1	Stochastic analysis of a heterogeneous micro-finite element
2	model of mouse tibia
3	Yongtao Lu ^{1,2} , Di Zuo ² , Junyan Li ³ and Yiqian He ^{1,2,*}
4 5	¹ State Key Laboratory of Structural Analysis for Industrial Equipment, Dalian University of Technology, Dalian, China
6 7	² Department of Engineering Mechanics, Dalian University of Technology, Dalian, China
8 9	³ Department of Design Engineering and Mathematics, School of Science and Technology, Middlesex University, London, UK
10	
11 12	
13	CORRESPONDING AUTHOR:
14	Yiqian He, PhD
15	State Key Laboratory of Structural Analysis for Industrial Equipment
16	Department of Engineering Mechanics
17	Dalian University of Technology
18	Dalian 116024, China
19	Phone: +86 13998576316
20	Email: <u>heyiqian@dlut.edu.cn</u>
21	
22	Number of words (Introduction to Discussion): 3033
23	Number of tables: 0
24	Number of Figures: 5

26 Abstract:

Finite element (FE) analysis can be used to predict bone mechanical environments that can be used for many important applications, such as the understanding of bone mechano-regulation mechanisms. However, when defining the FE models, uncertainty in bone material properties may lead to marked variations in the predicted mechanical environment. The aim of this study is to investigate the influence of uncertainty in bone material property on the mechanical environment of bone.

A heterogeneous FE model of a mouse tibia was created from micro computed tomography images. Axial compression loading was applied, and all possible bone density-modulus relationships were considered through stochastic analysis. The 1st and 3rd principal strains (ε_1 and ε_3) and the strain energy density (SED) were quantified in the tibial volume of interest (VOI).

The bounds of ε_1 , ε_3 , and SED were determined by the bounds of the density-modulus relationship; the bone mechanical environment (ε_1 , ε_3 , and SED) and the bone density-modulus relationship exhibit the same trend of change; the relative percentage differences caused by bone material uncertainty are up to 28%, 28%, and 21% for ε_1 , ε_3 , and SED, respectively. These data provide guidelines on the adoption of bone density-modulus relationship in heterogeneous FE models.

44

45 Keywords: bone mechanics, material uncertainty, density-modulus relationship,
46 stochastic analysis

47 **1. Introduction**

48 Micro computed tomography (μ CT) imaging has become an important tool to 49 reveal the detailed internal structure of bone, both ex vivo and in vivo [1, 2]. µCT 50 images can be used to generate micro-finite element (uFE) models, in which the 51 element size is on the order of micrometers, to investigate the mechanical behavior of 52 bone, the mechanism of bone mechano-regulation, and the strength of bone after 53 medical intervention [3-8]. Because the homogeneous μ FE models lack realistic 54 spatial variations in bone properties and exhibit limited accuracy, heterogeneous µFE 55 models with heterogeneous material properties have been adopted widely in previous 56 studies [9-11].

57 To generate heterogeneous μFE models of the bone from μCT images, the raw 58 CT attenuation values must first be related to the bone mineral density (BMD) and 59 subsequently converted to bone material properties using bone density-modulus 60 relationship. The relationship between CT attenuation values and BMD values can be 61 established by first scanning the calibration phantom that contains several rods with 62 known BMD values provided by the manufacturer, and subsequently fitting a linear 63 line to the scatter data of the CT attenuation and BMD values of the rods [12]. The 64 bone density-modulus relationship is typically obtained by first performing a 65 mechanical testing on the bone samples at the organ-level and subsequently relating 66 these bone mechanical properties to the various bone densities (apparent density, ash 67 density, etc.) [13]. Because the mechanical properties obtained from mechanical 68 testing are apparent values at the organ-level and the microstructures of bone varies 69 significantly among samples, large variations occur in the bone density-modulus 70 relationship [9-11, 14]. It is still unclear how these variations affect the predictions of

71 heterogeneous µFE models.

72 The detailed mechanical environment could provide important information for 73 the full-field validation of bone FE models [15] and for understanding the mechanical 74 signals driving bone adaptations [16]. For an example, it has been found that bone 75 sites with high strain energy density (SED) exhibit more activities of bone formation; 76 on the contrary, low SED leads to bone resorption [16, 17]. However, the uncertainties 77 in bone density-modulus relationships may affect the mechanical environment 78 predicted from FE models. To account for these uncertainties, many stochastic 79 analyses have been performed to assess the effect of variability in material property 80 on the mechanical environment predicted by FE models [18-21]. However, these 81 studies focused on either the peak values of the mechanical properties (maximal 82 principal strains, maximal principal stresses, etc.) or the apparent behavior (fracture 83 force, hardness, etc.) of the bone samples. The influence of variability in material 84 properties on the detailed mechanical environment (i.e., the distribution of the 1st and 3rd principal strains and the strain energy density) across the entire bone spatial space 85 86 is still unknown.

87 The aim of this study is to investigate the influence of uncertainty in bone
88 material property on the detailed mechanical environment of the bone using
89 heterogeneous μFE models and stochastic analysis.

90

91 **2.** Materials and methods

92 2.1 μCT image of mouse tibia and image processing

93 One entire right tibia dissected from a 12-week-old female C57Bl/6 mouse was
94 imaged using the *ex vivo* μCT imaging system (SkyScan 1172, Bruker, Belgium) with

95 the following setting: a voltage of 49 kV, a tube current of 179 µA, an exposure time 96 of 1180 ms, and an isotropic image voxel size of 4.3 µm. In preparation for generating 97 the FE models, the image datasets were processed based on the standard procedure 98 developed previously [22]. In brief, the tibia was placed back to its anatomic position, 99 i.e., its long (proximal-distal) axis was aligned along the z-axis approximately, and the 100 y-z plane passed through the central line of the articular surfaces of the medial and 101 lateral condyles (Fig. 1a and b). This step was to facilitate the application of the 102 compressive loading along the long axis of the mouse tibia. The image dataset was 103 subsequently transformed into the new position and resampled using the Lanczos 104 kernel, which is a low-pass filter and considered to be the "best compromise" among 105 several simple filters [23].

106 **2.2** Generation of heterogeneous finite element models of mouse tibia

107 The heterogeneous µFE model of mouse tibia was generated from the 108 transformed µCT images (Fig. 1b and d). In brief, the grayscale image dataset was 109 first smoothed with a Gaussian filter (convolution kernel [3 3 3], standard deviation = 110 0.65) and subsequently binarized into bone and background using a single threshold 111 value, i.e., 25.5% of maximal grayscale value (approximately 420 mg HA/cm³). 112 However, the tibia cannot be segmented completely using only one threshold value, 113 because the images includes other bones, such as the femur. Therefore, the tibia and 114 fibula were further segmented from other bones manually (Amira 5.4.3, FEI 115 Visualization Sciences Group, France). The tibial-fibula joint and the region of tibial 116 proximal growth plate were manually filled to allow for load transmission. From the 117 binarized tibia-fibula images, the µFE model with the element number of 1,944,774 118 was created by converting each bone voxel into an eight-node hexahedral element

119 mesh with the element type SOLID185 using an in-house developed Matlab code 120 (Matlab 2015a, The Mathworks, Inc. USA). The boundary condition was based on the 121 experimental setup used for the *in vivo* loading of the mouse tibia [7], i.e., all the 122 nodes on the concave articular surface of the distal tibia were coupled to a distal 123 reference point (RP), which is constrained in all degrees of freedom; the FE nodes at 124 the tibial plateau surface were coupled rigidly to a proximal RP, on which a load of 125 -11 N was applied [7] (Fig. 1c). Poisson's ratio for all the materials was set to 0.3. 126 The uncertainty of the bone's Young's modulus (E) was considered by selecting the 127 bone density-modulus relationship stochastically, which was an input of the FE 128 models (Fig. 1c and d). The details of this step are described as below.

129 **2.3** Stochastic selection of the bone density-modulus relationship

130 The uncertainty in Young's modulus (E) of the μ FE bone models was treated 131 through a stochastic analysis. First, after matching the anatomic sites, six 132 density-modulus relationships of the femur and tibia, which were typically adopted in 133 the literature [13, 14, 24], were reviewed and plotted (Fig. 2a). Here, the data on other 134 anatomic sites, such as the vertebra, were excluded, because these bones have 135 markedly different structures compared to the tibia and femur.

Subsequently, exponential density-modulus relationships were fitted to the mechanical testing data of the bone samples by adjusting the constants "a" and "b" in the exponential function (**Equation 1**). Because the μ FE model also included hollow structures of the bone (such as the tibia–fibula joint and the growth plate), which were generated by the manual filling of these regions in the image processing step, a lower threshold value of bone ash density of 0.4 g/cm³ was adopted in the density-modulus relationship to avoid unrealistically low moduli in the μ FE model. The modulus for

the elements with bone ash density less than 0.4 g/cm³ was set to 0.0104 MPa [13]. Meanwhile, some image voxels may have superficially high grayscale values owing to image noise, which lead to unrealistically high bone densities. Therefore, an upper threshold value of 1.2 g/cm³ was defined in the density-modulus relationship [13]. In summary, the exponential density-modulus relationship used in this study was formulated as below:

149
$$E = \begin{cases} 0.0104 & \rho_{ash} < 0.4 \\ a \times \rho_{ash}^{b} & 0.4 \le \rho_{ash} \le 1.2 \\ a \times 1.2^{b} & \rho_{ash} > 1.2 \end{cases}$$
(1)

150 where "a" and "b" are the two constants, *E* is Young's modulus (GPa), and ρ_{ash} is 151 the bone ash density (g/cm³). It is noteworthy that based on the conversion between 152 bone apparent and ash densities [14], the relationship between bone apparent density 153 and bone modulus can also be established and used for the investigations.

154 All possible bone moduli in the heterogeneous μFE models were considered by 155 adjusting the two constants ("a" and "b" in Equation 1) within the range covered by 156 the various bone density-modulus relationships reviewed (Figs. 2a and 2b). This was 157 implemented and realized in two steps: first, the intervals of "a" and "b" were 158 determined by initially selecting a relatively large interval and subsequently refined 159 by optimizing the two constants by the simplex method (Fig. 2b) [25]; next, in the 160 intervals calculated, "a" and "b" were selected stochastically based on the 161 transformation method [26], which has been proven to reduce the computation cost 162 effectively. It was found that when "a" changed from 10.22 to 12.07 and "b" changed 163 from 1.18 to 2.24, the exponential density-modulus function (Equation 1) covered the 164 full uncertain interval of the bone density-modulus relationships reviewed (Fig. 2b).

166 **2.4 Calibration of the bone modulus in the finite element models**

167 For each stochastic selection of "a" and "b," the calcium hydroxyapatite 168 (HA)-equivalent BMD was calculated at each µCT image voxel using the relationship 169 established through scanning the calibration phantom. In the present study, the 170 phantom with rod densities of 0.0 HA mg/cm³, 250.0 HA mg/cm³, and 750.0 HA mg/cm³ was used. The phantom was scanned using the same setting as used for 171 172 scanning the tibia. By calculating the image grayscale values at each rod of the 173 phantom, the relationship of $\rho_{HA} = 0.0059 \times Igray + 0.242$ (ρ_{HA} is the 174 HA-equivalent BMD, of units HA g/cm³; *Igray* is the image grayscale value) was 175 established to convert the image grayscale values to HA-equivalent BMD values. 176 After matching the phantom type and anatomic site, the density-conversion 177 relationship of $\rho_{ash} = 0.877 \times \rho_{HA} + 0.079$ was chosen to convert the 178 HA-equivalent BMD to bone ash density [14]. However, it is noteworthy that 179 variability exists in this conversion and its influence on the bone mechanical 180 environment requires further investigations. The modulus for each bone image voxel 181 was calculated using Equation (1) and subsequently mapped to the FE mesh using a 182 Matlab code developed in-house.

183 2.5 Finite element analysis and post-processing

Based on the stochastic selection algorithm of the transformation method, 11 values were selected for both "a" and "b" in their intervals, thus resulting in 121 bone density-modulus relationships and 121 FE models. The FE models were solved using ANSYS (Release 14.0.3, ANSYS, Inc.) on a workstation (Intel Xeon E-5-2670. 2.60 GHz, 256 GB RAM) using the formulation of a linear elastic constitutive model.

189 To investigate the influence of uncertainty in bone material property on the

190 mechanical environment of the mouse tibia, a volume of interest (VOI) was selected 191 in the FE models. The VOI started from the end of the proximal growth plate and 192 encompassed 80% of the tibial length (L), which was measured as the distance from 193 the most proximal pixel of the mouse tibia until the most distal pixel of the mouse 194 tibia, and is 17.82 mm for the tibia analyzed in the present study (Fig. 1d). To 195 quantify the results in the three-dimensional (3D) bone spatial space, the VOI was 196 partitioned into 20 compartments of equal length in the z-direction (Fig. 1e). Further, 197 the normalized length of VOI was defined, with the value of zero at the distal end of 198 the VOI. The 1st principal strain (ϵ_1), 3rd principal strain (ϵ_3), and SED were selected 199 as the parameters to describe the mechanical environment of the mouse tibia (Fig. 1e), 200 because ε_1 is likely linked to the bone opening fracture, ε_3 is the compressive strain 201 reflecting the primary loading scenario in the bone, and SED is highly correlated with 202 bone adaptations. The averaged values of ε_1 , ε_3 , and SED in the 20 compartments 203 were calculated and plotted against the normalized VOI length. The post-processing 204 of data in this manner is based on the previous findings where the mechanical values 205 are not reproducible at the image voxel level, but are reliable over a larger VOI [22]. 206 It was found that the bounds of ε_1 , ε_3 , and SED were determined by the upper and lower bounds of "a" and "b" and all different selections of "a" and "b" shared the 207 208 same upper and lower bounds. Therefore, to determine the bounds of ε_1 , ε_3 , and SED, 209 no further refinements on the selection of "a" and "b" were required.

210

```
211 3. Results
```

The occurrence frequencies corresponding to the softest and hardest bone material models are shown in **Fig. 3**. It was found that a lower bone stiffness led to

214 higher ε_1 , ε_3 , and SED. Further, 86% of the nodes in the hardest bone model exhibit 215 an ε_1 that is higher than 250 µ ε (a = 12.07, b = 2.24), compared to 89% of the nodes 216 in the softest bone model (a = 10.22, b = 2.24). Meanwhile, 83% of the nodes in the 217 hardest bone model exhibit an ε_3 lower than -250 µ ε (a = 10.07, b = 2.24), compared 218 to 89% of the nodes in the softest bone model (a = 10.22, b = 1.18). In summary, if 219 the softest bone model was used instead of the hardest bone model, 3% (= 89% - 86%) 220 occurrence of ε_1 were shifted above 250 µ ε and 5% (= 89% - 83%) occurrence of ε_3 221 were shifted below -250 με.

222 The material uncertainty-induced bounds of ε_1 , ε_3 , and SED across the tibial VOI 223 are presented in Fig. 4. It was found that when the bone density-modulus relationship 224 was changed in the FE models, the ε_1 , ε_3 , and SED across the tibial VOI exhibited the 225 same trend of change (Fig. 4). A lower bone stiffness (soft bone) led to an increased ε_1 , 226 an increased SED and a decreased ε_3 . It is noteworthy that the bounds of ε_1 and ε_3 227 were determined by different bone density-modulus relationships, i.e., the bounds of ε_1 were determined by $E = 10.22 \times \rho_{ash}^{2.24}$ and $E = 12.07 \times \rho_{ash}^{1.18}$, and the 228 bounds of ε_3 were determined by $E = 10.22 \times \rho_{ash}^{1.18}$ and $E = 12.07 \times \rho_{ash}^{2.24}$. 229

The relative percentage differences (defined as the difference between the maximal and minimal values divided by the minimal value) of these mechanical parameters across the tibial VOI are shown in **Fig. 5**. The relative percentage differences of ε_1 , ε_3 , and SED ranged from 8% to 28%, from 20% to 28%, and from 14% to 21%, respectively (**Fig.5**).

235

4. Discussion

237 The purpose of this study is to evaluate the effect of uncertainty in bone material

property on the mechanical environment of the bone using heterogeneous FE models and stochastic analysis. This study aims to provide guidelines on the adoption of bone density-modulus relationship in heterogeneous FE models.

241 Two major findings were revealed from this study. First, we found that if the 242 softest bone model was used instead of the hardest bone model, 3% occurrence of ε_1 243 were shifted above 250 $\mu\epsilon$, and 5% occurrence of ϵ_3 were shifted below -250 $\mu\epsilon$. This 244 affects the study of the bone mechano-regulation mechanism, which was first 245 proposed by Wolff and Frost [27, 28]. In particular, Frost's mechano-regulation theory 246 suggests that the local bone mass increases when the strain is above a certain upper 247 strain threshold, and decreases when the strain is below a certain lower strain 248 threshold [27]. Furthermore, it has been postulated that the local bone mass is not 249 responsive of the strain when it is within the interval encompassed by these lower and 250 upper thresholds, i.e., the "lazy zone" [29]. If -250 µE and 250 µE were set as the 251 lower and upper bounds of the "lazy zone" [30] respectively, this study implies that 252 approximately 8% (= 3% + 5%) of the bone voxels will become inactive in the bone 253 adaptation process if the hardest bone model, instead of the softest model, was used in 254 the heterogeneous FE models. Therefore, the uncertainty in bone material property 255 affects the quantification of mechanical stimulation signals of the bone, and is crucial 256 in the study of the bone mechano-regulation mechanism. Next, we found that owing 257 to the uncertainty in bone material property, the mechanical environment across the 258 mouse tibial VOI was changed by up to 28%, 28%, and 21% for ε_1 , ε_3 and SED, 259 respectively, thereby indicating the importance of assigning the appropriate bone 260 properties in studies such as the FE validation study. We also found that the bone 261 mechanical environment (ε_1 , ε_3 , and SED) and the bone density-modulus relationship

262 exhibited the same change trend. Therefore, using the bone density-modulus
263 relationship consistently for defining the bone property could be a feasible strategy in
264 parametric studies, such as evaluating the effect of medicine intervention on the bone
265 mechanical behavior [10].

266 It is noteworthy that in the present study, the magnitude of the load applied is 11 267 N [7] to engender 1200 µε at the medial midshaft of the tibia [31], and thus elicit an 268 osteogenic response in the mouse tibia [32]. Next, the bone density-modulus 269 relationships available in the literature are subject-specific and site-specific [33] 270 because the bone density-modulus relationships are derived from the mechanical 271 testing of organ-level specimens (e.g., vertebra) in previous studies [34, 35]. Hence, a 272 universal deterministic bone density-modulus relationship is required that poses a 273 significant challenge for future research. The universal relationship might be achieved 274 by the investigations at the bone tissue (microstructural) level. Once the tissue-level 275 relationship is developed, the accuracy of heterogeneous µFE models will be 276 increased significantly, because the mapping from bone density to modulus is defined 277 at the bone-tissue level in the heterogeneous µFE models. Furthermore, in the present 278 study, ε_1 , ε_3 , and SED were selected to describe the bone mechanical environment, 279 because ε_1 is likely to be linked to the mode I (opening) bone fracture, ε_3 is the 280 compressive strain reflecting the loading scenario performed in this study, and SED is 281 the resultant bone parameter containing information of both strain and stress that is 282 highly correlated with bone adaptations [20]. Additionally, a limitation in the present 283 study is that the FE analysis was only performed under the loading of axial 284 compression. Other complex loading scenarios, such as the three-point bending, are 285 not investigated. However, because axial compression was used widely in previous

286 preclinical studies of bone adaptations [7, 10], the results from this study can be 287 referred easily for a comparison.

288 In summary, uncertainty in the bone material property exhibited a marked effect 289 on the mechanical environment of the bone, thus implying that the bone 290 density-modulus relationship should be assigned appropriately in studies such as the 291 investigation of the bone mechano-regulation mechanism and FE validation. However, 292 the change trend in the bone mechanical environment is consistent with that of the 293 density-modulus relationship, thus suggesting that assigning bone bone 294 density-modulus relationships in the FE models consistently could be feasible for 295 parametric studies. This study provides guidelines on the adoption of the bone 296 density-modulus relationship in heterogeneous FE models.

297

298 **Conflict of interest**

299 The authors declare that there is no conflict of interest.

300

301 Funding

This study was supported by the National Natural Science Foundation of China (grant numbers 11702057, 11572077), the Chinese Fundamental Research Funds for the Central Universities (grant numbers DUT18LK19, DUT17LK11), the Open Fund from the State Key Laboratory of Structural Analysis for Industrial Equipment (grant number GZ1611), and the Liaoning Provincial Natural Science Foundation of China (grant number 2015020141). The raw μ CT used in the present study can be obtained upon the request to the corresponding author.

310 **References**

- Bouxsein ML, Boyd SK, Christiansen BA, Guldberg RE, Jepsen KJ, Mueller R.
 Guidelines for assessment of bone microstructure in rodents using
 micro-computed tomography. J Bone Miner Res 2010; 25(7): 1468-86.
- Lukas C, Ruffoni D, Lambers FM, Schulte FA, Kuhn G, Kollmannsberger P,
 Weinkamer R, Mueller R. Mineralization kinetics in murine trabecular bone
 quantified by time-lapsed in vivo micro-computed tomography. Bone 2013; 56:
 55-60.
- 318 3. de Bakker CM, Altman AR, Tseng WJ, Tribble MB, Li C, Chandra A, Qin L, Liu
 319 XS. μCT-based in vivo dynamic bone histomorphometry allows 3D evaluation of
 320 the early responses of bone resorption and formation to PTH and alendronate
 321 combination therapy. Bone 2015; 73: 198-207.
- 4. Lambers FM, Kuhn G, Weiqt C, Koch KM, Schulte FA, Mueller R. Bone
 adaptation to cyclic loading in murine caudal vertebrae is maintained with age
 and directly correlated to the local micromechanical environment. J Biomech
 2015; 48: 1179-87.
- 5. Lu Y, Boudiffa M, Dall'Ara, Liu Y, Bellantuono I, Viceconti M. Longitudinal
 effects of parathyroid hormone treatment on morphological, densitometric and
 mechanical properties of mouse tibia. J Mech Behav Biomed Mater 2017; 75:
 244-51.
- Brodt MD, Silva MJ. Experimental and finite element analysis of
 strains induced by axial tibial compression in young-adult and old female
 C57Bl/6 mice. J Biomech 2014; 47: 451-7.

- 333 7. Razi H, Birkhold AI, Zaslansky P, Weinkamer R, Duda GN, Willie BM, Checa S.
- 334 Skeletal maturity leads to a reduction in the strain magnitudes induced within the
 335 bone: a murine tibia study. Acta Biomater 2015; 13: 301-10.
- 336 8. Willie BM, Birkhold AI, Razi H, Thiele T, Aido M, Kruck B, Schill A, Checa S,
- Main RP, Duda GN. Diminished response to in vivo mechanical loading in
 trabecular and not cortical bone in adulthood of female C57Bl/6 mice coincides
 with a reduction in deformation to load. Bone 2013; 55: 335-46.
- 340 9. Easley SK, Jekir MG, Burghardt AJ, Li M, Keaveny TM. Contribution of the
 intra-specimen variations in tissue mineralization to PTH- and raloxifene-induced
 changes in hardness of rat vertebrae. Bone 2009; 46: 1162-9.
- Wang N, Rumney RM, Yang L, Robaye B, Boeynaems JM, Skerry TM, Gartland
 A. The P2Y13 receptor regulates extracellular ATP metabolism and the
 osteogenic response to mechanical loading. J Bone Miner Res 2012; 28(6):
 1446-56.
- 347 11. Yang H, Butz KD, Duffy D, Niebur GL, Nauman EA, Main RP. Characterization
 348 of cancellous and cortical bone strain in the in vivo mouse tibial loading model
 349 using microCT-based finite element analysis. Bone 2014; 66: 131-9.
- Lu Y, Engelke K, Glueer CC, Morlock MM, Huber G. The effect of in situ/in
 vitro three-dimensional quantitative computed tomography image voxel size on
 the finite element model of human vertebral cancellous bone.Proc Inst Mech Eng
 H 2014; 228(11): 1208-13.
- 354 13. Helgason B, Perilli E, Schileo E, Taddei F, Brynjofsson S, Viceconti M.
 355 Mathematical relationships between bone density and mechanical properties: A
 356 literature review. Clin Biomech 2008; 23(2): 135-46.

- 357 14. Knowles NK, Reeves JM, Ferreira LM. Quantitative computed tomography
 358 derived bone mineral density in finite element studies: a review of the literature. J
 359 Exp Orthoped 2016; 3: 36-52.
- 360 15. Costa MC, Tozzi G, Cristofolini L, Danesi V, Viceconti M, Dall'Ara E. Micro
 361 finite element models of the vertebral body: validation of local displacement
- 362 predictions. PLoS One 2017; 12(7): e0180151
- 363 16. Levchuk A, Zwahlen A, Weiqt C, Lambers FM, Badilatti SD, Schulte FA, Kuhn
 364 G, Mueller R. Large scale simulations of trabecular bone adaptation to loading
 365 and treatment. Clin Biomech. 2014; 29: 355-62.
- 366 17. Webster D, Wirth A, van Lenthe GH, Mueller R. Experimental and finite element
 analysis of the mouse caudal vertebrae loading model: prediction of cortical and
 trabecular bone adaptation. Biomech Model Mechanobiol 2012; 11: 221-30.
- 369 18. Berthaume MA, Dechow PC, Iriarte-Diaz J, Ross CF, Strait DS, Wang Q, Grosse
- 370 IR. Probabilistic finite element analysis of a craniofacial finite element model. J
 371 Theor Biol 2012; 300: 242-53.
- 372 19. Laz PJ, Stowe JQ, Baldwin MA, Petrella AJ, Rullkoetter PJ. Incorporating
 373 uncertainty in mechanical properties for finite element-based evaluation of bone
 374 mechanics. J Biomech 2007; 40: 2831-6.
- 375 20. Taddei F, Martelli S, Reggiani B, Cristofolini L, Viceconti M. Finite element
 376 modeling of bones from CT data: sensitivity to geometry and material
 377 uncertainties. IEEE Trans Biomed Eng 2006; 53: 2194-200.
- 378 21. Wille H, Rank E, Yosibash Z. Prediction of the mechanical response of the femur
 379 with uncertain elastic properties. J Biomech 2012; 45: 1140-8.

- 22. Lu Y, Boudiffa M, Dall'Ara, Bellantuono I, Viceconti M. Development of a
 protocol to quantify local bone adaptation over space and time: Quantification of
 reproducibility. J Biomech 2016; 49: 2095-99.
- 383 23. Turkowski K, Gabriel S. Filters for common resampling tasks. In: Glassner AS
- 384 (Ed.), Graphics Gems 1. Academic Press 1990; 147-65.
- 385 24. Keller TS. Predicting the compressive mechanical behavior of bone. J Biomech
 386 1994; 27: 1159-68.
- 387 25. Cerdà V, Cerdà JL, Idris AM. Optimization using the gradient and simplex
 388 methods. Talanta 2016; 148:641–8.
- 389 26. Hanss M, Klimke A. On the reliability of the influence measure in the
 transformation method of fuzzy arithmetic. Fuzzy Set Syst 2004; 143: 371-90.
- 391 27. Frost HM. Bone's mechanostat: a 2003 update. Anat Rec A discov Mol Cell Evol
 392 Biol 2003; 275: 1081-101.
- 393 28. Wolff J. Concerning the interrelationship between form and function of the394 individual parts of the organism. Clin Orthop Relat Res 1988; 228: 2-11.
- 395 29. Beaupre GS, Orr TE, Carter DR. An approach for time-dependent bone modeling
- and remodeling application: a preliminary remodeling simulation. J Orthop Res1990; 8: 662-70.
- 30. Geraldes DM, Phillips AT. A comparative study of orthotropic and isotropic bone
 adaptation in the femur. Int J Numer Method Biomed Eng 2014; 30: 873-89.
- 400 31. Fritton JC, Myers ER, Wright TM, van der Meulen MC. Loading induced
 401 site-specific increases in mineral content assessed by microcomputed tomography
 402 of the mouse tibia. Bone 2005; 36: 1030-8.

403	32. Sugiyama T, Saxon LK, Zaman G, Moustafa A, Sunters A, Price JS, Lanyon LE.
404	Mechanical loading enhances the anabolic effects of intermittent parathyroid
405	hormone (1-34) on trabecular and cortical bone in mice. Bone 2008; 43: 238-48.
406	33. Eberle S, Göttlinger M, Augat P. An investigation to determine if a single
407	validated density-elasticity relationship can be used for subject specific finite
408	element analyses of human long bones. Med Eng Phys 2013; 35: 875-83.
409	34. Morgan EF, Bayraktar HH, Keaveny TM. Trabecular bone modulus-density
410	relationships depend on anatomic site. J Biomech 2003; 36: 897-904.
411	35. Schileo E, Dall'Ara E, Taddei F, Malandrino A, Schotkamp T, Baleani M,
412	Viceconti M. An accurate estimation of bone density improves the accuracy of
413	subject-specific finite element models. J Biomech 2008; 41: 2486-91.
414	



417 Fig. 1. Schematic description of the image processing procedure. (a) The mouse tibia; 418 (b) the tibia was aligned along the global coordinate system; (c) and (d) the μ FE tibial 419 model and boundary conditions; (e) the volume of interest (VOI) was partitioned into 420 20 compartments and the mechanical environment of the bone was quantified in the 421 20 compartments.





424 Fig. 2. Determination of the uncertain interval for the bone density-modulus
425 relationships. (a) Fitting exponential functions to the density-modulus relationships
426 available in the literature; (b) the determined bone density-modulus interval.



429 Fig. 3. The influence of material uncertainty on the occurrence frequency of the 1st 430 principal strain, the 3rd principal strain, and the strain energy density. The plotted 431 curves of occurrence frequency are the ones with the hardest and softest bone 432 density-modulus relationships.



Fig. 4. The material uncertainty-induced bounds of the 1st principal strain, the 3rd principal strain, and the strain energy density across the tibial volume of interest (VOI), with the corresponding density-modulus relationships. The dotted data are the mean values in the 20 compartments across the tibial VOI.



440

441 Fig. 5. The relative percentage differences of tibial mechanical parameters across the 442 tibial volume of interest (VOI). Data are presented as the differences between the 443 maximal and minimal values divided by the minimal values in the 20 compartments 444 across the tibial VOI.