Porous architected biomaterial for a tibial-knee implant with minimum bone resorption and bone-implant interface micromotion

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ARTICLE INFO

Keywords: Porous biomaterial Tibial knee implant Bone resorption End-of-stem pain Multiscale mechanics Topology optimization

ABSTRACT

This investigation presents the numerical development of a fully porous tibial knee implant that is suggested to alleviate the clinical problems associated with current prostheses that are fully solid. A scheme combining multiscale mechanics and topology optimization is proposed to handle the homogenized analysis and property tailoring of the porous architecture with the aim of reducing the stiffness mismatch between the implant and surrounding bone. The outcome of applying this scheme is a graded lattice microarchitecture that can potentially offer the implant an improved degree of load bearing capacity while reducing concurrently bone resorption and interface micromotion. Asymptotic Homogenization theory is used to characterize the mechanics of its building block, a tetrahedron based unit cell, and the Soderberg fatigue criterion to represent the implant fatigue resistance under multiaxial physiological loadings. The numerical results suggest that the overall amount of bone resorption around the graded porous tibial stem is 26% lower than that around a conventional, commercially available, fully dense titanium implant of identical shape and size. In addition, an improved interface micromotion is observed along the tibial stem, with values at the tip of the stem as low as 17 µm during gait cycle and 22 µm for deep bend compared to a fully dense implant. This decrease in micromotion compared to that of an identical solid implant made of titanium can reasonably be expected to alleviate post-operative end of stem pain suffered by some patients undergoing surgery at the present time.

1. Introduction

Knee replacement implants currently used in total knee arthroplasty (TKA) are generally made of a fully solid material, such as Titanium alloys and Cobalt chrome (CoCr) (Carr and Goswami, 2009). Despite their improved characteristics when compared to late 1970s implants, existing knee stems feature homogeneous properties that fail to fulfill the complete set of mechanobiological requirements they are subjected to (Carr and Goswami, 2009). One of these is the end-of-stem pain, which is experienced by 7% of the patients undergoing primary TKA (Mihalko and Whiteside, 2015). In revision surgery, despite the adoption of modified long stems, end-of-stem pain has been reported to affect as many as 19% of the patients (Mihalko and Whiteside, 2015). The incidence of end-of-stem pain has been documented for both fully cementless and fully cemented stems with stable fixation such that 9% of patients with fully cemented stems suffer from end-of-stem pain (Barrack et al., 2004; Jasty et al., 1991; Rakotomanana et al., 1994).

Similarly, for patients receiving a press-fit, cementless stem, an incidence rate of 14% has been reported in the literature (Glenn et al., 2010). As an alternative, hybrid fixation of a stemmed tibial component has been proposed and has now become the standard of care. With this technique, the tibial plateau and the proximal metaphysis are cemented, whereas the stem is cementless and press-fit into the tibial diaphysis (Haas et al., 1995). Nevertheless, although this type of fixation offers a lower failure rate, pain at the stem tip has been reported by patients with a stemmed tibial component with hybrid fixation (Barrack et al., 2004, 1999; Haas et al., 1995). The etiology of this pain is most commonly attributed to the severe interface micromotion that develops between the stem tip and the surrounding native bone, a problem that results from the elastic modulus mismatch existing between them (Glenn et al., 2010). Recent efforts to reduce the risk of end-of-stem pain have included modification of the implant macrogeometry and use of advanced materials, such as composites and functionally graded solids (Bahraminasab et al., 2013, 2014; Comelto et al., 2010). Using...
advanced materials, an enhanced performance has been reported in
tibiofemoral articulation (Greenwald and Heim, 2005; McEwen et al.,
2005) and bone-implant interface. In a recent study, Completo et al.
(2012) postulated that fine-tuning the material properties of the
tibial stem tip to make it mechanically similar to that of the surrounding
cortical bone could reduce the incidence of post-operative pain. They
found that a polyethylene tip lowered the strain at the distal end of the
tibial stem, which could contribute to a reduction in the occurrence of
end-of-stem pain. Despite these modest improvements in trying to
eliminate end-of-stem pain in the proximal tibia, there has been no
modification of the tibial implant that has successfully dealt with the
modulus mismatch and micromotion that occurs between the stem and
the surrounding tibial bone (Glenn et al., 2010).

The second consequence of the elastic modulus mismatch between
the tibial implant and the surrounding host bone is bone resorption
secondary to stress shielding. The modulus mismatch between the tibial
component and the host bone results in periprosthetic bone loss in both
primary and revision TKA (Van Lenthe et al., 1997). There are studies in
the literature that have shown the potential of decreasing bone res-
orption by means of reducing the implant stiffness (Glenn et al., 2010;
Gefen, 2002; Huiskes et al., 1992; Shi et al., 2013). A reduction in
implant stiffness to the order of magnitude of the surrounding bone
tissue has been achieved in implants that use either composite materials
or porous solids with uniform porosity (Bahraminasab et al., 2014;
Karlsson et al., 2000; Mataassi et al., 2013). These attempts, however,
have shown only partial success. Only recently the tailoring of porosity
gradients in porous hip stems has been demonstrated effective in re-
ducing bone resorption secondary to stress shielding (Arabnejad
shielding around a TKA can predispose to a subsequent periprosthetic
fracture and decreases the amount and quality of bone at the time of
revision surgery. The contribution of material properties to stress
shielding in the proximal tibia was investigated in a computational
study, where a homogenous distribution of elastic properties was de-
signed to be tuned to the order of magnitude of the surrounding can-
cellous bone (Au et al., 2007). However, since the elastic properties
were assumed constant within the implant, the amount of stress
shielding could not be reduced to minimum and a global reduction of
bone resorption could not be achieved at the distal stem (Mataassi et al.,
2013). Enab (2012) found that customizing the elastic modulus within
the tibial tray could contribute to a 46% reduction in the stresses
transferred to the implant, thus resulting in a lower bone resorption.
This reduction can be achieved by varying the elastic modulus from
40 GPa at the top of the tray to 110 GPa downward. Other strategies
that deal with the resolution of the modulus mismatch between stan-
dard titanium implants and the surrounding tibia involve the use of
highly porous materials, such as tantalum foam (Bobyn et al., 2004).
Although its biocompatibility and high volumetric porosity can provide
an exceptional degree of compliance, tantalum foam has a quasi-uni-
form distribution of pores which has limited capacity in minimizing
bone resorption and interface micromotion simultaneously. Recently,
porous implants featuring uniform porosity have been introduced with
the aim of improving the performance of solid implants (Karlsson et al.,
2000; Mataassi et al., 2013; Paganias et al., 2012). Despite encouraging
results, these implants fall short in addressing post-operative complic-
ations, namely the concurrent reduction of bone resorption and in-
terface micromotion. Elastic property tailoring in hip stems was rec-
ognized already in the 1990s as a promising mean to reduce bone
resorption (Kuiper and Huiskes, 1992, 1997). Whereas those studies
focus on the use of functionally graded solids, more recently porous
materials with tailored cellular architecture have been proved suc-
cessful in reducing bone resorption secondary to stress shielding,
promoting bone ingrowth and providing implant stability (Arabnejad
This strategy has been recently adopted to design a hip replacement
implant that can facilitate osseointegration and concurrently minimize
bone resorption and bone-implant interface failure (Arabnejad Khanoki
and Pasini, 2012, 2013a, 2013b; Arabnejad et al., 2017). While this
work focused on the suppression of bone resorption in a hip implant, a
similar strategy can be extended to deal with stress shielding and mi-
cromotion that occur in present day stemmed tibial implants used in
TKA.

In this numeric investigation, we introduce a fully porous cement-
less stemmed tibial implant for primary and revision knee replacement
that has a 3D cellular architecture tailored to concomitantly minimize
interface micromotion and bone resorption, while satisfying clinical
strength and fatigue requirements. Section 2 of this study presents a
systematic methodology integrating multiscale mechanics and topology
optimization to tailor the material properties of the knee implant.
Section 3 describes the finite element model of the tibia and the pros-
thesis implanted into the tibia, while Section 4 details the calculation of
the mechanical properties of the implant building block. In Section 5,
we present the governing equations of a gradient-based topology opti-
mization scheme that is used to optimally design the 3D architecture of
the porous knee-implant. Finally, the performance of the proposed ti-
bial knee implant, in particular its interface micromotion and bone
resorption, is numerically assessed and compared to that of two base-
line concepts, a fully solid one that is commercially available, and a
fully porous tibial implant identical to the fully solid one, but with
uniform distribution of pores.

2. Methodology

In this work, we present a fully porous architectured biomaterial for a
tibial knee implant, specifically a stemmed tibial component, with
tailored properties and macro geometry identical to that of a com-
mercially available tibial component, as explained in detail in the fol-
lowing section. Fig. 1 briefly depicts the numeric strategy proposed
here. The mechanical properties of the building block are expressed as
a function of its relative density, and their optimal gradients are de-
termined via topology optimization starting from the tissue properties
of the tibial native bone of a patient.

The major aspects of the proposed methodology rest on the in-
tegration of multiscale solid mechanics and hierarchical optimal design
of materials (Coelho et al., 2011, 2008), as briefly summarized in the
steps below:

- CT scan data from a 38 year old male are used to create the nu-
merical model of his tibial geometry and bone tissue properties.
- An open cell, a Tetrahedron-based topology, is selected to modularly
build the interior architecture of the porous tibial stem. The high
strength characteristics of this unit cell ensure a minimum level of
fatigue and static resistance, as deemed necessary to resist the set of
repetitive loads the knee is subjected to Arabnejad et al. (2016).
- Asymptotic homogenization is used to calculate the elastic constants
of the unit cell, with characteristic length much smaller than the
implant, as a function of its relative density (Fang et al., 2005;
Zienkiewicz and Taylor, 2005).
- A first trial uniform distribution of relative density is assigned to the
implant and 3D finite element analysis is undertaken. From this
analysis, the stress and displacement regime over the geometry of
the prosthesis is obtained and used to formulate a multi constraint
topology optimization problem. On the microscale, the stress dis-
tribution over the lattice architecture is retrieved and used to de-
termine the implant safety factor (SF) under static and fatigue
conditions.
- To minimize bone resorption and interface micromotion around the
tibial stem, topology optimization is solved for maximum implant
compliance. The problem is also subjected to a set of inequality and
equality constraints, which include average porosity of the cellular
implant, along with safety factors for first cycle and infinite fatigue
Implant compliance is obtained from the calculation of the total strain energy of the implant. Implant displacements at each mesh element node along with unit cell homogenized properties are then used to build the macro stiffness tensor of each element. The gradient of the objective function is then obtained through partially taking the derivatives of the stiffness tensor components with respect to the relative density.

The design variables are grouped in the vector, $\mathbf{b}$, which collects the relative density of all elements and is updated via the standard optimality criterion. The optimization process continues until the attainment of the optimized distribution of relative density.

### 2.1. Finite element model

The three-dimensional (3D) finite element (FE) model of the tibial fixation is constructed through CT images from the left tibia of a 38 year old male with a body weight of 900 N, as shown in Fig. 2. The properties of bone are assumed to be isotropic, a simplification that does not lead to a considerable difference from the results that consider bone as orthotropic (Baca et al., 2008; Peng et al., 2006a). The effective Young’s modulus of each element is obtained through the apparent density of the corresponding element. The Hounsfield values (HU) obtained through the CT images are used to determine the apparent density of bone by using a linear interpolation between HU and apparent density. As a result, the effective elastic properties of bone are expressed as a function of the apparent density (Baca et al., 2008; Austman et al., 2008; Peng et al., 2006b). The procedure followed to compute the mechanical properties of the tibia is detailed in Appendix A (See Appendix Fig. A.1).

The macrogeometry of the stemmed tibial knee implant was taken from a NexGen component (Zimmer, Warsaw, IN) commercially available and made of solid titanium. The tibial dimensions were measured and the tibia was templated using NexGen templates to determine the appropriate size of the tibial component and the tibial stem. The component was sized to ensure the central portion of the tibial tray was externally rotated to lie over the medial 1/3 of the tibial tubercle while choosing the largest size possible that would cover the proximal tibia, 10 mm below the articular surface, but did not overhang. The tibial stem was chosen to fill the proximal tibial diaphysis. An offset tibial stem was required to permit anatomic positioning of the tibial tray and ensure centralization of the stem. This resulted in using a size 5 NexGen stemmed tibial implant with a 4 mm offset 100 mm stem extension, creating in a 145 mm offset stem. These implants were CT scanned to create the parameters used in this study. The identical design was then used to convert the solid stemmed tibial implant into the optimized 3D porous implant here proposed.

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**Fig. 1.** Flow chart illustrating the analysis and design scheme used to develop a graded cellular knee implant minimizing bone resorption and interface micromotion. Legend: mean stress: $\sigma_{ij}^m$, alternating stress: $\sigma_{ij}^a$, yield stress of tetrahedron cell unit: $\sigma_{ij}^y$, implant compliance: $c$, relative density of tetrahedron cell unit: $\rho$, minimum element density: $\rho_{\text{min}}$, maximum element density: $\rho_{\text{max}}$, displacement vector: $U$, stiffness matrix of tetrahedron unit cell: $K_i$, volume constraint: $f$, predefined volume of the implant: $V_0$, force vector: $F$, implant safety factor: $SF$, minimum allowable safety factor: $SF_{\text{min}}$. 

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**Fig. 3** shows the geometry of the intact and implanted bone along with the applied loads and boundary conditions. The distal end of the bone is fixed to avoid rigid body motion. As clinically recommended, the exterior of the implant stem is made of a thin shell that eases the implant removal during revision surgery. This ensures that in a postoperative situation no bone ingrowth occurs onto the stem, which is press fitted into the cortical bone. On the other hand, the implant tray is kept fully porous, and for this reason bone ingrowth can occur beneath it. To accurately capture interface stability, two distinct contact models are implemented in this work, one for the tray and one for the stem below. The tibial tray is assumed fully bonded to the underlying bone of the tibial plateau; this scenario represents the incidence of bone ingrowth onto the tray. One benefit of this choice is the reduction in the computational cost required for the stability analysis on a non-linear
frictional contact model (Viceconti et al., 2000). On the other hand, a frictionless contact is assumed for the tibial stem below the tray, where a thin solid shell covers the implant. This contact model represents the weak bonding between the smooth metal and the surrounding bone tissue (Viceconti et al., 2000; Hipp et al., 1985). At the bone implant interface, face to face contact elements with appropriate mesh type are used with results that show interfacial micromotion at an average error of about 10 µm (Viceconti et al., 2000). The entire stem surface is selected as the target member and the inner bone surface is selected to be the contact member. The Augmented Lagrangian method, where contact penetration is present but controlled to some degree, is used in the contact formulation and contact tractions are monitored at the Gauss points. A contact stiffness of 1 is considered at the contact regions. The time step control is set to be automatic, where the contact behaviour is reviewed at the end of each substep to ensure that neither drastic change in contact status nor excessive penetration occurs.

Four different load cases corresponding to the 20%, 30%, 40% of the gait cycle and deep-knee bend are all studied. The system of loads reported in Table 1 and adopted in this work represents the daily physiological loads experienced by a human. Obtained from in-vivo measurements on an instrumented knee joint, those values pertain to a subject with a male body weight of 1000 N and provide contact load measures with an error below 2% (Bergmann et al., 2014; Heinlein et al., 2009; Kutzner et al., 2010). The values in Table 1 are used with
equal weights to calculate an equivalent load pattern that represents concurrently walking and deep knee bend, and is here used for the implant design.

2.2. Homogenized mechanical properties

Mechanics and optimization of implant microarchitecture are here undertaken by treating the implant as a homogenized continuum, an assumption that allows to avoid detailed finite element simulations, which would be extensive and lengthy to handle (Arabnejad and Pasini, 2013). By doing so, we can focus on a representative volume element (RVE) of the implant, and consider its properties as those of a homogenized medium equivalent to the implant itself. Below is a description of the method used in this paper to calculate the stiffness tensor and implant fatigue resistance, which both depend on multiscale properties, specified at the macro and micro scale.

2.2.1. Elastic properties of unit cell

To obtain homogenized properties of the implant, we use asymptotic homogenization (AH) theory, which enables to calculate the stress and strain distribution developed within the cellular architecture, namely microstrain and microstress, from which the implant stiffness tensor can be obtained. The yield and ultimate strength of the lattice are also assessed for relative density $\rho \leq 0.3$ and used to capture the fatigue strength of the designed implant via the Soderberg fatigue criterion under multi axial stresses (Masoumi Khalil Abad et al., 2013).

The effective properties of the building block $E^{II}_{ijkl}$ are obtained by solving a local problem formulated on the RVE and defined as (Hollister and Kikuchi, 1992):

\[
E^{II}_{ijkl} = \frac{1}{|Y|} \int_Y E_{ijpm}M_{pmkl}dY
\]  

where $|Y|$ is the unit cell volume, $E_{ijpm}$ is the positive elastic tensor of the unit cell that varies between zero and the material elastic tensor corresponding to the voids and bulk material respectively. Furthermore, we define a structure tensor $M_{ijkl}$ that relates the local macro strains $\varepsilon_i$ to the micro strains as

\[
\varepsilon_i = M_{ijkl}\varepsilon_j
\]  

where $\delta_{ij}$ is the Kronecker $\varepsilon_i$, and $\varepsilon^{skl}_i$ is the microscopic stress corresponding to the component $kl$ of the macroscopic strain. Given that for this application small deformation and linear elasticity hold, the microscopic strain $\varepsilon^{skl}_i$ can be expressed as

\[
\varepsilon^{skl}_i = \int_Y E_{ijpm}C^{skl}_{ijm}(v)\tau_{pm}(u)du = \int_Y E_{ijpm}C^{skl}_{ijm}(v)\tau_d du
\]  

where $C^{skl}_{ijm}(v)$ denotes the virtual strain.

In three dimensions, six arbitrary macroscopic unit strains are required to construct the $M_{ijkl}$ matrix. The periodicity of the strain field is ensured by imposing periodic boundary conditions on the RVE edges (Hollister and Kikuchi, 1992; Hassani, 1996). As a result, the nodes of the opposite planes are set with identical displacement. Once the structure tensor, $M_{ijkl}$, is obtained, the homogenized stiffness tensor of the unit cell can be predicted by substituting the structure tensor $M_{ijkl}$ into (2a), (2b). The microscopic stress distribution, $\sigma_{ij}$, can be defined by

\[
\sigma_{ij} = E^{II}_{ijkl}M_{klmn}\varepsilon_{mn}
\]

Using the microscopic stress tensor and introducing the homogenized elastic tensor, $E^{II}_{ijkl}$, results in a simplified relationship between the microscopic stress distribution and the macroscopic stress tensor

\[
\sigma_{ij} = E^{II}_{ijkl}M_{klmn}(E^{III}_{lmnk})^{-1}\varepsilon_{mn}
\]

wherein $\sigma_{ij}$ is the macroscopic stress distribution applied to the RVE. If $\sigma_{ij}$ is the yield strength of the cell walls, then the yield surface of the unit cell can be written from (5) as

\[
\int_Y E^{II}_{ijkl}M_{klmn}\varepsilon_{mn}
\]

where $\sigma_{ij}$ is the von-Mises stress distribution within the unit cell caused by the applied macroscopic stress $\sigma_{ij}$.

Table 1

<table>
<thead>
<tr>
<th>No.</th>
<th>Load case</th>
<th>Force/N</th>
<th>Moment/N mm</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>$F_1$</td>
<td>$F_2$</td>
</tr>
<tr>
<td>1</td>
<td>Level walking</td>
<td>20% of gait cycle</td>
<td>-76.05</td>
</tr>
<tr>
<td>2</td>
<td></td>
<td>30% of gait cycle</td>
<td>-144.65</td>
</tr>
<tr>
<td>3</td>
<td></td>
<td>40% of gait cycle</td>
<td>-71.96</td>
</tr>
<tr>
<td>4</td>
<td>Level bending</td>
<td>Deep knee bend</td>
<td>75.8</td>
</tr>
</tbody>
</table>

2.2.2. Fatigue properties

Implant stiffness tailoring is undertaken against high cycle fatigue failure such that stress values generated within the material architecture do not exceed the yield strength of Ti6Al4V, the material the implant is made of. Hence, to obtain the fatigue surface of the implant, we multiply the RVE yield strength $\sigma_y$ by the ratio of the endurance limit $\sigma_e$ and yield strength $\sigma_y$ of the solid material. This gives

\[
\frac{\sigma_y}{\sigma_e} = \frac{\sigma_y^{max}}{\sigma_y^{min}}
\]

Once the fatigue surface is obtained, the Soderberg diagram, a conservative criterion, can be constructed to compute the fatigue safety factor, $S_{Fatigue}$, from the relation:

\[
\frac{\sigma_y}{\sigma_y^{max}} + \frac{\sigma_y}{\sigma_y^{min}} = \frac{1}{S_{Fatigue}}
\]

wherein, $\sigma_y^{max}$ and $\sigma_y^{min}$ denote the mean and alternating stresses respectively, given by:

\[
\sigma_y^{max} = \frac{\sigma_y^{max} + \sigma_y^{min}}{2}
\]

\[
\sigma_y^{min} = \frac{\sigma_y^{max} - \sigma_y^{min}}{2}
\]

With $\sigma_y^{max}$ and $\sigma_y^{min}$ being the multiaxial macroscopic stresses that cause, respectively, the highest and the lowest von-Mises stress within the unit cell. From Eqs. (7) and (8), the static safety factor of the RVE is expressed as:
\[
S_{E_{\text{active}}} = \frac{\sigma_{ij}^e}{\sigma_0}
\] (10)

The procedure above is used for the analysis of the tibia knee implant which is modelled with homogenized properties representing the tetrahedron-based cell made of Ti6Al4V (Parthasarathy et al., 2010).

The properties of Ti6Al4V here adopted (Young’s modulus: 120 GPa, Poisson’s ratio: 0.3, yield strength: 900 MPa, fatigue strength: 600 MPa at 10^7 cycles) are measured from mechanical testing of 3d-printed samples manufactured via Selective Laser Melting (Wycisk et al., 2013). SLM is the additive process that will be used for implant fabrication. In this work, we rely on the assumption that defects and imperfections emerging from the manufacturing process would not severely impact the mechanical properties (Arabnejad et al., 2017, 2016). Further work, currently undergoing, is required to include the influence of manufacturing deviations in the implant analysis.

3. Optimization scheme

A density-based topology optimization is here adopted to optimally tailor the material distribution in the stem of the tibial component (Liu and Tovar, 2014). The search for optimized density gradients is undertaken on an equivalent medium with homogenized properties obtained in Section 2.2.1 and expressed as a function of relative density (Fig. 4). A penalization scheme that penalizes elements with intermediate density values is used to create an implant with continuous density distribution (Zhou and Rozvany, 1991).

3.1. Problem formulation

Elastic stiffness tailoring aims at reducing the elastic modulus mismatch between implant microstructure and native bone, resulting in lower bone resorption. As such, to minimize bone-stem interface micromotion and bone resorption, the optimization problem is posed for minimum stiffness, which is equivalent to maximize implant compliance (Ridzwan et al., 2007), here expressed as

\[
c(\chi) = F^T U(\chi)
\] (11)

where \( F \) is the vector of the nodal forces applied to the implant and \( U(\chi) \) is the vector of nodal displacement. \( F \) can be expressed as

\[
F = K(\chi) U(\chi)
\] (12)

where \( K(\chi) \) is the total stiffness matrix of the implant. The problem is subjected to multiple constraints, e.g., on volume, density, and safety factor, and can be formally formulated as:

\[
\begin{align*}
\text{Find} & \quad x = [x_1, x_2, ..., x_n] \\
\text{Maximize} & \quad c(\chi) = F^T U(\chi) \\
\text{subject to} & \quad \nu(\chi) = x^T v - \nu \leq 0 \\
& \quad x \in \chi \quad \chi = \{x \in \mathbb{R}^n : x_{\text{min}} \leq x \leq x_{\text{max}}\} \\
& \quad SF_{\text{min}} < SF
\end{align*}
\] (13)

where \( \bar{x} \) is the density vector which is filtered through a basic filter function described in Appendix B, \( V \) is the total number of the elements that discretizes the implant domain, \( V = [v_1, ..., v_n] \) is the vector of the elements volume and \( \bar{v} \) is a prescribed volume of the tibial knee implant.

The optimization problem defined in (13) can be reformulated as a traditional problem of minimization by multiplying the objective function by \(-1\), which results in

\[
\begin{align*}
\text{Find} & \quad x = [x_1, x_2, ..., x_n] \\
\text{Minimize} & \quad c(\chi) = -F^T U(\chi) \\
\text{subject to} & \quad \nu(\chi) = x^T v - \nu \leq 0 \\
& \quad x \in \chi \quad \chi = \{x \in \mathbb{R}^n : x_{\text{min}} \leq x \leq x_{\text{max}}\} \\
& \quad SF_{\text{min}} < SF
\end{align*}
\] (14)

The derivative of the objective function can be determined as:

\[
\frac{\delta c(\chi)}{\delta x} = \sum_{i \in N_e} \frac{\delta c(\chi)}{\delta \bar{x}_i} \frac{\delta \bar{x}_i}{\delta x_i}
\] (15)

where \( \bar{x}_i \) represent the filtered density of element \( i \) and \( \frac{\delta \bar{x}_i}{\delta x_i} \) is defined by

\[
\frac{\delta \bar{x}_i}{\delta x_i} = \frac{H_{x_i} v_i}{\sum_{j \in N_e} H_{x_j} v_j}
\] (16)

where \( H_{x_i} \) is a weight factor matrix as described in Appendix B. From (11), \( \frac{\delta c(\chi)}{\delta x} \) can be defined by

\[
\frac{\delta c(\chi)}{\delta x} = -F^T U(\chi) \frac{\delta \chi}{\delta \bar{x}} = -U^T(\chi) K(\chi) \frac{\delta U(\chi)}{\delta \bar{x}}
\] (17)

where \( F^T \) and \( U^T(\chi) \) indicate the transpose matrix of \( F \) and \( U(\chi) \) respectively. Taking the derivative of (12) with respect to \( \bar{x}_i \) yields
4. Results and discussion

The methodology described in Section 2 is here applied for the design of a fully porous knee implant. Two sets of numeric results are herein presented. The first describes the micro-architecture of the proposed tibial implant, and the second its performance, namely bone resorption and bone-stem interface micromotion, which are then compared to those of the fully solid titanium tibial implant and a fully porous one with uniform relative density.

4.1. Implant architecture

Fig. 5 shows the von Mises distribution and optimal density distribution of the microstructure of the proposed tibial implant which features low porosity at the distal region where severe stresses are located. As can be observed, stress concentration appears beneath the tray due to the variation in implant macrogeometry. In addition, the applied moment in the frontal plane, which represents the Varus-Valgus movement of knee joint, leads to high stress values at the distal region. The tibial implant is fully porous with very low porosity in the tibial tray. In particular, unit cells with high value of relative density ranging 0.7–0.8 are located at regions with high stress in the tibial tray, which ensures a minimum level of fatigue resistance. Hence, underneath the tray as well as in the lower part of the stem, cells with high relative density are necessary to meet the level of fatigue resistance the implant should provide.

From the relative density distribution, the lattice microarchitecture is created through in-house mapping scripts. Fig. 5 shows that while the implant internal architecture is fully porous, a very thin shell is placed on the exterior of the stem only. This feature is designed to avoid bone ingrowth onto the stem and thus facilitate the process of implant replacement at the time of revision surgery. While the fully solid tibial implant shows fatigue strength 2 times higher than the cellular tibial implant, the latter has a safety factor of 3, which is well within the margin of safety for a biomedical device (ASTM F1800-12, 2012). If a further increase in the implant fatigue strength is desired, either a lattice with smooth cell geometry could be implemented (Masoumi Khalil Abad et al., 2013), or other part of the implant could be designed as fully dense.

4.2. Implant micromotion

The primary stability of the implant is crucial to the success of the total knee arthroplasty (TKA) (Small et al., 2016). An extensive body of the literature has demonstrated that the elastic modulus mismatch from the bone tissue to the implant is one main cause of the lack of stability at the bone-implant interface (Glenn et al., 2010; Completo et al., 2012). Although other factors such as the implant macrogeometry, the fixation technique used at the bone-implant interface, and the design platform of the implant play an important role in the development of interface micromotion (Completo et al., 2012; Small et al., 2016; Conlisk et al., 2012), in this work we address the elastic modulus mismatch. To do so, we perform a relative comparison between implants that share identical macrogeometry and prescribed clinical conditions, such as the fixation type at the bone implant interface. The differences in interfacial micromotion that we observe are therefore relative, and can be exclusively attributed to the distribution of elastic modulus that each tibial implant here examined features.

Figs. 6 and 7 illustrate the distribution of bone-stem interface micromotion at 30% of gait cycle and deep bend, respectively. The amount of micromotion is computed from the relative nodal sliding distance of mesh elements from bone and stem. Stable fixation at the bone-stem interface with micromotion between 20 and 50 μm provides a desirable range, since this degree of micromotion is known to be well tolerated by the periprosthetic bone, and in cases of porous implants, is

Fig. 5. (a) Von-Mises stress distribution; (b) optimum relative density distribution to ensure adequate fatigue resistance against daily cyclic loads; (c) graded cellular implant with tailored porosity in the stem ranging 0.3–0.7. A thin solid exterior (shown only partially in the figure) coats the stem, a clinically recommended feature introduced to ease implant removal at the time of revision surgery; (d) internal architecture of the tibial implant, where the thin shell is here omitted for a global visualization of the whole internal microstructure.
associated with bone ingrowth (Chong et al., 2010). This range of micromotion can also contribute to a notable reduction of the end-of-stem pain that arises from excessive interface micromotion. The percentage of surface area (SA) with micromotion below 50 µm is then assessed for the designed graded lattice implant and the two baselines, a fully dense tibial implant and a uniform porosity tibial implant, all made of Ti6Al4V.

Both the loading conditions of gait cycle and deep knee bend led to severe micromotion around the distal part of the stem, particularly around the tip of the stem. High values of micromotion can cause the patient to feel pain at this region. At 30% of gait cycle, the graded lattice tibial implant stem resulted in a maximum micromotion at the stem tip that is reduced by 17 µm and 10 µm when compared respectively to a fully solid and a uniform lattice tibial implant stem. Although the maximum micromotion is reduced by only 15% when compared to a fully solid implant, from a clinical point of view this result suggests a lower potential for postoperative end-of-stem pain in patients. As can be seen, the surface area with micromotion below 50 µm for the graded tibial implant (grey regions of the implant in Fig. 6c) is larger than that of the two other implants (Fig. 6a and b). This value of micromotion ensures an improved stability between the tray periphery and the distal part of the stem. During deep knee bend, a reduction in micromotion of the tip of the stem of 22 µm and 14 µm with respect to a fully dense titanium tibial implant and uniform porosity tibial implant were achieved. This amount represents a 14% reduction in the maximum micromotion at the stem tip for the graded lattice implant compared to the traditional solid implant. This provides the graded tibial implant with 76% prosthesis area with micromotion below 50 µm, values that are clinically stable.

Among the nodal displacement vectors, we now examine the displacement distribution at the mid (i) and distal areas (ii) of the implant, and plot the results in Fig. 8. Here, is the micromotion distribution visualized for the loading conditions of walking and deep bend. 50 Points around the stem surface have been specified with the first one

![Fig. 6. Interface micromotion distribution at 30% of gait cycle for a fully dense titanium implant (a), cellular implant with uniform relative density of 60% (b), and graded cellular implant (c). SA: surface area of the prosthesis (Chong et al., 2010).](image6)

![Fig. 7. Interface micromotion distribution at deep bend for a fully dense titanium implant (a), cellular implant with uniform relative density of 60% (b), and a graded cellular implant (c). SA: surface area of the prosthesis (Chong et al., 2010).](image7)
located at the anterior of the cross section and the remaining ones following a counter clockwise direction. The figure shows the relative sliding micromotion values with respect to the fully solid implant for deep bend and 30% of gait cycle. As can be seen, maximum micromotion occurs anteriorly for both loading conditions. The interface micromotion for the graded lattice implant is lower relative to both baseline implants at almost all the points. Although the relative reduction is small, this result offers a lower probability of local interface failure and end-of stem pain.

4.3. Bone resorption

To further investigate the improved performance of the implant here introduced, we compute its amount of bone loss and compare to that of a fully dense and a uniform porosity tibial implants ($\rho = 60\%$). Fig. 9 shows the results with bone resorption levels plotted around the tibial stems in the three cases. The results are obtained by computing the amount of underloaded bone (Arabnejad Khanoki and Pasini, 2012, 2013a, 2013b; Arabnejad et al., 2017). In particular, bone is assumed to start loosing its mass when the local strain energy ($U_i$) per unit of bone mass ($\rho$) averaged over ($n$) loading cases ($S = ((1/n) \sum_{i=1}^{n} U_i/\rho)$) in the postoperative situation is beneath the respective local strain energy of the implant preoperatively ($S < (1-s) S_{ref}$). $S_{ref}$ indicates the local strain energy of the intact bone and signifies a specialized threshold level or dead-zone for the bone to start degrading after implantation. As a result, the amount of resorbed bone mass can be expressed as

$$m_r(b) = \frac{1}{M} \int_{V} g(S(b)) \rho dV$$

(21)

where $M$ and $V$ are the mass and volume of the original bone and $g(S(b))$ is a resorptive function equal to unity if $S < (1-s) S_{ref}$ and otherwise is equal to zero. The value considered in this study for the dead-zone is 0.75 (Kuiper and Huiskes, 1992).

We observed that the computed amount of resorbed bone was consistent with the bone-stem stability. The less the bone-stem interface micromotion, the lower bone resorption. As can be seen, significant bone resorption appears proximally underneath the tray throughout the medial compartment of the tibia and propagates toward the lateral part. The concentration of bone resorption in the medial compartment can be attributed to the Varus-Valgus movement of the knee joint which causes a larger portion of the vertical load to be distributed over the medial part of the tibiae.

In Fig. 10, we compare the amount of bone resorption at four different regions of the implanted tibia. As previously described, bone resorption is maximum at the proximal region, whereas at the distal regions an overstressed bone results in bone formation (Ebrahimi et al., 2012). For the solid titanium stem, an overall bone resorption of 40% can be predicted with the greatest degree of resorption occurring in zone 3. Due to the higher compliance of the uniform lattice tibial implant, we observe an overall decrease of resorbed bone of 16%, i.e. two times lower than the solid tibial implant. The least amount of bone
resorption was seen around the graded cellular implant, with an overall decrease of 26%, 3 times less than that seen around the solid tibial implant. In summary with respect to the fully solid baseline tibial knee implant, Fig. 10 shows a reduction of bone resorption around the graded lattice tibial implant of 50% at zone 4, 76% at zone 3 54% at zone 2, and 18% at zone 1.

Recent technologies for additive manufacturing (AM), such as electron beam melting (EBM) and selective laser melting (SLM), bring versatile layer-by-layer processes that enable the fabrication of porous materials with tailored cellular architecture (Melancon et al., 2017). For example, SLM, a powder bed fusion technology, has been used to generate parts with improved mechanical, tribological and corrosion properties (Konda Gokulldoss et al., 2017). Recent works have demonstrated that AM can successfully build metallic lattice structures including porous implants with complex internal microarchitecture (Melancon et al., 2017; Vaezi et al., 2013; Balla et al., 2010; Moiduddin et al., 2017; Tunchel et al., 2016; Liu et al., 2017). In addition, AM facilitates the fabrication of cellular implants with tailored gradients of porosity and pore morphology that enables bone ingrowth (Sobral et al., 2011; Khoda et al., 2011). A fully porous hip implant featuring a lattice microstructure similar to the one presented in this paper has been recently manufactured through SLM and successfully tested in vitro (Arabnejad et al., 2017). Its graded microarchitecture features a minimum strut thickness of 200 µm and a maximum pore size of 800 µm, characteristics that are built through AM and are shared by the knee implant presented in this study (Arabnejad et al., 2016; Melancon et al., 2017). These previous works therefore demonstrate the feasibility of additively manufacturing the knee implant architecture herein reported.

Exploratory in nature, this numeric study holds some limitations that need to be addressed in the future. The first one is that a simplified loading system representing a high body weight has been used to perform the numerical analysis. Although this system of loadings describes

![Fig. 9. Distribution of bone resorption in knee prosthesis around (a) fully dense titanium implant; (b) cellular implant with uniform relative density of 60%; and (c) graded cellular implant. L: Lateral, M: Medial, A: Anterior, P: Posterior, AL: Anterolateral, AM: Anteromedial.](image)

![Fig. 10. Percentage of bone resorption with respect to a fully solid tibial implant here taken as a baseline for (i) graded cellular implant, and (ii) uniform cellular implant with relative density of 0.6.](image)
the worst-case scenario that can occur during normal daily activities, the real load scenarios that a knee joint undergoes are more complex. In particular, this work did not consider the Varus-Valgus movements that often result in a higher amount of load on the medial compartment than the lateral one. In addition, the magnitude of contact forces varies between ordinary activities and the phase of activity, thus resulting in a complex relationship between the flexion angle, the maximum joint load and the balance of medial to lateral load distribution for different activities. Another aspect that requires further investigation is the sensitivity of the results, e.g. the amount of interface micromotion, to variations in bulk material properties, bone properties, loading conditions and the contact model used at the bone-implant interface.

5. Conclusion

This study describes the computational design and numerical assessment of a stemmed tibial component with a fully porous stem with tuned tetrahedron lattice architecture. Although the results show an opportunity to improve the clinical performance of the current tibial implants, further experimental and numerical studies are required to validate the clinical applicability of the proposed design. The implant porosity has been optimally tailored to mitigate postoperative bone resorption and end-of-stem pain while satisfying the strength requirement necessary to sustain cyclic loadings. The mechanical properties of the implant are optimally tuned to bring about a concomitant reduction of bone resorption and interface micromotion. The numerical results suggest that the proposed concept could lead to a reduction of 17 µm and 21 µm in micromotion at the tip of the stem during gait cycle and deep knee bend, respectively. Clinically, this would be expected to alleviate the problem of end-of-stem pain that is not uncommon after total knee arthroplasty. In addition, the numerical results suggest a reduction in bone resorption of 26% with respect to an identical tibial knee implant made of solid materials. This level of performance would translate in decreased periprosthetic fractures and enhanced bone stock at the time of revision surgery.

Appendix A. Assigning bone material properties using computed tomography (CT) data

The heterogeneous material properties of the tibia are captured through computed tomography (CT) data obtained from a 38 year old male, provided by the Visible Human project (United States National Library of Medicine, Bethesda, MD). The radiographic density of the CT images quantified as Hounsfield Unit (HU) is used to calculate the local properties of the tibia. Since the relationship between the HU and bone density is monotonic, a linear relationship between the bone apparent density and the HU is adopted as shown in Figure (Peng et al., 2006a). The bone apparent density represents the density of solid bone excluding the density of the fluid mass, namely the density of blood. On the other hand, the bone effective density accounts for the fluid mass. At regions where there is no bone, the effective density would be about 1024 kg/m³, which represents the density of blood. However at these regions, the apparent density and the HU value are zero.

To obtain the mechanical properties of the tibia, the apparent density for each element of the finite element model is determined from the HU value measured from the CT data ranging from 0 HU to 1567 HU. This ensures that the density of the fluid has no contribution to the mechanical properties of the tibia. The maximum value of HU corresponds to the densest region of the cortical bone with an apparent density of 2000 kg/m³. The Young modulus of the tibia is then obtained using the relation

\[
\begin{align*}
E &= 1904 \rho^{1.64} \quad \rho < 0.95 \\
E &= 2065 \rho^{0.99} \quad 0.95 < \rho \\
v &= 0.3
\end{align*}
\]

where \(E\) is the elastic modulus of bone, and \(v\) is its Poisson’s ratio. The properties are assumed to be isotropic, a simplification that does not lead to a considerable difference from the results that consider bone as orthotropic (Baca et al., 2008; Peng et al., 2006a). This assumption contributes also to reduce the computational cost.

Fig. A.1. Linear relationship between Hounsfield number and both effective density and apparent density.

Appendix B. Filter density function

Early formulations of topology optimization problems generally yield instabilities in the optimal solutions that in turn affect result accuracy. A continuous density distribution is an asset for the implant strength while a discontinuous density distribution in the implant microstructure leads to stress concentration and compromise the implant strength thus increasing the probability of local failure. To avoid binary results (black and white
patterns) for the density distribution, we use the following filter density function (Sigmund and Petersson, 1998), to obtain the mechanical properties of each finite element mesh:

\[ \tilde{s}_i = \frac{\sum_{j \in N_i} H_{ij} v_j}{\sum_{j \in N_i} H_{ij}} \]  

(B.1)

wherein, \( N_i \) corresponds to neighborhood elements of element \( i \), with volume of \( v_i \).

\( H_{ij} \) is a weight factor matrix determined as follows:

\[ H_{ij} = R - \text{dist}(i, j) \]  

(B.2)

where \( R \) is the size of the neighborhood that is referred to the filter size and \( \text{dist}(i, j) \) is the distance between the element \( i \) and the center of the element \( j \). The neighborhood of an element is defined as

\[ N_i = \{ j \mid \text{dist}(i, j) \leq R \} \]  

(B.3)

Appendix C. Derivation of the stiffness tensor for the implant internal microstructure

To compute the sensitivity of the strain energy of the implant microstructure, as described in Section 3.1, the derivative of the stiffness tensor is required for the entire implant. We use here a direct stiffness approach to find the global stiffness tensor, where the implant is discretized into small elements, and the elastic tensor of each element is calculated before the global stiffness matrix, \( K \), assembly, where \( K \) is expressed as

\[ K = \int_\Omega B^T D_i B_i \, dV \]  

(C.1)

\( B \) is the strain-displacement matrix and \( D \) is the elastic tensor of element \( i \), and \( \Omega \) is the total volume of the implant. Since the strain-displacement matrix is independent of the design variables, the derivation of the stiffness matrix with respect to the design variables (relative density of each element) can be expressed as follows:

\[ \frac{\delta K}{\delta x_i} = \int_\Omega B^T \frac{\delta D}{\delta x_i} B_i \, dV \]  

(C.2)

Each mesh element corresponds to a tetrahedron-based cell that has three planes of symmetry; therefore 9 elastic constants are needed in the constitutive equations: 3 Young’s moduli \( E_x, E_y, E_z \), 3 Poisson’s ratios \( \nu_{yz}, \nu_{zx}, \nu_{xy} \) and 3 shear moduli \( G_{yz}, G_{zx}, G_{xy} \). The stiffness matrix of the unit cell can thus be expressed as

\[ D = \begin{bmatrix} \frac{1 - \nu_{yz} \nu_{zx}}{E_y E_z} & \frac{\nu_{yz} + \nu_{zx} \nu_{xy}}{E_y E_z} & \frac{\nu_{zx} + \nu_{xy} \nu_{yz}}{E_x E_z} & 0 & 0 & 0 \\ \frac{\nu_{yz} + \nu_{zx} \nu_{xy}}{E_y E_z} & \frac{1 - \nu_{yz} \nu_{xy}}{E_y E_z} & \frac{\nu_{zx} + \nu_{xy} \nu_{yz}}{E_x E_z} & 0 & 0 & 0 \\ \frac{\nu_{zx} + \nu_{xy} \nu_{yz}}{E_x E_z} & \frac{\nu_{zx} + \nu_{xy} \nu_{yz}}{E_x E_z} & \frac{1 - \nu_{zx} \nu_{xy}}{E_x E_z} & 0 & 0 & 0 \\ 0 & 0 & 0 & 2G_{yz} & 0 & 0 \\ 0 & 0 & 0 & 0 & 2G_{zx} & 0 \\ 0 & 0 & 0 & 0 & 0 & 2G_{xy} \end{bmatrix} \]  

(C.3)

Since AH is used to obtain the elastic constants of the unit cell across a range of relative densities, with results shown in Fig. 4, we can represent the elastic tensor of each element as a function of the element relative density and then use it to evaluate the derivative of the elastic tensor with respect to the design variables. The strain-displacement matrix is also derived by differentiating the displacements expressed through shape functions and nodal displacements. A 10 nodes isoparametric tetrahedron element is used to mesh the FE model of the implanted tibia. Hence, the strain-displacement matrix can be written as

\[ B = [B_1 \ B_2 \ B_3 \ B_4 \ B_5 \ B_6 \ B_7 \ B_8 \ B_9 \ B_{10}] \]  

(C.4)

With

\[ B_k = \begin{bmatrix} N_{k,x} & 0 & 0 \\ 0 & N_{k,y} & 0 \\ 0 & 0 & N_{k,z} \\ 0 & N_{k,y} & N_{k,z} \\ 0 & N_{k,z} & N_{k,y} \\ N_{k,x} & 0 & N_{k,z} \end{bmatrix} \quad k \in \{1, 2, \ldots, 10\} \]  

(C.5)

where \( N_{k,x}, N_{k,y} \) and \( N_{k,z} \) are the derivative of the shape functions with respect to the global coordinate system. The shape functions of a tetrahedron element with respect to the isoparametric coordinate system given by \( r, s \) and \( t \) (see figure below) can be defined as
\( N_1 = 2s^3 - s \)
\( N_2 = 2r^3 - t \)
\( N_3 = 2r^2 - r \)
\( N_4 = 2s^2 + 4rs + 4rt - 3r + 2s^3 + 4st - 3s + 2t^2 - 3t + 1 \)
\( N_5 = 4st \)
\( N_6 = 4rt \)
\( N_7 = 4rs \)
\( N_8 = 4s - 4rs - 4st - 4s^2 \)
\( N_9 = 4t - 4rt - 4st - 4t^2 \)
\( N_{10} = 4r - 4rs - 4rt - 4r^2 \)

To construct the strain-displacement matrix, the derivative of the shape functions with respect to the generalized coordinate system is computed via the chain rule as

\[
\frac{\partial N_i}{\partial \xi} = \frac{\partial N_i}{\partial x} \frac{\partial x}{\partial \xi} + \frac{\partial N_i}{\partial y} \frac{\partial y}{\partial \xi} + \frac{\partial N_i}{\partial z} \frac{\partial z}{\partial \xi} \tag{C.7a}
\]
\[
\frac{\partial N_i}{\partial \eta} = \frac{\partial N_i}{\partial x} \frac{\partial x}{\partial \eta} + \frac{\partial N_i}{\partial y} \frac{\partial y}{\partial \eta} + \frac{\partial N_i}{\partial z} \frac{\partial z}{\partial \eta} \tag{C.7b}
\]
\[
\frac{\partial N_i}{\partial \zeta} = \frac{\partial N_i}{\partial x} \frac{\partial x}{\partial \zeta} + \frac{\partial N_i}{\partial y} \frac{\partial y}{\partial \zeta} + \frac{\partial N_i}{\partial z} \frac{\partial z}{\partial \zeta} \tag{C.7c}
\]

Using (C.7a), (C.7b), (C.7c) results in

\[
\begin{bmatrix}
\frac{\partial N_1}{\partial \xi} & \frac{\partial N_1}{\partial \eta} & \frac{\partial N_1}{\partial \zeta} \\
\frac{\partial N_2}{\partial \xi} & \frac{\partial N_2}{\partial \eta} & \frac{\partial N_2}{\partial \zeta} \\
\frac{\partial N_3}{\partial \xi} & \frac{\partial N_3}{\partial \eta} & \frac{\partial N_3}{\partial \zeta} \\
\frac{\partial N_4}{\partial \xi} & \frac{\partial N_4}{\partial \eta} & \frac{\partial N_4}{\partial \zeta} \\
\frac{\partial N_5}{\partial \xi} & \frac{\partial N_5}{\partial \eta} & \frac{\partial N_5}{\partial \zeta} \\
\frac{\partial N_6}{\partial \xi} & \frac{\partial N_6}{\partial \eta} & \frac{\partial N_6}{\partial \zeta} \\
\frac{\partial N_7}{\partial \xi} & \frac{\partial N_7}{\partial \eta} & \frac{\partial N_7}{\partial \zeta} \\
\frac{\partial N_8}{\partial \xi} & \frac{\partial N_8}{\partial \eta} & \frac{\partial N_8}{\partial \zeta} \\
\frac{\partial N_9}{\partial \xi} & \frac{\partial N_9}{\partial \eta} & \frac{\partial N_9}{\partial \zeta} \\
\frac{\partial N_{10}}{\partial \xi} & \frac{\partial N_{10}}{\partial \eta} & \frac{\partial N_{10}}{\partial \zeta}
\end{bmatrix} = [J] 
\]

where \([J]\) is the Jacobian matrix. From (C.8), the Jacobian matrix of a 10 nodes tetrahedron element may be expressed as

\[
\begin{bmatrix}
\frac{\partial N_1}{\partial x} & \frac{\partial N_1}{\partial y} & \frac{\partial N_1}{\partial z} \\
\frac{\partial N_2}{\partial x} & \frac{\partial N_2}{\partial y} & \frac{\partial N_2}{\partial z} \\
\frac{\partial N_3}{\partial x} & \frac{\partial N_3}{\partial y} & \frac{\partial N_3}{\partial z} \\
\frac{\partial N_4}{\partial x} & \frac{\partial N_4}{\partial y} & \frac{\partial N_4}{\partial z} \\
\frac{\partial N_5}{\partial x} & \frac{\partial N_5}{\partial y} & \frac{\partial N_5}{\partial z} \\
\frac{\partial N_6}{\partial x} & \frac{\partial N_6}{\partial y} & \frac{\partial N_6}{\partial z} \\
\frac{\partial N_7}{\partial x} & \frac{\partial N_7}{\partial y} & \frac{\partial N_7}{\partial z} \\
\frac{\partial N_8}{\partial x} & \frac{\partial N_8}{\partial y} & \frac{\partial N_8}{\partial z} \\
\frac{\partial N_9}{\partial x} & \frac{\partial N_9}{\partial y} & \frac{\partial N_9}{\partial z} \\
\frac{\partial N_{10}}{\partial x} & \frac{\partial N_{10}}{\partial y} & \frac{\partial N_{10}}{\partial z}
\end{bmatrix}
\]

where \([(x_i, y_i, z_i) | i \in \{1, 2, \ldots, 10\}]\) are the coordinates of the element nodes. By substituting (C.8) into (C.5), the strain-displacement matrix is calculated. Once the strain-displacement matrix for each element is obtained, the gradient of the strain energy for the corresponding element is computed from (C.2). The sensitivity of the objective function \(\frac{\partial J}{\partial N}\) is assembled by the element sensitivity \(\frac{\partial J}{\partial N}\) to obtain the vector of strain energy differentiation for the whole implant microarchitecture. The sensitivity analysis is then implemented under the optimization scheme described in Section 3.1, so as to seek the optimum density distribution of the implant.

Appendix D. Convergence plot of the topology optimization scheme

As described in Section 3, a density-based topology optimization is used in this study to optimally tailor the density gradients of the implant. To ensure solution convergence, we resort to the optimality criteria methods (Bendsoe, 1995), in particular here we use the standard optimality criterion based on the Lagrangian function that benefits from knowledge on the physics and mechanics of the problem (Avinash Shukla, 2013). This method requires the calculation of the derivative of the objective function along with the derivative of the design constraints to update the design variables based on the initial guess (Avinash Shukla, 2013).

In this work, the derivative of the implant compliance – obtained as described in Section 3 – is used with the standard optimality criterion to find the optimum density distribution. As per the initial guess, a uniform relative density of 0.5 is assigned to the implant and the total strain energy of the implant is calculated. The optimization continues until the difference between two successive iterations is below 1%. As shown in the figure below, the objective function converges within 31 iterations.
Fig. D.I. The convergence plot of the topology optimization scheme.
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