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Finite Element Analysis to Model the Effects of Osteoarthritis in the Knee

INTRODUCTION

Significance and Background

Our Approach

Study Objective and Hypothesis

MATERIALS & METHODS

Modeling

Evaluation

- **Osteoarthritis (OA) is a degenerative disease that begins with the breakdown of the joint cartilage, and affects over half a million people globally [1,2]**
- **The presence of OA in the knee can lead to damage of the cartilage tissue, tendons, synovium, and bone, causing chronic pain, swelling, loss of motion, and may lead to excessive stress on the joint [1]**
- **Cartilage's avascular nature causes it to have a low regenerative capacity, because it does not have the blood vessels necessary to distribute the oxygen, nutrients, or white blood cells needed to stimulate healing [3]**
- **The subchondral bone (SB), calcified cartilage (CC), and articular cartilage (AC) provide mechanical support to the knee [4] (Fig. 1B)**
- **Pathological processes that affect the joints morphology or joint components, lead to changes in the biomechanics of the knee, and has been correlated to the presence of OA [1]**
- **Computational modeling with finite element analysis is used to stimulate and study complex systems with models. These models contain variables that characterize the system being studied, and can be adjusted to study the response from system [8]**
- **Understanding how knees with OA experience the stresses and strains from the compressional forces applied throughout everyday use is integral to engineering cartilage that is capable of replacing damaged tissue**

- **The objective of this study is to use finite element analysis to model the subchondral bone and calcified cartilage interface, in order to quantify and analyze how the morphology of the interface, and presence of osteoarthritis affects the stress and strain distributions experienced by the knee.**
- **We hypothesize that the pathological changes experienced by knees with osteoarthritis will result in changes to how compressional loads are distributed across the calcified cartilage and subchondral bone interface, and will affect how and where the knee experiences these stresses and strains**

- **Develop specific and generalized models of the SB and CC in FEBio to conduct finite element analysis to determine if there are differences in the stress and strain distributions for healthy and OA**
- **Identify the regions of maximum stress and strain, and the magnitudes of those stresses and strains**

- **SB and CC interface to analyze if the stress and strain distributions vary between each group Generalized Models: n = 1**
	- **The generalized models were created using averaged, subject specific data from the histology samples (Fig. 3A,B) where the dimensions of the model, and the shape and frequency of the undulations (Fig. 6) of the SB and CC interface were determined through histomorphometric analysis**

Subject Specific Models: n = 3

• **The subject specific models were created using 2D histology images of the**

- **osteochondral interface of healthy and OA subjects to generate the 3D models in FEBio** • **In order to analyze how the morphological differences between healthy and OA models affects the stress and strain distributions, the Young's Modulus, Poisson's ratio, and density between both groups were fixed**
- **Once the dimensions, material properties, and mesh size were determined, FEBio was used to generate the models**

- **After the generalized and subject specific models were generated, a compression test was performed at 4 MPa on all models and the stress and strain colormap's were outputted, showing how the compressional load was distributed in each model**
	- **The 4 MPa compressional load was applied to top surface of model, the magnitude was determined was chosen based on physiological loading experienced in the knee when walking [7]**
	- **A confined compression test was performed, where the top and bottom of the model were fixed in all directions and all sides were fixed in direction of they are perpendicular to (Fig. 4)**
- **After the stress and strain colormaps were generated, the node IDs (Fig 5), their associated stress values, and the x, y, z positions were extracted from FEBio and imported into Excel to conduct further analysis on the stress distribution**

be conducted

REFERENCES ACKNOWLEDGMENTS The authors would like to acknowledge the funding and support from the Amazon Summer Undergraduate Research Experience (SURE) Fellowship and NSF DGE-2036197 (JZL), NIH 5R01AR073529 (HHL) [1] J.H. Abidi. BMC Musculoskelet. Disord. 2023;(175-182) [2] WHO. Osteoarthritis. 2023:(1) [3] Lou-Ren C.StatPearls. 2022;(1) [4] Lin-Fen H. Braddom's Rehabilitation Care: A Clinical Handbook. 2018;(208-213) [5] A. Zelink Dev Biol. 2020;8(3-4) 2020;8:(3-4) [7] H. Yoshida. Thr Dimen Dyn hi con area press dist. 2006;39(11) [8] NIH. Computational Modeling. 2020:(1)

Subject Specific Models:

- **healthy and OA groups**
- **Maximum stress is applied**

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- **Further analysis of how fatigue loading affects the stress and strain distributions, along with the rate of deformation on the SB and CC interface, should**

