The influence of osteoporotic bone structures of the pelvic-hip complex on stress distribution under impact load

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Purpose: The aim of this study was to determine the effect of bone mineral density (BMD) on the stress distribution in pelvic-hip complex (PHC) model which included bone structures and soft tissues. Bone mass changes in osteoporosis and osteopenia were considered in this analysis. In addition, the relations between force direction and stress distribution causing PHC fractures were determined.

Methods: This paper presents the development and validation of a detailed 3D finite element model with high anatomical fidelity of the PHC and BMD changes in trabecular and cortical bones, modelled based on CT scans. 10 kN loading was induced on a model consisting of 8 ligaments, the pelvis, sacrum, femur in front and side directions.

Results: For validation, the results of this model were compared to physiological stress in standing position and previous results with high-energy crashes under side impact load. Analysis of side-impact indicated the influence of BMD on femoral neck fractures, acetabular cartilage and sacroiliac joint delaminations. Front-impact analysis revealed the inferior pubic ramus, femoral neck fractures and soft tissue injuries, i.e., acetabular cartilage and symphysis pubis in osteoporosis and osteopenia.

Conclusions: The elaborated PHC model enables effective prediction of pelvis injuries in high-energy trauma, according to Young-Burgess classification, and the determination of the influence of BMD reduction on pelvis trauma depending on force direction. The correlation between BMD and stress distribution causing varying injuries was determined.

Key words: pelvic-hip complex (PHC), bone mineral density (BMD), finite element analysis (FEA), osteoporosis

1. Introduction

Osteoporosis is classified as one of the main civilization diseases and characterized by low bone mineral density and changes in compact bone microstructures. In 2010, the estimated prevalence of new diagnosed osteoporosis among American community was 8 mln women and 2 mln men, according to Burge et al. [4]. By 2020, the new osteoporotic diagnosis are projected to increase to 14 mln [21]. Osteoporotic changes in bone cause reduction in overall skeleton strength, which leads to excessive sensitivity to low-energy fractures, mainly located in hip bone (72%), vertebrae (6%) and pelvis (5%). More than 1.5 mln fractures per year is a result of osteoporosis, which makes it a global, social and economical problem [4]. The most common injuries (60%) apply to fractures of the femoral neck and pelvis in patients aged 85 and over [4], [10], [13].

The occurrence and development of osteoporosis are associated with many factors, such as age, progressive bone demineralization or abnormal hormone levels [24]. A lack of estrogen in blood serum of post-menopausal women, which prevents the absorption and utilization of calcium, makes older woman most affected by osteoporosis. Due to the decrease in levels of estrogen, which is responsible for the management of the body calcium, more women, especially during and after menopause, suffer from osteoporosis [4], [8].

Osteoporotic changes are characterized by indirect and non-uniform decrease in bone density, particu-
larly in trabecular bone. The annual rate of bone loss in women aged 38 and over has been calculated to be 0.3 percent per year, in postmenopausal women to be 1.2 percent, whereas in men to be 0.3–0.5 percent. This is due to the release of calcium and magnesium from bone, resulting in decreasing in trabecular bone thickness and bone volume loss [21]. The severity of osteoporosis depends on bone resorption activity and local biomechanical factors.

The development of osteoporosis significantly affects the decrease of the mechanical parameters of bone structures. According to Burr et al. [5], the decrease of collagen in bone tissue results in a strength loss up to 50%. Experimental studies employed on the sectional preparations pointed out that the average Young’s modulus for normal bone tissue has value of 16.586 GPa (range 67–86 years), however for an osteoporotic bone it is much smaller, i.e., 11.554 GPa (range 67–91 years) [7]. According to Mazurkiewicz et al. [17] these value for people aged 53–76 is lower, and amounts to 15.7 GPa. Furthermore, tensile strength (from 117 MPa to 95.1 MPa) and yield strength (from 80.8 MPa to 75.8 MPa) tend to decrease with bone mineral density loss. Based on reports elaborated by the World Health Organization (WHO) and the National Osteoporosis Foundation (NOF), bone mineral density (BMD) higher than 833 mg/cm² for physiological bone, in the range 833–648 mg/cm² for osteopenia, and less than 648 mg/cm² for osteoporosis has a significant influence on bone mechanical properties [17]. Experimental research carried out by Rosholm et al. [20] confirmed the above classification and indicated the average BMD in age group 59–84 diagnosed with osteoporosis as 410 mg/cm². The lower elastic modulus (233.6 MPa), yield strength (5.3 MPa) and tensile strength (6.6 MPa) were measured for reduced BMD (380 mg/cm²) and located in cancellous bone of femoral head [27]. Demineralization process affects also the deterioration of elastic shear modulus (from 310 MPa to 247 MPa) and yield strength (from 3.3 MPa to 2.5 MPa) [14].

In view of the complicated geometric structure of pelvic-hip complex (PHC) and the problem to determine the elastic/visco-plastic material parameters of osteoporotic bone, previous numerical analyses were limited to modeling particular joint, i.e., lumbar vertebrae [15], [19], or hip [2], [9], [25].The aim of the analysis carried out by Pitzen et al. [19] was to determine the critical forces causing disruption of bone structures in the lumbar spine. Numerical model was included in structure of the vertebral body consisting of cancellous and cortical structures, ligaments: anterior and posterior longitudinal, interspinous, yellow, supraspinous and articular cartilages. More detailed model was elaborated by Liang et. al. [15] and included a structure localized in T12-L1 spine, i.e., cortical and trabecular structures of vertebrae, the presence of the nucleus pulposus, the annulus fibrosus and various ligaments. Bauer et. al [2] characterized trabecular bone with varying radiological density (0, 70, 140 and 210 mg/cm³ by Hounsfield scale) by uniaxial compressive strength until axial force of 25–243 N, using the following material parameters: Young’s modulus of 10 GPa, Poisson’s ratio of 0.3. The analysis [9] carried out by Newton–Raphson method treated the femur as a homogeneous material with a Young’s modulus equal to 10 GPa, Poisson’s ratio of 0.4 and a density of 200 mg/cm³.

Presented differences in biomechanical properties of bones characterized by abnormal BMD indicate varying stress/strain distribution under impact load, what may cause completely specific injuries than numerical analysis performed hitherto. Many numerical models of pelvic-hip complex have been elaborated and used to determine the stress distribution and the critical force in these structures under side impact load so far. According to Dawson et al. [6], the impact force which induced fracture of the pelvic bone was 8.6 kN (impact force – 129.2 N/s) on side impact. The region, which was the first to fracture was right pubic ramus, suggesting that frontal sections of pelvis are most susceptible to injury. The minimum force required to cause the left ischium and pubic rami fractures was 10.28 kN at an impulse of 154.2 N/s (impact speed 44.6 km/h). Fractures of the left and right ischium, pubic rami, left iliac fossa, and acetabulum are predicted under a peak force of 15.55 kN. Majumder et al. [16] indicated that the force causing the above-mentioned fractures was 16.98 kN. The superior pubic ramus has been damaged in the direction of the impact.

There are no models used in FEM analysis of osteoporotic changes in PHC in the literature. Characteristics carried out by Verhulp et al. [26] refer only to the stress distribution evaluation in the femur during a side falls. Forces in the range from 1 to 5 kN were applied to the pelvic-hip contact area and directed toward the center of the head. The elaborated model took into account the heterogeneity of bone – Young’s modulus of the cortical and cancellous was 22.5 GPa and 15 Gpa, respectively, bone mineral density was 0.917 and 0.976 g/cm² in the neck and trochanter of healthy bone and 0.496 and 0.656 g/cm² in the osteoporotic femurs, with Poisson’s ratio of 0.3. The maximum stress occurred in the cortical bone in the femoral neck and the value of the critical force that caused its breaking was equal to 3.29 kN.
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Up-to-date research indicates a significant role of osteoporotic changes in the mechanical properties of bone and its susceptibility to fracture. In addition, the significant prolongation of treatment time in patients with reduced mineral bone density was confirmed. It is, therefore, essential to know the mechanism of osteoporotic bone fracture and stress distribution in the osteoarticular system, not only in single bone analyzed until now, allowing the development of new crash protection features provide greater levels of injury protection. The aim of this study was to determine the effect of bone mineral density on the stress distribution in pelvic-hip complex which consisted of the bone and soft tissues. Changes in bone mineral density characteristic of osteoporosis and osteopenia were taken into account in this analysis. In addition, the relationship between direction of exerted force and stress characteristic result in bone fracture was investigated in some detail.

2. Materials and methods

2.1. The FE model of PHC

A detailed PHC model was elaborated using a computed tomography (CT) scans of a 25-year-old patient (woman, height: 1.64 m, weight: 52 kg) without osteoporotic changes in bones (T-score < −1), performed using an 8-row spiral CT with an accuracy of 2.5 mm. Bone structures were meshed with 8-node tetrahedral finite elements (245262 elements), because tetrahedral element instead of cubic or hexahedral were adopted to represent a smooth surface which accurately depict stress in the concentration areas with the use of local grid densification.

Homogeneous orthotropic properties derived from the literature are assigned to the PHC model (Table 1). Herein we use the relationship between Young’s modulus and bone mineral density for each bone according to Eqs. (1)–(4), where $E_{\text{cort}}$ – Young’s modulus of cortical bone, $E_{\text{FEMURtrab}}$ – Young’s modulus of trabecular femur bone, $E_{\text{SACRUMtrab}}$ – Young’s modulus of trabecular sacrum bone, $E_{\text{PELVIStrab}}$ – Young’s modulus of trabecular pelvic bone, $\rho$ – bone density [11]:

$$E_{\text{cort}} = 3.981 \rho^{2.39}$$  \hspace{1cm} (1)

$$E_{\text{FEMURtrab}} = 1.310 \rho^{1.4}$$  \hspace{1cm} (2)

$$E_{\text{SACRUMtrab}} = 1.540 \rho - 0.058$$  \hspace{1cm} (3)

$$E_{\text{PELVIStrab}} = 2.0173 \rho^{2.46}$$  \hspace{1cm} (4)

The shear modulus of cortical and trabecular bones used in this analysis were calculated according to the Eq. (5), where $G$ – shear modulus, $E$ – Young’s modulus and $\nu$ – Poisson’s ratio as listed in Table 1.

$$G = \frac{E}{2(1+\nu)}$$  \hspace{1cm} (5)

Due to the lack of experimental data, pelvic ligaments were modeled as one-dimensional line elements, which act as linear springs where mechanical

| Table 1. Elastic constants of cortical and trabecular bone of pelvic-hip complex modeled as orthotropic material for healthy bone, bone with osteopenia and osteoporosis, where cort – cortical bone, trab – trabecular [11] |
|---------------------------------|-----------------|-----------------|-----------------|
| Density [g/cm³] | Normal Cort | Normal Trab | Osteopenia Cort | Osteope
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<tr>
<td>E1</td>
<td>6983</td>
<td>2029</td>
<td>4136</td>
<td>239</td>
<td>150</td>
<td>875</td>
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<tr>
<td>E2</td>
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<td>2029</td>
<td>4136</td>
<td>239</td>
<td>150</td>
<td>875</td>
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<tr>
<td>E3</td>
<td>18155</td>
<td>3195</td>
<td>10752</td>
<td>259</td>
<td>180</td>
<td>712</td>
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<td>$\nu_{12}$</td>
<td>0.4</td>
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<td>$\nu_{23}$</td>
<td>0.25</td>
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<td>$\nu_{31}$</td>
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<td>Shear Modulus [MPa]</td>
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<tr>
<td>G12</td>
<td>4888</td>
<td>1420</td>
<td>2895</td>
<td>167</td>
<td>105</td>
<td>613</td>
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<tr>
<td>G23</td>
<td>4364</td>
<td>1268</td>
<td>2585</td>
<td>149</td>
<td>94</td>
<td>547</td>
</tr>
<tr>
<td>G31</td>
<td>11347</td>
<td>1997</td>
<td>6720</td>
<td>162</td>
<td>113</td>
<td>445</td>
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properties are shown in Table 2 [3]. The elaborated PHC model consists of the ligaments responsible for keeping the weight of the trunk and upper limbs, providing stability and attenuate forces on the lower extremities and the spine: anterior and posterior sacroiliac ligaments. Additional ligaments responsible for preventing the apex of sacrum from tilting farther back: sacrotuberous and sacrospinous ligaments.

The hip capsules on the left and right sides were modelled as the part of the acetabulum and provided connection of the hip and femur bones. The material properties for these capsules were estimated based on the tangent modulus determined by Hewitt et al. [12]. The Young modulus was calculated as the average stress measured from 0 to 80% of strain in the capsule. In the result, the mean value was assumed to be the same as for articular cartilage material in the acetabulum.

### 2.2. Loading and boundary conditions

The elaborated PHC model (Fig. 1) maps the sitting position of a driver/passenger resulting from ischial tuberosities support (Fig. 1A and B). Additional

![Fig. 1. The PHC model and boundary conditions of performed analysis: A, B – ischial tuberosities support, C – iliac crest support, D – physiological load of human body, E – impact load](image-url)
ally, the iliac crest support was applied where the seatbelts are fastened (Fig. 1C). The model also takes into account the physiological load value 500 N acting perpendicularly to the base of sacrum and imitating the human body mass (Fig. 1D).

To constrain rigid body motion, the model translation was fixed in the horizontal direction (at the top of the sacrum bone by load of 500 N and at the bottom of the ischial tuberosities by fixed support) and in the vertical direction (at the iliac crest surface, by fixed support). Thus, no degrees of freedom remained. This assumption could be applied due to the analysis of principal stretches not taking into account the displacement of anatomical structures.

We also analyzed the stress distribution under impact load as a force value 10 kN – force causing first fracture in pelvic complex [6], [16], [26] loading in two directions (Fig. 1E):

- SIDE IMPACT – load acting perpendicularly to the greater trochanter of the femur
- FRONT IMPACT – load acting perpendicularly to the cross-section of femur

The above analysis was carried out using ANSYS 15.2. software.

3. Results

3.1. The PHC model validation

Validation of the elaborated PHC model (Fig. 1) was achieved through the simulation of physiological stress distribution during sitting. To this end, the ischial tuberosities were supported and the load was imposed vertically onto the sacral promontory. The stress distribution under an axial compressive load of 500 N is shown in Fig. 2.

Fig. 2. Physiological stress distribution in pelvic complex under load of 500 N imposed onto the sacral promontory during sitting

The stress concentrated mainly on the surface of sacroiliac joint, lunate surface of the acetabulum, in particular on rami inferior ossis pubis, where the maximum stress value 22 MPa was localized.

The PHC model was also validated by the stress distribution under side and front impact load of 15 kN concentrated in greater trochanter and on cross-section of femur, respectively. The simulation results from the PHC model are shown in Fig. 3, with the maximum stress and fracture (stress > 135 MPa) occurring in ramus of ischium under forces acting in both directions. Other fractures are localized in pubic

Fig. 3. Stress distribution of pelvic-hip complex under front (A) and side (B) impact load of 15 kN acting he cross-section of femur and on the greater trochanter, respectively
rami. In the front impact case, the fractures of femur neck were additionally observed.

### 3.2. FE model boundary condition analysis

The stress distribution on the pelvis under side and front impact load (10 kN) was performed using the elaborated and validated PHC model in which bones varies by bone mineral density. Obtained results are shown in the following charts: pelvic bone (Fig. 4), femur (Fig. 5), and sacrum (Fig. 6).

The analysis of the stress distribution in the pelvic bone (Fig. 4) showed the highest stress concentration in the body of the ischium and its fractures in each analyzed case under side and front impact load. The fracture of the body of the pubis was recorded only in bone with osteopenic and osteoporotic changes under front impact load. Stresses exceeding ultimate stress at failure (135 MPa) were observed in the bodies of the

![Fig. 4. Stress distribution in pelvic bone with varying bone mineral density under front and side impact having value of 10 kN](image1)

![Fig. 5. Stress distribution in femur with varying bone mineral density under front and side impact having value of 10 kN](image2)
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ischium and pubis on side impact load, regardless of bone mineral density. Maximum stresses concentrated in the lower part of the head of the femur (Fig. 5) and the neck of the femur in the case of front impact load. Stresses exceeding the ultimate strength of bone were calculated for osteoporotic bone under two directions forces (max. stresses: 162 MPa). Forces of value 10 kN acting on the greater trochanter in side impact and cross-section of femur in front impact do not lead to fracture of sacrum bone, regardless of its BMD (Fig. 6).

The numerical PHC model has been widely applied to analyze the behavior of cartilages under compressive stress (Table 3). We have also reported an acetabular labrum fracture during side impact and sacroiliac joint destruction under front impact.

4. Discussion

Osteoporosis is classified as one of the diseases of civilization, also called the “epidemic of the 21st century”, being not only a health problem, but also a socio-economic problem of today’s world. Apart from the development of pharmacological treatment of osteoporosis taken to prevent or slow down the bone degradation, it is also important to understand the
mechanism of osteoporotic bone fractures, to reduce or eliminate risk factors. Research and past numerical analyses have been related to the influence of bone mineral density on mechanical properties, development of computer system to aid diagnosis of osteoporosis or traumatological analysis of low-energy trauma (i.e., falls). The authors attempted to elaborate a numerical model of pelvic-hip complex with regard to bone mineral density and analyze the mechanism of osteoporotic fractures under impact load acting in different directions.

The failure criterion adopted for the numerical analysis in the present work is the ultimate stress for cortical bone: 135 MPa and for cartilage: 40 MPa [18].

Using the PHC model (Fig. 1), the predicted stress distribution in pelvic girdle during sitting (Fig. 2) is in accordance with analogical analysis performed in literature [28]. Zhou et al. [28] indicated the maximum stress concentration in greater sciatic notch. Stresses occur also in sacroiliac joint and acetabulum regions including ischium bones. Stress concentration regions are almost identical with our results. The difference in maximal stress obtained using the PHC model, i.e., 22 MPa, compared to 1.1 MPa described in literature [23], arises from varying mechanical parameters of two types of bone tissue used in our model. The influence of one- and two-phase biomechanical models of bone tissue were analyzed [17], [14], [26] and proved the two-phase model much more accurate [11]. The application of trabecular bone with reduced strength parameters (Table 1), compared to the cortical one, have an influence on higher stresses concentrated in bones. In addition, the FE model elaborated by Zhou et al. [28] contained the fifth lumbar vertebra and did not include the ligaments which cause stress distribution to the lower part of the pelvic-hip complex.

The second stage of validation was carried out for high-energy accidents using forces of 15 kN. According to research [6], [16], forces acting in two directions – side and front cause total destruction of pelvis and allow to compare these fractures with Tile, Young and Burgess classification [1]. Material properties of bone and soft tissue were used for normal BMD (T-score $>-2.5$). Tensile stress values and stress distribution are similar to the one studied within a single bone structure, without taking into account the soft tissue and the articular joint [2], [9], [15], [19], [25]. On the other hand, estimation of pelvic injuries using finite element method was performed only for bone without bone mineral changes [6], [16], [26]. Fractures of the ischium and head of the femur fractures were observed under side and front impact of 10 kN in model of pelvic girdle bone with normal BMD (T-score $<-1$). The changes in stress distribution of pelvis with force directions were demonstrated. Side impact loads cause stress concentration in the ischium and pubis bones (Fig. 4), in the neck and head of the femur (Fig. 5) and in the base of sacrum (Fig. 6), however, under front impact stress concentration was localized in superior ischial ramus (Fig. 4), the head of the femur (Fig. 5) and sacral foramina (Fig. 6). These stress distributions are also in accordance with Young and Burgess classification [1]. Performed numerical research [6], [16] also indicate the stress concentration in the above-mentioned regions under impact load.

Inclusion of bone mineral density and mechanical properties characteristic of osteopenia (Table 1) influence the destruction of pelvic-hip complex. The bone density set at 740.5 mg/cm$^2$ in the PHC model corresponds to T-score of $-1.75$, an half of BMD range for osteopenia (T-score $-1$ to $-2.5$). Tensile stress values and stress distribution are similar to the one studied for bone within the normal range of BMD. That is, only the advanced states of osteopenia affect the formation and extent of fractures. Significant changes are observed in the FE analysis of model consisting of osteoporotic bone (i.e., T-score $>-2.5$). The stress distribution in PHC under side impact load is changed with BMD loss. Impact load acting on the greater trochanter caused the head of the femur to fracture (Fig. 5) and total degradation of the left acetabulum (Table 3). This results in lower stress in the pelvic bone, concentrating in the ischium (Fig. 4) and its further transfer to the sacrum bone, mainly via the sac-
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5. Conclusions

The stress distribution in pelvic-hip complex in diseases such as osteopenia and osteoporosis is investigated under impact loading condition. A bone mineral density changes model is used in the FE simulation of the PHC to examine the extent and sequence of pelvic fracture and soft tissue influence on stress transmission under varying force directions. The elaborated PHC model with varying BMD in each bone and multiple ligaments and cartilages described as inhomogeneous orthotropic material. The detailed PHC model was validated by comparison to the model described in literature, experimentally validated back during physiological standing and under side impact load.

Executed numerical analysis showed significant effect of bone mineral density on stress distribution in the PH complex under impact load. It was also demonstrated that the force directions affect the stress distribution and not independently of BMD concentration in the same regions. The maximum stress concentration in the femur (in the neck of the femur for side impact and in the head and neck of the femur for front impact) and its transfer through the joint causing pelvic bone, cartilages and symphysis pubic were confirmed.

The mechanism of damage evolution coupled with yielded criterion is considered as one of the unclear subjects in failure analysis of cortical bone materials and still debatable. The elaborated complex model can be useful in further simulation of PHC fractures depending on interaction between drugs and fracture, hormone influence on bone strength and implant loosening mechanism in osteoporotic patients.

Acknowledgments

This investigation was supported by a research grant, as a part of the project DOBR-BIO/22/13149/2013: “Improvement of safety and protection of soldiers on missions through military-medical and technical operations” sponsored by the National Centre for Research and Development in Poland.

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