Simulation on the internal structure of three-dimensional proximal tibia under different mechanical environments

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Abstract

Background: Bone can adjust its morphological structure to adapt to the changes of mechanical environment, i.e. the bone structure change is related to mechanical loading. This implies that osteoarthritis may be closely associated with knee joint deformity. The purposes of this paper were to simulate the internal bone mineral density (BMD) change in three-dimensional (3D) proximal tibia under different mechanical environments, as well as to explore the relationship between mechanical environment and bone morphological abnormity.

Methods: The right proximal tibia was scanned with CT to reconstruct a 3D proximal tibia model in MIMICS, then it was imported to finite element software ANSYS to establish 3D finite element model. The internal structure of 3D proximal tibia of young normal people was simulated using quantitative bone remodeling theory in combination with finite element method, then based on the changing pattern of joint contact force on the tibial plateau in valgus knees, the mechanical loading was changed, and the simulated normal tibia structure was used as initial structure to simulate the internal structure of 3D proximal tibia for old people with 6° valgus deformity. Four regions of interest (ROIs) were selected in the proximal tibia to quantitatively analyze BMD and compare with the clinical measurements.

Results: The simulation results showed that the BMD distribution in 3D proximal tibia was consistent with clinical measurements in normal knees and that in valgus knees was consistent with the measurement of patients with osteoarthritis in clinics.

Conclusions: It is shown that the change of mechanical environment is the main cause for the change of subchondral bone structure, and being under abnormal mechanical environment for a long time may lead to osteoarthritis. Besides, the simulation method adopted in this paper can more accurately simulate the internal structure of 3D proximal tibia under different mechanical environments. It helps to better understand the mechanism of osteoarthritis and provides theoretical basis and computational method for the prevention and treatment of osteoarthritis. It can also serve as basis for further study on periprosthetic BMD changes after total knee arthroplasty, and provide a theoretical basis for optimization design of prosthesis.

Keywords: Bone remodeling, Three-dimensional, Proximal tibia, Finite element, Valgus
Background

Osteoarthritis is a chronic joint disease, and severe osteoarthritis can cause knee-joint pain or malfunction. Total knee arthroplasty is needed to restore its function in serious cases. Its incidence and prevalence are rising with aging. It has serious influence on the life quality of the elder [1]. The typical radiographic features of knee osteoarthritis include degeneration of cartilage, subchondral bone sclerosis and osteophyte formation. It is thought that subchondral bone changes after cartilage degenerates, so much attention is paid to the change of cartilage and the treatment of osteoarthritis. However, recent studies showed that subchondral bone plays an important role in the development of osteoarthritis. The subchondral bone changes in the early stage of osteoarthritis and leads to the degeneration of articular cartilage, which may be the initial cause of osteoarthritis [2,3]. So it is important to investigate the roles of subchondral bone for the prevention and treatment of osteoarthritis.

In many cases, mechanical loading on subchondral bone appears to play a crucial role in the changes of BMD distribution of subchondral bone since bone tissue is living material that has the functionality to adapt to mechanical loading in terms of its mass and architecture. This attribute is known as functional adaptation, and it includes both modeling and remodeling processes [4]. Bone modeling happens simultaneously with bone growth, and its primary role is to control the structure and shape of bone during the growing period. Bone remodeling always exists in the whole life and its primary role is to renew and adjust the bone structure to adapt to the mechanical loading change [5]. The numerical simulation of bone functional adaptation behavior showed that the adaptation behaviors of bone tissue at different mechanical levels were different and mechanical stimulus was an important factor for the change of bone mass and architecture [6]. If varus or valgus deformity of the knee joint occurs, it will change the lower limb alignment, as well as local mechanical loading on the tibial plateau, thus results in the joint under abnormal mechanical environment. These changes break the mechanostat of bone tissue. So when knee joint deformity occurs, bone tissue will adjust its structure and bone mineral density (BMD) distribution of proximal tibia to adapt to this abnormal mechanical loading which may be one of the causes of osteoarthritis. Animal experiments proved that the former varus or valgus deformity was associated with the occurrence of knee osteoarthritis [7]. Afterward, some longitudinal studies on patients during early stage osteoarthritis found that if varus or valgus deformity occurred before osteoarthritis, it might be one of the causes of osteoarthritis [8]. As a result, the abnormal loading may be the cause of osteoarthritis [9]. And for patients already with osteoarthritis, knee joint deformity can aggravate the progress of disease.

When varus or valgus deformity of the knee joint happens, the loading acted on the tibial plateau will change and result in changes of BMD in proximal tibia, and it could cause the ratio of medial BMD to lateral BMD at the tibial plateau (M:L BMD Ratio) to become larger or smaller, which can increase the risk of knee osteoarthritis [10]. Thus BMD can be used an index for measuring bone quality [11]. In clinics, radiography is often used to detect BMD, but at least 30 percent change of BMD is required before it can be detectable radiographically [12]. So it is impossible to detect changes of BMD during early osteoarthritis with radiography accurately, nor to predict it. Quantitative bone remodeling theory in combination with finite element method cannot only be used to simulate bone structure efficiently, but also analyze the change of BMD caused
by the change of loads and explore the relationship between mechanical environment and osteoarthritis, thus provide theoretical basis for prevention and diagnosis of early osteoarthritis.

The important aspect of bone remodeling simulation is quantitative bone remodeling theory, in which mathematical formula is utilized to quantitatively describe the remodeling process of bone tissue. One of the fundamental theories of bone remodeling is the theory of adaptive elasticity proposed by Cowin et al. [13], Hegedus et al. [14] and Firoozbaksh et al. [15]. It was based on general continuum mechanics principles. Hart et al. [16] combined this theory with finite element model. Afterwards Weinans et al. [17], Xinghua et al. [18,19] developed the bone density adaptation algorithm, in which the strain energy density (SED) was used as mechanical stimulus and the bone internal structure was described by the apparent density. Another representative quantitative bone remodeling theory was proposed by Huiskes et al. [20]. The behaviors of osteoclasts resorption and osteoblasts formation were separately described. The co-effects of osteoclasts and osteoblasts can simulate the growth, adaptation and maintenance of trabecular [20,21]. With the development of computer technology, the minimum element size of the model used for simulation can be as small as 50 microns. From the previous bone remodeling studies, it is shown that the complex structure of trabecular bone can be simulated [22-24].

Some researchers successfully simulated the change of internal bone structures quantitatively, e.g. Weinans et al. simulated the change of periprosthetic BMD in a two-dimensional (2D) proximal femur with prosthesis and analyzed the effects of different prosthetic materials and fixation methods on distribution of periprosthetic BMD [25]. Similarly, the periprosthetic BMD change of ankle and tibia after ankle arthroplasty were simulated, and bone loss caused by stress shielding was quantitatively analyzed [26]. The periprosthetic BMD in 2D proximal tibia with tibial implant was also simulated and the bone loss caused by four different fixation methods of long stem prosthesis was studied [27]. Later, the BMD change in proximal tibia after long-stemmed total knee arthroplasty with and without bisphosphonates treatment were simulated, and the role of bisphosphonates treatment for mitigating bone loss of proximal tibia was investigated [28]. In addition, Zhu et al. developed a high-order nonlinear equation of bone remodeling, and the internal density distribution and external shape of vertebral body were simulated [18]. The effects of functionally gradient materials on periprosthetic BMD after implanting dental prosthesis were also simulated [29,30]. But these simulations were mostly based on 2D model. Using 2D model to represent three-dimensional (3D) structure has its limitation since 2D model cannot accurately describe the actual structure of bone and its loading condition. So it is very necessary to establish 3D models and simulate internal structures of 3D bone models using quantitative bone remodeling theory in combination with finite element method.

There are also some investigations regarding the simulation of internal structures of 3D bone models. For example, Marsik et al. established a 3D finite element model of proximal femur and simulated its internal structure, and their simulation results were consistent with real proximal femoral structures [31]. 3D finite element models of femur, tibia and mandible were also established to simulate the internal BMD distributions [32]. The effect of different prosthesis fixation methods for periprosthetic bone resorption after total knee arthroplasty with 3D tibial models were simulated and analyzed [33].
Sharma et al. modeled the intact 3D finite element models of glenoid and glenoid prosthesis-bone. They simulated not only the BMD distribution, but also the influences of different materials and fixation methods of prosthesis on periprosthetic BMD [34]. Although some 3D bone remodeling simulations were successful, similar studies on proximal tibia are few. There are even fewer investigations on the relationship between the change of mechanical environment and knee osteoarthritis based on 3D bone remodeling simulation.

Accordingly, this paper aimed to simulate 3D internal structure of proximal tibia under different mechanical environments and to explore the relationship between the change of mechanical environment and the internal structure change of proximal tibia. This study may help to better understand the mechanism of osteoarthritis and provide theoretical basis for early diagnosis and risk prediction for knee osteoarthritis in clinics, thus can serve as theoretical basis for further study on periprosthetic bone resorption after total knee arthroplasty and optimal design of prosthesis.

Methods

In this paper, the bone remodeling control equation proposed by Weinans et al. [17] in combination with finite element method was utilized to simulate the internal structure of 3D proximal tibia with fixed shape. The load applied on proximal tibia included joint contact force and ligament forces. In order to validate the method in this paper and investigate the influence of mechanical environment on the BMD of proximal tibia, the internal structure of proximal tibia of young people under normal mechanical environment was simulated first. The BMD of the selected regions of interest (ROIs) were obtained and compared with clinical measurements to verify if the method adopted in this paper can accurately simulate BMD distribution of 3D proximal tibia. Then the loading condition on the tibial plateau in valgus knee was changed according to the changing pattern of joint contact force, and the change of BMD in proximal tibia of old people with 6° vaglus deformity was simulated.

Bone remodeling control equation

The control equation of bone remodeling process is as follows [17]:

\[
\frac{d\rho}{dt} = B\left(\frac{U_a}{\rho-k}\right) \quad 0 < \rho \leq \rho_{cb}
\]

\[
U_a = \frac{1}{n} \sum_{i=1}^{n} U_i
\]

Where \(\rho\) is apparent density, which is used to describe the characteristics of internal structure of bone. \(U_a\) is apparent strain energy density. \(U_a/\rho\) is strain energy density per unit mass, which is used as mechanical stimulus. \(B\) and \(k\) are constants. \(\rho_{cb}\) is the maximal bone mineral density, which is usually chosen the density of cortical bone. \(U_i\) is apparent strain energy density for loading case \(i\). \(n\) is the total number of loading cases. In this paper, the constant \(B\) is taken as \(0.05 \text{(g/cm}^3\text{)}^2 \text{(MPa \times time-unit)}^{-1}\). The reference stimulus \(k\) is \(0.14 \text{ J/g}\). The maximal BMD is \(1.92 \text{ g/cm}^3\) and the Poisson’s ratio is 0.3.
The piecewise function proposed by Zhu et al. is used to describe the relationships between elastic modulus and apparent density according to cancellous bone microstructural models and bone physiology as follows [35]:

\[
E = \begin{cases} 
1007 \times \rho^2 & \rho \leq 0.25 \\
255 \times \rho & 0.25 < \rho \leq 0.4 \\
2972 \times \rho^2 - 933 \times \rho & 0.4 < \rho \leq 1.2 \\
1763 \times \rho^3 & \rho > 1.2 
\end{cases}
\] (2)

Wherein, the units of elastic modulus and apparent density are MPa and g/cm³, respectively. This expression is more specific in describing the relationship between apparent density of cancellous bone and elastic modulus.

The flow chart of bone structure simulation using quantitative bone remodeling theory in combination with finite element method is shown in Figure 1 [36].

The iteration will stop when the bone remodeling process reaches equilibrium condition. According to the physiological mechanism of bone remodeling, bone remodeling equilibrium condition is as follows:

\[
d\rho/dt = 0, \text{ i.e. } U/\rho = k \text{ or } \rho = \rho_{\text{min}} \text{ or } \rho = \rho_{\text{max}} 
\]

Where \( \rho_{\text{min}} \) is taken as 0.01 g/cm³ and \( \rho_{\text{max}} = \rho_{cb} \). In this paper \( \rho_{cb} = 1.92 \text{ g/cm}^3 \).

**Finite element model**

*The 3D finite element model of proximal tibia for young normal people*

The 3D finite element model of proximal tibia for young normal people is derived from the CT scan data of a right proximal tibia of a patient. This patient is male, 34 years old with the height of 173 cm and the body weight of 70 kg. The left tibia of this patient is fractured, but the right knee is healthy. The images were reconstructed into 3D model of proximal tibia in MIMICS software, and then imported into ANSYS software to mesh. The finite element model is shown in Figure 2. This model contains bone tissue in proximal tibia, but the structure of articular cartilage and other soft tissues are not included. This model includes 176708 tetrahedral elements and 30920 nodes. The average element size is 2.0490 mm. It was shown that the initial density had little effect on the simulated results [37]. Thus the initial density is chosen as 0.8 g/cm³ in this paper.

![Figure 1 Iterative feedback mechanism of bone remodeling simulation in combination with finite element method [36].](http://www.biomedical-engineering-online.com/content/12/1/130)
The proximal tibia model is constrained distally and the loads applied on it consist of joint contact force, anterior shear force produced by joint contact force, medial collateral ligament force and anterior cruciate ligament force. According to the anatomical structure of knee joint, there are four main ligaments on the knee joint, i.e. medial collateral ligament, lateral collateral ligament, anterior cruciate ligament, and posterior cruciate ligament. In addition to the lateral collateral ligament that is attached to fibula, the other three ligaments are all attached to proximal tibia [38,39]. Shelburne et al. used a 3D model of lower limb to calculate the joint contact force and ligament forces on the normal knee in a complete gait cycle [40]. Posterior cruciate ligament was found unloaded during stance. Thus the ligament forces that applied on proximal tibia in this paper include medial collateral ligament force and anterior cruciate ligament force (see Figure 2). The magnitudes of loads applied on proximal tibia are the mean value of 0%, 10%, 20%, 30%, 40%, 50% and 60% gait cycle loading [40]. The loads applied on the proximal tibia, i.e. joint contact force, shear force, anterior cruciate ligament force and medial collateral ligament force are 1233.3 N, 102.04 N, 139.3 N and 6.4 N, respectively.

The joint contact force acted on tibial plateau, and the medial and lateral tibial plateau bore about 55% and 45% of it [41]. The direction of joint contact force is perpendicular to the surface of meniscus and the part of the tibial plateau not covered by the meniscus. Actually the medial and lateral tibial plateau is covered by medial and lateral meniscus, respectively. The coverage area of medial and lateral tibial plateau by meniscus is about 64% and 84%, respectively, and this proportion remains unchanged during growth [42]. The main role of meniscus is to bear the forces acted on tibial plateau and can carry about 40-60% of the total force [42]. The applied loads of ligaments include the medial collateral ligament force and the anterior cruciate ligament force. The positions of ligament forces acted on proximal tibia are the attached region of ligaments, and the load directions are consistent with ligament orientations [43,44].

**The 3D finite element model of proximal tibia for old people with 6° valgus deformity**

We simulated the internal BMD distribution of proximal tibia with 6° valgus deformity. The reason was that Hulet et al. measured BMD of proximal tibia for the patients who
needed treatment due to knee pain, in which 22 cases were valgus deformity of the knee joint and the valgus angle was larger than 4° [12].

The loadings acted on tibial plateau are changed due to the change of lower limb alignment in valgus knees. In normal knees, the connection between femoral head and ankle center is through the center of knee joint, and the loading acted on articular surface are relatively uniform. But when valgus deformity of the knee joint occurred, the lower limb alignment was through lateral tibial plateau, the load center would offset to the lateral tibial plateau and in the corresponding plateau the loading increased [45]. The resultant pressure on medial and lateral plateau in knees with 5° valgus deformity is similar to that in normal knees, while the distribution of loadings acted on the medial and lateral tibial plateau was changed [46]. So in this paper, we assumed that the resultant joint force was unchanged when knee valgus deformity occurred, and the lateral and medial tibial plateau accounted for about 66.5% and 33.5% of joint contact force, respectively. We also simulated the change of internal structure of proximal tibia in valgus knees from uniform initial density. The initial density is 0.8 g/cm³.

**BMD measurement**

In order to verify the simulation method and compare with clinical observations, in this paper, four ROIs in proximal tibia are selected to analyze and compare the BMD. These four ROIs are throughout the tibial plateau from front to back. The selection methods are the same with the actual clinical region selection methods in the literature. The coronal plane diagram for these four selected ROIs of proximal tibia is shown in Figure 3. Figure 3 shows the CT-images of right proximal tibia of a patient, which is used not only to mark the locations of ROIs, but also to compare with the simulated results in Results section. ROI1 and ROI2 are the same as the measuring regions selected by Hudson et al. in clinics [47]. Taken fibular head as reference, the height of ROIs is one-half the distance from the fibular head to the superior border of the cortical plate, and its width is one half of tibial plateau. The ROIs located beneath the medial and lateral tibial plateau are named ROI1 and ROI2, and the coronal plane diagram of these two ROIs is shown in Figure 3a. ROI3 and ROI4 are the same as the measuring regions selected by Hulet et al. in clinics [12]. The distance from the fibular head to the superior border of the cortical plate is about 11 mm in reference [48], while it is 14 mm in our model, the width of tibial plateau is equally divided into 14 parts and these regions are named as R1-R14 successively, where R2, R3, R4 constitute ROI3 in medial tibia, and R11, R12, R13 constitute ROI4 in lateral tibia. The coronal plane diagram of ROI3 and ROI4 is shown in Figure 3b.

**Results and discussion**

**Results**

Simulation result of the internal structure of 3D proximal tibia for young normal people using quantitative bone remodeling theory in combination with finite element method is shown in Figure 4. Simulation result of internal structure of 3D proximal tibia for old people with 6° valgus deformity is shown in Figure 5.
Figure 3 The coronal diagram of the four selected ROIs according to the measuring regions in clinics. (a) ROI1 and ROI2. (b) ROI3 and ROI4.
Figure 4 The simulation result of 3D proximal tibia for young normal people. (a) Top view. (b) Right view. (c) Posterior view. (d) Cross-sectional view of the coronal plane.
Figure 5 The simulation result of 3D proximal tibia for old people with 6° valgus deformity.

(a) Top view. (b) Right view. (c) Posterior view. (d) Cross-sectional view of the coronal plane.
The simulated normal proximal tibia structure of young people is similar with the real proximal tibia structure (Figure 3), which can be seen from the following three aspects:

(1) Seen from Figure 4a-c, there is a layer of cortical bone covering the whole tibia, and the cortical bone is relatively continuous and completed. This is consistent with the real proximal tibial cortical bone structure in Figure 3;

(2) From Figure 4d it can be seen that there is medullary cavity within distal tibia. This characteristic is also consistent with the real proximal tibial structure in Figure 3;

(3) Beneath the tibial plateau there is cancellous bone, and the BMD in medial region is bigger than that in lateral region. It is in agreement with the measurements in clinics [49].

From the above results it can be seen that the typical characteristics of the simulation results of proximal tibia structure for young normal people are consistent with real tibia.

The average BMDs of the four ROIs and the whole proximal tibia for young normal people and old people with 6° valgus deformity simulation results are listed in Table 1. From the simulation results of proximal tibia of young normal people it can be seen that the average BMDs in ROI1 and ROI2 are 1.23 g/cm³ and 1.03 g/cm³, and the M:L BMD Ratio is 1.19. While for the same ROIs, the proximal tibial BMDs of 30 young healthy people were measured by Hudson et al. [47], and the obtained average M:L BMD Ratio was 1.20 ± 0.10. Thus it can be seen that the simulated M:L BMD Ratio in this paper is consistent with that measured by Hudson et al. [47]. It shows that the BMD distribution of 3D proximal tibia for young normal people we simulated is consistent with real proximal tibia. Thus it appears that the method used in this paper can simulate and predict the BMD distribution of 3D proximal tibia accurately.

From the simulation result of old people with 6° valgus deformity it can be seen that the average BMDs of ROI3 and ROI4, and the M:L BMD Ratio are 0.908 g/cm³, 1.033 g/cm³ and 0.879, respectively. While the BMDs of proximal tibia of 22 osteoarthritis patients who had severe knee pain with mean age of 71 years old and valgus deformity degree larger than 4° for the same ROIs were measured by Hulet et al. [12] with the average BMDs of ROI3 and ROI4 0.827 ± 0.198 g/cm³ and 0.939 ± 0.229 g/cm³, and the M:L BMD Ratio 0.878-0.886. It can be seen that under such loading condition, the BMD of medial proximal tibia decreases and that of lateral region increases, i.e. the M:L BMD Ratio is very small. This result is consistent with the clinical observation data of knee osteoarthritis patients with valgus deformity, which means that when valgus deformity of the knee occurs, the BMD distribution of proximal tibia changes accordingly under that abnormal mechanical environment for a long period.

**Discussion**

In this paper, the internal structure of 3D proximal tibia was simulated using bone remodeling theory in combination with finite element method. The internal structure of 3D proximal tibia of young normal people was simulated to validate the method used in this paper. Comparing the simulated results in Table 1 with the observations in the literature, it can be seen that the average BMD ratio of ROI1 to ROI2 in the simulated result of internal structure of 3D proximal tibia for young normal people
## Table 1 The average BMD obtained by simulation and measurement in the literature

| Simulation results of proximal tibia for young normal people | Simulation results of proximal tibia for old people with 6° valgus deformity | Measurement results of young people obtained by Hudson et al. [47] | Measurement results of old people with valgus deformity >4° obtained by Hulet et al. [12] |
|-------------------------------------------------------------|--------------------------------------------------------------------------------|------------------------------------------------------------------|------------------------------------------------------------------------------------------|
| The average BMD of ROI1 (g/cm³)                            | 1.23                                                                          | 0.89                                                             | 0.827 ± 0.198                                                                            |
| The average BMD of ROI2 (g/cm³)                            | 1.03                                                                          | 1.10                                                             | 0.939 ± 0.229                                                                            |
| The average BMD of ROI3 (g/cm³)                            | 1.246                                                                         | 0.908                                                            |                                                                                            |
| The average BMD of ROI4 (g/cm³)                            | 0.871                                                                         | 1.033                                                            | 0.879 ± 0.198                                                                            |
| The average BMD of the whole (g/cm³)                       | 1.343                                                                         | 1.185                                                            | 1.20 ± 0.10                                                                               |
| ROI1/ROI2                                                  | 1.19                                                                          | 0.81                                                             |                                                                                            |
| ROI3/ROI4                                                  | 1.431                                                                         | 0.879                                                            | 0.878-0.886                                                                              |

The average BMD of young normal people and old people with 6° valgus deformity in the selected ROI and the whole proximal tibia was calculated, and the measurement result in the same zone was listed.

*ROI* region of interest, BMD bone mineral density.
is consistent with the result of young healthy people in the same area measured by Hudson et al. [47]. Hence with the method used in this paper the BMD distribution of 3D proximal tibia can be simulated more accurately.

After the simulated results were verified, the loading applied on the proximal tibia was changed according to the changing pattern of loading when valgus deformity occurred. Then the simulated result of internal structure of proximal tibia for old people with 6° vaglus deformity was obtained. The average BMD ratio of ROI3 to ROI4 was in the range that measured in the same region of old people with greater than 4° vaglus deformity who need treatment because of knee joint pain [12]. It means that when the loads change, the internal structure of proximal tibia is consistent with clinical observations on osteoarthritis patients, that is, under the mechanical environment of valgus knees, the BMD distribution of proximal tibia has changed. It shows that mechanical environment changes the BMD distribution under cartilage in proximal tibia, and the change of mechanical environment is one of the causes for bone structural abnormality. If knee joint deformity occurred, it may lead to the change of loads acted on the tibial plateau, and that change can break the primary mechanostat, which will enhance the bone remodeling of proximal tibia and increase the risk of osteoarthritis.

In clinics, the bone mass and distribution of cancellous bone in proximal tibia are thought to be closely associated with age and gender, and BMD of proximal tibia would reduce with ageing for males, as well as females [50]. In Table 1, the BMDs of ROI1, ROI3 and the whole proximal tibia obtained in our simulations for young normal people are all greater than old people with valgus deformity. The BMDs of ROI1, ROI3 and the whole decease by 27.64%, 27.127% and 11.765% for old people in comparison with those of young people. These results show that with ageing, the BMD of proximal tibia became smaller. This pattern is consistent with clinical observations. But in ROI2 and ROI4, however, the average BMD of young normal people is smaller than old people with valgus deformity. The reason may be that we simulated the BMD distribution of proximal tibia in this paper when valgus deformity of the knee occurs, the joint contact force is not changed in valgus knees, but its distribution changes with the load center on tibial plateau moving from medial to lateral region [45], resulting in lateral tibial plateau bears more load, and the BMDs of ROI2 and ROI4 beneath lateral tibial plateau increase. Even so, the average BMD of proximal tibia of old people in ROI2 and ROI4 is still smaller than the whole BMD of proximal tibia of young people; the BMD changing pattern is in agreement with the loading that acted on tibial plateau. In addition, from Figures 4d and 5d we can see that the cortical thickness of proximal tibia of young normal people is thicker than that of old people with 6° valgus deformity.

The specific features of osteoarthritics such as joint space narrowing, osteophyte and hardening are closely related to extreme M:L BMD Ratio, and the M:L BMD Ratio is proportional to medial joint space narrowing and inversely proportional to lateral joint space narrowing [10]. If M:L BMD Ratio is too low, it can increase the possibility of lateral joint space narrowing, lateral osteophyte formation and hardening. When the varus or valgus deformity of the knee joint occurs, the axial alignment of the lower extremity changes and load center will be moved from the center of tibial plateau to medial or lateral tibial plateau [45]. This changes the distribution of loads acted on tibial plateau, thus affects the internal structure of proximal tibia, i.e. BMD of the proximal tibia would increase with the increasing load on its corresponding plateau,
which leads to the change of M:L BMD Ratio. Thus the M:L BMD Ratio can be used as a diagnosis index for osteoarthritis in clinics. Another direct application of the method used in this paper is to accurately simulate the change of BMD of proximal tibia caused by load changes when varus or valgus deformity occurred, so it can provide theoretical basis for prevention and early diagnosis of osteoarthritis.

In summary, the change of BMD in proximal tibia is closely related to mechanical environment, and mechanical environment is the direct reason for the change of internal structure of proximal tibia. If keen joint abnormalities occur, being under this abnormal mechanical environment for a long period may lead to osteoarthritis. The proximal tibial model we used in this study didn’t contain the articular cartilage. That is why the degeneration of articular cartilage was not discussed. But the effect of changes of mechanical properties of articular cartilage was included in the change of loading. In this paper the relationship between mechanical environment and bone morphological structure is mainly studied. Although the effect of articular cartilage cannot be investigated directly, it was shown that the subchondral bone structure and the degree of articular cartilage damage were closely related [51]. The change of BMD occurred after varus deformity may imply that the articular cartilage has been damaged already and moreover, the damage of articular cartilage will cause the uneven loading distribution on tibial plateau, which will aggravate the change of subchondral bone. From the above result it can be seen, the change of mechanical environment directly lead to the change of bone morphological structure. The mechanism of osteoarthritis is analyzed from the influence of mechanical environment on bone structure, which helps to better understand the mechanism of osteoarthritis.

In order to verify the convergence of the model utilized in this paper (model 1), the same proximal tibia was also meshed with a smaller or a larger element size, i.e. the average element size of this two model is 1.6000 mm (model 2) and 2.6706 mm (model 3), respectively, while the average element size of model 1 used in this paper is 2.0490 mm. These three models contain 176708, 360789 and 80353 tetrahedral elements, respectively. Under the same loading and boundary conditions, these three models were analyzed in ANSYS software. The detailed information and results are listed in Table 2. It is found that the relative difference of average element equivalent stress, average equivalent strain and average SED between model 1 and model 2 are 0.01296%, 1.4467% and 4.5333%. Although the relative difference of element size between these two models is 28.0625%, all the relative differences in finite element results are within 5%. The differences

|                              | Model 1 | Model 2 | Model 3 | Relative difference between model 1 and model 2 (%) | Relative difference between model 1 and model 3 (%) |
|------------------------------|---------|---------|---------|-----------------------------------------------------|-----------------------------------------------------|
| Average element size (mm)    | 2.0490  | 1.6000  | 2.6706  |                                                     |                                                     |
| Number of elements           | 176708  | 360789  | 80353   |                                                     |                                                     |
| Average element equivalent stress (MPa) | 0.7717 | 0.7716  | 0.8451  | 0.01296                                            | 9.5115                                              |
| Average element equivalent strain | 0.0006774 | 0.0006676 | 0.0007312 | 1.4467                                              | 7.9421                                              |
| Average element SED (MPa)    | 0.0004125 | 0.0003938 | 0.0004731 | 4.5333                                              | 14.6909                                              |

SED strain energy density.
in results are relative small, but decreasing the element size will cause the number of ele-
ments increasing and consuming too much time to compute. Comparing the finite element
results of model 3 with model 1, the relative differences are 9.5115%, 7.9421% and
14.6909%, respectively. It is shown that the relative differences between model 3 and model
1 are large. From the above analyses it is shown that the element size of the model used in
this study is small enough to guarantee the accuracy of the results.

The simulation method in this paper has wide applications. For example, it can be
used for further computational simulation on the change of periprosthetic BMD due to
stress shielding after total knee arthroplasty or other joint replacement. And the
method can also simulate the change of periprosthetic bone mass for different
prosthesis material or fixation methods, which provides a theoretical basis for the
optimal design of prosthesis material and suitable selection of prosthesis fixation methods.
By changing the control equations of bone remodeling process and introducing biological
factors, the change of BMD for menopausal women may be further simulated. The
method used in this paper can quantitatively simulate the BMD change due to the
change of mechanical environment accurately and rapidly. It also solves the problem in
clinics that too much time cost on observing the change of BMD in human body, and
makes up the disadvantage that some experimental methods cannot be applied directly to
human body.

Conclusions
Bone morphological abnormity is closely related to its abnormal mechanical environment,
thus the knee joint deformity would be the cause of osteoarthritis, and the method used
in this paper can be used as an effective method to predict the BMD change of proximal
tibia under different mechanical environments to help diagnose osteoarthritis in clinics.
Another important conclusion is that the M:L BMD Ratio can be used as another criteria
to measure the structural change caused by osteoarthritis on specific site.

Abbreviations
3D: Three-dimensional; 2D: Two-dimensional; BMD: Bone mineral density; ROI: Region of interest; M:L BMD Ratio:
The ratio of medial BMD to lateral BMD at the tibial plateau.

Competing interests
The authors declare that they have no competing interests.

Authors’ contributions
JF established the finite element model, carried out the numerical simulations and drafted the manuscript. HG
conceived of the study design and helped to draft the manuscript. LYK participated in the numerical simulations.
DZ provided the CT images. All authors read and approved the final manuscript.

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References
1. Woolf AD, Pfleger B: Burden of major musculoskeletal conditions. Bull World Health Organ 2003, 81:646–656.
2. Goldring SR: Role of bone in osteoarthritis pathogenesis. Med Clin North Am 2009, 93:25–35. xv.
3. Quasnicka HL, Anderson-Mackenzie JM, Bailey AJ: Subchondral bone and ligament changes precede cartilage degradation in guinea pig osteoarthritis. Biomech Eng Online 2006, 43:389–397.

4. Frost HM: Seeking genetic causes of "osteoarthritis": insights of the Utah paradigm of skeletal physiology. Bone 2001, 29:407–412.

5. Popoda P, Priemel M, Rueger JM, Amling M: Bone remodeling: new aspects of a key process that controls skeletal maintenance and repair. Osteoporos Int 2005, 16:518–524.

6. Gong H, Zhu D, Gao JZ, Lv LW, Zhang XZ: An adaptation model for trabecular bone at different mechanical levels. Biomed Eng Online 2010, 9:32.

7. Tetsworth K, Paley D: Malalignment and degenerative arthropathy. Orthop Clin North Am 1994, 25:367–377.

8. Sharma L, Song J, Felton DT, Cahué S, Shamiyeh E, Dunlop DD: The role of knee alignment in disease progression and functional decline in knee osteoarthritis. JAMA 2001, 286:188–195.

9. Guiltak F: Biomechanical factors in osteoarthritis. Best Pract Res Clin Rheumatol 2011, 25:815–823.

10. Lo GH, Zhang Y, McLennan C, Niu J, Kiel DP, McLean RR, Allabadi P, Felton DT, Hunter DJ: The ratio of medial to lateral tibial plateau bone mineral density and compartment-specific tibifemoral osteoarthritis. Osteoarthr Cartilage 2006, 14:984–990.

11. Petersen MM: Bone mineral measurements at the knee using dual photon and dual energy X-ray absorptiometry. Acta Orthop Scand 2000, 71(suppl):2–31.

12. Huilet C, Sabater JP, Souquet D, Locker B, Marcelli C, Velpau C: Distribution of bone mineral density at the modelling and clinical osteoarthritis. Calcified Tissue Int 2002, 71:315–322.

13. Cowin SC, Hegedus DH: Bone remodeling: theory of adaptive elasticity. J Elasticity 1976, 6:313–326.

14. Hegedus DH, Cowin SC: Bone remodeling II: small strain adaptive elasticity. J Elasticity 1976, 6:337–352.

15. Firoozabadi K, Cowin SC: An analytical model for paupels functional adaptation mechanism in bone. J Biomech Eng 1981, 103:247–252.

16. Hart RT, Davy DT, Heiple KG: A computational method of stress analysis adaptive elastic materials with a view toward application in strain induced remodeling. J Biomech Eng 1984, 106:342–350.

17. Weinans H, Huiskes R, Grootenboer HJ: The behavior of adaptive bone remodeling simulation models. J Biomech 1992, 25:1425–1441.

18. Zhu XF, Gong H, Zhu D, Gao BZ: A study of the effect of non-linearities in the equation of bone remodeling. J Biomech 2002, 35:951–960.

19. Zhu XH, Gong H, Gao BZ: The application of topology optimization on the quantitative description of the external shape of bone structure. J Biomech 2005, 38:1612–1620.

20. Huiskes R, Ruimernan R, Van Lente HT, Janssen JD: Effects of mechanical forces on maintenance and adaptation of form in trabecular bone. Nature 2000, 405:704–706.

21. Ruimernan R, Van Rietbergen B, Hilbers P, Huiskes R: A 3-dimensional computer model to simulate trabecular bone metabolism. Biomech 2003, 40:315–320.

22. Jang IG, Kim FY: Computational simulation of trabecular adaptation progress in human proximal femur during growth. J Biomech 2009, 42:573–580.

23. Jang IG, Kim FY: Computational study of Wolff’s law with trabecular architecture in the human proximal femur using topology optimization. J Biomech 2008, 41:2353–2361.

24. Tsukuba K, Suzuki Y, Yamada T, Hojo M, Makinouchi A, Adachi T: Computer simulation of trabecular remodeling in human proximal femur using large-scale voxel FE models: approach to understanding Wolff’s law. J Biomech 2009, 42:1088–1094.

25. Weimans H, Huiskes R, Grootenboer HJ: Effects of material properties of femoral hip components on bone remodeling. J Orthop Res 1992, 10:845–853.

26. Bouguecha A, Weigel N, Behrens BA, Stukenborg-Colsman C, Waizy H: Numerical simulation of strain-adaptive bone remodeling in the ankle joint. Biomed Eng Online 2011, 10:58.

27. Nyman JS, Hazewoold SJ, Rodrigo JJ, Martin RB, Yeh OC: Femoral stem and knee arthroplasty with interlocking screws: a computational bone adaptation study. J Orthop Res 2004, 22:51–57.

28. Nyman JS, Rodrigo JJ, Hazewoold SJ, Yeh OC, Martin RB: Predictions on preserving bone mass in knee arthroplasty with biphosphonates. J Arthroplasty 2006, 21:106–113.

29. Lin D, Li Q, Li W, Swain M: Bone remodeling induced by dental implants of functionally graded materials. J Biomed Mater Res B 2010, 92:430–438.

30. Lin D, Li Q, Li W, Swain M: Dental implant induced bone remodeling and associated algorithms. J Mech Behav Biomed 2009, 2:410–432.

31. Marck F, Kikla V, Chup H: Remodelling of living bone induced by dynamic loading and drug delivery-numerical modelling and clinical treatment. Math Comput Simulat 2010, 80:1278–1288.

32. Perez MA, Forrester P, Dobllare M, Garcia-Aznar JM: Comparative analysis of bone remodeling models with respect to computerized tomography-based finite element models of bone. Comput Method Biomech 2010, 13:71–80.

33. Chong D, Hansen UN, van der Venne R, Verdonschot N, Amis AA: The influence of tibial component fixation techniques on resorption of supporting bone stock after total knee replacement. J Biomech 2011, 44:946–954.

34. Sharma GB, Debvisi RE, McMahurn PJ, Robertson DO: Effect of genoid prosthesis design on genoid bone remodeling: adaptive finite element based simulation. J Biomech 2010, 43:1653–1659.

35. Zhu XH, Gong H, Bai XF, Wang FR: Application of subsectional relationship between elastic modulus and apparent density in the structural simulation of proximal femur. Chin J Biomed Eng 2003, 22:250–257.

36. Weimans H, Huiskes R, Grootenboer HJ: Effects of fit and bonding characteristics of femoral stems on adaptive bone remodeling. J Biomech Eng 1994, 116:393–400.

37. Marzban A, Canavan P, Warner G, Vaziri A, Nayeb-Hashemi H: Parametric investigation of load-induced structure remodeling in the proximal femur. J Eng Med 2012, 226(6):450–460.

38. Woo SL, Abramowitch SD, Kliger R, Liang R: Biomechanics of knee ligament: injury, healing, and repair. J Biomech 2006, 39:1–20.
39. Blackburn TA, Craig E: Knee anatomy: a brief review. Phys Ther 1980, 60:1556–1560.
40. Shelburne KB, Tony MR, Pandy MG: Muscle, ligament, and joint-contact forces at the knee during walking. Med Sci Sport Exerc 2005, 37:1948–1956.
41. Zhao D, Banks SA, D’Lima OD, Colwell CW, Fregly BJ: In vivo medial and lateral tibial loads during dynamic and high flexion activities. J Orthop Res 2007, 25:593–602.
42. Clark CR, Ogden JA: Development of menisci of the human knee joint: morphological changes and their potential role in childhood meniscal injury. J Bone Joint Surg Am 1983, 65:538–547.
43. Staubli HU, Rauschning W: Tibial attachment area of the anterior cruciate ligament in the extended knee position: anatomy and cryosections in vitro complemented by magnetic resonance arthrography in vivo. Knee Surg Sport Tr A 1994, 2:138–146.
44. Liu F, Yue B, Gadikota HR, Kozaneck M, Liu WJ, Gill TJ, Rubash HE, Li G: Morphology of the medial collateral ligament of the knee. J Orthop Surf Res 2010, 5:69–77.
45. Bruns J, Volkmer M, Luessenhop S: Pressure distribution at the knee joint. Arch Orthop Traum Surg 1993, 133:12–19.
46. Harrington U: Static and dynamic loading patterns in knee joints with deformities. J Bone Joint Surg Am 1983, 65(2):247–259.
47. Hudson D, Royer T, Richards J: Bone mineral density of the proximal tibia relates to axial torsion in the lower limb. Gait Posture 2007, 26:446–451.
48. Le Graverand MP, Mazzuca S, Lassere M, Guermazi A, Pickering E, Brandt K, Peterfy C, Cline G, Nevitt M, Woodworth T, Canaghian P, Vignon E: Assessment of the radioanatomic positioning of the osteoarthritic knee in serial radiographs: comparison of three acquisition techniques. Osteoarthr Cartilage 2006, 14(suppl A):A37–A43.
49. Li MG, Nilsson KG: The effect of the preoperative bone quality on the fixation of the tibial component in total knee arthroplasty. J Arthroplasty 2000, 15:744–753.
50. Khodadadyan-Klostermann C, Von Seebach M, Taylor WR, Duda GN, Haas NP: Distribution of bone mineral density with age and gender in the proximal tibia. Clin Biomech 2004, 19:370–376.
51. Bobinac D, Spanjoli J, Zoricic S, Marc I: Changes in articular cartilage and subchondral bone histomorphometry in osteoarthritic knee joints in humans. Bone 2003, 32:284–290.