Fatigue Simulation of Human Cortical Bone using Non-Homogeneous Finite Element Models to Examine the Importance of Sizing Factors on Damage Laws

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ABSTRACT

Finite element modeling has become a powerful tool in orthopedic biomechanics, allowing simulations with complex geometries. Current fatigue behavior simulations are unable to accurately predict the cycles to failure, creep, and damage or modulus loss even when applied to a bending model. It is thought that the inhomogeneity of the models may be the source of the problem. It has also been suggested that the volume size of the element will affect the fatigue behavior. This is called a stressed volume effect. In this thesis non-homogeneous finite element models were used to examine the effects of “sizing factors” on damage laws in fatigue simulations.

Non-homogeneous finite element models were created from micro computed tomography (CT) images of dumbbell shaped fatigue samples. An automatic voxel meshing technique was used which converted the CT data directly into mesh geometry and material properties.

My results showed that including these sizing factors improved the accuracy of the fatigue simulations on the non-homogeneous models. Using the Nelder-Mead optimization routine, I optimized the sizing factors for a group of 5 models. When these optimized sizing factors were applied to other models they improved the accuracy of the simulations but not as much as for the original models, but they improved the results more than with no sizing factors at all. I found that in our fatigue simulations we could
account for the effects of stressed volume and inhomogeneity by including sizing factors in the life and damaging laws.
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Chapter 1: Introduction

The quantification of the mechanical behavior of healthy bone is a necessity for the study of implant failure, femoral neck fractures, stress fractures and senile fractures. Fatigue testing on bone can help define empirical relationships and methods that predict the mechanical behavior and properties (Cotton, Winwood et al. 2005). These methods can then simulate implants in an attempt to increase the fixation life in the surrounding bone (Taylor and Tanner 1997). Stress fractures that are common in athletes and military recruits, occur in healthy bone under high cyclic loading situations; while senile fractures occur in diseased bone (Donahue, Sharkey et al. 2000).

Numerous studies have investigated the fatigue behavior in bone (Carter and Caler 1983; Zioupos, Wang et al. 1996; Zioupos, Currey et al. 2001; Cotton, Winwood et al. 2005). These studies have examined cycles to failure vs. stress, damage vs. stress and have shown bone can experience a loss of the material stiffness, loss of strength, and accumulation of permanent strain during fatigue. Damage in bone can be defined as loss of the mechanical properties reducing the usefulness. The damaged state could be found by measuring the permanent changes in the mechanical behavior during a damaging event.

Computed tomography (CT) uses x-ray equipment to obtain cross-sectional images of the body. CT images map different attenuation coefficients, which differentiate bones, organs and other tissues. The information from a CT image can be used to model specific geometry and modulus of bone.
Finite element (FE) modeling has become a powerful tool in orthopedic biomechanics, allowing simulations with complex geometries to be solved. Advanced material behavior has been simulated including; fatigue behavior (Taylor, Verdonschot et al. 1999), implant failure (Taylor and Tanner 1997) and fracture location prediction (Keyak, Rossi et al. 2001). Even though finite element analysis is widely used, the results can vary due to meshing techniques used before solving the problem (Viceconti, Bellingeri et al. 1998).

This thesis describes the first FE simulation of fatigue behavior in non-homogeneous samples of human cortical bone. Previous fatigue behavior simulations used homogeneous FE models of human cortical bone samples. The simulation’s predicted modulus degradation and permanent strain data were in close agreement with the experimental data, both in terms of the trends and absolute magnitudes (Taylor, Verdonschot et al. 1999). This simulation technique did not accurately predict the cycles to failure, creep or damage when applied to a model representing cancellous bone. It was suggested that the inhomogeneity of the model could be the source of the problems (Taylor, Cotton et al. 2002). It has also been proposed the fatigue behavior of bone like many other materials is dependent on the size of the test specimen called stressed volume or “sizing effects” (Taylor and Kuiper 2001). In our simulation non-homogeneous FE models of cortical bone samples were created using an automatic voxel meshing technique, where CT images were used as source information for the geometry and material properties. The primary goal of this thesis was to examine the effects of incorporating “sizing factors” ($K_N$, $K_D$, and $K_{cr}$) into the life and damaging laws to account for sizing effects and inhomogeneity to achieve improved fatigue simulations for
non-homogeneous FE models. Where, the life and damaging laws are the empirical equations from Cotton, Winwood, et al. (2005).

This chapter presents background information relevant to the of fatigue behavior simulation of cortical bone. The mechanical behavior of cortical bone during \textit{in vitro} fatigue testing is described. FE meshing techniques, models, and simulations relevant to this research are discussed.

\section{1.1 Bone}

There are two major classifications of bone: cortical and cancellous bone. Cortical, or compact bone represents nearly 80\% of the skeletal mass and forms a structural shell around most every bone in the body (Carter and Hayes 1977). It is stiff with a longitudinal elastic modulus between 14-20 GPa (Taylor, O'Reilly et al. 2003) and a low porosity of 5-30\% (Carter and Hayes 1977; Cowin 2001). Cancellous, or trabecular bone, while only representing 20\% of the skeletal mass represents 80\% of bone volume. Cancellous bone is less dense and has a porosity ranging from 30-90\% (Carter and Hayes 1977). Cancellous bone forms low weight interior scaffolding, which to helps bone maintain its shape by absorbing compress forces.
1.2 Role of Fatigue in Orthopedic Problems

Cortical bone when subjected to cyclic loading exhibits a reduction of elastic modulus, accumulation of permanent strain for non zero mean stresses, and failure at stresses below the static strength, which are all indicative of fatigue damage. Fatigue damage can contribute to femoral neck fractures (Keyak, Rossi et al. 2001), stress fractures (Valimaki, Alfthan et al. 2005) and failure of implants (Taylor and Tanner 1997).

Uniaxial Fatigue Testing

A servo-hydraulic machine is usually used to perform fatigue tests. The fatigue specimens are clamped in the grips of a testing machine equipped with a water bath chamber maintained at body temperature, 37˚C. The load and displacement are recorded with time, and the load is converted into stress \( \sigma \), by dividing by the initial cross-sectional area. The engineering strain values, \( \varepsilon \) are produced from the movement of the actuator or extensometer. A machine-generated loading waveform is used to cycle specimens at a cycle frequency of 2 Hz between a near zero load level (Zioupos, Wang et al. 1996; Zioupos, Wang et al. 1996). Other examples of uniaxial fatigue testing can be seen in (Carter and Hayes 1977; Pattin, Caler et al. 1996; Lee, O'Brien et al. 2000).

Cycles to Failure

The traditional approach to studying the cycles to failure of bone usually involves the stress-life (\( \sigma-N \)) approach (Carter and Hayes 1977), where specimens are cycled at
various stress levels in order to determine the number of cycles to failure at a specific stress. From this data a $\sigma$-$N_f$ diagram is made, in which the stress is plotted against the number of cycles to failure. When both values are plotted on log scales, the distribution is roughly linear and has been expressed as a power law relationship,

$$N_f = A\sigma^B$$  \hspace{1cm} (1.1)

where $A$ and $B$ are constants from the data (Basquin 1910). Damage has been expressed as the remaining life of the material. In this case the damage, $d$, is thought to increase inversely proportional to the total cycles to failure, $N_f$,

$$d = n / N_f$$  \hspace{1cm} (1.2)

$n$ represents the number of cycles experienced; hence in this case damage represents a fraction of the remaining life.

**Modulus degradation**

Bone, when subjected to a uniaxial tensile load exhibits microscopic losses in continuity leading to microcracks which decrease the longitudinal stiffness. The damage can be interpreted as the ratio of the damaged area, which no longer carries the load to the total load, or as the remaining life of the material (Kachanov 1986). When observing damage as a functional loss of stiffness it can be expressed as,

$$d = 1 - E_i / E_o$$  \hspace{1cm} (1.3)

where, in fatigue studies, $E_i$, is the current secant modulus (Zioupos, Wang et al. 1996). The undamaged modulus, $E_o$, can be defined as the initial tangent modulus of the first load cycle (Zioupos, Currey et al. 1994; Courtney, Hayes et al. 1996).
Loss of Strength

Strength degradation is thought to have more clinical importance as a damage state measure than stiffness, but characterizing strength degradation is difficult because there is no easy way to define an undamaged strength value. It would be necessary to destroy the samples to compare the failure event across a population to evaluate the change in strength (Cowin 2001).

Cracking

It has been shown that during in vitro fatigue testing, fractures of machined bone specimens are the result of a progressive accumulation of micro damage and micro cracks that form ahead of catastrophic macrocracks. This microdamage contributes to the formation of stress fractures and fragility fractures (Schaffler et al. 1995). The resistance of any material to fatigue failure is a function of its resistance to either the initiation or propagation of cracks, or both.

Creep

Bone, like many other engineering materials, exhibits a three stage creep (Figure 1.1) (Carter and Caler 1983). The primary stage is a rapid concave-down curve followed by a slower, constant rate, called the second stage; and then a rapid concave-up curve, called the third stage. The degree of modulus degradation and permanent strain, $\varepsilon_{pn}$, accumulation has been shown to be dependent on the magnitude and sign of the applied stress or strain. It has been shown for a zero to tensile (0- T) cyclic load test that the
permanent strain fits the functional form and magnitude of creep, which can be seen in (Cotton, Zioupos et al. 2003). Figure 1.2 from Cotton, Zioupos et al. (2003) shows the accumulation of percent permanent strain vs. the number of cycles from 53-year-old female fatigue sample with an applied normalized stress of 0.0049 (4900 με). The curve contains the three portions that are typical to the functional form of creep.

Figure 1.1: Basic functional form of creep: (A) Primary stage, rapid concave-down section, (B) Secondary stage, slower almost constant rate, (C) Third Stage, rapid concave-up before failure.
1.3 Finite Element Analysis in Orthopedics

FE models allow the complex geometries and properties of bone to be represented. FE modeling of bone provides a valuable tool for examining behavior. FE models can be created by transforming a CT image into a solid model and then meshing the solid model, and then material properties are applied from the CT image. FE models can also be produced by direct or automatic meshing, where the geometry and material properties are taken directly from the CT image.
Computed Tomography Images

Computed tomography describes a procedure in which x-ray measurements are made at multiple angles around an object and are then used to reconstruct a section, or slice, of the object. Each local volume of interest, the size of which depends on scanner resolution, is called a voxel. The CT numbers measured for a voxel are related to the x-ray attenuation coefficient of the material in that volume. This allows differentiation between materials with different attenuation coefficients. CT images can provide accurate information on bone geometry since the attenuation coefficient of bone tissue is much higher than the one of the surrounding soft tissues which results in contrasting edges. It has also been shown that the numbers in the CT images can be related with the mechanical properties of bone tissues (Carter and Hayes 1977; Taddei, Pancanti et al. 2004). A three-dimensional scan is formed by combining multiple adjacent slices. The CT image is essentially a three-dimensional array in which CT numbers are stored (Goodenough 2000).

Finite Element Meshes Created From Solid Models

Once a solid model has been created from CT data there are various methods for creating FE meshes using various commercial FE software packages. A more detailed description and comparison of three techniques and the software used to perform the meshing can be found in Viceconti, Bellingeri et al. (1998). FE meshes created from solid models produce smooth surface geometries.
Finite Element Meshes Created directly from CT images

Keyak, Meagher et al. 1990 developed a method that converts voxel information obtained from CT images directly into elements. The apparent material density, $\rho$ which is dominated by the mineral content, is calculated through a linear relationship (McBroom, Hayes et al. 1985). Bone mineral content shows up well in CT images because of the high attenuation coefficients (Taddei, Pancanti et al. 2004). The elastic modulus for each voxel is then be calculated by using a power-law relationship such as,

$$E_o = A \rho^B \text{ MPa} \quad (1.4)$$

where $A$ and $B$ are experimentally derived constants. This equation can be rewritten as,

$$\log E_o = \log A + B \log \rho \quad (1.5)$$

the relationship between $E_o$ and $\rho$ is linear when plotted on log-log scale. The regression slope for the data is $B$ and the intercept is $\log A$ (Carter and Hayes 1977).

This method uses a simplified element structure, a regular eight-node brick, to define the model. The geometry is created by stacking the contours of each layer of the CT scan. These brick elements do not accurately model the surface geometry of the bone surface. The accuracy of the bone geometry can be increased by reducing the size of the elements (Keyak, Meagher et al. 1990; Keyak, Fourkas et al. 1993). Figure 1.3 shows a mesh created from a solid model and a mesh by an automatic voxel technique, as described by the above procedure. The meshes in Figure 1.3 were created using the same geometry (Viceconti, Bellingeri et al. 1998).
1.4 Numerical Models

Taylor, Verdonchot et al. (1999) developed a technique to simulate the tensile fatigue behavior of human cortical bone samples. In their study they assumed damage, $d$, to be: (i) a linear function of the life fraction $d = n / N_f$; (ii) zero at the beginning of the analysis rising to a value of 1 at failure. The number of load cycles to failure, $N_f$ at a given stress level was determined using a power law,

$$\log(N_f) = -28.23 \log(\sigma / E_o) - 58.43$$

(1.6)
where $E_0$ is the initial Young’s modulus of the specimen and $\sigma$ is the applied tensile stress. During Fatigue the current modulus, $E_i$ and permanent strain, $\varepsilon_{pn}$ were assumed to be non-linear functions of the damage and were chosen to match experimental fatigue data.

\[ E_i = a(d)^3 + b(d)^2 + c(d) + E_0 \]  
\[ \varepsilon_{pn} = e(d)^f \]

where $a$, $b$, $c$, $e$, and $f$ are material dependent, linear functions of the normalized stress level.

\[ a = 2.346(\sigma / E_o) - 19032 \]  
\[ b = -2.567(\sigma / E_o) + 21737 \]  
\[ c = -1.425(\sigma / E_o) + 4320 \]  
\[ e = 0.931(\sigma / E_o) - 3395 \]  
\[ f = 1.52 \times 10^{-4}(\sigma / E_o) - 0.525 \]

where all fatigue data was taken from Zioupos, Wang et al. (1996).

Cotton, Winwood et al. (2005) relate the cycles to failure $N_f$, damage rate $\Delta d / \Delta n$ and creep rate $\Delta \varepsilon_{cr} / \Delta n$ to the normalized stress range $\sigma / E^*$, where $\sigma$ is the applied stress, during a zero-tension (0-T) fatigue test.

The damage in the $i$th cycle, $d_i$ was defined as the fraction loss in the modulus and was expressed as,

\[ d_i = 1 - E_i / E_o \]

where $E_o$ is the initial modulus. A regression analysis was performed for damage from 10% to 90% of the sample life against the cycle number. During this portion of the life...
the damage tends to be linear and the slope of the regression line represents the damage rate per cycle.

The creep per cycle was calculated using the same method. It was found that the cycles to failure, \( N_f \), creep rate, \( \Delta \varepsilon_{cr} / \Delta n \) and damage rate \( \Delta d / \Delta n \) could be expressed as a function of the normalized stress (Cotton, Winwood et al. 2005),

\[
N_f = A_f \left( \frac{\sigma}{E} \right)^{B_f}, \quad \log A_f = -27.9 \quad B_f = -13.5 \quad (1.15)
\]

\[
\frac{\Delta \varepsilon_{cr}}{\Delta n} = A_{cr} \left( \frac{\sigma}{E} \right)^{B_{cr}}, \quad \log A_{cr} = 33.9 \quad B_{cr} = 17.1 \quad (1.16)
\]

\[
\frac{\Delta d}{\Delta n} = A_d \left( \frac{\sigma}{E} \right)^{B_d}, \quad \log A_d = 35.5 \quad B_d = 17.0 \quad (1.17)
\]

1.5 Simulating Fatigue using FEA

Taylor M., Verdonschot et al. (1999) simulated the tensile fatigue behavior of homogeneous human cortical bone samples by implementing their above numerical model with FE analysis.

At the beginning of each iteration the FE model was loaded and the stresses within each element calculated. The number of remaining cycles to failure, \( n_r \) for each element was calculated using,

\[
n_r = N_f (1 - d) \quad (1.18)
\]

where \( N_f \) is calculated from Equation 1.6. The element with the lowest number of cycles to failure remaining within the model was then determined. To simulate damage and creep an iterative solution was implemented. An iteration can represent 1 to tens of thousands loading cycles. The number of loading cycles, \( \Delta n \), for the current iteration was based on a chosen percentage of the lowest cycles to failure within the model. The
change in damage, $\Delta d$, within each element was calculated and added to the total damage, $d$. If the damage of an element reached a value of 1, the element was removed causing the load to be redistributed to surrounding elements. The modulus loss was calculated using Equation 1.7, and permanent strain using Equations 1.8, 1.12, 1.13. At the start of the next iteration the material properties for each element were updated and this procedure was repeated until failure of the sample occurred.

This approach was used to predict the cycles to failure, modulus degradation and the permanent strain. The corresponding FE models were assumed isotropic, homogeneous and linear elastic with only one quarter of the specimen being modeled due to symmetry. Simulations of eight cortical bone samples were run and compared to the experimental data from Zioupos, Wang, et al. (1996). They found that the results of the FEA simulations were in close agreement with the experimental data (Taylor, Verdonschot et al. 1999).

### 1.6 Stressed Volume

Taylor, D. and Kuiper (2001) proposed the fatigue behavior of bone like many other materials is dependent on the size of the test specimen used. It is known that larger components will tend to fail more easily by fatigue than small components subjected to the same stress levels due to a weakest-link phenomenon: where weak regions exist from pre existing micro cracks. The life of a given specimen is controlled by the properties of the weakest region. This causes shorter lives for larger components since a large stressed volume has a higher probability of containing a weak region. This also causes scatter in
the lives of components of the same size (Taylor and Kuiper 2001). The number of cycles to failure, \( N_f \) is described as,

\[
N_f = A \Delta \sigma^B
\]

(1.19)

Taylor, D. and Kuiper (2001) showed the fatigue strength of bone is affected by the sample volume and that larger bone samples tend to have lower fatigue strengths. The number of cycles to failure for specimens of different volumes can be predicted using,

\[
\frac{\Delta \sigma_o}{\Delta \sigma^*} = \left( \frac{-V_{so}}{V_s + V^*} \right) \ln (0.5)^{\frac{1}{B}} = K_n
\]

(1.20)

where \( V_s \) is the stressed volume and \( V_{so} \) the original specimen volume. \( V^* \) is a constant added to compensate for microstructures or for stressed volumes below 10 mm\(^3\). \( B \) and \( \Delta \sigma^* \) are constants that express the degree of scatter in the data.

1.7 Summary of Research Conducted in this Thesis

FE meshes of human cortical bone fatigue samples were created through an automated voxel meshing technique in order to determine the effects of adding “sizing factors”; \( K_N \), \( K_D \), and \( K_{cr} \) to fatigue simulations. I took the existing simulation from Taylor M., Verdonschot et al. (1999) and added the sizing factors to the life, damage and creep laws, which are empirical equations (Equations 1.15-1.17) from Cotton, Winwood, et al. (2005).

Micro CT images of ten dumbbell fatigue samples were used as source data to derive the geometry and non-homogeneous material properties for each FE mesh. The micro CT images were randomly divided into two groups; “Sample group 1” (SMP1 or
optimization) and “Sample group 2” (SMP2 validation). The automatic meshing technique produced a mesh size of 0.28 mm face length for every cubic element for both sample groups, which would be used for the majority of the analysis performed. Additional mesh sizes of 0.21 and 0.14 mm were produced for the SMP1 group; these meshes will be used to check the effects of mesh size on model properties and the fatigue simulations. The meshes were imported into MSC MARC (MSC software, Santa Ana, CA) to create FE models. Boundary conditions and loads were applied to simulate in vitro fatigue testing conditions.

The SMP1 group was used to determine optimal sizing factors for each FE model in the group using the Nelder-Mead optimization. FEA simulations would then be compared with 1) predicted values of the fatigue behavior computed from Equations 1.15-1.17, 2) FEA simulations of the current stressed volume approach which only includes the $K_N$ and 3) the previous method which does not include any sizing factors. The accuracy of the results were computed using an average percent error between the FEA simulations and the empirically predicted values for the cycles to failure, $N_f$, damage rate, $\Delta d/\Delta n$ and creep rate, $\Delta \epsilon_c/\Delta n$. The SMP2 group was used to evaluate the success of the sizing factors on other models. FEA simulations were run at a different mesh size to examine the effects of different mesh size on our determined values.
Chapter 2: Materials and Methods

In this chapter, the materials and methods used in the finite element simulation of the fatigue behavior—cycles to failure, damage per cycle and creep per cycle are described. The CT images used as data sources are presented in Section 2.1. Section 2.2 describes the processes used to develop the models used in the simulations. The method of simulating the fatigue behavior is shown in Section 2.3. Descriptions of the analyses performed can be found in Section 2.4.

2.1 CT Images

CT images for ten fatigue samples originated from the Department of Oral Growth & Development, Queen Mary, and University of London. The dumbbell shaped cortical bone samples were machined at Cranfield University, Shrivenham, UK from the femur of a 74 year old female cadaver. All were longitudinally oriented; with a cross section of approximately 5.3 mm² and a gauge length of 10 mm. The scans had a resolution of 14.95 μm and the image sizes varied depending on the sample and are shown in Table 2.1.

The CT images were randomly divided into two groups (Table 2.2), “Sample group 1” (SMP1) group and a “Sample group 2” (SMP2) group. The SMP1 group was used to determine the optimal sizing factors implemented in the simulations to increase accuracy of fatigue on each individual model. The SMP2 group was used to check the effectiveness of the sizing factors found on other models. Table 2.2
Table 2.1: Micro CT image data

<table>
<thead>
<tr>
<th>CT name</th>
<th>$x$</th>
<th>$y$</th>
<th>$z$</th>
<th>File size</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pz1tm</td>
<td>380</td>
<td>240</td>
<td>912</td>
<td>83.17</td>
</tr>
<tr>
<td>Pz2tm</td>
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<td>320</td>
<td>912</td>
<td>122.57</td>
</tr>
<tr>
<td>Pz3tm</td>
<td>370</td>
<td>370</td>
<td>912</td>
<td>124.85</td>
</tr>
<tr>
<td>Pz4tm</td>
<td>370</td>
<td>370</td>
<td>912</td>
<td>124.85</td>
</tr>
<tr>
<td>Pz5tm</td>
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<td>400</td>
<td>912</td>
<td>102.14</td>
</tr>
<tr>
<td>Pz6tm</td>
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<td>400</td>
<td>912</td>
<td>87.55</td>
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<td>240</td>
<td>912</td>
<td>96.31</td>
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<tr>
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<td>Pz10tm</td>
<td>420</td>
<td>390</td>
<td>912</td>
<td>149.39</td>
</tr>
</tbody>
</table>

Table 2.2: CT images randomly divided into groups

<table>
<thead>
<tr>
<th>“Sample group 1” (SMP1)</th>
<th>“Sample group 2” (SMP2)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CT name</td>
<td>Model name</td>
</tr>
<tr>
<td>Pz1tm</td>
<td>SMP1-1</td>
</tr>
<tr>
<td>Pz3tm</td>
<td>SMP1-2</td>
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<tr>
<td>Pz5tm</td>
<td>SMP1-3</td>
</tr>
<tr>
<td>Pz6tm</td>
<td>SMP1-4</td>
</tr>
<tr>
<td>Pz8tm</td>
<td>SMP1-5</td>
</tr>
</tbody>
</table>

2.2 Model Development

Finite element models of the human cortical bone fatigue samples were created using the CT images as source information. The process of converting a CT image into a finite element model required: 1) image processing, 2) creating meshes through an automatic voxel technique and 3) applying loads and boundary conditions. Figure 2.1 shows an outline of the modeling process used to create the finite element models.
Image Processing

The first step in creating the meshes was rotating the CT images in MRICro (Chris Rorden, University of South Carolina, Columbia, SC). The rotations were performed to simplify the surface geometry of the models once the meshes were created and imported into MARC. The CT images were rotated to get the longitudinal axis of the CT images vertical; which represents the z-axis in MARC. Rotations were also performed to align
with the front and side of the samples with the x-z and y-z plane in MARC. A 3-D rendering of the desired orientation for the samples is shown in Figure 2.2. Figure 2.3 shows sample xPz3tm before and after rotations have been performed. Once the CT images were rotated the meshes could be created.

Figure 2.2: 3-D rendering of a SMP1-1: Showing desired orientation before automatic meshing to simplify surface geometry.
Automatic Meshing

The FE meshes were constructed using a voxel based automatic meshing technique similar to the procedure described by Keyak et al. (1990). The FE meshes were generated using two programs, VOXSUB and VOXMSH (Laboratorio di Tecnologia Medica of Istituti Ortopedici Rizzoli, Bologna – Italy). The first program VOXSUB takes the CT data and creates a sub-sampled dataset by combing voxels. The user defines input such as the number of voxels that will be combined in x, y and z-direction. For example if the user defined input is set to two in each direction then two
voxels each in the x, y and z-direction will be combined forming a larger “super voxel” now containing eight voxels. This sub-sampling generates a set of super voxels. Each super voxel is then assigned a CT number obtained as an average of the composing voxels CT number.

The user defined input was set to combine 20 voxels in each of the three orthogonal directions giving an element edge length of 0.28 mm which I will call our “normal” mesh for both the SMP1 group and the SMP2 group. This normal mesh would be used in most of the analysis performed. To test the effects of the mesh size on a model’s modulus, surface geometry and fatigue behavior simulation two more meshes were created for the SMP1 group. To create these additional meshes the user defined inputs were set to 15 and 10 creating edge lengths of 0.21 (fine) and 0.14 (super fine) mm respectively.

The second program VOXMSH converts the averaged CT number for each super voxel into the tissue apparent density, $\rho$, using the relationship,

$$\rho = \frac{\text{brightness}}{\kappa} \times 1.037 \text{ g/cm}^3 \quad (2.1)$$

where the brightness ranged from 0-255 and $\kappa$, a scale factor determined from the calibration of the scanner. The scale factors are shown below for each sample in Table 2.3.

Table 2.3: Scale factors for CT images

<table>
<thead>
<tr>
<th>CT Image</th>
<th>xPz1tm</th>
<th>xPz2tm</th>
<th>xPz3tm</th>
<th>xPz4tm</th>
<th>xPz5tm</th>
<th>xPz6tm</th>
<th>xPz7tm</th>
<th>xPz8tm</th>
<th>xPz9tm</th>
<th>xPz10tm</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\kappa$</td>
<td>130</td>
<td>130</td>
<td>120</td>
<td>130</td>
<td>130</td>
<td>120</td>
<td>75</td>
<td>130</td>
<td>130</td>
<td>130</td>
</tr>
</tbody>
</table>
The modulus for each super-voxel was then calculated using an empirical relationship from Carter and Hayes (1977),

\[ E_o = 3790 \rho^3 \text{ MPa} \]  

(2.2)

this relationship was determined from compression data and assumed valid for tensile modulus (Keyak, Meagher et al. 1990). We used this relationship because it placed the modulus values of the models inside the reported values of cortical bone 14-20 GPa (Taylor, O'Reilly et al. 2003).

Once the material properties were assigned the super-voxels were converted into linear, isotropic, regular 8-node finite element bricks. An element and its nodes were created only if the modulus was above 1000 MPa. Those elements whose modulus was lower than 1000 MPa were ignored. The FE mesh was then created by placing nodes in a cube lattice, represented by a three dimensional array in the software. This process was repeated until all the elements and their properties for the model are created and stored in a neutral PATRAN (MSC software, Santa Ana, CA) format.

**Loads and Boundary Conditions**

The PATRAN files of each mesh were imported into MARC where the boundary conditions and loads were applied. Boundary conditions were applied to the bottom of the model to simulate a real test where the sample would be clamped lower than the model; this allowed movement in the transverse planes. The bottom of each model was fixed in place by first setting the nodal displacements to zero in the \( z \)-direction. Then the nodal displacement for the center node on the bottom of the model, point A in Figure 2.4, was
fixed in all directions. Next the $x$-displacements were set to zero for the nodes on line BC in Figure 2.4. Finally the $y$-displacements were set to zero for the nodes on line DE in Figure 2.4. This keeps the model from sliding in the axial plane.

![Diagram of nodal constraints for bottom of finite element models to emulate in vitro fatigue testing](image)

The loading force was applied to each model to approximate an actual test where the sample is loaded above the finite element model. This was achieved by placing a point load at the center node of the top of each sample and then “tying” the remaining nodes on the top center node. “Tying” in MARC applies identical displacements for all tied nodes. In this case the nodes were tied in the $z$-direction and left free to move independently in the $x$ and $y$-directions.
2.3 Simulating Fatigue Behavior

To fatigue the samples, a cyclic load was implemented within MARC where an iteration consisted of three “steps.” Each step represents an increment within MARC. During the first step, “load”, the force or load is applied, the load is then held during the second step, “hold”, allowing creep to occur; finally during the third step, “unload” the load is removed. The loading steps for a cycle are shown in Figure 2.5. In reality steps one and two would occur simultaneous but are dived into two separate parts for computational ease. Each iteration may represent more than one cycle depending on the damage (refer to Taylor M., Verdonschot, et al. 1999).

![Figure 2.5: The three loading steps during an iteration; load, hold and unload](image)

The fatigue behavior was simulated within MARC using a set of user subroutines written in FORTRAN by Taylor M., Verdonschot, et al. (1999) and later updated by
Taylor M., et al. (2002). MARC input files containing the model geometry; material properties, boundary conditions and loading conditions was also used.

The user subroutine `HOOKLW` is called at the beginning of every load step and sets the modulus, $E_i(k)$ for every element, where $i$ represents the current iteration. The index $k$ represents both the integration point and the element number, because I used a reduced integration element with one integration point per element. During the first increment each element’s modulus is set to the initial modulus, $E_o(k)$. These are the moduli that were computed from the CT data and stored in the MARC input file. The damage, $d(k)$, is set to zero because no loading or damage has occurred yet, and we assume no initial damage. The damage during a loading cycle will be computed later in the program and used to compute and update the modulus.

Next the user subroutine `ELEVAR` is called. This user subroutine first finds the absolute maximum value of the three principal stress values, $\sigma_{\text{prin}}(k)$ using the utility routine `PRINCV` within MARC for each element. Once the maximum principle stress is found for each element it is then normalized using the initial modulus,

$$\bar{\sigma}(k) = \frac{\sigma_{\text{prin}}(k)}{E_o(k)}$$

(2.3)

Once the normalized stress is calculated the cycles to failure for each element can be calculated using,

$$N_f(k) = A_f \left( \frac{\bar{\sigma}(k)}{K_N} \right)^{B_f}$$

(2.4)

where $K_N$ is the “life sizing factor” added to account for sizing effects and inhomogeneity and $A_f = \log -27.9$ and $B_f = -13.3$ from Cotton, Winwood et al. (2005). The minimum number of cycles to failure, $\left(N_f\right)_{\text{min}}$ within the whole model is then found based on a
predetermined percentage of the lowest number of cycles. A more detailed explanation can be found in Taylor M., Verdonschot, et al. (1999).

Next the total number cycles run, \( n_i \), and damage, \( d_i \), were calculated using the subroutine \textsc{UPDATVAR}. The number of cycles for the current increment, \( \Delta n \), were calculated based on the minimum number of cycles to failure, where \( \Delta n \) is the maximum of \( (N_f)_{\text{min}} \) or one. Then the total number cycles run updated is using,

\[
n_i = n_{i-1} + \Delta n
\]

The increase in damage for each element was calculated using,

\[
\Delta d(k) = \Delta n \times A_D \left( \frac{\sigma(k)}{K_D} \right)^{B_D}
\]

where \( A_D = \log 35.5 \) and \( B_D = 17.0 \) are from Cotton, Winwood et al. (2005) and \( K_D \) is the “damaging sizing factor.” The damage is updated using,

\[
d_i(k) = d_{i-1}(k) + \Delta d(k)
\]

The creep for each integration point was then calculated with the user subroutine \textsc{CRPLAW}. The change in creep for the iteration, \( \Delta \varepsilon_{cr} \), and the total creep, \( \varepsilon_{cr} \), for each element was found using,

\[
\Delta \varepsilon_{cr}(k) = A_{cr} \left( \frac{\sigma(k)}{K_{cr}} \right)^{B_{cr}} \Delta n
\]

\[
\varepsilon_{cr}(k) = \varepsilon_{cr}(k)_{i-1} + \Delta \varepsilon_{cr}(k)
\]

where \( A_{cr} = \log 33.9 \) and \( B_{cr} = 17.1 \) from Cotton, Winwood et al. (2005) and \( K_{cr} \) is the “creep sizing factor.”
Once the creep was calculated the load was released during the third loading increment. At the beginning of the next iteration before the load was applied, the HOOKLAw was called again to update the modulus and damage for every element, where the new modulus is defined as,

$$E_i(k) = (1 - d_i(k))E_o(k)$$  \hspace{1cm} (2.10)

The damage within each element was checked to see if the damage exceeded a value of 1. If an element had reached or exceeded a value of 1 then the corresponding elements modulus was reduced to 10 MPa. This essentially removes the element from the analysis which caused the stresses to be redistributed to other surrounding elements. The procedure was repeated until the sample failed. A detailed description of the simulation can be found in Taylor M., Verdonschot, et al. (1999).

### 2.4 Analysis Performed

**Computing the Model’s Initial Modulus**

The initial modulus, \(E_o\) for a sample will be different from the individual moduli of the elements. The initial modulus, \(E_o\), represents the modulus for the whole model. The initial modulus, \(E_o\), for all models was computed by applying a 420 N force, \(F\) to the models and running the fatigue simulation through at least one iteration, where an iteration consists of the load, hold and unload step. In an actual fatigue test multiple cycles would be run with a small load so no damage would occur and the initial modulus
would be calculated from this data. Once the load was applied the elastic strain, $\varepsilon$ was found using,

$$
\varepsilon = \frac{(d_z^{\text{load}} - d_z^{\text{unload}})}{g l}
$$

(2.11)

where $d_z^{\text{load}}$ and $d_z^{\text{unload}}$ are displacements averaged from eight nodes at the top of the gage length, $gL$ taken from the end of the load and unload steps respectively. The normal and super fine models had a gauge length of 9.8 mm and the fine models had a gauge length of 9.87 mm. The model’s initial modulus, was the found using Hooke’s Law,

$$
E_o = \frac{F}{A \varepsilon}
$$

(2.12)

where the cross sectional area, $A$ of the sample and was taken from the CT image by averaging the area at slices 256, 456 and 656.

**Calculating Damage and Creep Rates**

The predicted fatigue behavior was computed using the empirical equations (Equations 1.15-1.17) Cotton, Winwood, et al. (2005). The simulated life, $N_f^{\text{sim}}$, damage, damage rate, $(\Delta d/\Delta n)^{\text{sim}}$ and creep rate $(\Delta \varepsilon_{cr}/\Delta n)^{\text{sim}}$ were calculated from the FE analysis. The displacements at the unload step could be used to calculate the permanent strain/creep in the model using,

$$
\varepsilon_{cr} = \frac{(d_z^{\text{unload}})}{g l}
$$

(2.13)

Then the creep per cycle can be found by dividing by the number of total cycles that occurred during the cycle. The damage rate is found by first calculating the elastic strain
for the current cycle, using Equation 2.11. Then the current modulus for the cycle and is calculated using,

\[ E_i = \frac{F}{A\varepsilon} \]  \hspace{1cm} (2.14)

The damage was then calculated using,

\[ d_i = 1 - \frac{E_i}{E_o} \]  \hspace{1cm} (2.15)

The damage rate was then calculated by dividing the damage by the number of cycles run.

**Nelder-Mead Numerical Optimization to find \( K_N, K_D, \) and \( K_{cr} \)**

I needed a direct search method to optimize a set of parameters to minimize a non-linear function. It was necessary to implement this optimization routine, since my results were only numerical and would not provide any derivative information. I decided to optimize the sum of the squares of the relative errors between the predicted (Equations 1.15-1.17) and simulated fatigue behaviors.

The Nelder-Mead optimization is a simplex pattern search used to find a local minimum of a function of several variables using only function values, without any derivative information. For \( n \) variables, the simplex contains \( n+1 \) vertices. For our case the simplex is a tetrahedral. The method compares function values at the vertices of the simplex. Let \( \Phi_k(x_k) \) be the function that is to be minimized, where \( x_k = (K_n, K_D, K_{cr}) \),
The function value at the worst vertex is rejected and replaced with a new vertex. A new simplex is formed and the search is continued. The process generates a sequence of simplexes, for which the function values at the vertices get smaller and smaller. The sizes of the simplexes are reduced and the coordinates of the minimum point is found.

The simulated cycles to failure, \( (N_f)_{\text{sim}} \) simulated damage rate, \( (\Delta d/\Delta n)_{\text{sim}} \) and the simulated creep rate, \( (\Delta \varepsilon_{cr}/\Delta n)_{\text{sim}} \) are calculated from the FE simulations as described above. The predicted cycles to failure, \( (N_f)_{\text{pre}} \) predicted damage rate, \( (\Delta d/\Delta n)_{\text{pre}} \) and the predicted creep rate, \( (\Delta \varepsilon_{cr}/\Delta n)_{\text{pre}} \) are calculated using Equations 1.15-1.17.

The Nelder-Mead algorithm described below uses a set of constants in the calculation of various points throughout the algorithm, they are,

\[
\rho = 1, \quad \chi = 2, \quad \gamma = \frac{1}{2}, \quad \text{and} \quad \sigma = \frac{1}{2}
\]

To start, four vertices for the initial tetrahedral are chosen and the function values are computed and ranked from one to four, \( \Phi(x_1) \leq \Phi(x_2) \leq \Phi(x_3) \leq \Phi(x_4) \) one having the lowest value and four having the worst. Next the reflection point, \( x_r \) is calculated and evaluated,

\[
x_r = (1 + \rho)\bar{x} - x_4
\]

where, \( \bar{x} = \sum_{i=1}^{n} \frac{x_i}{n} \) is the centroid of the best points. If \( \Phi_1 \leq \Phi_r < \Phi_3 \) then accept
reflection point $x$, to replace $x_4$ and move on to the next iteration. If $\Phi_r < \Phi_1$
calculate and evaluate the expansion point $x_e$,

$$
x_e = (1 + \rho \chi)x - \rho \chi x_4
$$

(2.18)

If $\Phi_e < \Phi_r$, accept $x_e$ and stop the iteration, otherwise accept $x_r$. If $\Phi_r \geq \Phi_3$ then a
contraction must be performed. If $\Phi_3 \leq \Phi_r \leq \Phi_4$ then an outside contraction, $x_{co}$ must
be calculated and evaluated. If $\Phi_r \geq \Phi_4$ then perform an inside contraction, $x_{ci}$ where.

$$
x_{co} = (1 + \rho \gamma)x - \rho \gamma x_4
$$

(2.19)

$$
x_{ci} = (1 - \gamma)x + \gamma x_4
$$

(2.20)

If the function value for the corresponding contraction is better than the reflection point,
the contraction is accepted and the next iteration can started. If the function values of the
corresponding contractions are worse than the reflection than a shrink must be performed.
When performing the shrink a new simplex is formed by keeping $x_1$ and replacing the
other vertices with,

$$
v_i = x_1 + \sigma(x_i - x_1)
$$

(2.21)

where $i=2, 3, 4$. Evaluate $\Phi$ at the new vertices of the simplex and order them as before,
$\Phi_1 \leq \Phi_2 \leq \Phi_3 \leq \Phi_4$. Once the ordering is complete the next iteration can be started.
Chapter 3: Results

In this chapter, results from the FE simulations are presented. Section 3.1 describes the models generated and their properties. The optimization of the SMP1 (optimization) group is shown in section 3.2. The FEA Fatigue simulations and comparisons are shown in section 3.3.

3.1 Models Generated

Three models were created for each sample in the SMP1 group. All models were created with 8 node cube elements, where each node had three degrees of freedom. The models for the SMP1 group had element edge lengths of 0.28, 0.21 and 0.14 mm and were named “normal”, “fine” and “super fine” respectively. The number of nodes and elements for each model in the SMP1 (optimization) group are shown below in Table 3.1. Only a normal model for each sample in SMP2 (verification) group was produced. These model properties are shown in table 3.2. The normal, fine and super fine models for SMP1-5 is shown in Figure 3.1.

<table>
<thead>
<tr>
<th>Model</th>
<th>Normal</th>
<th></th>
<th>Fine</th>
<th></th>
<th>Super Fine</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>SMP1-1</td>
<td>4638</td>
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<td>10757</td>
<td>12817</td>
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<td>40029</td>
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<td>10643</td>
<td>12720</td>
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<td>40223</td>
</tr>
<tr>
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<td>5928</td>
<td>11075</td>
<td>13166</td>
<td>36109</td>
<td>40779</td>
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<td>6111</td>
<td>10914</td>
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<td>39905</td>
</tr>
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</table>
Table 3.2: Model properties for SMP2 group used for verification

<table>
<thead>
<tr>
<th>Model</th>
<th>elements</th>
<th>nodes</th>
</tr>
</thead>
<tbody>
<tr>
<td>SMP2-1</td>
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<tr>
<td>SMP2-5</td>
<td>5001</td>
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</tr>
</tbody>
</table>

Figure 3.1: FE models for SMP1-5 showing effects of element sizes on surface geometry: (A) Superfine mesh with element edge length of 0.14 mm, (B) Fine mesh with element edge length of 0.21 mm, (C) Normal mesh with element edge length of 0.28 mm.

To demonstrate mesh convergence, moduli of all models in the SMP1 group are plotted against the number of elements (figure 3.2). It can be seen that the size of the element affects the model’s modulus. The modulus increased as the mesh size decreased. All the moduli lay within the reported 14-20 GPa range for cortical bone (Taylor, O'Reilly et al. 2003). Figure 3.3 shows all the model’s moduli for the SMP2 group.
Figure 3.2: SMP1-normal model’s moduli verse the number of elements in the mesh.

Figure 3.3: Sample modulus values for the SMP2 group models used for verification.

The inhomogeneity of typical samples can be seen in Figure 3.4; the SMP1 group normal mesh models are shown. Histograms of the SMP1 group (Figure 3.5) show the distribution of the elements properties within the SMP1 group. All the models are quite
similar. This is not unexpected since all the samples were taken from the same donor. The inhomogeneity of the model was affected by the mesh size. Differences in material distributions when different meshes were used are shown in Figure 3.6.

Figure 3.4: Non-homogeneous material distribution for SMP1 group.
Figure 3.5: Element property distribution through SMP1 group.

Figure 3.6: Cross section showing material property distribution at height $z = 7.6 \text{ mm}$ for sample SMP1-5; (A) Super fine mesh, (B) Fine Mesh, (C) Normal Mesh
3.2 Nelder-Mead Optimization

The Nelder-Mead optimization was implemented to find the optimal $K_N$, $K_D$, and $K_{cr}$ values for the normal samples in the SMP1 (optimization) group using Equation 2.16, the sum of the squares of the relative error of the fatigue behavior, where we were comparing the predicted (computed from Equations 1.15-1.17) and FEA simulated values. During the optimization and fatigue simulation the samples were loaded to fail around 1000 cycles. The load, predicted cycles to failure, predicted damage per cycle and predicted creep per cycle are shown below in Table 3.3. Figure 3.7 shows $\Phi$ versus the increment of the optimization for the SMP1 group as $\Phi$ converges towards zero. The values found for $K_N$, $K_D$, $K_{cr}$ and $\Phi$ for the normal mesh models in SMP1 group are shown in Table 3.4.

<table>
<thead>
<tr>
<th>Model</th>
<th>Force, N</th>
<th>$N_F$</th>
<th>Damage per cycle</th>
<th>creep per cycle</th>
</tr>
</thead>
<tbody>
<tr>
<td>SMP1-1</td>
<td>420</td>
<td>1107</td>
<td>8.86E-05</td>
<td>1.35E-06</td>
</tr>
<tr>
<td>SMP1-2</td>
<td>420</td>
<td>1068</td>
<td>9.26E-05</td>
<td>1.41E-06</td>
</tr>
<tr>
<td>SMP1-3</td>
<td>385</td>
<td>1021</td>
<td>9.82E-05</td>
<td>1.49E-06</td>
</tr>
<tr>
<td>SMP1-4</td>
<td>445</td>
<td>982</td>
<td>1.03E-04</td>
<td>1.57E-06</td>
</tr>
<tr>
<td>SMP1-5</td>
<td>370</td>
<td>1008</td>
<td>9.98E-05</td>
<td>1.52E-06</td>
</tr>
</tbody>
</table>
When the fatigue simulations were run for the normal mesh models in the SMP1 group without sizing factors, the cycles to failure were greatly under-predicted while the damage per cycle and creep per cycle were over-predicted. An example of this is
shown in Figure 3.8 for SMP1-5. In this FEA simulation the model failed at 100 cycles when it was loaded to fail at around 1000 cycles.

Once the optimized sizing factors were applied to the simulations, the behavior was brought into closer agreement (Figure 3.9). A simulation was also run where only
the life sizing factor $K_N$ was applied while the other sizing factors were not to see if it was necessary to include sizing factors on the damage and creep laws. This represents the current stressed volume approach, proposed for bone by Taylor, D. and Kuiper (2001). The cycles to failure (Figure 3.10), damage per cycle (Figure 3.11) and creep per cycle (Figure 3.12) for each simulation is shown below. The average percent error of the fatigue behavior is shown below (Table 3.5) for the simulations and was computed by taking an average of the sum of the simulated percent errors against the predicted fatigue behavior. The average for the group is also shown to try and display an overall trend. It is apparent that the simulations that included the optimized sizing factors greatly increased the accuracy of the simulations. The current stressed volume approach slightly improves the simulations.
Figure 3.9: Comparison of predicted and simulated results. Predictions computed from Equations 1.16 and 1.17 and FEA simulation are results for SMP1-5 normal mesh using optimized sizing. (A) Creep vs. number of cycles, (B) Damage vs. number of cycles.
Figure 3.10: Cycles to failure comparisons; predicted value from empirical Equation 1.15, FEA simulation using optimized sizing factors $K_N, K_D, K_C$, FEA simulation using only life sizing factor $K_N$, FEA simulation without any sizing factors.

Figure 3.11: Damage per cycle comparisons; predicted value from empirical Equation 1.15, FEA simulation using optimized sizing factors $K_N, K_D, K_C$, FEA simulation using only life sizing factor $K_N$, FEA simulation without any sizing factors.
Figure 3.12: Creep per cycle comparisons; predicted value from empirical Equation 1.15, FEA simulation using optimized sizing factors $K_N, K_D, K_{cr}$, FEA simulation using only life sizing factor $K_N$, FEA simulation without any sizing factors.

Table 3.5: Average percent errors between FEA simulations and predictions for the SMP1 group

<table>
<thead>
<tr>
<th></th>
<th>$K_N, K_D, K_{cr}$</th>
<th>$K_N$ only</th>
<th>No sizing factors</th>
</tr>
</thead>
<tbody>
<tr>
<td>SMP1-1</td>
<td>15.95%</td>
<td>116.73%</td>
<td>70.26%</td>
</tr>
<tr>
<td>SMP1-2</td>
<td>9.14%</td>
<td>61.72%</td>
<td>405.57%</td>
</tr>
<tr>
<td>SMP1-3</td>
<td>5.71%</td>
<td>9.80%</td>
<td>67.98%</td>
</tr>
<tr>
<td>SMP1-4</td>
<td>8.67%</td>
<td>347.58%</td>
<td>132.55%</td>
</tr>
<tr>
<td>SMP1-5</td>
<td>3.53%</td>
<td>14.01%</td>
<td>97.79%</td>
</tr>
<tr>
<td>Average</td>
<td>8.60%</td>
<td>109.97%</td>
<td>154.83%</td>
</tr>
</tbody>
</table>

3.4 Validation of $K_N, K_D, K_{cr}$

To evaluate the success of the sizing factors found for the normal mesh models in the SMP1 group three sets of $K_N, K_D, K_{cr}$ were run for the SMP2 group, a “low”,
“mean” and a “high”. The $K_N$ values were used to choose the low and high values because most fatigue applications are life related. The mean was averaged from all values. The sizing factors, $K_N$, $K_D$ and $K_{cr}$ used for these evaluations are shown below in Table 3.6. A simulation was also run where no sizing factors were implemented to set a base for comparisons. The results from model SMP2-5 were excluded because it would break at the bottom of the sample after one cycle. This is similar to a sample breaking at the grips in a real test. The cycles to failure (Figure 3.13), damage per cycle (Figure 3.14) and creep per cycle (Figure 3.15) for each simulation is shown below. The average percent errors for the simulations are shown in Table 3.7. It can be seen that the high simulation improved the fatigue behavior the best.

<table>
<thead>
<tr>
<th>Case</th>
<th>$K_N$</th>
<th>$K_D$</th>
<th>$K_{cr}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>$K_{low}$</td>
<td>1.074543</td>
<td>0.999085</td>
<td>0.983091</td>
</tr>
<tr>
<td>$K_{mean}$</td>
<td>1.21903</td>
<td>1.037101</td>
<td>1.012522</td>
</tr>
<tr>
<td>$K_{high}$</td>
<td>1.175586</td>
<td>1.08647</td>
<td>1.036767</td>
</tr>
<tr>
<td>No Sizing Factors</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
</tbody>
</table>
Figure 3.13: Comparison of cycles to failure for SMP2 group; predicted cycles to failure calculated from equation 1.15, FEA simulation using "low" sizing factors, FEA simulation using "mean" sizing factors, FEA simulation using "high" sizing factors, FEA simulation using no sizing factors.
Figure 3.14: Comparison of damage per cycle for SMP2 group; predicted cycles to failure calculated from equation 1.15, FEA simulation using “low” sizing factors, FEA simulation using “mean” sizing factors, FEA simulation using “high” sizing factors, FEA simulation using no sizing factors.
Figure 3.15: Comparison of creep per cycle for SMP2 group; predicted cycles to failure calculated from equation 1.15, FEA simulation using “low” sizing factors, FEA simulation using “mean” sizing factors, FEA simulation using “high” sizing factors, FEA simulation using no sizing factors.

Table 3.7: Average percent errors for SMP2 group

<table>
<thead>
<tr>
<th></th>
<th>K_LOW</th>
<th>K_MEAN</th>
<th>K_HIGH</th>
<th>No Sizing Factors</th>
</tr>
</thead>
<tbody>
<tr>
<td>SMP2-1</td>
<td>2548.10%</td>
<td>145.53%</td>
<td>36.78%</td>
<td>2332.19%</td>
</tr>
<tr>
<td>SMP2-2</td>
<td>916.73%</td>
<td>40.73%</td>
<td>30.16%</td>
<td>891.48%</td>
</tr>
<tr>
<td>SMP2-3</td>
<td>116.38%</td>
<td>66.80%</td>
<td>27.92%</td>
<td>120.34%</td>
</tr>
<tr>
<td>SMP2-4</td>
<td>60.89%</td>
<td>15.37%</td>
<td>61.22%</td>
<td>82.05%</td>
</tr>
<tr>
<td>Average</td>
<td>910.52%</td>
<td>67.11%</td>
<td>39.02%</td>
<td>856.52%</td>
</tr>
</tbody>
</table>
Effects of Element Size on Sizing Factors.

To test the effect of mesh size on the sizing factors, model SMP1-5 using the fine mesh was run using the sizing factors optimized from the SMP1-5 normal mesh. Using the sizing factors from the normal mesh on the fine mesh gave an average percent error of 73.69% from the predicted behaviors. An optimization was performed on the fine mesh (figure 3.16) to compare the sizing factors computed, the values are shown in Table 3.8. Running the fine model with sizing factors from the normal model improved the behavior (Figures 3.17, 3.18 and 3.19) but the optimized results for the fine model produced improved averaged behavior by 56% over the sizing factors optimized from normal mesh.

Figure 3.16: Φ vs. optimization iteration number for the SMP1-5 fine mesh model, showing convergence
Table 3.8: Sizing factors and average percent errors for SMP1-5 fine mesh

<table>
<thead>
<tr>
<th>Simulation</th>
<th>$K_N$</th>
<th>$K_D$</th>
<th>$K_{cr}$</th>
<th>Average errors</th>
</tr>
</thead>
<tbody>
<tr>
<td>No Sizing</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>107.22%</td>
</tr>
<tr>
<td>Normal Optimized</td>
<td>1.074543</td>
<td>0.999085</td>
<td>0.983091</td>
<td>73.69%</td>
</tr>
<tr>
<td>Fine Optimized</td>
<td>1.132407</td>
<td>1.056481</td>
<td>1.007407</td>
<td>17.39%</td>
</tr>
</tbody>
</table>

Figure 3.17: Comparison of cycles to failure for the SMP1-5 fine mesh: Predicted cycles to failure from Equation 1.15, FEA simulation using sizing factors optimized from the normal mesh, FEA simulation using sizing factors optimized from the fine mesh, FEA simulation using no sizing factors.
Figure 3.18: Comparison of creep per cycle for the SMP1-5 fine mesh: Predicted cycles to failure from Equation 1.16, FEA simulation using sizing factors optimized from the normal mesh, FEA simulation using sizing factors optimized from the fine mesh, FEA simulation using no sizing factors.

Figure 3.19: Comparison of damage per cycle for the SMP1-5 fine mesh: Predicted cycles to failure from Equation 1.17, FEA simulation using sizing factors optimized from the normal mesh, FEA simulation using sizing factors optimized from the fine mesh, FEA simulation using no sizing factors.
Chapter 4: Discussion

This chapter discusses the outcomes of the fatigue simulations. The significance of automatic meshing is observed in Section 4.1. Element size is discussed in Section 4.2, showing the importance of these properties. The damage accumulation is discussed in section 4.3. Section 4.4 discusses the effectiveness of the sizing factors. Finally, a summary of the important conclusions is given in Section 4.5.

4.1 Automatic meshing

CT images provide an excellent source for model geometry and material properties for bone. The images I used were micro CT images with a resolution of 14.95 µm. The resolutions of normal scanners do not capture this fine of a detail. The automated voxel meshing technique provided an easy method of generating meshes of our samples quickly. The technique has been criticized (Viceconti, Bellingeri et al. 1998) because it does not accurately model curved surface geometries due to the use of regular 8–noded cube elements. This was not a major concern of ours because the surfaces of our samples were mostly flat. Further, because our samples were loaded in uniaxial tension, we do not get the high surface stresses as seen in bending. It was shown in Figure 3.1 that while the element size did affect the surface geometry slightly, this did not effect my simulations as I could see with the slight variation in sample modulus. Alternative techniques of solid modeling followed by automated 3D meshing are time consuming and would have added months to the project to create all the FE meshes used.
The computed model modulus lay within the reported 14-20 GPa range for cortical bone (Taylor, O'Reilly et al. 2003).

4.2 Element size effects

It was shown (Figure 3.2) that as the element size decreased the model stiffness increased. Usually in finite element models the stiffness is over predicted by a coarse mesh. The reason for our results are that the edge elements are softer in the more coarse meshes, because when the super voxels near the edges are being created by averaging in more voxels with lower CT numbers are include and averaged. This can be seen in Figure 4.1; where the blue represents a lower modulus, C represents the normal mesh; B represents the fine mesh, and A shows the super fine mesh. The amount of weaker elements decreases as the mesh size is refined. Furthermore the relationship between voxel brightness and modulus is nonlinear. I saw slight variations in modulus, however due to simulation time constraints, I concentrated on the normal mesh size.
Figure 4.1: Cross section showing material property distribution at $z = 7.6 \text{mm}$ for SMP1-5; (A) Super fine mesh, (B) Fine Mesh, (C) Normal mesh

4.3 Damage in the Finite Element Simulations

During the FEA simulations I noticed that some of the models initially experienced little or zero damage and then the damage would experience a huge increase (Figure 3.8 (B) and 3.9 (B)). In real tests the damage accumulates more gradually. One possible cause for this jump could be the use of the coarse mesh. As one element fails there is a large reduction in the surface area.

The use of clinical scanners would cause the use of coarse meshes due to the fact that they are unable to produce scans with such a fine resolution as I used. I think that this further work must be performed to explain the sudden jumps in the damage seen in some of the simulations.
4.4 Sizing Factors Effectiveness

I was able to show that by applying the sizing factors to the damage laws within the fatigue simulation, I was able to improve the accuracy of the simulated behavior. By using a numerical optimization I was able to find a set of sizing factors for each individual model in the SMP1 optimization group normal mesh. The average percent error for the group was 9%; while the current stressed volume approach yielded an average error of 110%

When the optimized sizing factors were tested for effectiveness on other models in the SMP2 or verification group, the high and mean simulations improved the behaviors, with average percent errors of 39% and 67% respectively. The errors in the SMP2 group were overall higher than in the optimized group but they still showed improved simulated behavior over the previous methods where no sizing factors, or sizing factor applied only to the life law were used.

The optimal sizing factor for the SMP1-5 normal mesh was tested on the SMP1-5 fine mesh model to check the effects of the mesh size on the sizing factors. Applying the normal sizing factors to the SMP1-5 fine mesh model improved the behavior and gave an average percent error of 74%. Running the optimization to find the better sizing factors for the fine mesh gave: $K_N = 1.132407$, $K_D = 1.056481$ and $K_{cr} = 1.007407$. All these values are greater than the normal sizing factors found for SMP1-5. Running the simulation with the new sizing factors gave an average percent error of 17%. This shows that the size of the mesh affects the sizing factors for a model.
4.5 Summary

The novel approach of this thesis applies sizing factors to creep, damage and life laws within fatigue simulations. The automatic voxel meshing technique allowed us to easily produce non-homogeneous FE models using micro CT images as source information for geometry and material properties. I showed that applying these sizing factors to simulations of non-homogeneous FE models improved the accuracy of the simulation. I also showed that the use of all three sizing factors is more effective than just using the life sizing factor of the current stressed volume approach (Taylor and Lee 2003). I used Nelder-Mead optimization to find optimal sizing factor for individual models. I showed that these optimized values could be used on other models, still improving the simulations. Finally, I showed it was more effective to perform a new optimization for different models and mesh sizes.
Chapter 5: References


