A Design Method for FES Bone Health Therapy in SCI

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Abstract

FES assisted activities such as standing, walking, cycling and rowing induce forces within the leg bones and have been proposed to reduce osteoporosis in spinal cord injury (SCI). However, details of the applied mechanical stimulus for osteogenesis is often not reported. Typically, comparisons of bone density results are made after costly and time consuming clinical trials. These studies have produced inconsistent results and are subject to sample size variations. Here we propose a design process that may be used to predict the clinical outcome based on biomechanical simulation and mechano-biology. This method may allow candidate therapies to be optimized and quantitatively compared. To illustrate the approach we have used data obtained from a rower with complete paraplegia using the RowStim (III) system.

Key Words: FES, spinal cord injury, osteoporosis, bone stress, FES rowing

Material and Methods

Here we illustrate the design process using our RowStim(III) system with 4 channels of electrical stimulation (Quads + Hamstrings) as described in [2]. The rowing machine used was based on the fixed stretcher Concept 2 model (D) ergometer. The FES rowing subject RG and co-author was 57yrs, 75 kg male, 1.72m height with a T4 AIS (A) injury sustained 11 years previously. The data was acquired in 2012 and RG started FES rowing in 2003 and has maintained 2-3 30 min sessions per week since then. Following an initial 10 minute warm-up the subject was asked to row as hard as he could for approximately 20 strokes. The trials analysed had an indicated mean power of 60W on the Concept2 display.

The USA Model Spinal Cord Injury System report fracture incidences due to osteoporosis of 14% at 5 years, 28% at 10 years, and 39% at 15 years post injury, based on outpatient studies, and a prevalence of 25-46%. The causes of fractures are often unknown or are associated with relatively low energy trauma and have an associated 78% increase in mortality risk.1 The repetitive pattern of limb loading is generally accepted as a major mechanical stimulus determining BMD as illustrated by the Stanford bone mineral density index (BDI) BDI = (n,β,GRF)m[10] where n is the number of loadings per day, β is a subject-specific scale factor accounting for differences in body weight, GRF is the ground reaction force, m is an empirical exponent that can be thought of as weighting factor for the relative importance of load magnitude and the number of daily loading cycles.2,3 Dudley-Javoroski and Shields (2008) clinically determined that FES ankle plantar-flexion contractions (30 contractions per min with about 8,000 per month for 3yrs) resulted in compressive loads approximately 1.5 times BW were required to attenuate BMD loss in SCI.4 Previous studies indicate modest BMD gains that appear localized to regions of expected higher bone stress.5 However, it is important to know the 3D distribution of bone mineral such as can be obtained clinically using imaging techniques such as pQCT or high resolution pQCT.6 In 1964 Harold Frost proposed a regulatory mechanism for bone adaptation to loading - the mechanostat. More recently, Frost provides evidence for genetically determined stress thresholds for bone reabsorption, maintenance and remodelling of bone.10 Here, we propose a design process for FES bone health therapies that involves calculating the 3D distribution of microstress in the bone of interest – here we use the tibia of a paraplegic subject. A beam model of the tibia, scaled to the subject, will be used. This model will then be used to determine the regions of the tibia where osteogenesis is expected when using a FES bone health therapy – for example, FES rowing.
The handle forces were measured using an in-line strain gauge transducer. The normal foot stretchers were replaced by cantilevers attached to two floor mounted OR6-7 force plates as shown in Fig 1(a). Kinematic and force data was acquired using a Vicon12 MX camera system running Nexus software.

The next step was to develop a beam model scaled to the tibial dimensions of RG. Stress distribution was calculated across the tibial bone model, assuming the tibia was fixed at the knee joint and forces were applied at the distal end of the tibia (i.e. a cantilever). Cross sectional measurements of the tibia were obtained from the literature.\(^6\) Capozza \textit{et al} (2008)\(^6\) calculated, the periosteal perimeter, cortical thickness and second moment of inertia (in the longitudinal, lateral-medial and anterior-posterior axis) from 18 pQCT scans of the entire tibia at intervals equivalent to 5% of the bone’s length. This data was combined with lateral, compressive and bending forces obtained from FES rowing, to calculate the stress along the tibial bone. The perimeter of each pQCT slice was used to calculate the external radius. Using the values for cortical thickness the internal radius was calculated. The area of each section was obtained by assuming a hollow cylinder. To calculate the total stress we used standard moment and stress equations,\(^5\) i.e.

$$\sigma_T = \sigma_{\text{comp}} \pm \sigma_{\text{lat}} \pm \sigma_{\text{bend}}$$

$$\sigma_T = \frac{F_{\text{comp}}}{A} \pm \frac{M_{\text{lat}}r_{\text{out}}}{I_{\text{LM}}} \pm \frac{M_{\text{bend}}r_{\text{out}}}{I_{\text{PA}}}$$

$$M_{\text{lat}} = l * F_{\text{lat}} \quad \text{and} \quad M_{\text{bend}} = l * F_{\text{bend}}$$

Where \(r_{\text{out}}\) is the outer radius, \(I_{\text{LM}}\) and \(I_{\text{PA}}\) are the second moment of inertia in the lateral-medial and posterior-anterior axes respectively and \(l\) is the length of the tibia.
Results and Discussion

We then applied the Frost threshold stress $\text{MESm}_{10}$, i.e. bone’s genetically determined modeling threshold strain range, in and above which modeling usually turns on to strengthen a bone. Where $\text{MESm}_{10}$, 1000–1500 microstrain; $\sim 20$ MPa, or $\sim 2$ kg/mm$^2$. Applying the threshold at 20MPa can be seen in Figure 3. The fringing occurring at the edge of the threshold is an artefact due to the graphical colour interpolation and has no physical significance.

The biomechanical calculation of joint contact forces, as shown in figure 1(b), exceed the 1.5 times body weight threshold proposed in Dudley-Javoroski S, Shields RK (2008). However, these plots not indicate how the loading is distributed throughout the bones such that regions of osteogenesis can be predicted. The 3D stress analysis allows detailed visualization of the stress applied to the tibia. Figure 2 is a typical single frame taken from a movie at the point of peak force and viewed from one perspective. Matlab software was written to enable 3D rotation to facilitate visual analysis. In this example, it is revealed that in no regions are unsafe levels of stress applied and that the Frost $\text{MESm}_{10}$ threshold is exceeded in an extensive region of the tibia where one might expect osteogenesis. It also reveals regions that are under stressed such as the distal region and the posterior plateau. NB: here the cross sectional tibial measurements were obtained from healthy subjects. To the author’s knowledge there are no corresponding data available for the SCI tibia. It is therefore possible that some of these measurements such as the thickness of the cortical bone and moment of inertia, would change in SCI subjects – this is an area where further radiographic studies are required to determine a model SCI tibia. The analysis was based simple beam theory using a point load applied to the distal tibia constrained as a cantilever so deflections, forces and boundary conditions are approximations.

The actual stress patterns illustrated can only be regarded as approximations but should be indicative of gross distributions and trends. However, at this stage the analysis presented here serves mainly to illustrate the proposed design technique. This provides a quantitative tool to generate hypotheses and further optimize the therapeutic application. For example, a number of experimental parameters could be manipulated. For example, in addition to the quadriceps, during the drive phase, it was expected that stimulating the ankle plantar flexors, in phase with the quadriceps during drive would extend distally the region that is above the $\text{MESm}_{10}$ threshold 20MPa. However, in practice, figure 4 illustrates that this was not the case, with our beam model in which the ankle joint was simplified, however, the above threshold region around the condyle was extended, indicating that stresses in the distal tibia are similar the 4-Ch FES case and a little more widely distributed in the proximal region. Other parameters could then be experimentally explored in a stepwise design and development process. For example, electrically stimulating the hip extensors or changing the rowing style (e.g. the coordination between handle force and seat position during the drive phase) or the type of rowing ergometer used, e.g. floating versus fixed stretcher types. At each step in the design process the above stress analysis is performed and compared with alternatives to check progress towards a solution. Thus relatively few subjects would be involved in finding a solution that provides the best distribution of stress. This would support the follow on clinical trials, involving larger groups of subjects, to validate the predictions. The personalized beam model may be extended to other bones that are compromised by osteoporosis. The beam model is computationally very efficient providing results in the order seconds on a Windows i7 PC. The present beam model may
underestimate the ankle region, due to the tibia, ankle joint and foot being represented by a single cantilever - this we plan to improve. The stress distribution modelling may be further extended, with greatly increased computation time, by using a finite-element model derived from high resolution pQCT scans. The stress distribution model may be improved by adjusting thresholds that better predict actual clinical scan data and further extended by including a porosity stage that may provide a deeper understanding of the basic mechano-transduction, poroelastic fluid low and cellular signaling.9 The proposed technique may also be extended using simulated force data derived from biomechanical models of the therapeutic activity – in this case it may even be possible to design and evaluate a candidate FES bone therapy in-silico prior to clinical trials. This is illustrated in Figure 5 where the bi-lateral force actions of the gluteus maximus muscles were biomechanically simulated and added to the quadriceps actions during the drive phase. This produced a slightly wider distribution of stresses above 20 Mpa. These modest changes due to additional extensors. To demonstrate a more dramatic change in distribution we simulate a floating stretcher type rower by putting the acceleration of the center of mass of the upper body to zero (Figure 6).7 This produced a much reduced distribution of supra threshold stress.

In conclusion, the approach appears to offer a quantitative design tool and method to assess potential bone therapy benefits for FES assisted exercise activity.

Author’s Contribution
All authors contributed equally.

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Conflict of Interest
The authors declare no conflict of interests.

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