An Investigation of the Fluid Structure Interaction in Articular



Engineering and Physical Sciences Research Council Cartilage across Disparate Scales

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Clinical Background

- Articular cartilage (AC) is found at opposing surfaces in mammalian joints. It provides a smooth bearing surface, promoting low friction articulation, facilitating continuous operation under relative motion.
- A lack of cells within AC renders a low capacity for intrinsic healing or repair, leaving it prone to degeneration.
- High clinical demand for cartilage repair, with
 8.75 million people aged 45 and over having sought treatment for osteoarthritis (a disease developed when cartilage breaks down) in the UK [1]. Which is predicted to cost the NHS around £118.6 billion in the next decade [2].

Modelling Articular Cartilage

- AC has a biphasic structure, with a deformable matrix of collagen, immersed in an interstitial fluid. The complex tissue composition spans a range of length scales (Fig.1).
- The micro-structure of the tissue contains a collagenous network of cells with variable permeability across a stratified composition.



Fig.1: Multi-scale structure of AC (adapted from [3]).

• Current single-scale constitutive models do not capture the physical interactions at both the fibre and material scale.

 This project aims to create a coupling between the micro- and macro-scales to map the pore-scale effects onto the material-scale response.

Scientific Aims and Objectives

Project Aim

Couple **an immersed fibre network** (micro-scale) model with a **poroelastic continuum mechanical** (macro-scale) model to create a novel multi-scale model of articular cartilage and its mechanical response.

Objectives:

- Refine computational set up to create an efficient, accurate multi-scale solver.
- Complexify the model to incorporate a stratified
 composition and damage modelling to improve
 clinical understanding of damaged tissue.



Representative Computational Results

- Multi-scale model uses COMSOL-MATLAB Live Link. Micro- and Macroscales coupled using Heterogeneous Multi-scale Methods.
- Currently, the continuum-continuum macro-scale model undergoes a 1D, unconfined compression in the negative y-direction (Fig2a).



- Next, include **fibrous network micro-scale model** using extensional shear.
- Replace continuum solid with a bond-diluted triangular **spring network**.
- Challenges: fibre model is inherently oscillatory, calculating non-constant parameters, e.g. the Biot Willis coefficient.

Model Validation

• Current **multi-scale continuum-continuum model** has been validated against a **single scale model** (e.g. Fig 3a).



Fig.3 Types of validation for the multi-scale model.

- Experimental compression and indentation testing of cartilage samples will be used to validate the model (Fig 3b).
- Atomic Force Microscopy to understand the fibre-structure of the tissue for validation and improvement of the fibre-model
- Clinical Collaboration will be used to optimize a biological model (Fig 3c).

Model Refinement and Optimization

- o Increasing model complexity will require computational optimization.
- **Meta-modelling,** for example GPTIPS is being explored to speed up multiscale model simulations.
- Machine Learning implemented to select the most pertinent data points.
- **Model efficiency** will be improved with efficient data storage.

References

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