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An Immediate Use Dialysis Graft Designed to Prevent Needle-Related Complications

Disclaimer: The InnAVasc Vascular Access Graft has not received regulatory clearance & is therefore not currently available for commercial use

Disclosure

Shawn M. Gage, PA-C

I have the following conflicts of interest to report:

- ✓ Co-inventor of InnAVasc technology
- ✓ Ownership in InnAVasc
- ✓ Shareholder in InnAVasc
- ✓ Employed by InnAVasc
- ✓ Receipt of grants/research support from NIH and NSF

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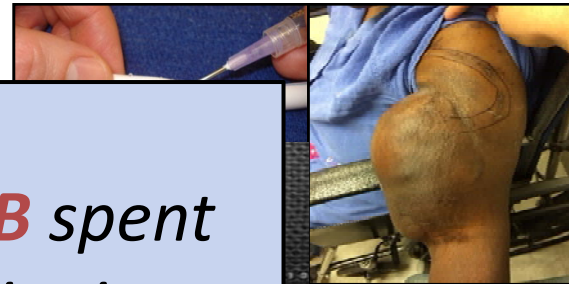
Current dialysis grafts have many pitfalls

Not Reliable

Lack of Durability



Preventable Complications



*Contribute to the **\$2.9B** spent annually on malfunctioning vascular access*

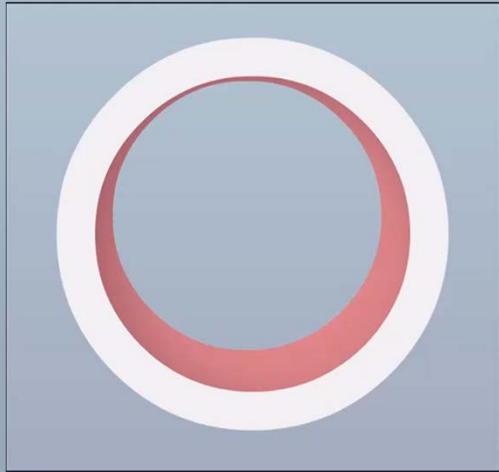
Challenging



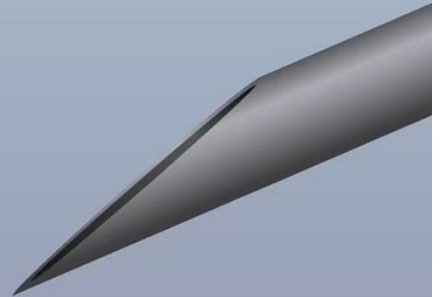
Intervention Required

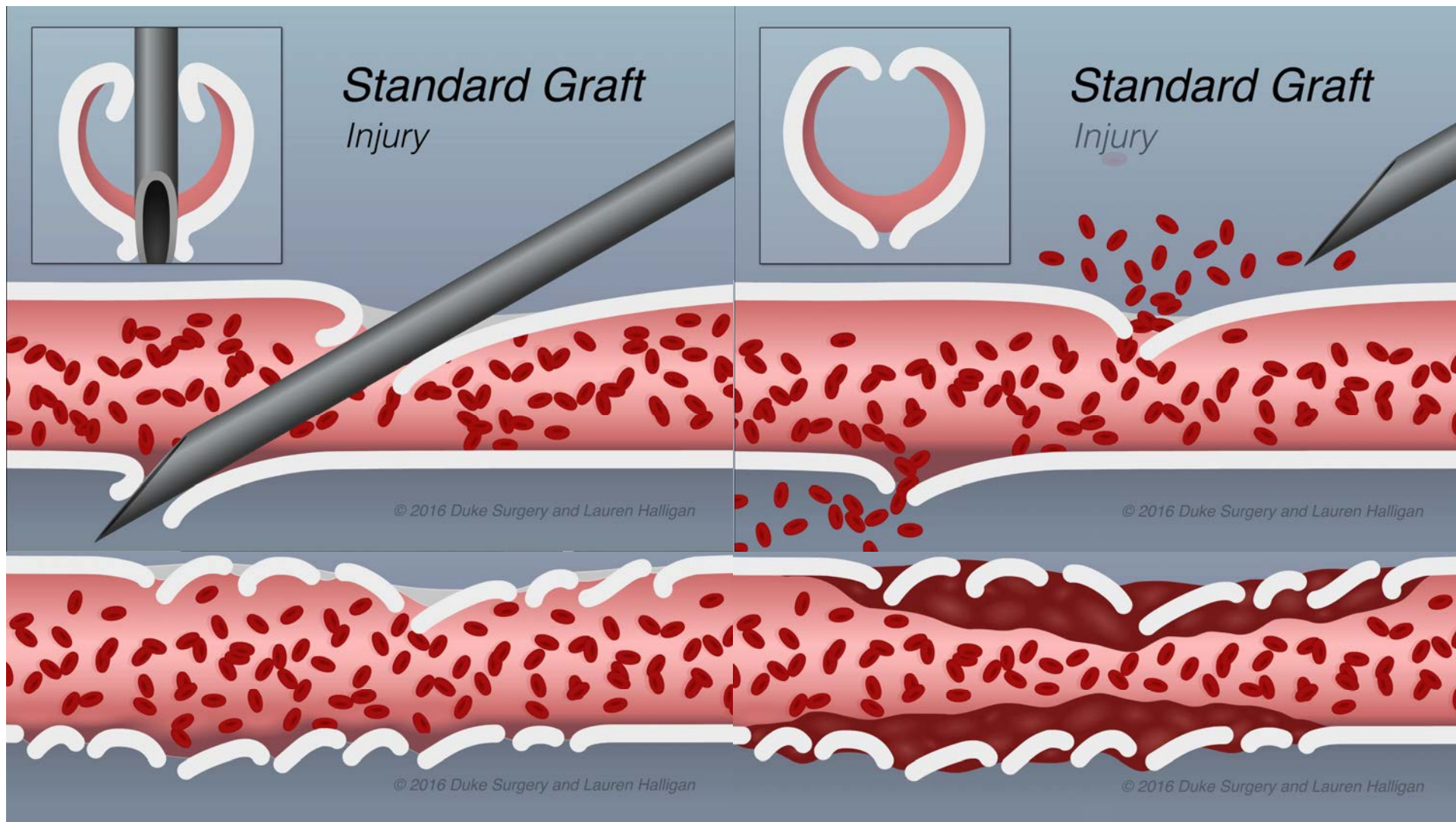


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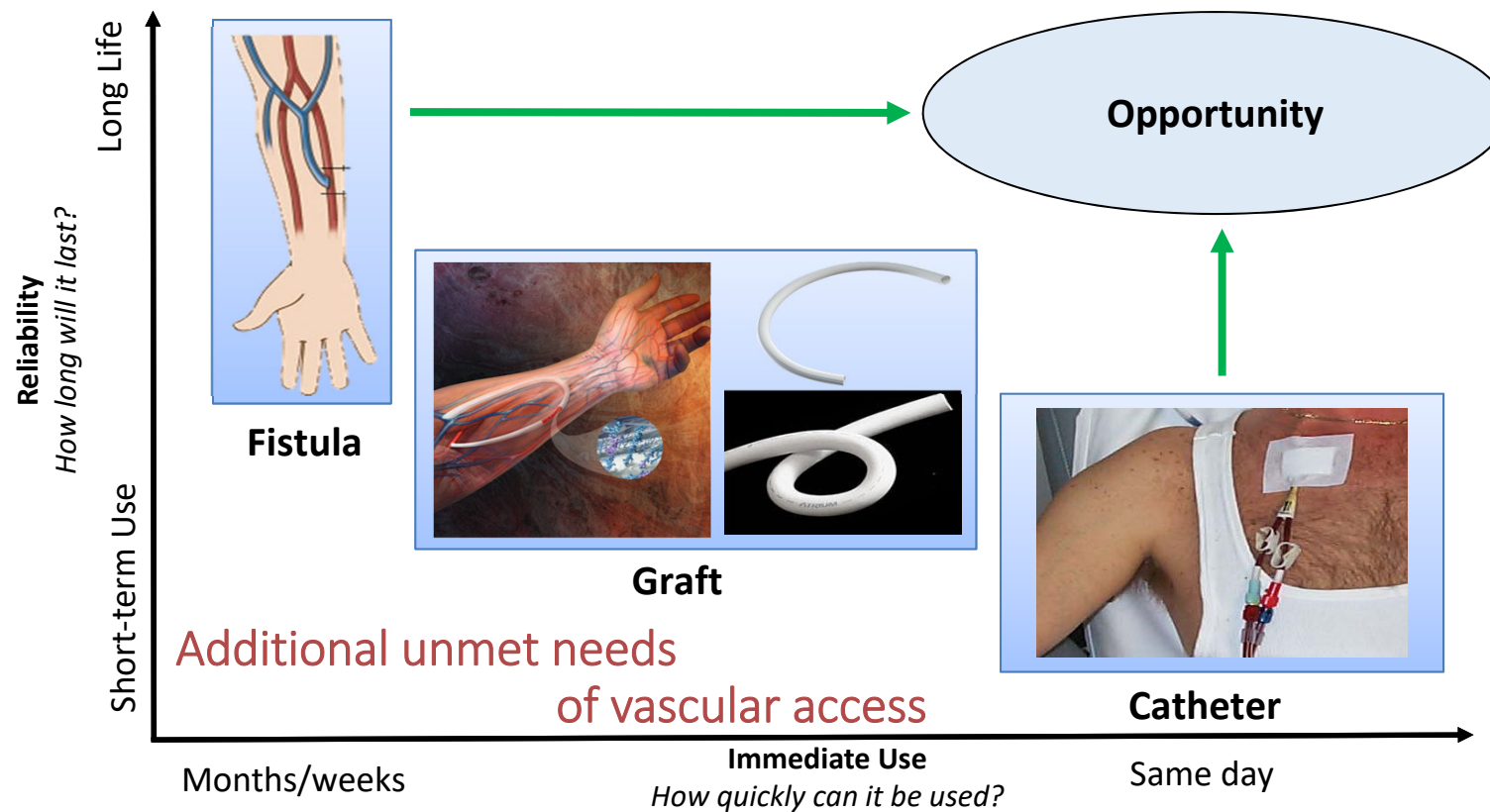
Standard Graft





What is needed?

A **reliable** access that can be used **immediately**

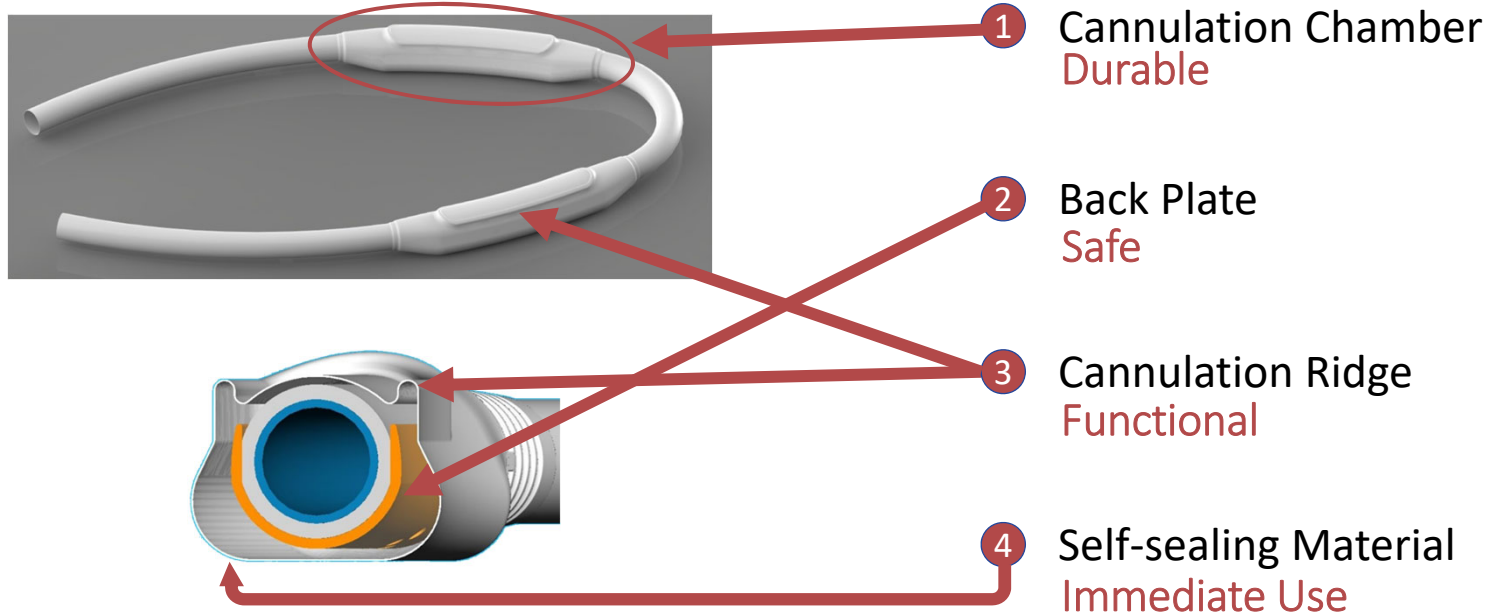


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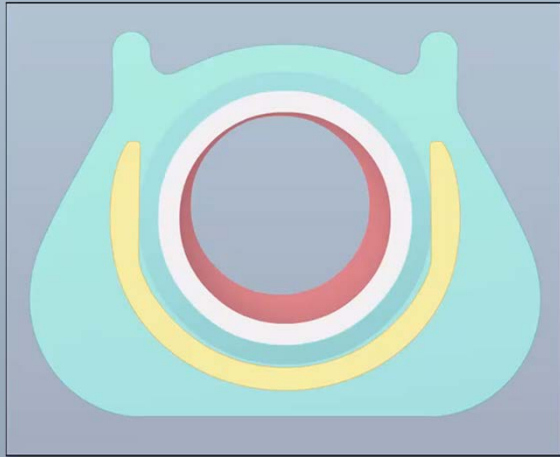
Key InnAVasc device features



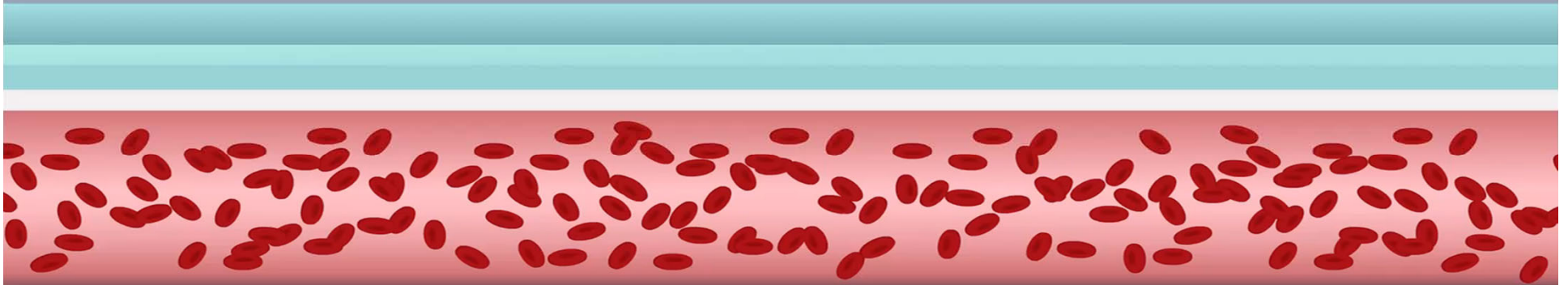
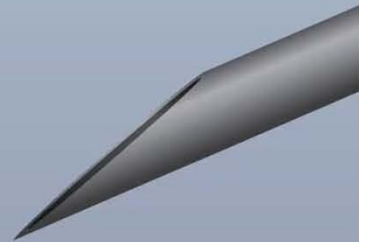
Designed to address complications faced with current grafts



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INNAVASC *Dialysis Device*



Current Clinical Feasibility Study



- 7 clinical sites in the US
- Enrollment began Jan 2, 2019
- 26 patients enrolled/implanted
 - 13 Looped, 13 Straight



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Chamber Performance to Date

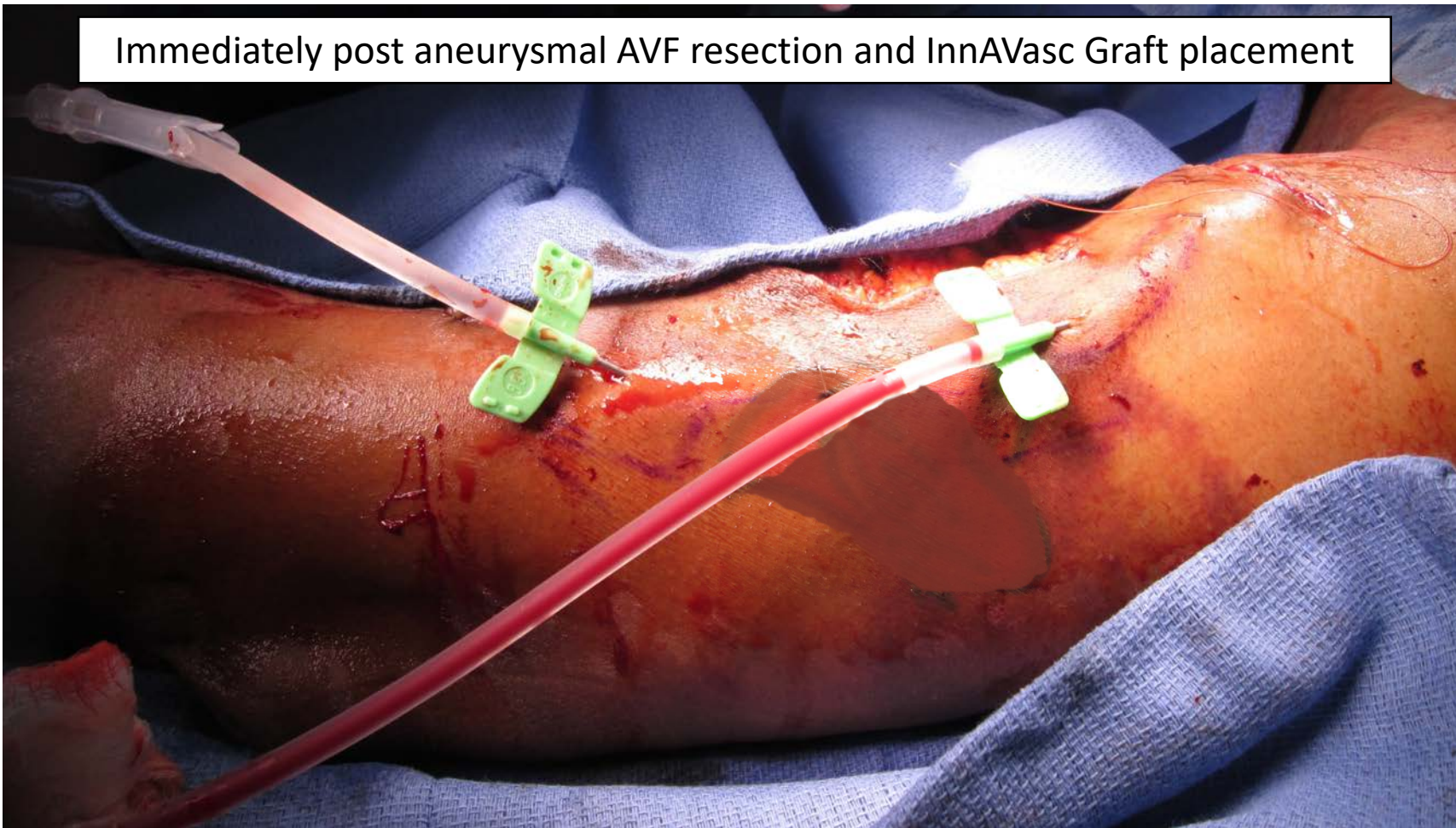
- Half of subjects have had graft used for HD POD#1
- Majority have had the graft used within 2-7 days
- Percutaneous interventions have been performed through the chamber within 1 week of implant
- No needle-related complications have been reported to date
- First 3 subjects have reached 4 months of follow up

Aneurysmal AVF – Difficulty with cannulation – No perm cath

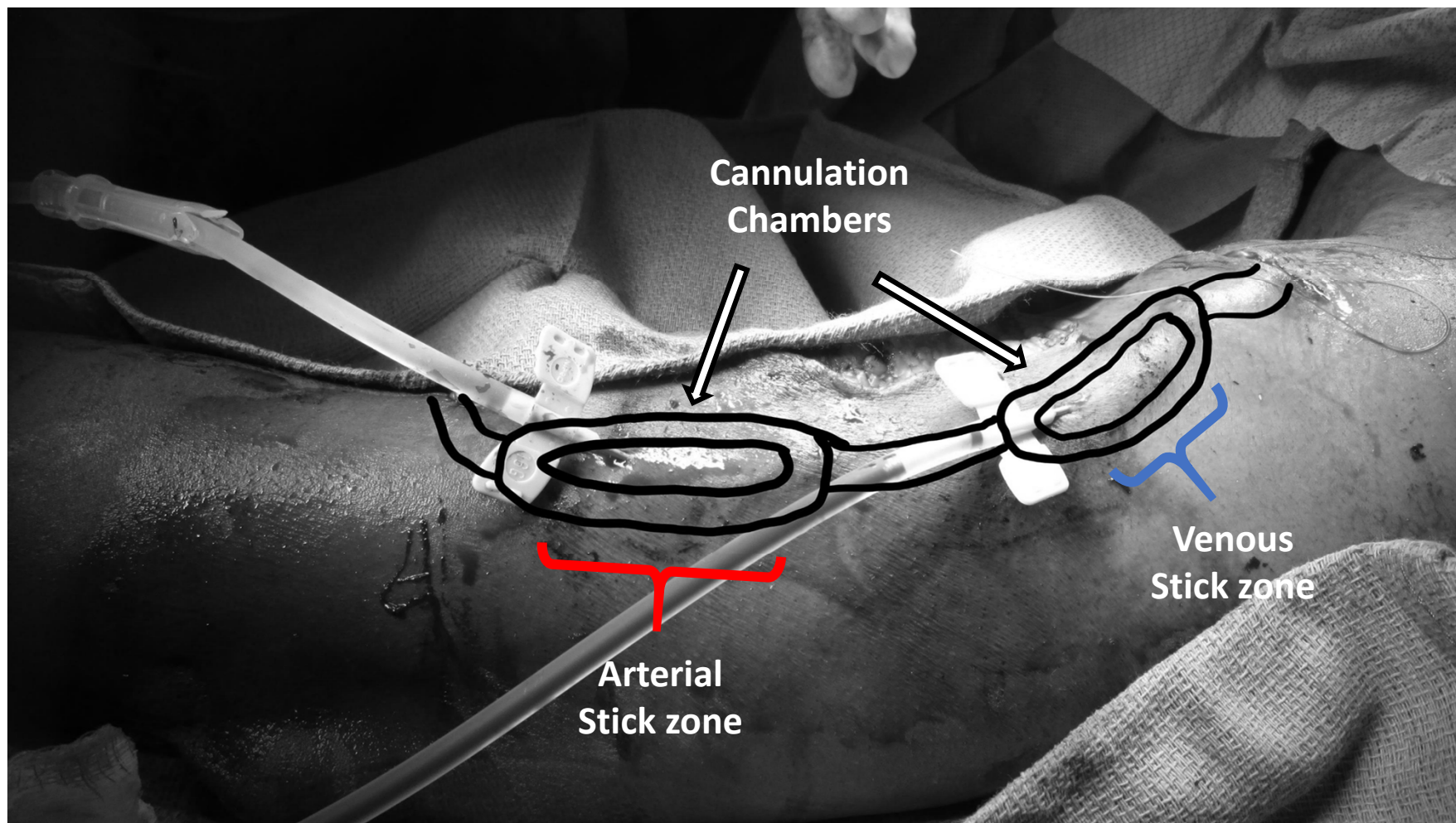


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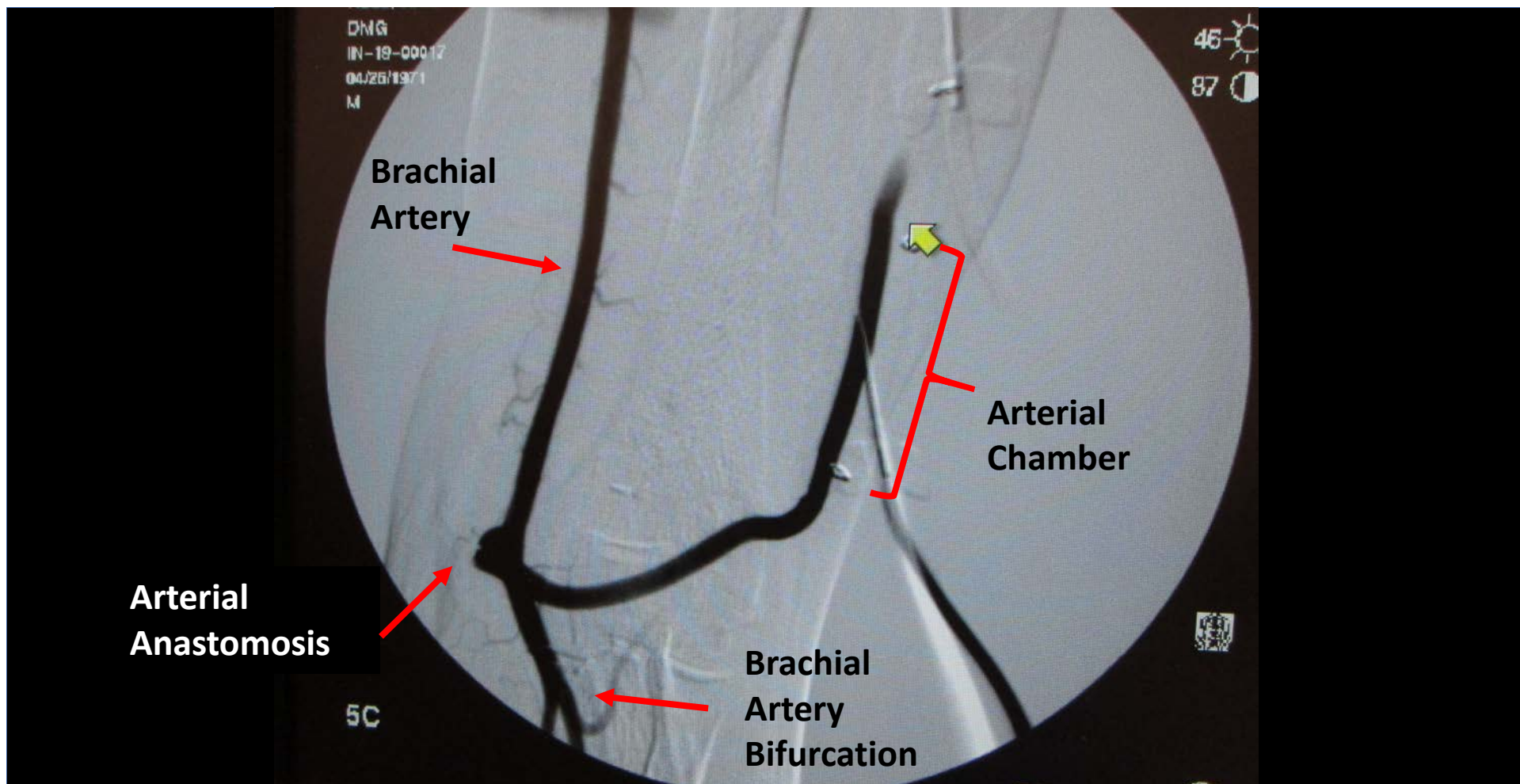
Immediately post aneurysmal AVF resection and InnAVasc Graft placement



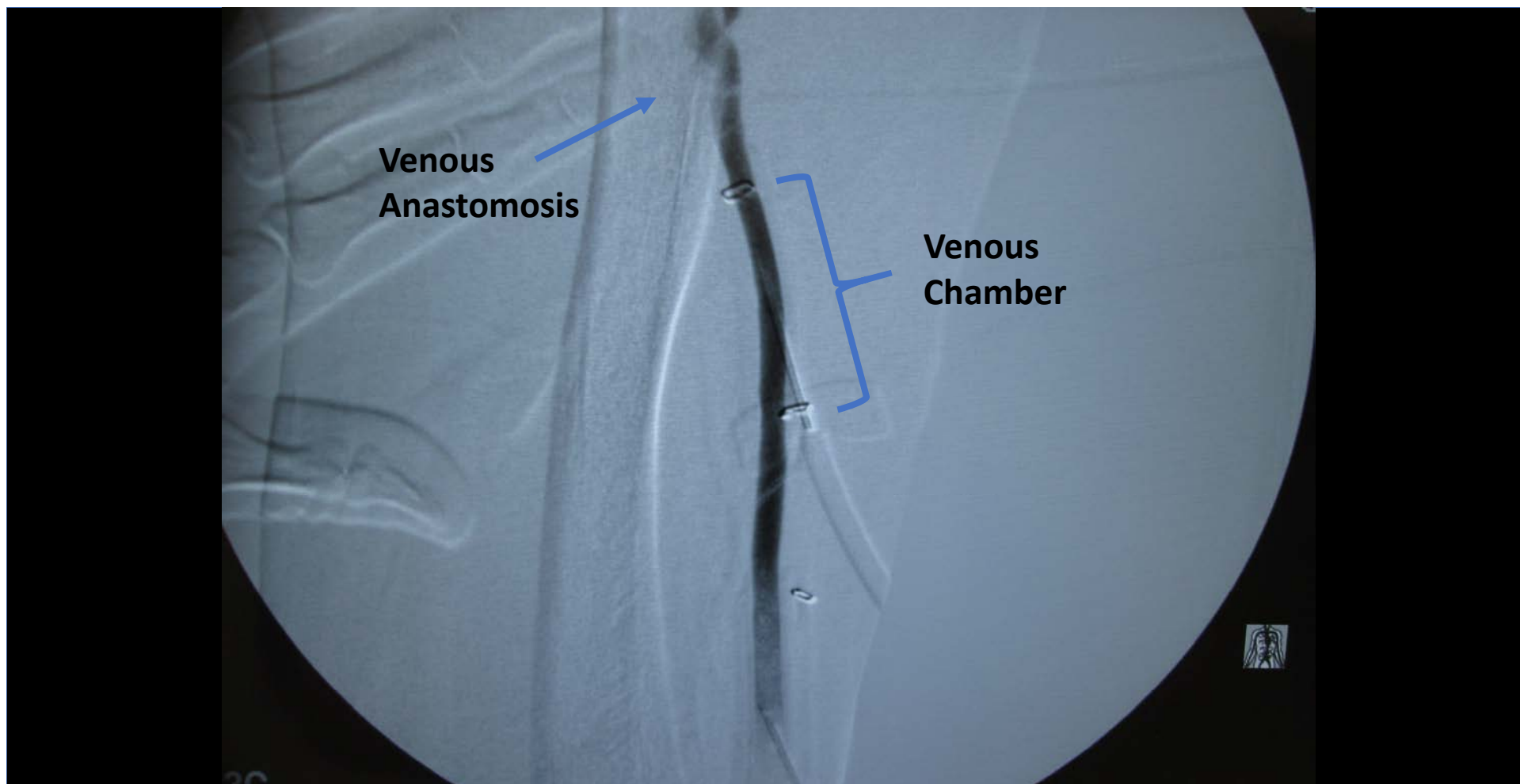
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Immediate Post-op



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Large, Aneurysmal AVF replaced with InnAVasc Graft



Used POD#1

NO
Perm Cath
Required

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Conclusion

- Immediate access with the InnAVasc Graft appears feasible and safe
- Appears to be working to prevent needle related injuries/problems
- Appears to be working to promote reliable access zones for cannulation
- Early percutaneous access for interventions appears feasible and safe
 - Accommodates standard sizes 6F-8F
- Information from this study will inform ongoing development



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