How I Approach The Swollen Arm

Renal Access Session, VS, 28th November 2018

Mr Paul Gibbs Renal Transplant and Access Surgeon Wessex Kidney Centre QAH, Portsmouth





I have no financial disclosures relevant to this talk.

What I will talk about

- Aetiology
- Investigations
- Simple balloon fistuloplasty
- Surgical Options
- Stents Grafts
- DCBs

What I won't talk about

- Central venous pathology
- HeRO Graft
- Surfacer

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Aetiology

- CVC usage
- Graft-venous anastomosis
- Swing points
 - Cephalic arch
 - BBFTs



- Previous damage to the vein (lines, phlebotomy....)
- Flow effects (valves, aneurysms, myointimal hyperplasia)

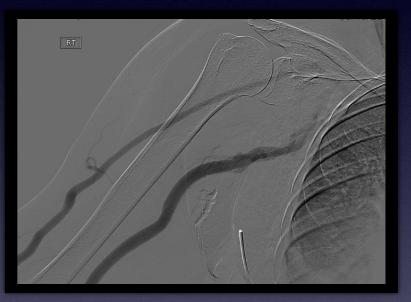
What I will talk about

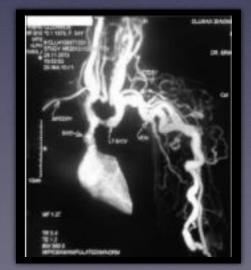
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Investigations

- Duplex USS
- Fistulogram
- CT Venography
- MR Venography

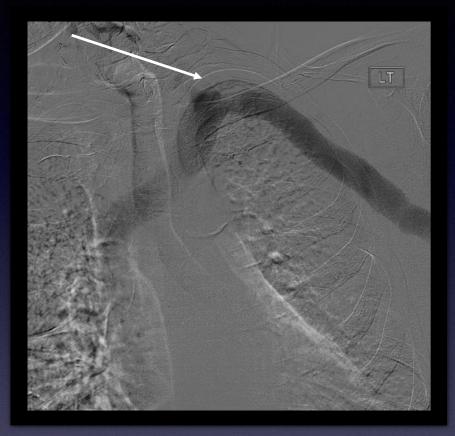


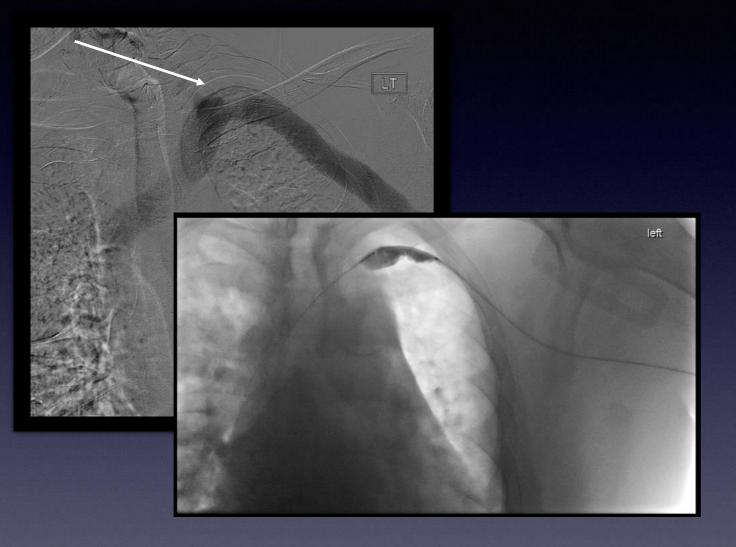


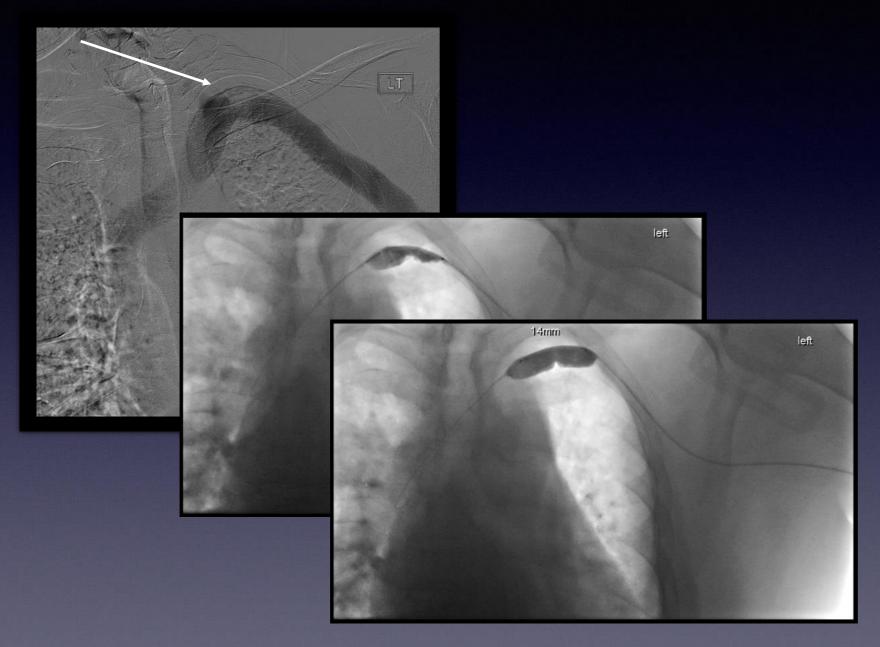


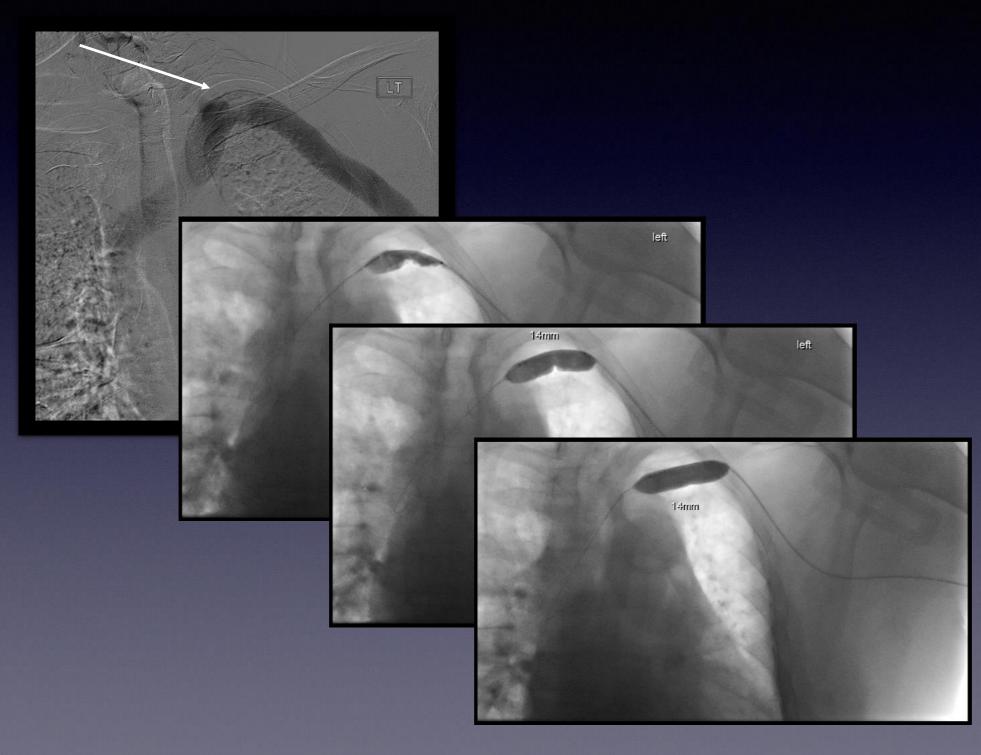
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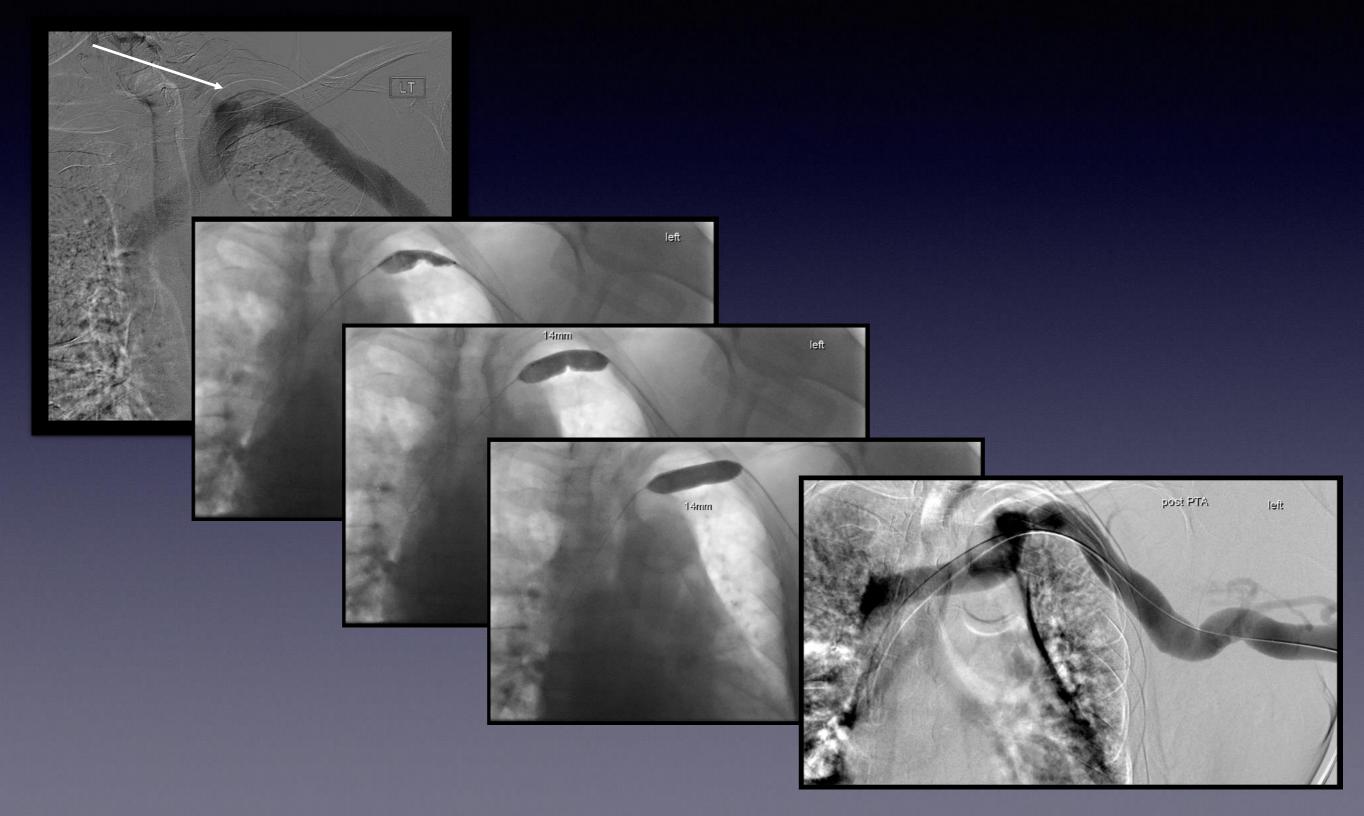
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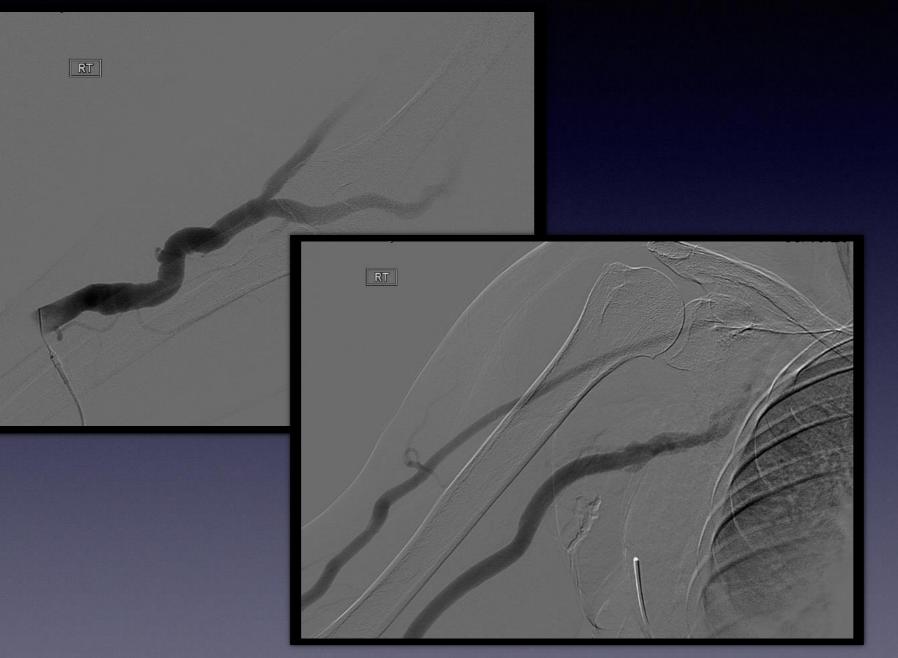
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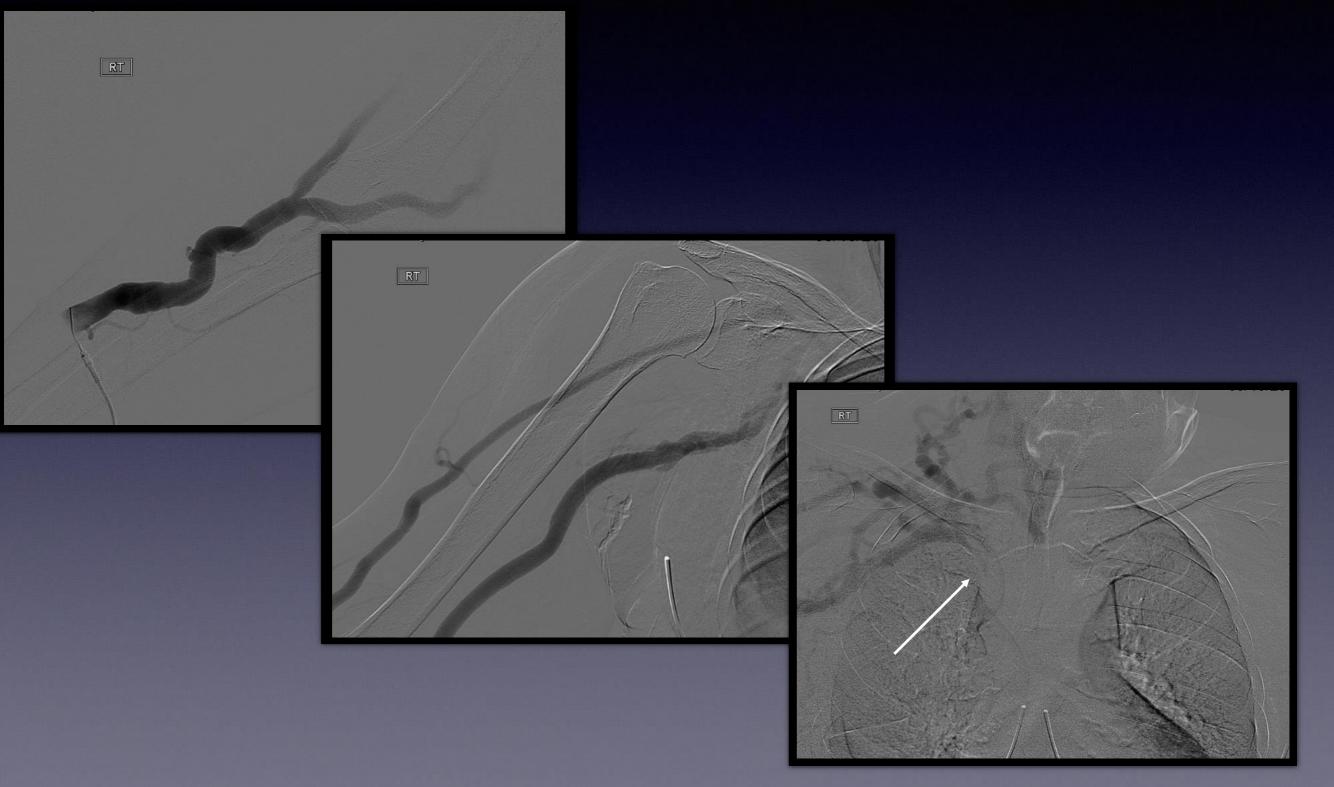
Ligation



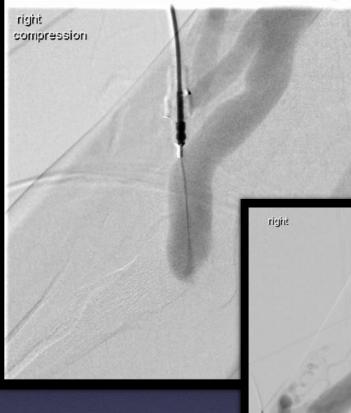
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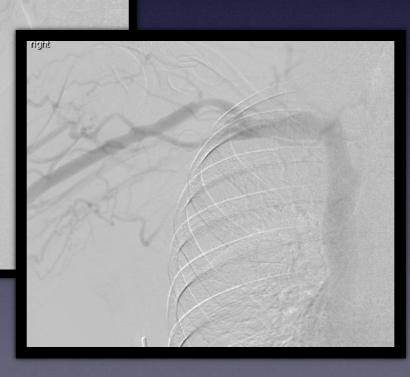


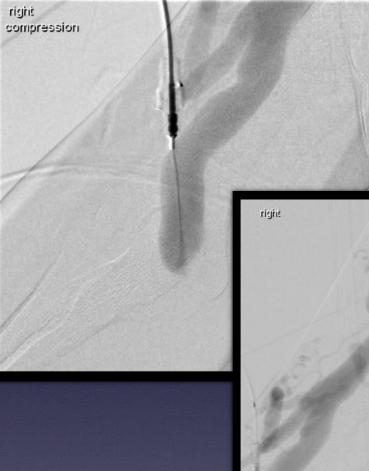


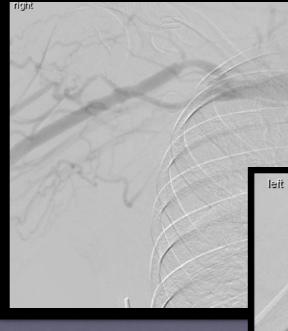


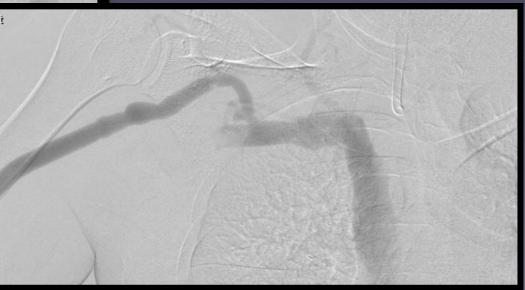








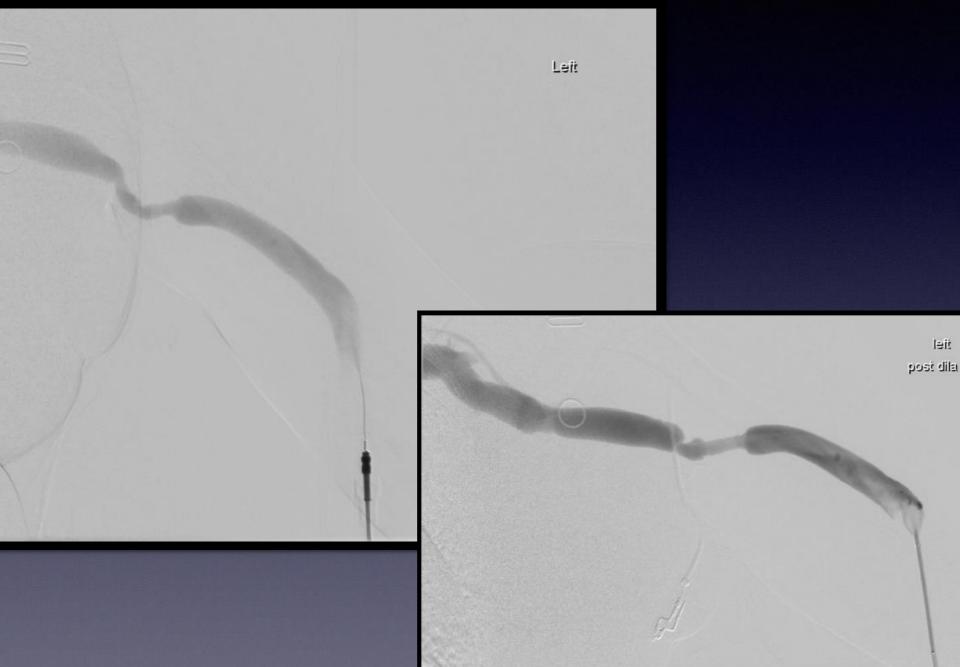




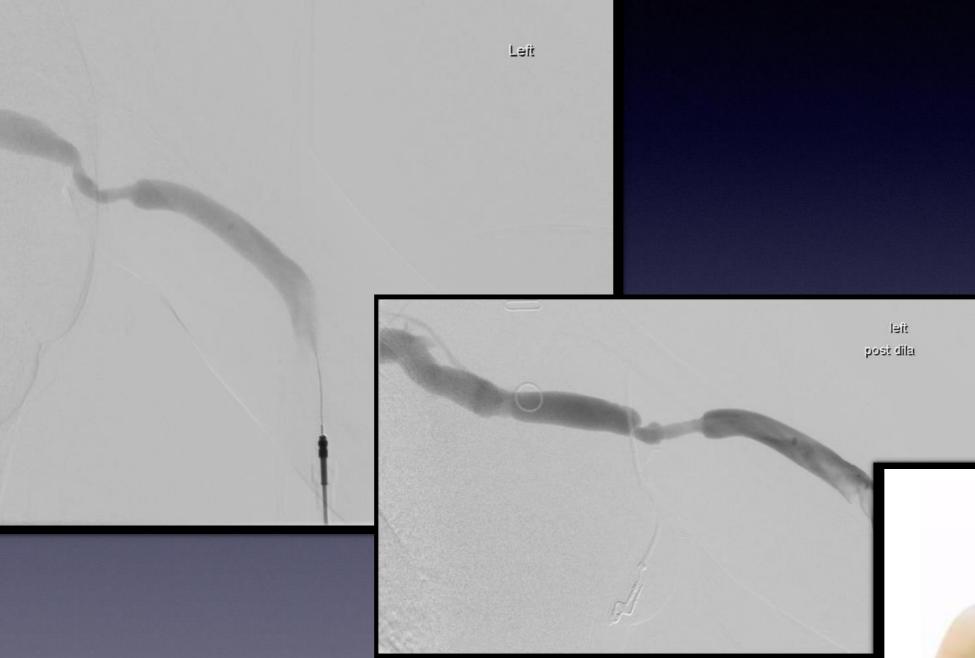
Surgical Patch Plasty

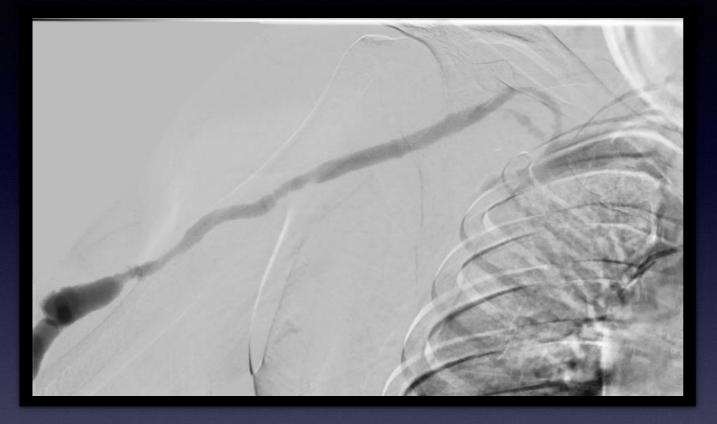


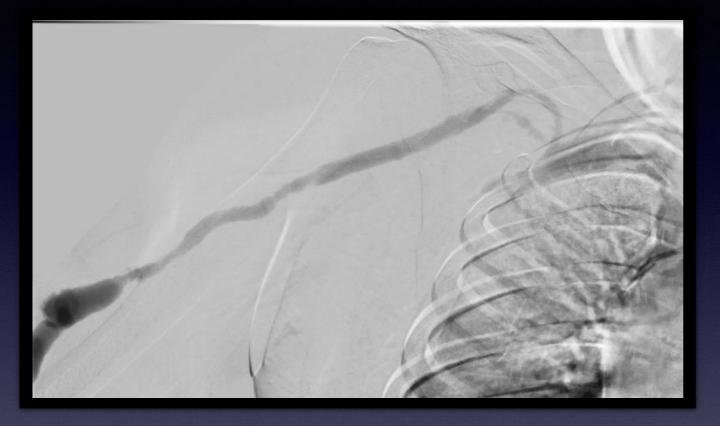
Surgical Patch Plasty



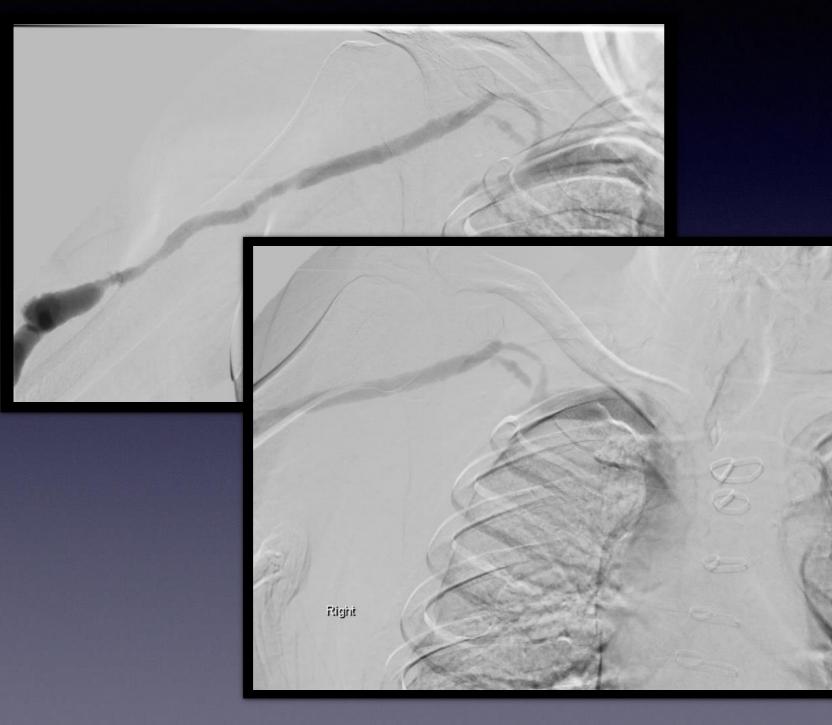
Surgical Patch Plasty



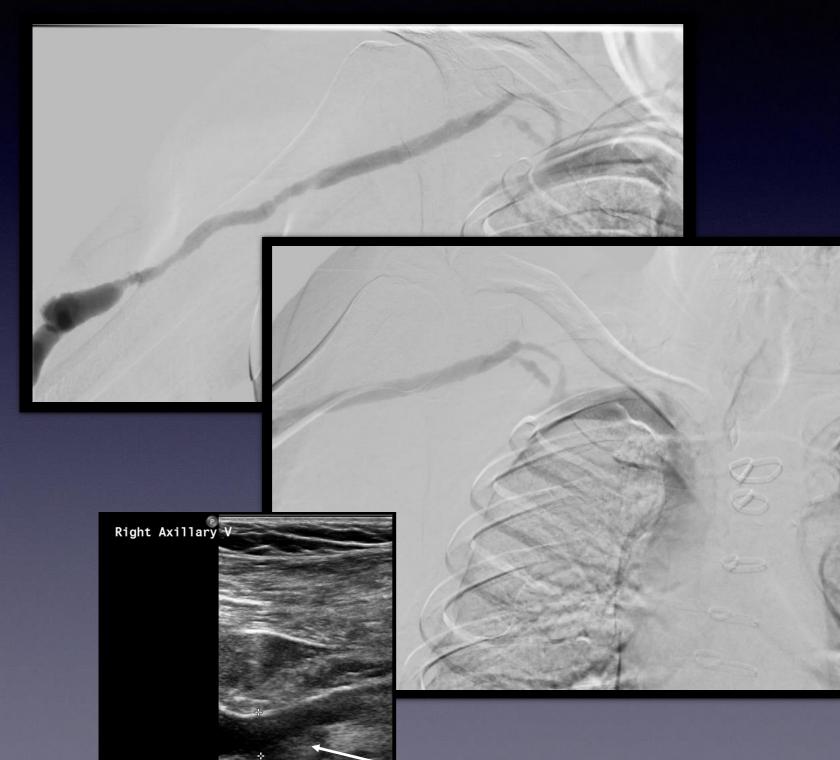




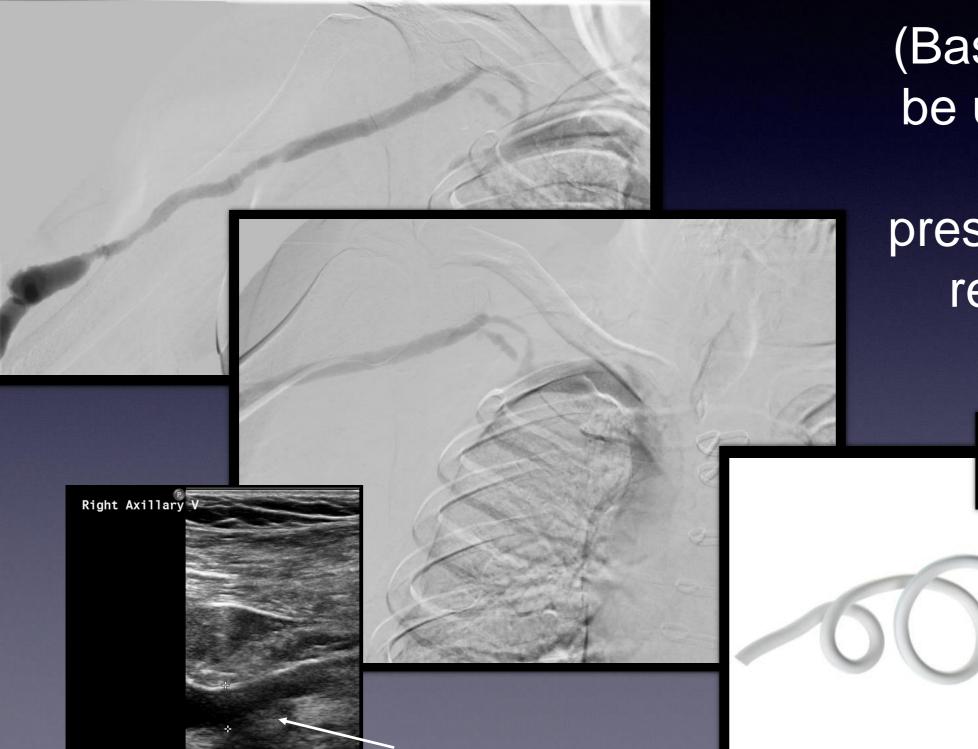
(Basilic vein can be used as well and may preserve venous real estate)



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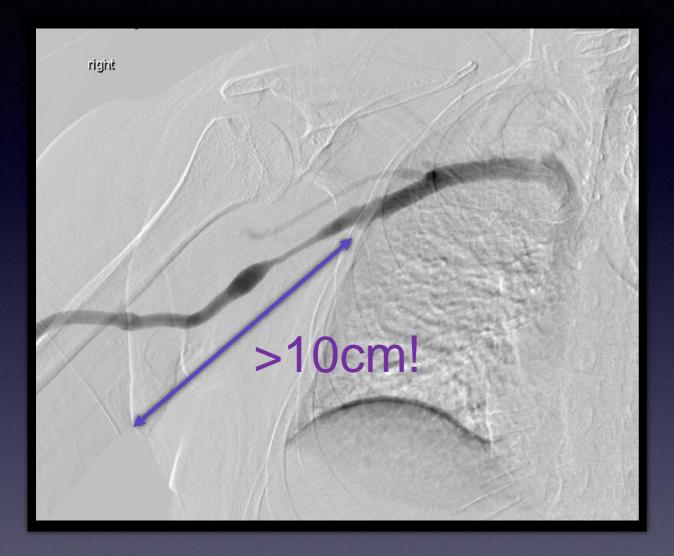


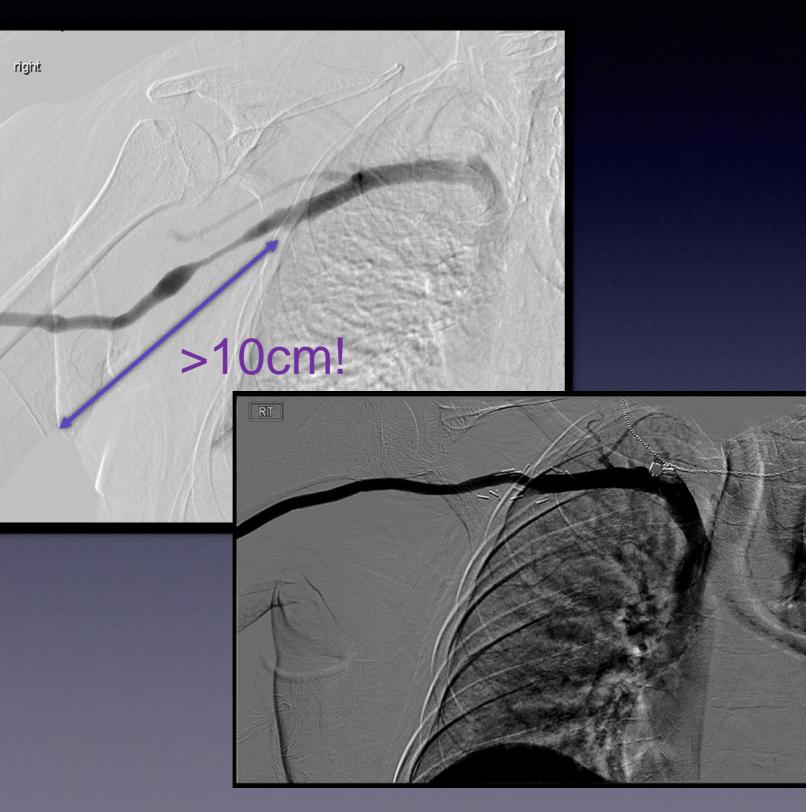
(Basilic vein can be used as well and may preserve venous real estate)

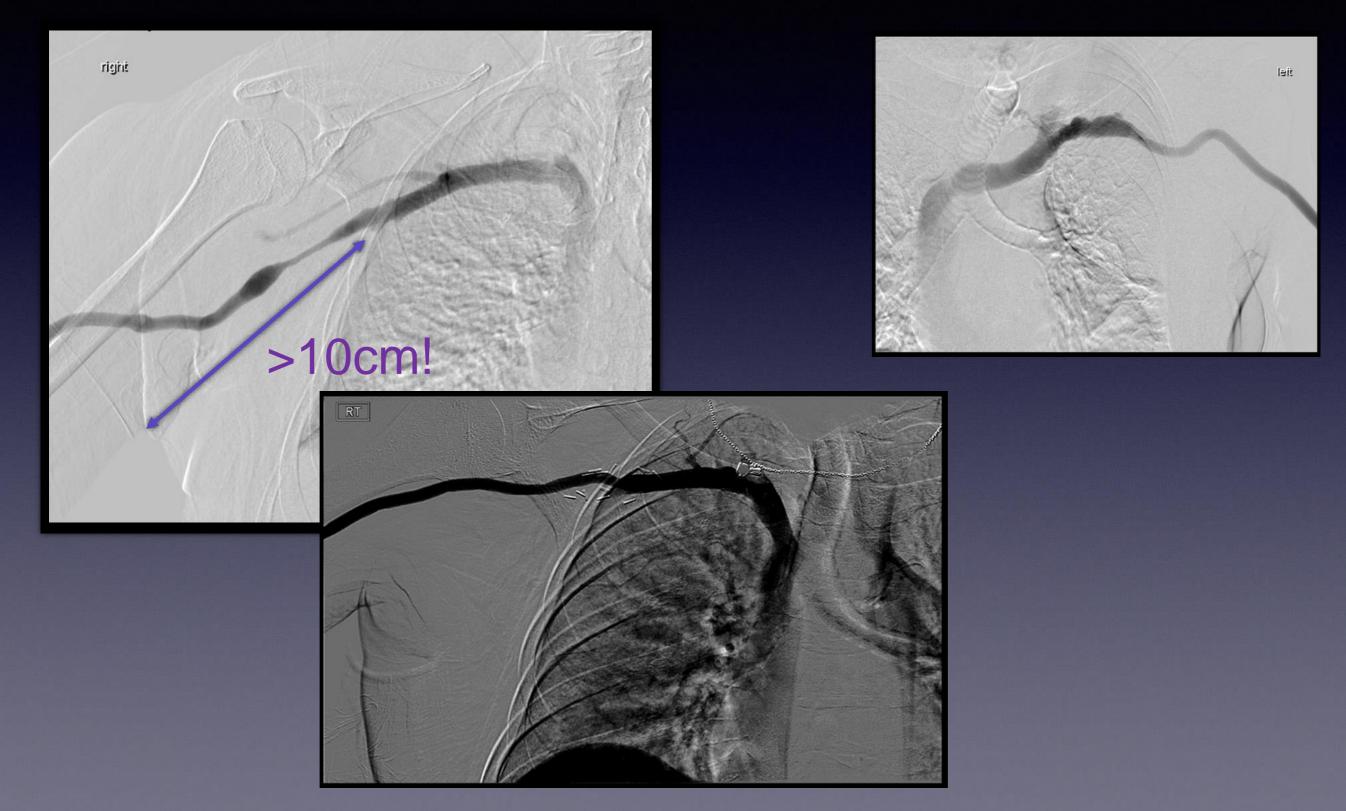


0.957 cm

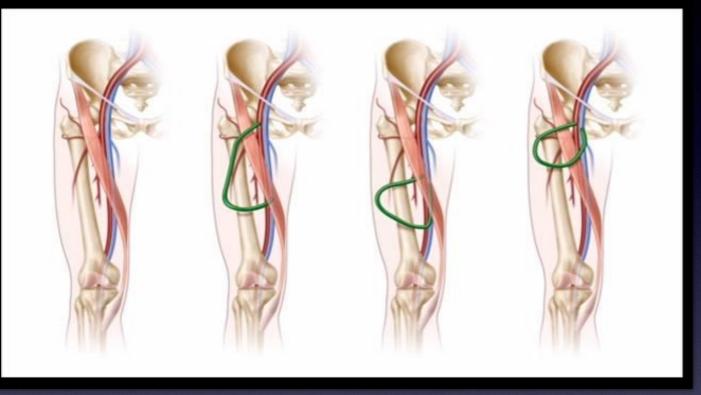








PTFE Leg Loops

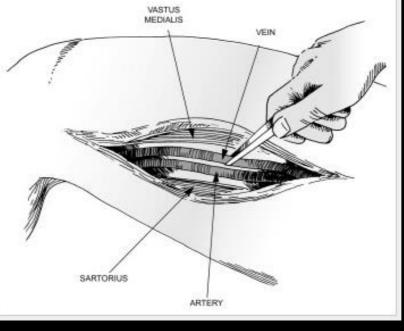


PTFE Leg Loops







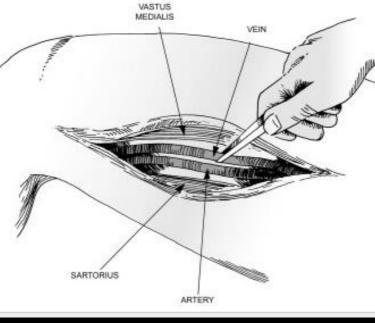


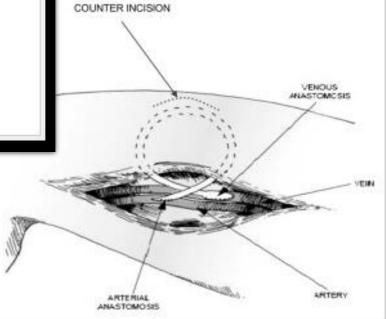
PTFE Leg Loops











PTFE Leg Loops

The Journal of Vascular Access

Good long-term patency: 10-year follow-up using the mid-thigh adductor loop arteriovenous graft

Samantha J McEwan, Hannah Maple, Paul J Gibbs

Conclusion:

Autogenous venous access remains the perceived gold standard for patients requiring dialysis for end stage renal failure, despite some published data reporting poor outcomes. We have shown that adductor loop arteriovenous grafts can be a reliable, safe and long-term alternative in those whom fistula formation is not possible and may have a role earlier in the patient journey than previously thought, as a result of good patency and lower complications.



Lower Limb AVFs

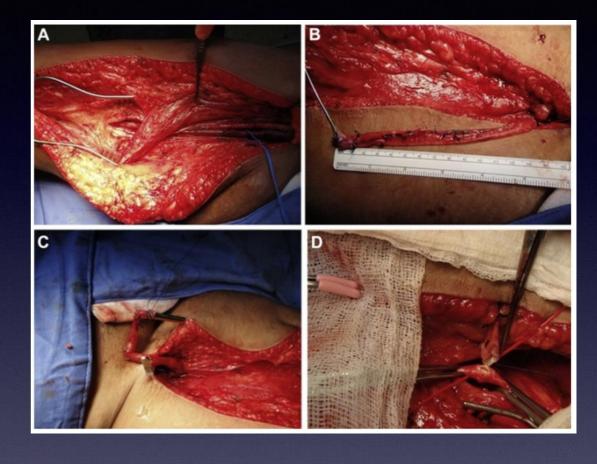
Annals of Vascular Surgery Available online 14 August 2018

The Transposed Femoral Vein Fistula: The Native Choice in Desperate Vascular Access

Rebecca Lefroy ¹ $\stackrel{\otimes}{\sim}$ ¹, Nikesh Dattani ¹, Mariane Reyes ², Sriram Rajagopalan ¹, Jack Fairhead ¹, Anthony Jaipersad ¹, Lorraine Corfield ¹, John Asquith ³, Michael W. Greenway ⁴, Arun Pherwani ¹

Conclusions

Our patient group showed good fistula patency at 1 year and did not experience any incidence of ischemic steal syndrome. We believe this to be due to careful preoperative patient assessment and meticulous surgical technique. Our experience suggests that such procedures should be performed by surgeons with vascular expertise wherever possible to reduce the incidence of complications.





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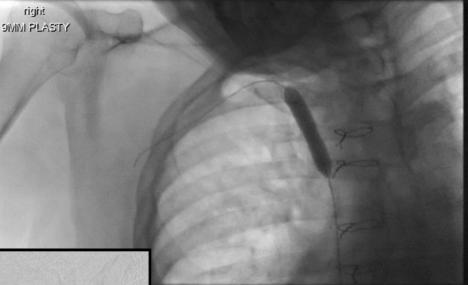








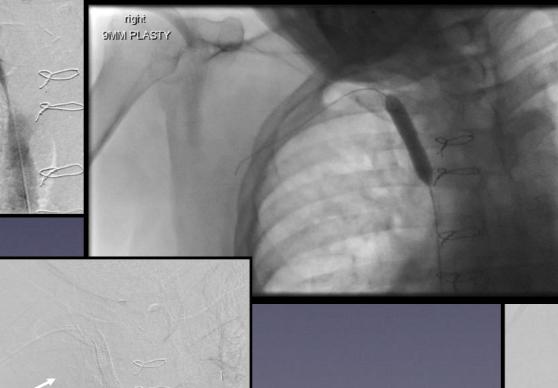






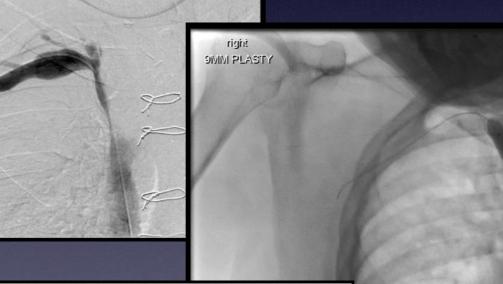


right POST PLASTY









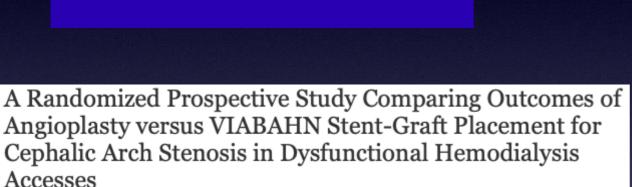






PROVEN TO INCREASE TIME TO NEXT INTERVENTION

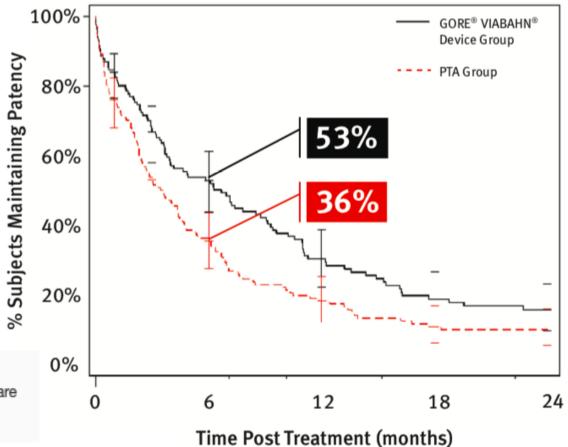
Kaplan-Meier Analysis of the Target Lesion Primary Patency



Dheeraj K. Rajan, MD, FRCPC 🖾 🖂, Abigail Falk, MD

Purpose

To determine if postintervention cephalic arch stenosis (CAS) primary patency and access circuit patency are superior with the VIABAHN stent graft compared with angioplasty at 3, 6, and 12 months.



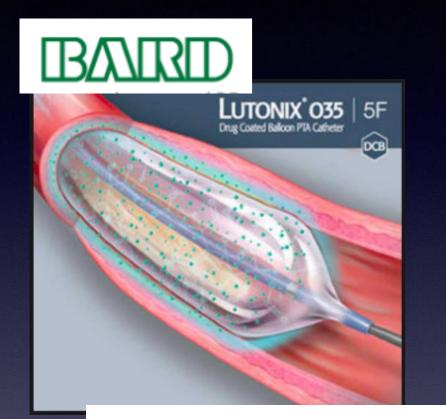
Conclusions

Treatment of CAS with the VIABAHN stent graft appears to provide statistically superior primary patency rates compared with balloon angioplasty.

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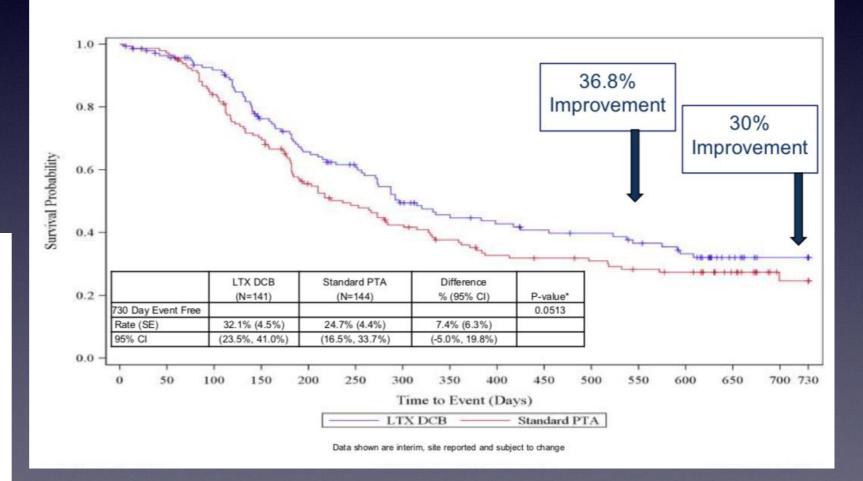
Drug Coated Balloons



Features

- First and only drug coated balloon approved for use in dysfunctional/stenosed dialysis fistulae
- Shown to enable longer AV fistula function due to increased time to first reintervention compared to standard angioplasty
- 71.4% Primary Patency in the LUTONIX [®] AV Clinical Study at 6 months
- 31.3% fewer reinterventions than PTA at 6 months in the LUTONIX [®] AV Clinical Study
- · Demonstrated a safety profile that is as safe as PTA

Lutonix AV Clinical Trial



Thank You



