

A Novel Biological Dermal Matrix for Skin Reconstruction

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1. Introduction

Skin and soft tissue defects from trauma, burns, or surgical resection remain a major clinical challenge. Biological dermal substrates support reconstructive surgery by providing a scaffold for neovascularization and wound healing, but accessible alternatives with comparable performance are still needed.

We developed a novel dermal substrate from rabbit skin, a sustainable agri-food by-product, preserving a collagen-elastin network and ECM organization similar to human dermis. Its performance was benchmarked against two commercial substitutes, Matriderm® (MedSkin) and NovoSorb® (PolyNovo).

3. Results and Discussion

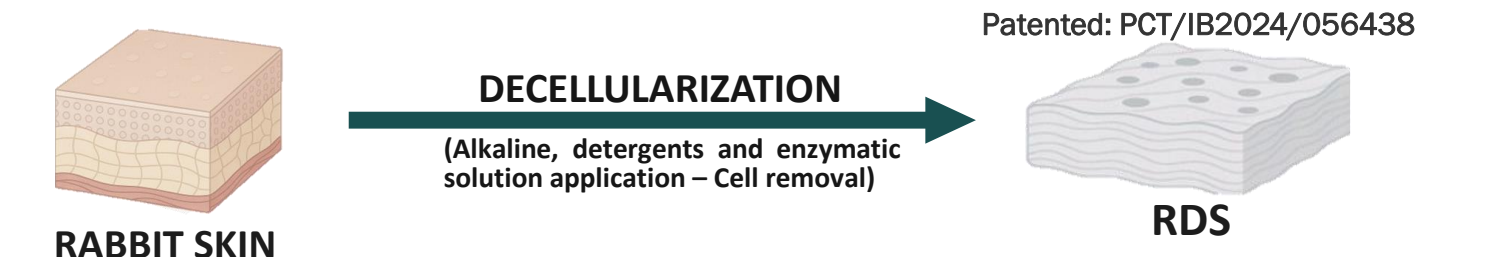
Product Characterization

RDS Composition	
ECM Component	Concentration (µg/ mg of dry tissue)
Collagen	
Soluble	59,6 ± 5,47
Unsoluble	188,2 ± 72,3
Total Collagen	248,9 ± 77,8
Elastin	22,1 ± 7,45
GAGs	0,20 ± 0,13
DNA	66,8 ± 11,0 ng/ mg of dry tissue

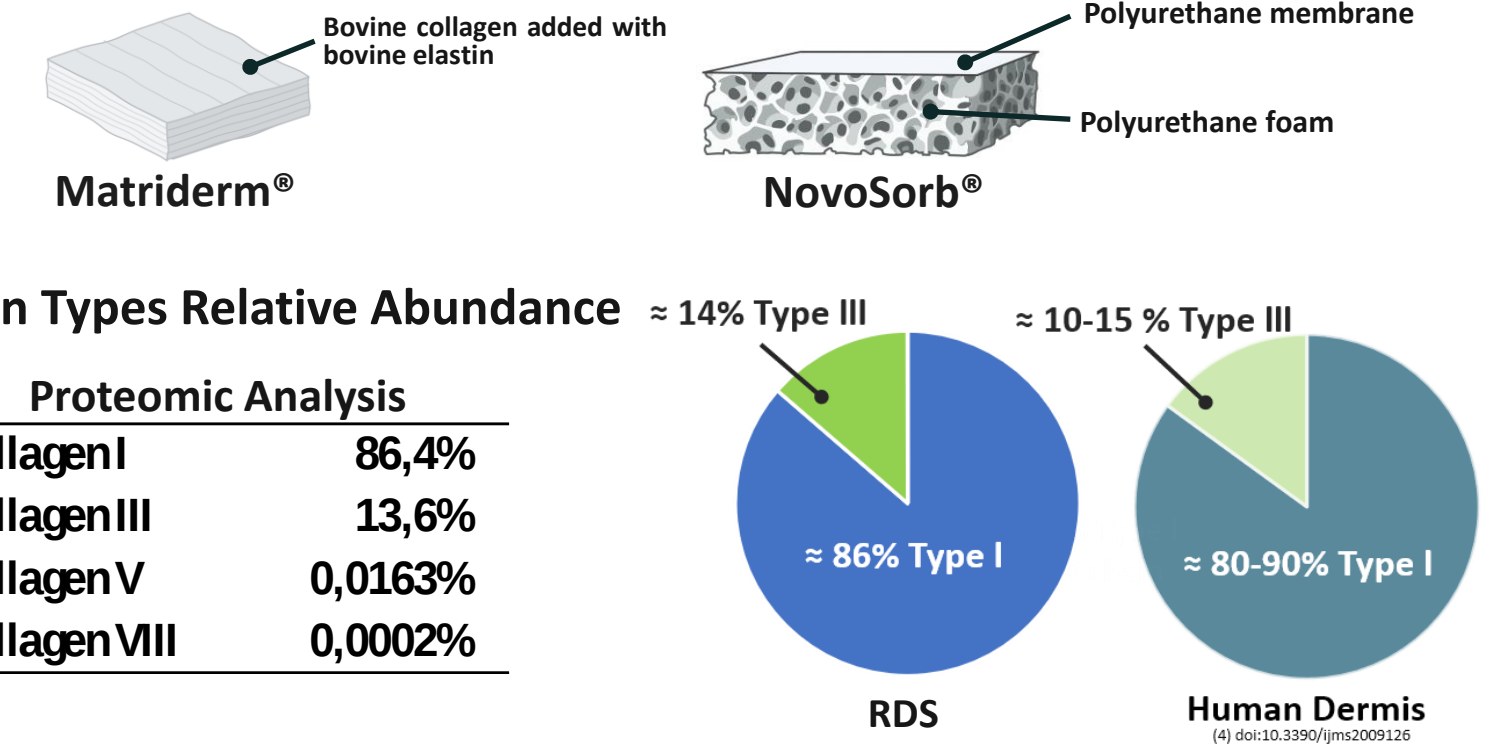


2. Materials and Methods

Production of rabbit skin-derived dermal substrate (RDS)



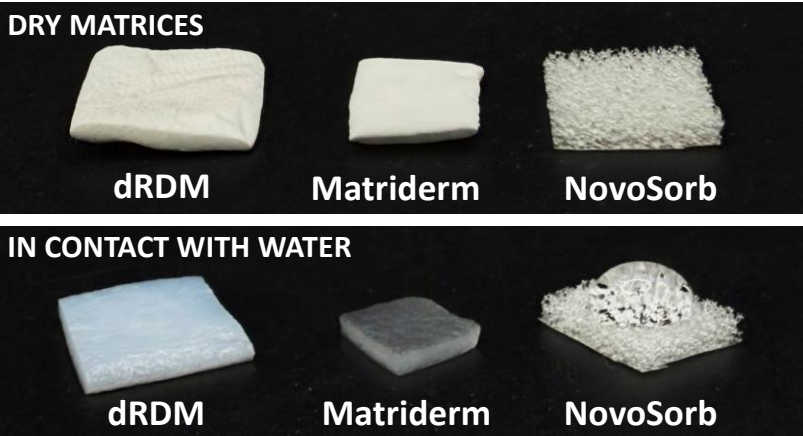
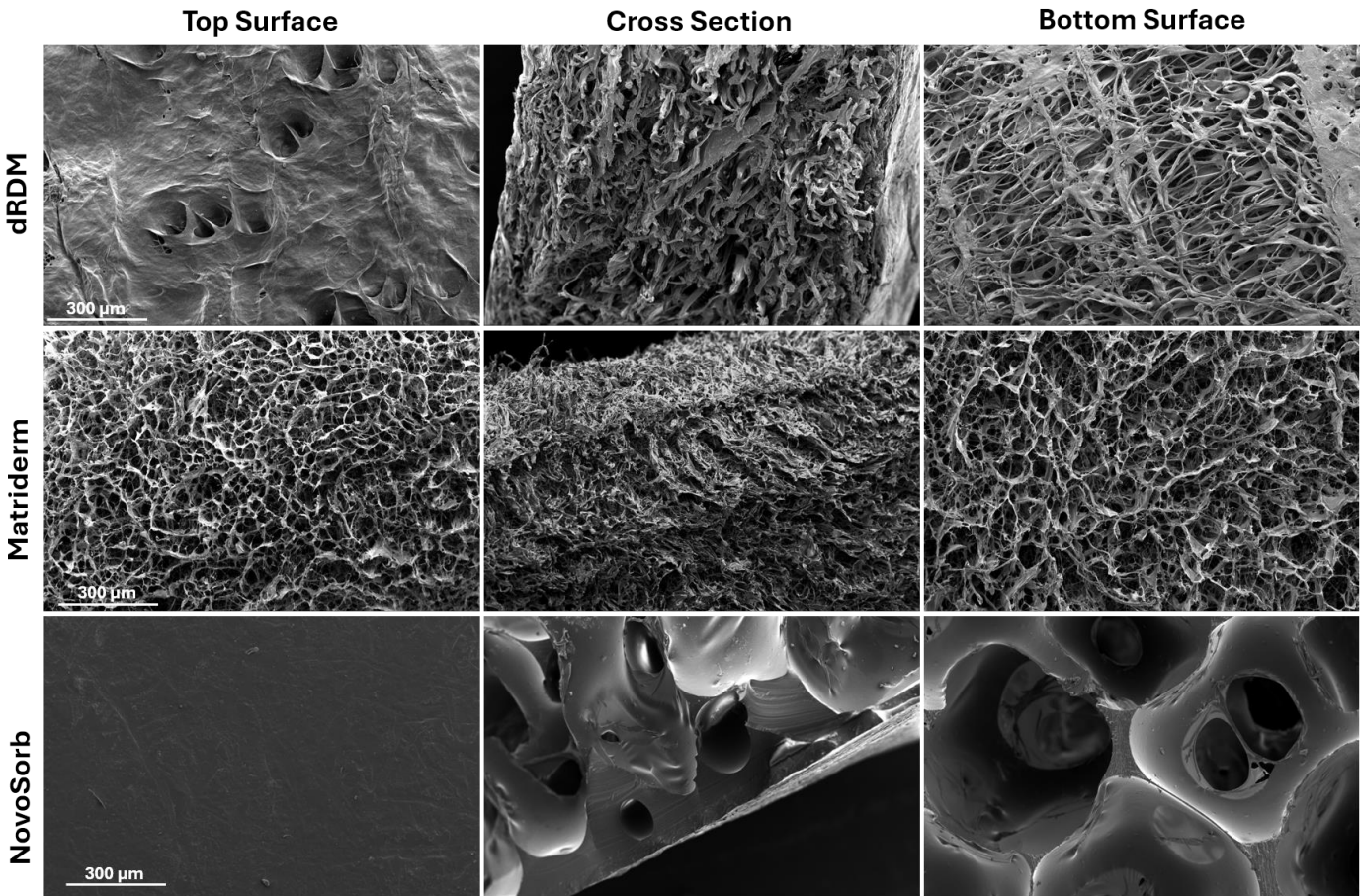
Benchmarked against commercial matrices:



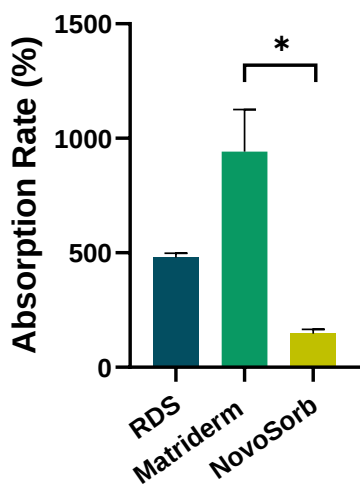
RDS is composed mainly of collagen (predominantly Type I and III, similar to human dermis) and elastin, confirmed by proteomic analysis. DNA is successfully removed, and histology confirms well-preserved collagen fibers with no cells or cellular debris.

Benchmarking with Matriderm® and NovoSorb®

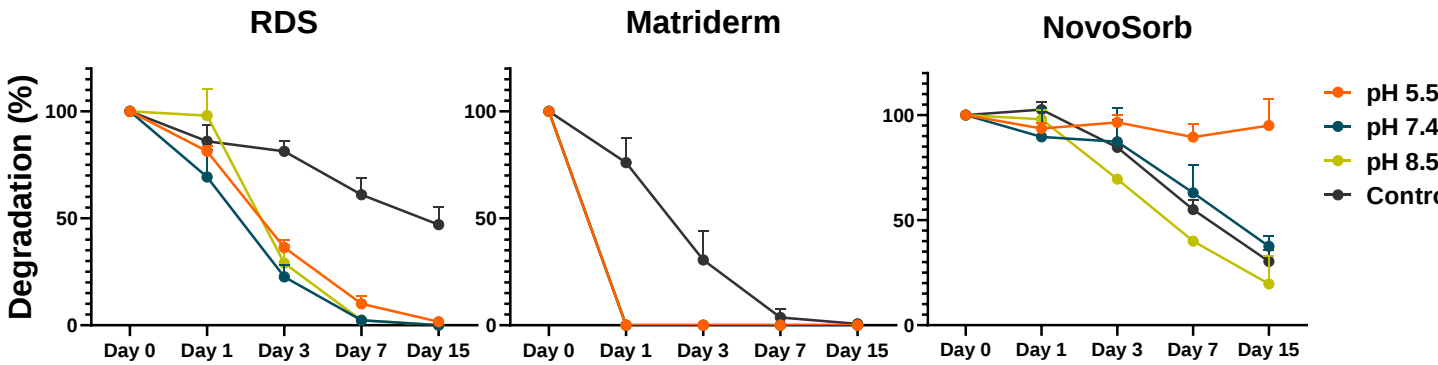
Scanning Electron Microscopy



Swelling Properties



Degradation Assay with Simulated Wound Fluid



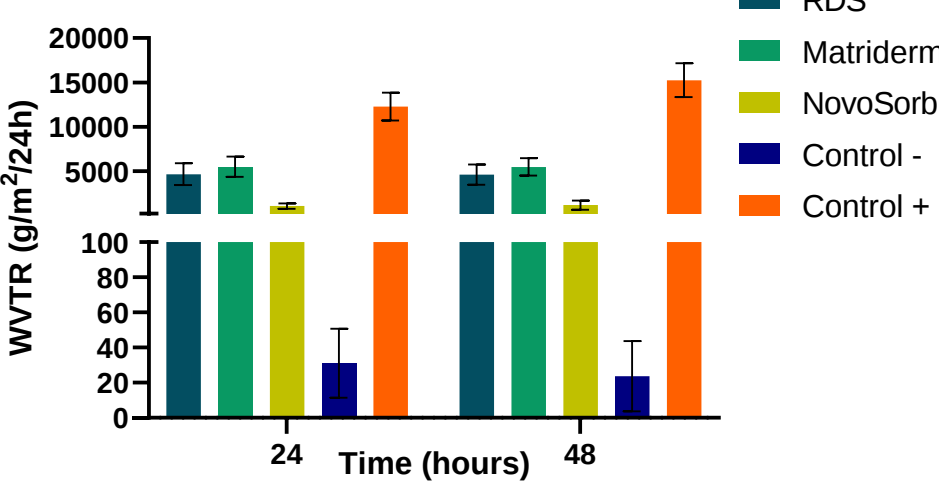
Morphology: RDS and Matriderm are both collagen-fiber structure, but RDS presents a more organized fiber structure while Matriderm presents more random fibers. RDS preserved basal membrane supports epidermis formation.

Permeability: RDS and Matriderm's porous network enables water/nutrient exchange while NovoSorb's more hydrophobic polyurethane film limits this, hindering regeneration.

Degradation: RDS degrades gradually, providing prolonged tissue support, whereas Matriderm degrades rapidly and NovoSorb degrades very slowly, potentially compromising regeneration.

Cell support: RDS and Matriderm both sustain cell growth for 30+ days while NovoSorb revealed reduced cellular performance

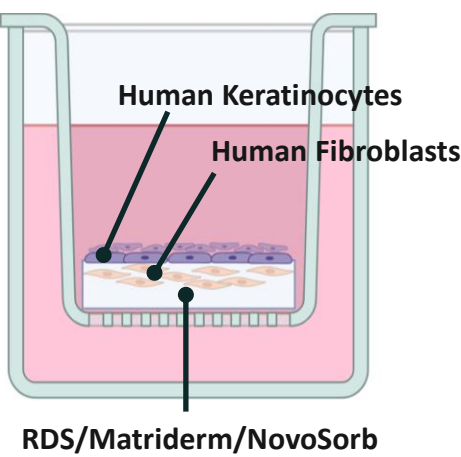
Poster WVTR



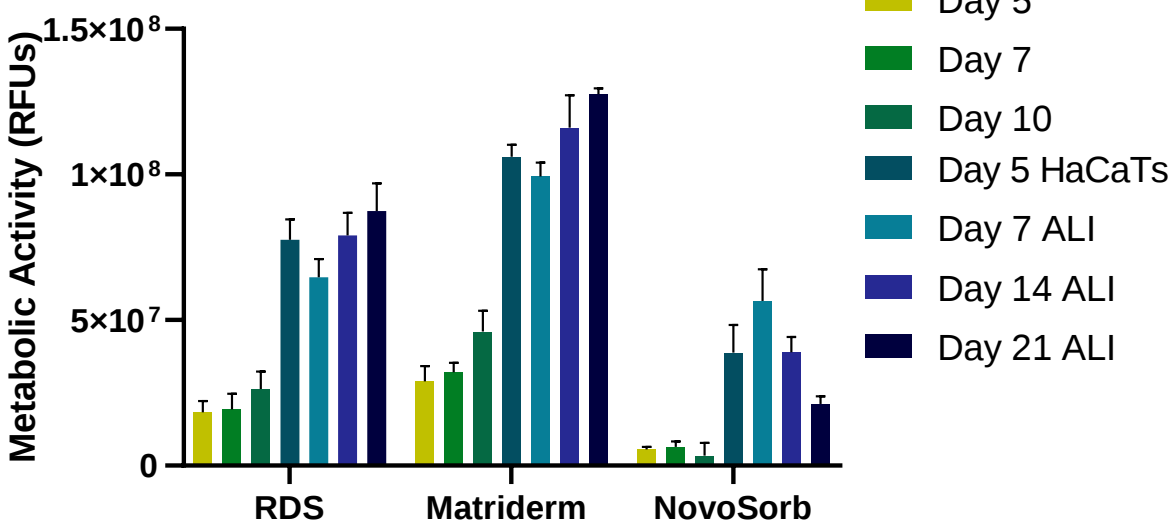
4. Conclusion

The RDS shows a collagen-elastin architecture comparable to native dermis with efficient decellularization. It matches Matriderm's permeability and cell-support performance while offering a more controlled degradation, outperforming NovoSorb overall. These findings support dRDM as a sustainable, effective alternative for dermal reconstruction.

Dermo-Epidermal Model



Dermo-Epidermal Support Capacity



Acknowledgments

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