# Northwestern CENTER FOR ADVANCED REGENERATIVE ENGINEERING



# Introduction

- Cardiovascular diseases are a leading cause of mortality and morbidity worldwide [1], often necessitating the replacement of damaged blood vessels with prosthetic vascular grafts.
- Conventional vascular grafts suffer from limitations such as poor long-term patency [2] and the inability to provide real-time information about graft performance.



Atherosclerosis



**Neointimal** Hyperplasia



Calcification

• Conductive polymers have been shown to promote tissue regeneration [3] and modulate vascular cell phenotype [4].



Sulfonated poly(3,4-ethylenedioxythiophene)

**Goal:** Develop a novel generation of vascular grafts that are biocompatible AND electrically conductive.

**HYPOTHESIS**: Vascular grafts coated with S-PEDOT will demonstrate increased conductivity and support healthy vascular cell phenotype



# Engineering Conductive Vascular Grafts with S-PEDOT for Next Generation Vascular Prostheses

Taylor K. Brown<sup>\*1</sup> & Rachel Daso<sup>\*1</sup>, Joshua Tropp<sup>1</sup>, Calvin L. Chao<sup>2</sup>, Caitlyn Dang<sup>2</sup>, Aurea del Carmen<sup>2</sup>, Jonathan Rivnay<sup>#1</sup>, Bin Jiang<sup>#1,2</sup> <sup>1</sup>Department of Biomedical Engineering, Northwestern University, Chicago, IL; <sup>2</sup>Department of Surgery, Northwestern University Feinberg School of Medicine, Chicago, IL \*co-first authors; #co-corresponding authors

### 1. S-PEDOT can be polymerized in situ to coat collagen sponges and decellularized aortas.

### S-EDOT <sup>1</sup>H NMR



1% S-PEDOT

S-PEDOT Loading Efficiency



### 2. S-PEDOT-modified collagen sponges and aortas demonstrate increased conductivity.



Implantation







3. Human aortic smooth muscle cells (HAoSMCs) and human umbilical vein endothelial cells (HUVECs) adhere and survive on S-PEDOT-modified collagen sponges.



# Results





S









**IMPACT:** These bioelectronic vascular grafts will pave the way for smart vascular prostheses with built-in bio-sensing capabilities, addressing the shortcomings of current vascular graft technologies.

Scanning Electron Microscopy (Hitachi S-4800) performed at the EPIC facility of Northwestern University's NUANCE Center, supported by NSF ECCS-2025633, the IIN, and NSF DMR-2308691 • Confocal Microscopy (Nikon AXR) imaging was performed at the Northwestern University Center for Advanced Microscopy, supported by NCI CCSG P30 CA060553 awarded to the Robert H Lurie Comprehensive Cancer Center • Funding from Center for Advanced Regenerative Engineering RE-Training Program: NIH T32-EB031527.

[1] Martin, S. S. et al. *Circulation* **2024.** DOI: 10.1161/CIR.0000000000001209 [2] Pashneh-Tala, S.; MacNeil, S.; Claeyssens, F. Tissue Eng. Part B Rev. 2016. DOI: 10.1089/ten.teb.2015.0100 [3] Petty, A. J.; Keate, R. L.; Jiang, B.; Ameer, G. A.; Rivnay, J. J. Chem. Mater. 2020. DOI: 10.1021/acs.chemmater.0c00767

[4] Rowlands, A. S.; Cooper-White, J. J. *Biomaterials* 2008. DOI: 10.1016/j.biomaterials.2008.07.052 [5] Yano, H.; Kudo, K.; Marumo, K.; Okuzaki, H. Sci. Adv. 2019. DOI: 10.1126/sciadv.aav9492



**M Northwestern** Medicine<sup>®</sup> Feinberg School of Medicine

# Results

4. S-PEDOT-modified aortas implanted subcutaneously show nuclear infiltration but no infection, seroma, or delayed wound healing.

### Acknowledgements & Funding

### References