

Use of recombinant osteopontin (SPP1) protein for prevention of foreign body response

Researchers at Stanford University have found that recombinant osteopontin (SPP1) protein reduces foreign body response (FBR) and thereby facilitates successful integration and function of implantable devices.

FBR is an immune response in which detected foreign material is encapsulated in dense fibrotic scar tissue. While this response is crucial for protecting the body from bioactive foreign substances, it can be a problem when implants are intentionally placed in the body for treatment. FBR against these implants can lead to complications such as malfunction, infection, soft tissue disfigurement, and pain. Acellular dermal matrix (ADM) has been used as a coating on implants to effectively attenuate FBR, despite an incomplete understanding of the underlying mechanisms. Given that ADM is expensive and not easily scalable, it is important to identify alternatives by uncovering its mechanisms of action.

Stanford researchers have discovered that SPP1, upregulated by ADM, is necessary and sufficient for FBR attenuation. When a recombinant-SPP1-loaded hydrogel was placed adjacent to an implant, the fibrotic capsule was thinner compared to the conventional FBR. Additionally, the connective tissue architecture within the fibrotic encapsulation was comparable to that derived from ADM-coated implants. Directly managing FBR's main underlying driver SPP-1 could lead to significant improvements in biocompatibility of implantable devices.

Figure

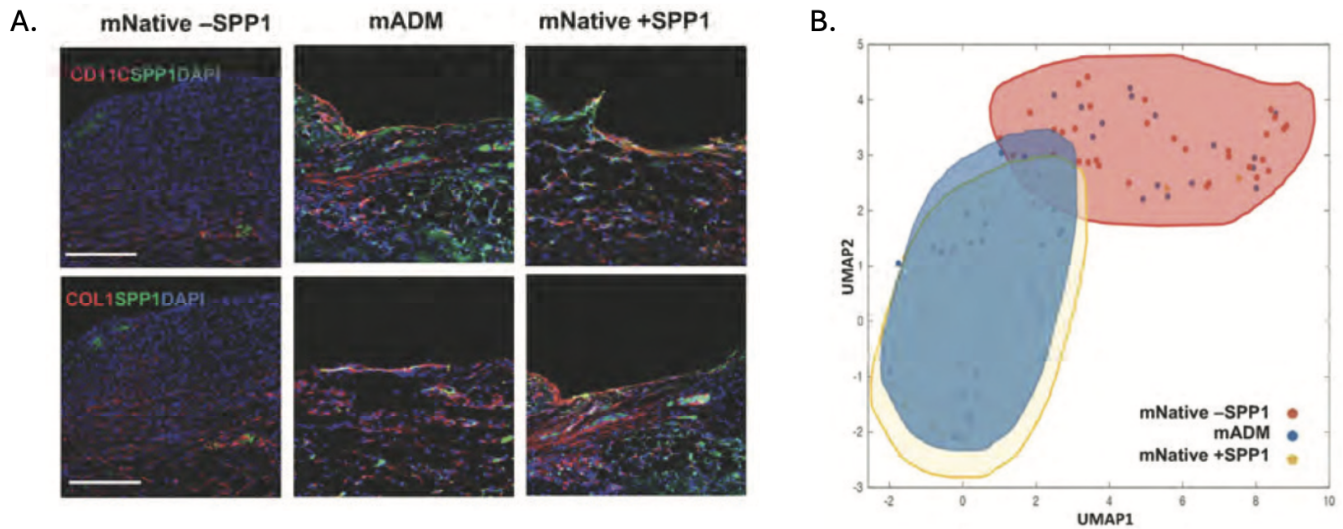


Figure Description: **A)** A comparison of fibrotic capsule thicknesses among different conditions. **B)** UMAP plot showing similarities in ADM and SPP1 treatment.

Stage of Development

In vivo data

Applications

- Biomedical implants
- Biomedical devices

Advantages

- Cost effective
- Scalable

Publications

- Griffin, M., Tevlin, R., Parker, J., Liang, N., Guo, J., Valencia, C., Morgan, A., Kuhnert, M., Momeni, A., Wan, D., & Longaker, M. T. (2024). [Acellular dermal matrix modulates monocyte transcriptome in mice and humans to attenuate foreign body response](#). *Plastic & Reconstructive Surgery-Global Open*, 12(S4), 42-43.

Innovators

- Michelle Griffin
- Norah Liang
- Jennifer Parker
- Ruth Tevlin
- Emily Meany
- Michael Longaker
- Arash Momeni
- Derrick Wan
- Eric Appel

Licensing Contact

Seth Rodgers

Licensing Manager, Life Sciences

[Email](#)