

Quantifying postburn immune responses: Using a computational model to predict scars early

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Introduction

- **Persistent** and **dysregulated inflammation** is a key driver of scar outcome in burn survivors, contributing to the development of hypertrophic (A) or normotrophic scars (B) during healing.

Aim

- To develop a **computational model** that simulates the **mechanistic link between inflammatory profiles** and the **proliferative phase of burn wound healing**.

Research questions

- Which **inflammatory markers** in human post-burn are **predictors of scar outcomes** during the remodeling phase of wound healing?
- Can **early-stage inflammatory profiles** accurately predict the development of hypertrophic scars during the proliferative phase?

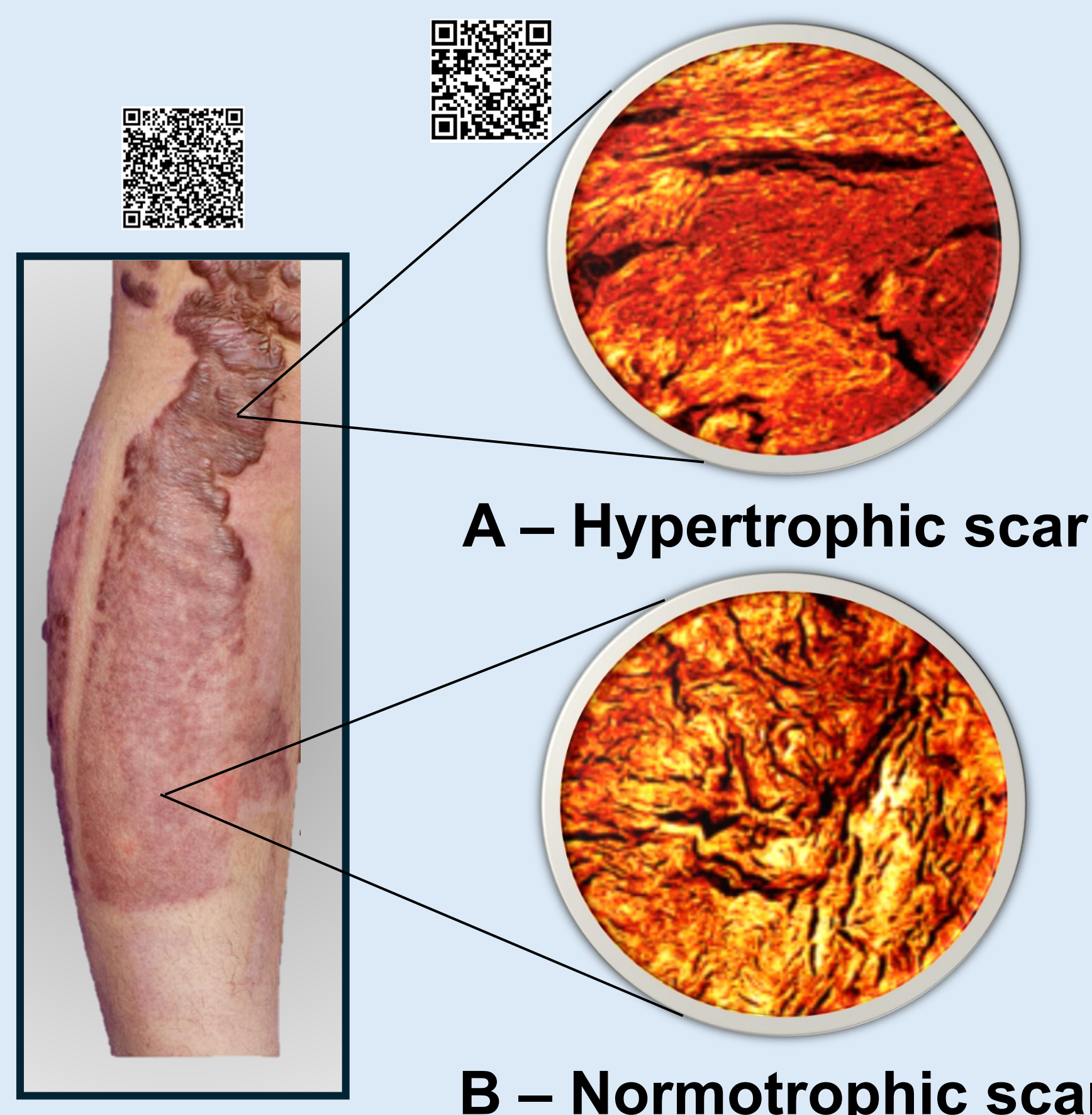


Figure 1 – Hypertrophic scar and normotrophic scar on a leg and the corresponding microscopic collagen deposition, A and B, respectively. Sources for each figure accessible through the QR code.

Methods

Human data was used in a computational model

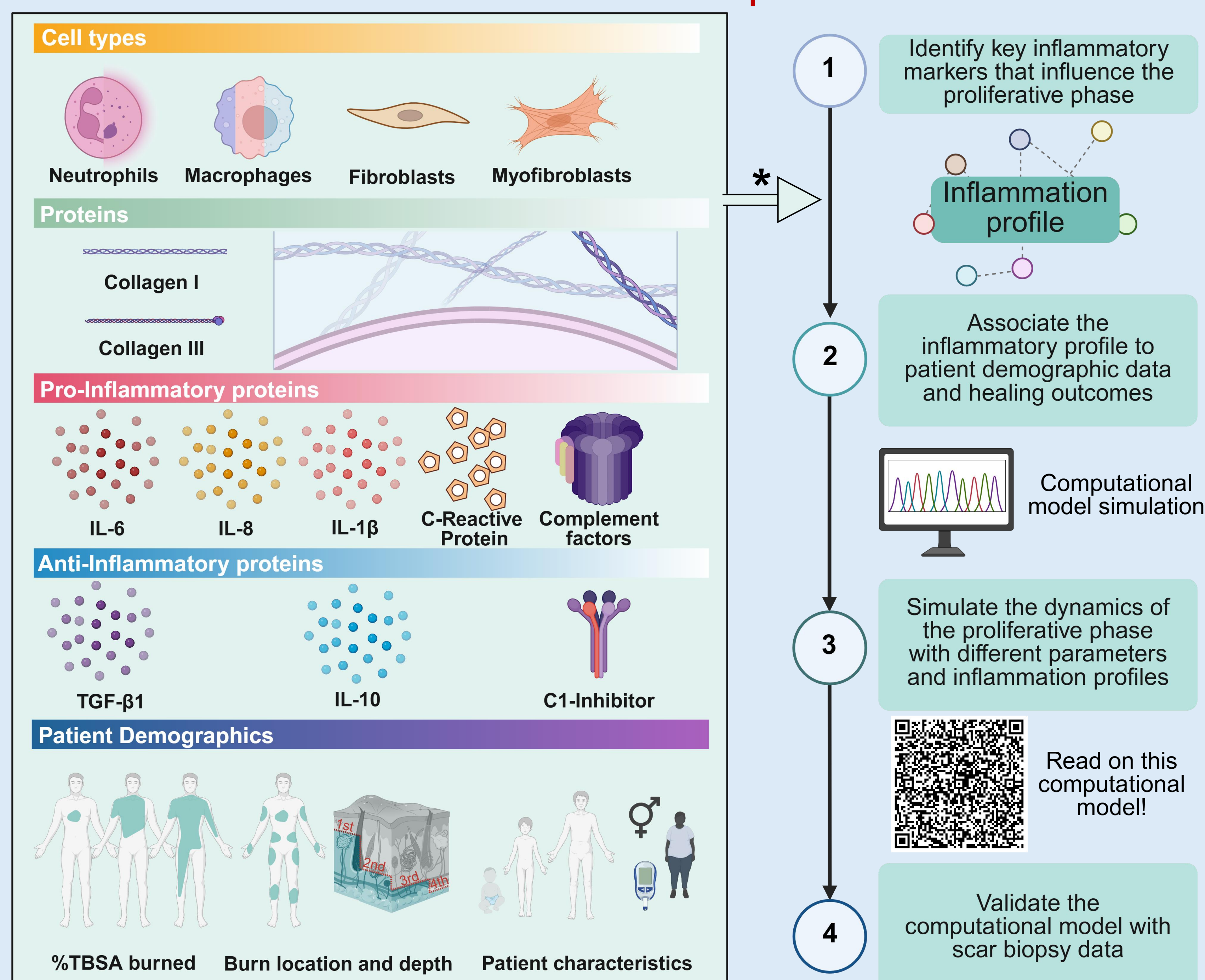


Figure 2 – Overview of clinical data collected from burn patients, including various cell types, extracellular matrix proteins, pro-/anti-inflammatory markers, and patient demographics (left). The proposed methodology (right) outlines how this data informs computational modelling for early prediction of scar outcomes during the proliferative phase. Collected data supports generation of patient-specific inflammation profiles, while reported outcomes serve for model validation. A hypothetical version of the model can be accessed via the QR code.

Preliminary Results

Prediction of the scar outcome

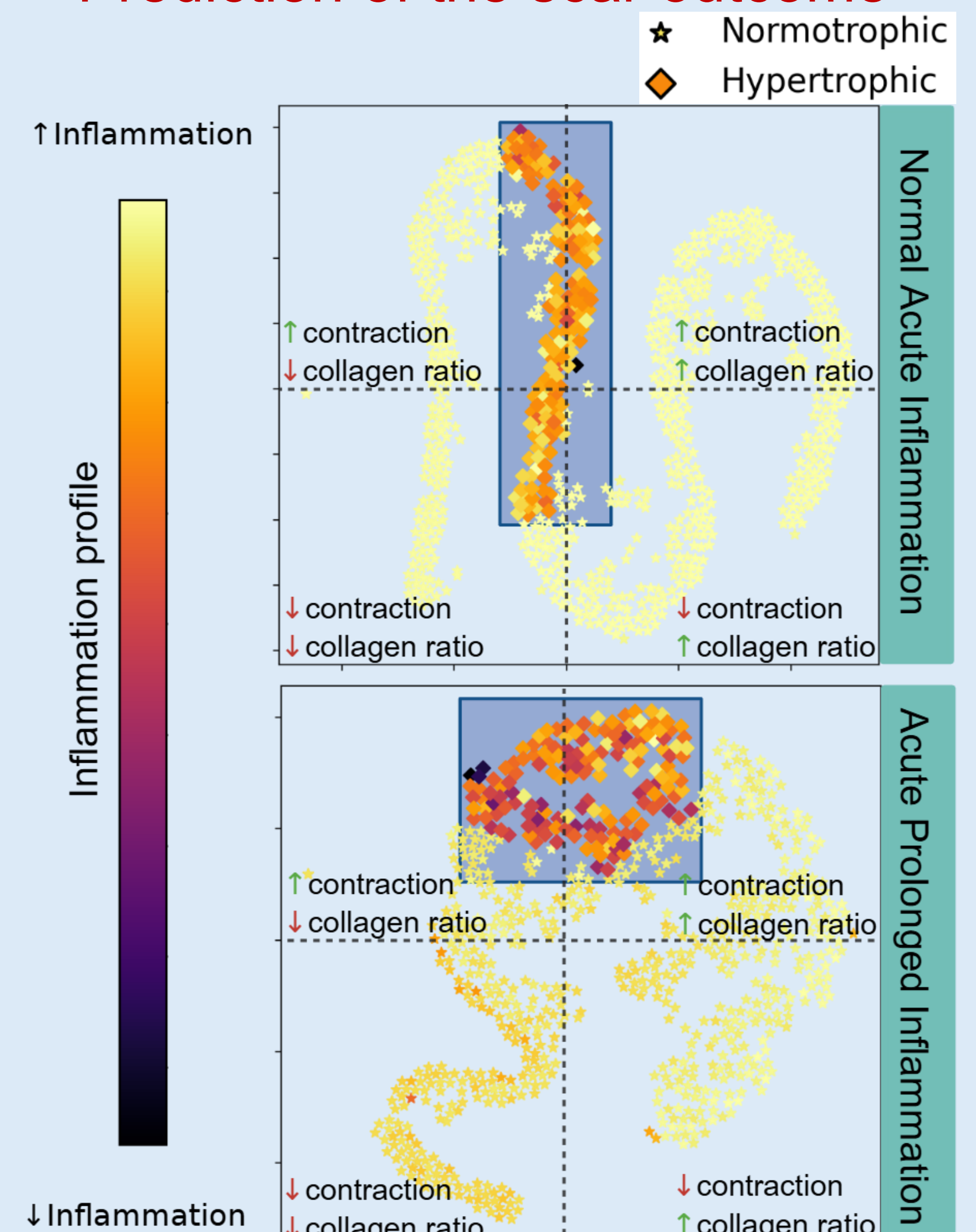


Figure 3 – Prediction of 1000 virtual scar outcomes under two scenarios, based on myofibroblast activity, collagen I/III ratio, and inflammation profile (defined by the ratio of pro- to anti-inflammatory cells and cytokines). The blue area indicates the cluster of hypertrophic scars. Plot quadrants categorize outcomes according to contraction and collagen ratio.

Discussion

- **Contraction** and **collagen ratio**, alongside **inflammation**, are strong predictors of scar outcomes
- The model successfully **simulates diverse scar outcomes** using a simplified mechanistic framework
- Key predictors for the remodelling phase outcomes are **still being identified** as the model undergoes continued development
- The model is currently **undergoing validation with human data (Step 4)**
- While **effective for generating population-level predictions**, the **current data lacks the specificity** required for personalized patient outcomes

About me

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