Research Article

Association between P2X7 Receptor Polymorphisms and Bone Status in Mice

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Macrophages from mouse strains with the naturally occurring mutation P451L in the purinergic receptor P2X7 have impaired responses to agonists (1). Because P2X7 receptors are expressed in bone cells and are implicated in bone physiology, we asked whether strains with the P451L mutation have a different bone phenotype. By sequencing the most common strains of inbred mice, we found that only a few strains (BALB, NOD, NZW, and 129) were harboring the wild allelic version of the mutation (P451) in the gene for the purinergic receptor P2X7. The strains were compared by means of dual energy X-ray absorptiometry (DXA), bone markers, and three-point bending. Cultured osteoclasts were used in the ATP-induced pore formation assay. We found that strains with the P451 allele (BALB/cJ and 129X1/SvJ) had stronger femurs and higher levels of the bone resorption marker C-telopeptide collagen (CTX) compared to C57Bl/6 (B6) and DBA/2J mice. In strains with the 451L allele, pore-formation activity in osteoclasts in vitro was lower after application of ATP. In conclusion, two strains with the 451L allele of the naturally occurring mutation P451L, have weaker bones and lower levels of CTX, suggesting lower resorption levels in these animals, which could be related to the decreased ATP-induced pore formation observed in vitro. The importance of these findings for the interpretation of the earlier reported effects of P2X7 in mice is discussed, along with strategies in developing a murine model for testing the therapeutic effects of P2X7 agonists and antagonists upon postmenopausal osteoporosis.

1. Introduction

In the past decade several reports have shown that P2-receptor signalling plays a central role in bone physiology [1–5]. Bone cells express several types of P2 receptors [6], allowing them to respond differently to nucleotides, depending on the types of nucleotides present, their concentration, and the duration of exposure [7, 8]. Prolonged exposure to high agonist concentrations initiates the formation of large pores in the membrane mediated by the P2 subtype P2X7, a feature often assessed by ATP-induced dye uptake [7, 9]. Activation of the P2X7 receptor results in changed cellular morphology and membrane blebbing [10], which initiates necrotic and apoptotic mechanisms in macrophages [11]. There are several studies directing a role to the P2X7 receptor in mediating ATP-induced apoptosis in other cell types [7] and accordingly increased osteoclast numbers have been found in mice with ablation of the P2X7 receptor (P2X7−/− mice) [3]. In humans the single nucleotide polymorphism Glu496Ala is associated with abolished pore formation activity of the P2X7 receptor [12] and with decreased ATP-induced apoptosis of osteoclasts in vitro [13]. The P2X7 receptors are expressed in both osteoclast precursors and resorbing osteoclasts [8, 14, 15], and therefore, in addition
to activating the apoptotic pathway, the P2X7 receptor could play a role in osteoclast development [16–18] and activation [19].

In calvarial cells in vitro activation of P2X7 receptors increases expression of osteoblast markers, enhances mineralization, and induces membrane blebbing [20, 21]. The effects of P2X7 activation could also be mediated through the activation of the cytokine interleukin β (IL-1β) on bone cells or on cells adjacent to the bone compartment, resulting in a systemic effect on bone [22]. In macrophages from people carrying two C alleles of the Glu496Ala (A1513C) polymorphism, the processing of pro-IL-1β is impaired, leading to decreased levels of mature serum IL-1β.

The ablation of the P2X7 receptor generated by Solle (Pfizer P2X7−/−) in mice, led to reduced total bone mineral content (BMC), as a result of increased trabecular bone resorption, decreased periosteal circumference of the femur, reduced periosteal bone formation [3], and impaired response to mechanical loading [4]. Another murine model, with ablation of the P2X7 gene (GSK P2X7−/−), was generated by Chessell and his group [23] and has a dissimilar bone phenotype, showing increased cortical thickness in the tibial shaft, but no changes in total BMD [15]. The contradicting observations have been attributed to the dissimilar sample sizes, methods of the gene knockout, and different genetic backgrounds of the inbred strains used to generate the mice.

That the genetic background of the two P2X7−/− strains is important was shown in the study by Adriouch et al. [24] describing the presence of a naturally occurring mutation in the murine P2X7 gene. A thymine to cytosine change was found at nucleotide position 1352 (T1352C) in the B6 genome, but not in BALB/c and outbred mice. The mutation results in a change in the amino acid sequence (proline to leucine) at position 451 (P451L), in the cytoplasmic tail of the P2X7 receptor [24]. By transfecting HEK cells with constructs of both genotypes, they found that ATP-induced pore formation was reduced by approximately 50% in cells carrying the mutated 451L allele [24]. In murine thymocytes the P451L mutation affects apoptosis acting through the ATP-induced pore formation [25]. The reduced responsiveness of the P2X7 receptors to ATP in mice with 451L could have led to underestimation of the effects of the P2X7−/− upon bone and other parameters. Pfizer P2X7−/− mice were generated on 129/Ola × C57Bl/6 (B6) × DBA/2 genetic backgrounds, and maintained on the B6 × DBA/2 background [3, 26]. The GSK P2X7−/− mice were maintained on B6 background, but originate from a B6/129 hybrid [15, 23].

Two papers have described the expression of a novel splice variant (called P2X7-k) for the rodent P2X7 [27, 28]. The splice variant was first found in lymphocytes of the GSK P2X7−/− mice by Taylor et al. [27]. Nicke et al. reported shortly after that P2X7-k shows higher agonist sensitivity and slower deactivation than the normal P2X7 receptor complexes [28]. The alternative exon of the P2X7-k splice variant includes the intracellular N-terminus and ~80% of the first transmembrane domain, and thereby escapes gene inactivation in the GSK P2X7−/− mice. However, the expression of the splice variant P2X7-k is tissue-specific, and not expressed on the plasma membrane of osteoclasts from BALB/cj P2X7−/− [29].

None of the current P2X7−/− models are quite suitable for studying the therapeutic potential of the agonist acting through P2X7 in ovariectomized mice. First, the impaired ATP response in P451L mutated cells could mask the “true” effect of agonists and antagonists. Second, some of the founder strains included in the genetic background of these animals lacks response upon ovariectomy. The inbred mouse strains traditionally used for bone research represent different skeletal phenotypes during development and aging [30], which could be of importance in the interpretation of the precise role of P2X7 receptors in bone remodelling.

In the present study, the major goal was to investigate the distribution of the P451L alleles in inbred strains of mice. Furthermore, we wanted to investigate the bone status of these animals in order to select a suitable strain with the preferred genetic background (i.e., carrying the P451 allele) for crossing the P2X7−/− genotype into.

2. Materials and Methods

2.1. Sequencing of P2X7. A single-coding mutation (T1352C) in the murine P2X7 gene resulting in a change from proline to leucine at position 451 (P451L) has earlier been reported in four major strains of inbred mice [24]. To examine the distribution of the two alleles of T1352C (P451L) among laboratory mice genomic DNA was analyzed by sequencing exon 13 in the murine P2X7 gene. DNA from 20 different inbred strains (C3H/HeJ, C3H/HeJcrI, NZB/B1NJ, NZW/J, SJL/J, AKR/J, BALB/cjJ, BALB/cByJ, BALB/cAnNcr, 129J, 129X1/SvJr, Swr/J, C57L/J, C57BL/10J, DBA/1J, DBA/2J, Sm/J, NOD, Ddy/cI, and Calb/Rk) and from 2 strains of outbred (Mus caroli/Ei and M. spretus) was obtained from The Jackson Laboratories (Bar Harbor, ME). The B6 originates from Charles River (Germany). The Danish Pest Infestation Laboratory (Lyngby, Denmark) donated ten wild Mus musculus domesticus, from different locations. Earpieces and tail parts were used for DNA isolation with the QIAamp DNA Blood Mini Kit (Qiagen), which was used as template in PCR reactions with the following primers recognizing exon 13: F-Act TGA GGG GTT GTC ATT GC and R-TCC AAG GGA AGC TGT ATT GTG giving an amplicon of 595 bp. For each PCR product two sequencing reactions were prepared with the specific sequencing primers (F-TGC TGA TGG GTC and R-TCC AAG GGA AGC TGT ATT GTG) in the sequencing reaction with the BigDye Terminator v3.1 Cycle Sequencing Kit (Applied Biosystems) as described by Ohlendorr et al. 2007 [13], and loaded on the ABI PRISM 3100 Genetic Analyzer. Duplicate amplification and sequencing reactions were run per strain to confirm the sequences.

2.2. Animals. Female mice of nine inbred strains (C3H/HeJ, NZB/B1NJ, SJL/J, AKR/J, BALB/cjJ, 129X1/SvJr, Swr/J, C57L/J, and Dba/2J) were obtained at eight weeks of
age from Jackson Laboratories (Bar Harbor, ME). The B6 originated from Charles River (Germany).

2.3. Study Protocol. The Danish Animal Welfare Council approved all animal procedures in advance (protocol: 2002/561–634). Female mice at the age of 120 days were starved overnight before euthanized by CO₂. Blood was collected into 5 mL syringes by cardiac puncture. Serum was collected and stored at −80°C for later measurements of bone markers. The animals were scanned on a PIXImus (Lunar Corporation, Madison, WI) densitometer.

2.4. Bone Mineral Measurements and Body Composition. Bone mineral measurements and body composition of the animals were determined on the PIXImus densitometer. Animals were fixed in a standard position, and measurements were performed sequentially, with duplicate determinations. Intra assay CV was 0.47% and inter assay CV 0.73%. Due to the large mineral content in the skull, it was excluded from the calculations.

2.5. Bone Formation and Resorption Markers. To investigate possible differences in bone formation markers between the genotypes, osteocalcin was measured in serum samples in duplicate using the Mouse Osteocalcin RIA Reagents from Biomedical Technologies, Inc. (Stoughton, MA), following the protocol supplied with the reagents. Inter assay CV was 12% and intra assay CV was 6%.

Bone resorption as expressed by fragments of type I collagen in mouse serum (s-CTX) was measured in duplicate using the RatLaps Elisa Assay (C-telopeptide collagen type I fragment Assay) developed by Nordic Bioscience Diagnostics (Herlev, Denmark) and following the procedure supplied with the kit. Inter assay CV was 14.8% and intra assay CV was 9.2%.

Alkaline phosphatase activity was measured in duplicate mouse serum using the Alkaline Phosphatase Reagent Kit (Sigma). Alkaline phosphatase activity was measured directly on the serum in multi well plates, using a slight modification of the standard clinical chemistry procedure. Serum replicates were diluted with alkaline buffer solution and substrate solution was added to each well, and the plate was incubated at 37°C for 30 min. Finally 2.0N NaOH was added to each well to stop the reaction, before the absorbance was measured on a plate reader at 405 nm. Inter assay CV was 5.9% and intra assay CV was 2.4%.

2.6. Bone Strength Measurements. On the day of sacrifice the mouse femurs were collected, cleaned for tissue, wrapped in saline gauze, and frozen at −20°C until biomechanical testing. The strength of the femur was measured by the 3-point-bending test on a Lloyd Instruments compression device (Lloyd Instruments, Fareham, UK), performed after rehydrating the femur in a saline solution at room temperature, as described earlier [31].

2.7. Bone Histomorphometry. To investigate histologic and morphometric changes in four different inbred strains histomorphometric analyses were performed, as described earlier [31]. In short, the total spines and tibias were collected and fixed in 70% ethanol at 5°C. The following indices were determined; bone volume in percentage of total volume (BV/TV%), cortical thickness (C.Th, μm), trabecular thickness (Tb.Th, μm), and eroded surface as percentage of bone surface (ES/BS%). Under fluorescent light the mineralizing surfaces were determined as percentage of bone surface (MS/BS%). Further, mineral appositional rate (MAR, μm/day) was calculated.

2.8. In Vitro Dye Uptake Assay. Osteoclasts were isolated from bone marrow from the long bones of 3–4 weeks old female mice of the strains; BALB/cJ, 129X1/SvJ, DBA, and B6 (10 mice per strain) as described by Wu et al. [32]. Bone marrow monocyte/macrophage lineage precursors were seeded in α-MEM (10% FCS, 100 U/mL penicillin, 100 μg/mL streptomycin, 2 mM L-glutamine, 50 ng/mL M-CSF, and pH = 7.35–7.4) and plated with a density of 2 × 10⁶ cells/cm². More than 91% of both mononuclear and multinucleated cells in these cultures stained positively for TRAP (tartrate resistant acid phosphatase) using the leukocyte acid phosphatase kit (Sigma).

To investigate pore formation in osteoclasts from the four strains, as described for osteoblasts by Ke et al. [3], the medium was changed after 7 days in culture to a HEPES/glucose (20/10 mM) divalent cation-free buffer and the osteoclasts ability to take up the dye YO-PRO (Molecular Probes) upon ATP (200 nM) stimulation was assessed. After 9 minutes Hoechst 33342 (5 μg/mL, BioWhittaker) was added to the culture for another minute to reveal unstained nuclei in the osteoclasts. Only multinucleated osteoclasts (with 3 or more nuclei) were counted on a Leica BZ:03 Microscope and the number of YO-PRO containing cells was compared to the number of total osteoclasts in ten randomly chosen fields of sight. Control cultures were incubated with YO-PRO and Hoechst, but not ATP. Inter assay CV was 9.8% and intra assay CV was 5.4%.

2.9. Resorption Assay. Osteoclasts were isolated and cultured as described above for the dyeuptake assay, but seeded in 96-well plates on bovine bone slices (Nordic Bioscience, Herlev Denmark) or glass-cover slips. After 10 days of culture the osteoclasts on glass-cover slips were stained for TRAP and counted by visible inspection of ten randomly chosen fields of sight. Only TRAP positive cells with more than 3 nuclei were quantified as multinucleated osteoclasts. The cells were removed from the bone slices and the resorption pits were stained with hematoxylin, prior to quantification with the C.A.S.T. (Computer Assisted Sterological Toolbbox) Grid system on Olympus Bx51. The results were calculated as area resorbed in percentage of total area. Inter assay CV was 0.67% and intra assay CV was 0.45%.

2.10. Statistics. Statistical analyses were performed using the SPSS software, v.11.5. Standard parametric and nonparametric tests were used. For the comparison of results from the different strains the one-way ANOVA was used, with
The distribution of the strains in two groups with different allelic versions of the P451L mutation. Strains written in italic are outbred or mice from wild populations. *Confirming the data also shown by Adriouch et al. [24].

| Strains with P451 | Strains with 451L |
|------------------|------------------|
| Mus musculus*    | C57BL/6J (B6)*   |
| Mus caroli*      | C57BL/6NCrl      |
| BALB/cByJ*       | DBA/1*           |
| BALB/cAnNCrl*    | DBA/2*           |
| NOD*             | C3H/HeJ          |
| NZW*             | C57L/J           |
| 129*/            | NZB/BINJ         |
| 129X1/SvJ        | SIL/J            |
|                  | SWR/J            |
|                  | DDDY/J           |
|                  | SM/J             |
|                  | CALB/RkJ         |

3.1. Distribution of the P451L Mutation in the Inbred Strains. The distribution of the two alleles of the P451L mutation among common strains of laboratory mouse strains was determined in 21 strains and sublines by sequencing genomic DNA. Only mice from wild populations, outbred strains and a few of the inbred strains, which included the four major strains, BALB, NOD, NZW, and 129, had the P451 allele, hereafter called the original genotype of the P451L mutation. The rest of the examined strains and sublines had the mutated 451L allele (Table 1).

3.2. Bone Phenotypes in Relation to P451L Mutation. Ten inbred strains frequently used in bone related studies, were examined by DXA, three-point bending and serum concentration of bone markers to reveal the bone phenotype. As summarized in Table 2 there were significant differences in all measured parameters between groups, but only few associated to the different allelic versions of the P2X7 P451L mutation. With the genetic background of the two existing P2X7−/− mice [23, 26] in mind, the inbred strains of 129X1/SvJ and BALB/cJ (P451) were compared with B6 and DBA/2 (451L) with respect to bone status and these background strains were further examined by histomorphometric analysis, resorption activity, and in vitro dyeuptake assay.

3.3. Bone Status of the Background Strains. When focusing on BMD/BMC 129X1/SvJ had significantly (P < 0.001) higher whole body bone mineral (BMD and BMC) compared to the other three strains (Figure 1(a)). In the load-bearing region, femoral BMD and BMC were higher in the strains 129X1/SvJ and BALB/cJ than B6 and DBA/2 (P < 0.001. Figure 1(b)). The ultimate way of determining bone quality is to test the ability of bone to resist mechanical forces. Determined by a three-point bending test, the femoral bone strength was different among the strains of mice (Table 2). Of the four background strains 129X1/SvJ and BALB/cJ were more resistant to mechanical forces at the femur than both DBA/2 and B6 strains (P < 0.001, Table 2, Figure 1(c)).

In vivo bone markers are summarized in Table 2. The serum concentration of the bone formation markers, alkaline phosphatase and osteocalcin, were not associated with the P451L mutation in the P2X7 gene. The BALB/cJ strain had significantly (P < 0.001) higher s-CTX than all the other strains (Figure 1(d)). 129X1/SvJ mice had significantly higher s-CTX than the B6 mice, but not the DBA/2 (Table 2).

3.4. Histomorphometric Analysis in Relation to the P451L Mutation. Bone volume (%) in the vertebrae was higher in 129X1/SvJ than in the BALB/cJ and the B6 (P < 0.001), whereas BALB/cJ and B6 did not differ significantly (Figure 2(a)). No significant difference was found in the Tb.Th between the different strains. The cortical thickness (C.Th) for BALB/cJ (102.1 μm) was not significantly different from B6 (93.0 μm) and DBA/2 (104.4 μm). The mean C.Th was 121.5 μm for the 129X1/SvJ strain, was significantly higher than in the three other strains. Neither MAR nor MS/BS showed any significant differences between the four strains. However ES/BS% showed that the 129X1/SvJ strain had a significantly higher ES/BS% than B6, and BALB/cJ, B6, and DBA/2 (Figure 2(b)).

3.5. Ex Vivo Bone Resorption Assay. The area of resorption pits per bone slice was calculated as percentage of total area in 10 mice from each strain. BALB/cJ and 129X1/SvJ osteoclasts showed 19.1% and 18.7% area resorbed in comparison to 93.6 and 97.4 for B6 and DBA/2, with no significant difference (P = 0.478 between BALB/cJ and DBA/2). The number of TRAP-stained osteoclasts did not differ significantly. BALB/cJ and 129X1/SvJ had 79.5 and 83.7 TRAP-positive osteoclasts in comparison to 93.6 and 97.4 for B6 and DBA/2, with no significant difference (P = 0.108 between BALB/cJ and DBA/2).

3.6. In Vitro Dye Uptake Assay. To elucidate the cellular bases of these changes, we examined the response to ATP stimulation in a dye uptake assay, comparing the pore formation in 4 cultures from 10 mice in each of the following strains; B6, BALB/cJ, DBA/2, and 129X1/SvJ. Half of the cell cultures were incubated with dye, but without ATP stimulation. These cultures were used as controls to determine the spontaneous background staining, compared to the total number of cells in the assay. A fraction of the osteoclasts in the un-stimulated control cultures showed spontaneous dye uptake, with no significant difference between the strains. Interestingly,
TABLE 2: Bone parameters and concentration of *in vivo* bone markers presented as means (±SD). Number of animals in each strain is indicated at the top of the table (n). BMD, BMC, and bone area, in whole body or femoral region were determined by DXA scanning. An electronic digital caliper measured femoral length, before bone strength of the femur was determined by a 3-point bending test. *In vivo* bone markers were measured on serum using commercial available kits. One-way ANOVA was performed to test differences between groups and post hoc Bonferroni corrections, using 0.05 as the significance level. Simple descriptive of data is presented as means and standard deviations (SD). Significant difference between the 129X1/SvJ, BALB/cJ, B6, and DBA/2 animals at the *P* < 0.05 level is extracted from Bonferroni multiple comparison analysis and indicated with asterisks when different from the three other strains, with A when different from 129X1/SvJ, with B when different from B6, with C when different from BALB/cJ, and with D when different from DBA/2.

| Sample size (n) | P451L genotype | 129X1/SvJ | BALB/cJ | B6 | DBA/2 | SWR/J | C57L/J | AKR/J | SL/J | NZB/BINJ | C3H/HeJ |
|----------------|----------------|------------|----------|----|--------|--------|--------|-------|------|----------|----------|
| 15             | P              | L          | L        | L  | L      | L      | L      | L     | L    | L        | L        |
| 14             | P              | L          | L        | L  | L      | L      | L      | L     | L    | L        | L        |
| 16             | P              | L          | L        | L  | L      | L      | L      | L     | L    | L        | L        |
| 14             | P              | L          | L        | L  | L      | L      | L      | L     | L    | L        | L        |
| 14             | P              | L          | L        | L  | L      | L      | L      | L     | L    | L        | L        |
| 15             | P              | L          | L        | L  | L      | L      | L      | L     | L    | L        | L        |
| 15             | P              | L          | L        | L  | L      | L      | L      | L     | L    | L        | L        |
| 15             | P              | L          | L        | L  | L      | L      | L      | L     | L    | L        | L        |
| 15             | P              | L          | L        | L  | L      | L      | L      | L     | L    | L        | L        |
| 15             | P              | L          | L        | L  | L      | L      | L      | L     | L    | L        | L        |

**Bone Parameters**

|                  | 129X1/SvJ | BALB/cJ | B6 | DBA/2 | SWR/J | C57L/J | AKR/J | SL/J | NZB/BINJ | C3H/HeJ |
|------------------|-----------|---------|----|-------|-------|--------|-------|------|----------|----------|
| BMD (g/cm²)      | 0.0562±0.0020 | 0.0513±0.0018 | 0.0502±0.0022 | 0.0468±0.0004 | 0.0477±0.0013 | 0.0579±0.0030 | 0.0576±0.0021 | 0.0548±0.0007 | 0.0525±0.0015 | 0.0563±0.0018 | 0.001 |
| BMC (g)          | 0.535±0.045 | 0.471±0.028 | 0.458±0.031 | 0.358±0.029 | 0.394±0.035 | 0.548±0.042 | 0.593±0.046 | 0.459±0.025 | 0.465±0.064 | 0.558±0.036 | 0.001 |
| Area (cm²)       | 9.53±0.53 | 9.19±0.30 | 9.13±0.40 | 7.65±0.41 | 8.26±0.54 | 9.46±0.35 | 10.29±0.60 | 8.38±0.40 | 8.83±0.60 | 9.90±0.50 | 0.001 |
| BMD in femur region (g/cm²) | 0.0798±0.0039 | 0.0764±0.0021 | 0.0655±0.0033 | 0.0639±0.0028 | 0.0650±0.0024 | 0.0786±0.0009 | 0.0782±0.0030 | 0.0774±0.0004 | 0.0740±0.0036 | 0.0826±0.0037 | 0.001 |
| BMC in femur region (g) | 0.0439±0.0044 | 0.0402±0.0028 | 0.0364±0.0026 | 0.0281±0.0027 | 0.0307±0.0021 | 0.0464±0.0004 | 0.0466±0.0029 | 0.0391±0.0030 | 0.0378±0.0037 | 0.0439±0.0036 | 0.001 |
| Area of femur region (cm²) | 0.552±0.035 | 0.529±0.031 | 0.558±0.027 | 0.443±0.035 | 0.476±0.031 | 0.595±0.046 | 0.599±0.026 | 0.507±0.032 | 0.511±0.053 | 0.33 | 0.001 |
| Femur length (mm) | 15.25±0.36 | 15.06±0.64 | 15.53±0.23 | 14.34±0.45 | 14.29±0.30 | 15.79±0.40 | 16.00±0.33 | 14.08±0.28 | 15.90±0.45 | 16.11±0.64 | 0.001 |
| Strength/Max. load femurs (N) | 31.5±3.2 | 24.6±2.7 | 16.3±1.5 | 18.7±1.2 | 16.7±1.5 | 28.0±2.4 | 30.5±2.8 | 20.0±2.4 | 23.3±3.0 | 33.9±4.2 | 0.001 |

*In vivo* bone markers:

|                  | 129X1/SvJ | BALB/cJ | B6 | DBA/2 | SWR/J | C57L/J | AKR/J | SL/J | NZB/BINJ | C3H/HeJ |
|------------------|-----------|---------|----|-------|-------|--------|-------|------|----------|----------|
| Osteocalcin (ng/mL) | 40.6±9.48 | 55.9±11.78 | 38.4±10.89 | 71.7±8.97 | 36.4±8.75 | 72.3±11.25 | 51.0±10.20 | 48.4±8.06 | 63.2±11.38 | 0.001 |
| ALP (nmol/mL)    | 266.8     | 314.2    | 260.2 | 317.6 | 272.5  | 179.7  | 190.9  | 213.7 | 299.4    | 270.6    | 0.001 |
| RatLaps-Telopeptide collagen (mg/mL) | 14.28±5.61 | 19.09±4.40 | 9.81±1.86 | 12.13±1.88 | 10.75±1.62 | 9.37±2.53 | 7.07±2.62 | 7.15±1.83 | 8.83±4.02 | 8.76 | 0.001 |
Figure 1: Bone parameters in the 129X1/SvJ, BALB/c, B6, and DBA/2 inbred strains. Significant difference between the strains at the \( P < 0.05 \) level is indicated with asterisks when different from the three other strains, with A when different from 129X1/SvJ, with B when different from B6, with C when different from BALB/cJ, and with D when different from DBA/2. P451L genotype indicated as P or L at each strain. (a) In BALB/c and B6 whole body BMDs were not significantly different from each other, but significantly lower than 129X1/SvJ and BALB/cJ. BALB/cJ had higher BMD than DBA. (b) The femoral BMD in 129X1/SvJ and BALB/cJ were significantly higher than B6 and DBA/2. The latter were not significantly different from each other. (c) The femoral strength assessed by a three-point bending test showed significantly higher strength in 129X1/SvJ femurs. BALB/c had stronger femurs than the DBA and B6 strains had. The latter two were not significantly different from each other. (d) The serum concentration of the resorption marker s-CTX-I, showed significantly higher resorption in the BALB/cJ strain and lowest in the B6 strain.

4. Discussion

The P451 allele of the P451L mutation in P2X7 was present in different wild mice populations, in the outbred strains Mus caroli/Ei and M. spretus and the four inbred strains, BALB, NOD, NZW, and 129 (Table 1). In the bone phenotypic study, two of the strains we characterized, had the natural version (P451) of the mutation, the other eight strains had the 451L version. We found great diversity in all the examined bone phenotypic parameters, but only a few were
associated to the P451L mutation. The strain 129X1/SvJ had higher BMD, stronger femurs and higher bone resorption in bone histomorphometric analysis, compared to BALB/cJ, DBA/2 and B6. BALB/cJ had stronger femurs with higher BMD and higher levels of the bone resorption marker s-CTX than DBA/2 and B6.

We found strong femurs with high BMD in the P451 mice. Mechanical stress of load-bearing parts of the body could induce release of ATP in the bone compartment. Mechanical stimulation of bone formation has been shown to be partly mediated through the P2X7 receptor [4]. However, we did not detect any significant differences in bone formation markers associated to the P451L alleles. Perhaps the local effect of ATP stimulation in the femurs cannot be detected on the systemic bone markers in the serum samples. The histomorphometric analysis was only of the vertebrae, where higher BV/TV% was found in the P451 mice.

In the bone phenotypic part of the study we found that strains with the naturally occurring mutation 451L in the gene for P2X7 had lower resorption than strains with the wild type allele (P451). Agonists activating P2X7 in low concentrations can initiate osteoclast formation [33, 34]. Significant differences in the number of osteoclasts after 10 days of culture were not detected, but bear in mind that osteoclast culture was done without agonist stimulation. The absence of an effect of the P451L mutation on osteoclast development in vitro could also be due to the lack of systemic hormones, mechanical stimulation, and lack of accessory cells in the culture system.

The next step was to investigate if there were any differences in resorption activity in vitro though we did not detect a significant difference in resorbed area there appeared to be a tendency to higher resorption in cultures with osteoclasts derived from P451 strains. The duration of resorption was not investigated in this study, but keeping in mind that the P2X7 receptor is involved in apoptosis of osteoclasts [13], there could be differences in the length of the osteoclasts’ resorption phase related to the P451L allelic versions.

In humans, the polymorphism called Glu496Ala abolishes the pore formation activity of the P2X7 receptor [12] and has been associated with decreased ATP-induced apoptosis of osteoclasts in vitro [13]. A tendency towards more...
TRAP-positive osteoclasts was seen in the 451L osteoclast cultures. Apoptosis was not directly addressed in this study, but in vitro osteoclasts derived from the examined strains showed differences in the ATP-induced dye uptake assay. The examined mutation P451L is in the C-terminal part of P2X7, which is thought to be involved in the initiation of the pore formation, by activating pannexin-1 [35, 36]. In this manner the 451L mutated P2X7 receptors could blunt the pannexin-1 dependent dye uptake, and explain the reduced dye uptake we found in B6 and DBA/2 osteoclasts.

The physiological importance of the pore formation remains to be elucidated, but besides having a role in apoptosis it could be related to IL-1β processing and its release through pannexin-1 [37, 38]. Interleukin-1β has been reported to affect both bone resorption and bone formation dependent on the duration of the administration [39]. Interleukin-1β has also been shown to induce osteoclast precursor proliferation and differentiation, and increases bone resorption [40]. This suggests that IL-1β induces bone resorption initially, but that the long-term effect increased bone turnover, which corresponds to our data.

In macrophages from people carrying two C alleles of the Glu496Ala (A1513C) polymorphism, the processing of pro-IL-1β is impaired, leading to decreased levels of mature serum IL-1β. Following the theory that the reduced activity of the cytoplasmic tail of P2X7 (measured by pore formation) has significant importance in determining the bone cells’ response to ATP, the mechanical stimulation of bone cells in strains with the mutation should be impaired, and thereby the release of mature IL-1β reduced. Then the level of IL-1β induced resorption would be decreased in strains with the mutation and bone remodelling would favour bone formation, and that correlates with the decreased levels of resorption measured in vivo in B6 and DBA/2.

Background strains of both the previously described P2X7−/− mice harbor the 451L allele of the examined mutation in P2X7. The observed differences in the two P2X7−/− murine models cannot be addressed to the P451L mutation in P2X7 found in the background strains, since both models are maintained on genetic background harboring the 451L allele. However, the different bone status of DBA/2 and B6 may affect the direction of the effect of the ablation of P2X7. The observed lack of mechanical stimulation upon bone formation in the Pfizer P2X7−/−, would probably not have been found in a pure B6 background since the DBA/2 shows increased remodeling compared to the B6 mice. The best reason for the different observations will be the escape of gene knockout due to the splice variant P2X7-k [28] in the GSK P2X7−/− model [23]. Even though no expression of the splice variant P2X7-k has been found in osteoclasts [29], and GSK P2X7−/− osteoblasts shows no dye uptake, the splice variant P2X7-k could still be found in other cells in the bone environment.

The underlying goal was to find a candidate strain of mice that could be used as new genetic background for the P2X7−/− mice, in the search for the therapeutic potential of P2X7 agonists and antagonists in a murine model for postmenopausal osteoporosis. Four strains were determined to carry the P451 allele by sequencing, however the NOD mice develop type 1 diabetes (during which P2X7 expression change [41]) and the NZB mice get autoimmune anaemia. Only two strains with the P451 allele of the mutation were left, namely, 129X1/SvJ and BALB/cJ. In this study we found that DBA/2 and B6 had low BMD (totally and regionally in the femur) and had a low bone strength, which could make it difficult to detect changes in the parameters. Besides carrying the P451 allele, high baseline BMD, relatively strong bones, and high trabecular bone volume should characterize the selected strain. All that fits well with the bone phenotype of 129X1/SvJ, but mice from this strain show no cancellous bone loss upon oestrogen depletion, as has been reported in postmenopausal women and oestrogen depleted rats [42, 43]. In contrast, BALB/cJ has relatively strong bones and high BMD in the femur, high Tb.Th number, and respond as predicted upon ovariectomy [44].

In conclusion, we find that the P451 strains have stronger femurs and higher BMD, but the cellular basis could not be established in this study. The major finding in this study is that the genetic background could be of significant importance when determining the effect of the P2X7 ablation and that an optimal strain for studying the bone-specific effect of the P2X7 ablation could be the BALB/cJ mice.

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