term improvement and in some cases (low EDSS) completely resolve. This suggests that these particular "MS" symptoms may be more accurately categorized as related to venous obstruction.
 - Early-term follow-up of functional mobility (high EDSS) is not conspicuously changed from preprocedure.



What are the procedural complications of endovascular intervention (PTA, stent placement, etc.) for venous obstruction?



Complications and Untoward Effects

- Bleeding: groin hematoma, excessive bruising, intra-cranial hemorrhage/death
- Stent migration
- Headache, nausea, ear plugging, neck/throat discomfort
- Trapezius pain, shoulder weakness and decreased ROM due to CN XI (accessory) pressure "injury"
- Re-narrowing or obstruction of treated segment
- No stroke, MI, worsening of existing symptoms



The Association of Chronic Cerebro-Spinal Venous Insufficiency (CCSVI) and Multiple Sclerosis Summary Analysis

- Extracranial venous obstruction
 - Lesion site is non-specific (dural sinus, jugular, brachiocephalic, azygous veins alone or in combination)
 - Lesion etiology is non-specific (congenital/hereditary, osseous impingement, arterial compression, post-inflammatory, arachnoid granulation, etc., alone or in combination)

