

Bone and Cellular Immune System of Multiparous Sows are Insensitive to Ovariectomy and Nutritive Calcium Shortage

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Key words

- osteoporosis
- osteoimmunology
- pig

Abstract

Research in osteoporosis, which is a complex systemic disease, demands suitable large animal models. In pigs, most research has been done in growing minipigs, which probably are not ideal models for postmenopausal osteoporosis. Therefore, our aim was to analyze the effects of ovariectomy (OVX) and nutritive calcium shortage on multiparous Large White sows. 32 animals were randomly assigned to 4 groups in a cross design with OVX vs. sham and physiological calcium supplementation (0.75% calcium) vs. dietary calcium shortage (0.3% calcium). The observation period was 10 months with blood sampling every 2 months for hematological, immunological, and biochemical bone

marker measurements. At the termination of the experiment, animals were sacrificed. Samples of trabecular bone of distal radius, proximal tibia, and sixth lumbar vertebra were subjected to micro-computed tomography imaging and ashed afterwards. Dual X-ray absorptiometry scans of the proximal femora were performed with prepared bones being placed in a water bath for mimicking soft tissue. Analyses of bone marker and cytokine profile kinetics, distribution of leukocyte subpopulations, and morphometrical and densitometrical analyses showed no evidence of any impact of OVX or calcium shortage. In conclusion, the skeleton of adult sows of a conventional breed is seemingly protected from effects of OVX and calcium shortage.

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Introduction

Osteoporosis is a complex systemic disease [1]. In vitro models are not sufficient to analyze systemic effects, thus, there is a great need for suitable animal models. Rodent models, which have several advantages in terms of laboratory management, are well established and have been widely used in osteoporosis research. However, the US Food and Drug Administration demands also the use of large animal models, mainly due to biomechanical issues, besides rats in preclinical testing of antiosteoporotic substances with an experimental time frame of 12 months when using rats and 16 months when using larger species. According to FDA regulations, valid animal models have to develop an osteoporotic phenotype either spontaneously or after ovariectomy (OVX), which mimics postmenopausal estrogen deficiency [2].

Pigs as well as sheep are among the most frequently used large animal species in osteoporosis research. Although there are a lot of similarities

of diverse porcine organ systems to their human analogues, the pig's usefulness as an osteologic model species is still not entirely clear. However, even though porcine femoral compact bone is predominantly plexiforme, it is converted to well-developed osteonal bone earlier than in sheep [3]. Peak bone mass is obtained with an age of 2–3 years. The main body of investigations in this area of research was performed using growing minipigs, which, however, might not appropriately reflect the situation of the postmenopausal osteoporotic woman due to their juvenile age and dwarfism. On the other hand, minipigs achieve sexual maturity earlier than conventional pigs and thus OVX may induce the desired phenotype earlier than in conventional sows. OVX in 10 months old minipigs resulted in a 6% decrease in bone mineral density (BMD), 15% in bone volume (BV), and 13% in trabecular number, and an increase of 15% in trabecular separation after 6 months, whereas OVX in combination with a mild nutritive calcium shortage (0.75% Ca²⁺), which had been started already at

an age of 4 months, led to a 10% reduction in vertebral BMD and significant increases in final erosion depth and vertebral marrow star volume [4]. A study investigating multiparous sows being fed a standard diet (1.5% Ca²⁺) showed moderate and transient increases in plasma PTH, calcitriol and bone-specific alkaline phosphatase (BAP) levels over a time span of 1 year after OVX [5]. However, no significant changes concerning bone chemistry and histomorphometry were observed. Consequently, we were interested in evaluating the impact of OVX in combination with pronounced calcium shortage on bone metabolism, bone microstructure, and immunological parameters in multiparous conventional sows in contrast to the minipigs analyzed by other authors. Analysis of immunological parameters was included as osteoporosis as a systemic disease is known to be correlated with a proinflammatory reactivity of the immune system [6].

Materials and Methods



Animals and group assignment

32 multiparous Large White sows aged 33.5±9.6 months in the mean and all of them approximately 2 months post partum and thus not lactating for over 5 weeks were allocated equally to 4 groups to compare the effects of OVX and dietary calcium shortage: (I) OVX, 0.75% Ca²⁺, (II) OVX, 0.3% Ca²⁺, (III) sham-OVX, 0.75% Ca²⁺, and (IV) sham-OVX, 0.3% Ca²⁺. Ovariectomy was performed from the left flank under full anesthesia with an IV bolus of ketamine and azaperone. Post-surgical treatment was done with enrofloxacin, metamizole, and local oxytetracycline spray. Sows were housed as groups of 5 or 6 in a separate stable under standard conditions. Diets were mainly based on barley, soy, and lignocellulose and produced by a farm animals' feed company (Biomim, Herzogenburg, Austria). One diet contained the usually supplemented calcium levels for nongestating sows, that is, 0.75%, and the other one was a low calcium diet containing 0.3%. For detailed composition of diets see • Table 1. Blood samples were collected by venipuncture of the jugular vein every 2 months. All animal experiments were approved by the institutional and the governmental ethics committees. The experiment was terminated after 10 months.

Biochemical bone markers and hormones

Serum levels of following bone metabolism markers were determined by commercially available ELISA kits, which have been validated also for pigs [7,8]: receptor activator of nuclear factor-κB ligand (ampli-sRANKL ELISA, Biomedica, Vienna, Austria), calcitriol (25-OH-Vitamin D direct ELISA, Immunodiag-

nostik, Bensheim, Germany), PTH (Porcine Intact PTH ELISA, Immotopics, San Clemente, CA, USA), bone specific alkaline phosphatase (Metra[®]BAP ELISA, Qidel Corporation, San Diego, CA, USA), osteocalcin (Metra[®]Osteocalcin ELISA, Qidel Corporation), CACP (Metra[®]CACP ELISA, Qidel Corporation), pyridinoline (Metra[®]Serum PYD ELISA, Qidel Corporation), and crosslaps (Serum CrossLaps[®] ELISA, Immunodiagnostic Systems, Frankfurt/Main, Germany). For measurement of serum estrogens, a home made enzyme immunoassay for the detection of estron and 17β-estradiol was applied [9]. The low detection limit of this test of 2 pg of estrogens/ml is superior to the one of commercial systems.

Hematological and immunological analyses

Hematological analyses were performed out of EDTA-blood using an ADVIA[®]120 (Siemens Health Care Diagnostics, Deerfield, IL, USA) with veterinary software adapted for pigs. Cytokine levels were determined at both the genomic as well as the protein levels. For preparation of total RNA out of peripheral blood mononuclear cells (PBMCs), the pellet was suspended in 1.2 ml of hemolysis buffer (140 mM NH₄Cl, 17 mM Tris, pH 7.2) and incubated at 37 °C for 15 min. Samples were then centrifuged and supernatant was removed. Pelleted white blood cells were resuspended in 1 ml of TriReagent[®] (Molecular Research Center, Inc., Cincinnati, USA) and frozen at -80 °C until analysis. Total RNA was extracted in accordance to the manufacturers' protocol. Integrity, quantity, and contamination with genomic DNA were analyzed on the Agilent BioAnalyzer 2100 (Agilent Technologies, Palo Alto, USA) using the RNA6000 Nano LabChip[®] kit (Agilent). 1 µg of total RNA was used to synthesize cDNA using Superscript[™] II RNase H-reverse transcriptase (200 U/reaction; Invitrogen, Carlsbad, USA) and anchored oligo-dT-primers (3.5 µM final concentration). To check the generation of amplifiable cDNA in the reverse transcription, a conventional PCR step was performed using GAPDH specific primers. The profiling of the expressions of the cytokine genes interleukin (IL)-1, IL-2, IL-4, IL-6, IL-10, interferon (IFN)-γ, tumor necrosis factor (TNF)-α, as well as inducible NO synthase (iNOS) and heme oxygenase (HO)-1 and the 3 housekeeping genes cyclophilin, GAPDH, and β-actin was performed by means of real-time triplex PCR, using TaqMan[®] probes and specific primer pairs on the iCycler iQ[®] (Bio-Rad, Hercules, USA) as described by others [10].

Intracellular cytokine expressions of IL-1β, IL-2, IL-4, IL-6, IL-10, IL-12p35, TNF-α, and IFN-γ were measured in PBMCs as described previously using a FACSARIA[®] flow cytometer (Becton Dickinson, San Jose, CA, USA) [11]. Subpopulations of white blood cells including monocytes, plasmacytoid dendritic cells, B cells, naïve, activated, and memory T helper cells, regulatory T cells, cytotoxic T lymphocytes, γδ-T cells, and natural killer cells were discriminated and quantified by FACS. Antibody clones used for distinguishing cell markers as well as surface marker co-expression profiles used for identification of antigen presenting cells and lymphocyte subpopulations were the same as used by Sipos et al. [11].

µCT and DXA analyses of selected bone sites

The following bone sites were considered for µCT imaging and analysis: sixth lumbar vertebra (L6), proximal tibia, and distal radius. Bones were frozen at -80 °C until preparation. After freeing bones from soft tissue, a transverse section was obtained from each site of interest and from each animal (diamond band saw, 300 CP, Exakt GmbH, Germany). Then, one trabecular bone cylinder of 8 mm in nominal diameter and 12 mm in nominal

Table 1 Composition of diets (per kg)

	Low-calcium diet	Physiological diet
ME (MJ)	11.1	11.1
Crude protein (%)	13.0	13.0
Crude fat (g)	22.0	22.0
Crude fiber (g)	102.0	102.0
Vit A (IU)	9900	9900
Vit D ₃ (IU)	1320	1320
Vit E (mg)	70	70
Ca (g)	3.0	7.5
P (g)	3.8	6.0
Na (g)	1.7	1.7
Cu (mg)	16.5	16.5

length was extracted from each section using a diamond coated coredrill. The biopsies were immersed in saline solution, freed of air bubbles by application of vacuum and scanned with 12 µm resolution using a µCT 40 system (ScancoMedical AG, Switzerland) with 70 kV, 114 mA, and 200 ms integration time. The region of interest was restricted to an inner cylinder with 7 mm diameter and 11.5 mm length via contouring to exclude peripheral artefacts. Morphological analysis of each biopsy was performed with the inbuilt software tools provided by the manufacturer (IPL). The images were filtered using a Gaussian filter (sigma=0.7, support=1) and segmented using a global threshold value corresponding to 20% of the maximum gray value intensity. A standard evaluation protocol was used to compute the histomorphometric parameters from the segmented images. 2 biopsies containing a significant part of the growth plate were excluded from the analysis.

DXA scans of femoral neck, trochanter, and the total region including head, neck, and trochanter were performed using a Lunar iDXA® bone densitometer with prepared femora being placed in a water bath for mimicking soft tissue.

Bone ash analysis

The volume of each trabecular bone biopsy used for µCT analysis was calculated out of 5 consecutive measurements of height and diameter. Ashing was performed for 24 h at 650 °C and ash weight was measured after a consecutive cooling down phase at room temperature for another 24 h.

Statistics

Statistical evaluation was performed by PASW-Statistic Software, version 17.0. 2. All variables were tested for normal distribution. As most of them fulfilled the requirements for parametric evaluation, ANOVA was applied for group comparisons. In cases where normal distribution was excluded, a Kruskal-Wallis test was used. For post hoc analysis, a Bonferroni test was applied.

Results



Hematology and cellular immune parameters

Hematological analyses including FACS-based analysis of the distribution of antigen-presenting cell and lymphocyte subpopulations as well as cytokine profile kinetics revealed no meaningful trends over the course of the trial. Also, no differences could be observed between the 4 groups at either time point. Instead, cytokine titers or cell numbers of immunocyte subpopulations were found within the respective reference ranges [11] (data not shown). They only slightly undulated from one time point of measurement to the next and thus gave no evidence of

any OVX- or calcium shortage-induced change in hematological or immune status.

Biochemical bone markers and hormones

Mean values and standard deviations of investigated biochemical bone markers of the last sampling session are given in ◻ Table 2. As with hematological and cellular immune parameters, levels were only slightly undulating without any trend of either increasing or decreasing titers for each of the parameters over time. Results of group 3, which was the control group, can be considered as physiological reference values.

Serum estrogens (estron and 17β-estradiol) were measured at baseline, 1 month after OVX, and at the termination of the experiment. At baseline, estrogens were detectable in only 13 out of 32 animals, giving evidence of 13 sows being in estrus at that time point. 1 month post OVX, estrogens were detectable in 6/16 ovariectomized sows as compared to 10/16 non-ovariectomized animals. At the end of the experiment, only 1 ovariectomized sow had a detectable estrogen titer (4.6 pg/ml), whereas estrogen titers of 7 non-ovariectomized sows were exceeding the detection limit.

Morphological and densitometrical analyses

Also morphological as well as densitometrical analyses gave no evidence of any impact of OVX or calcium shortage on bone microstructure or density (◻ Table 3, 4). To better demonstrate the uniformity of trabecular bone microstructure of each anatomical site between groups, 2D µCT images of all investigated bone sites of reference animals of each group are given in ◻ Fig. 1.

Discussion and Conclusions



To date, mostly minipigs are used as large animal biomedical model species. Pure bred minipigs are expensive and have distinct ontogenetic and physiologic features when compared to conventional pigs [12]. These include earlier sexual maturity and a chondrodystrophic phenotype, which is common to most minipig strains. Additionally, most experiments in osteoporosis research were performed with growing minipigs, which may be a drawback when trying to extrapolate data to the situation of postmenopausal women. Therefore, our aim was to investigate the suitability of multiparous conventional sows as a model in osteoporosis research. We analyzed the effects of calcium shortage and OVX over a time span of 10 months. We chose this time frame because changes in bone metabolism as well as bone mass and architecture due to OVX and calcium shortage could have

Table 2 Biochemical bone marker levels acquired at the termination of the experiment (10 months post OVX)

	Group 1 (OVX)		Group 2 (OVX, low Ca)		Group 3 (Sham)		Group 4 (Sham, low Ca)		p-Value
	Mean	SD	Mean	SD	Mean	SD	Mean	SD	
sRANKL (pmol/l)	0.7	0.6	0.6	0.7	0.5	0.5	0.4	0.6	0.89
BAP (E/l)	13.5	4.2	16.5	2.3	14.7	5.7	15.7	5.2	0.55
OC (ng/ml)	194.7	139.3	304.2	88.6	236.2	150.4	192.7	106.8	0.34
CICP (ng/ml)	40.0	38.3	71.2	109.7	32.9	53.2	88.7	74.2	0.54
PYD (nmol/l)	8.5	0.9	8.0	0.8	8.5	0.4	8.4	0.6	0.55
Crosslaps (ng/ml)	1.5	0.9	2.0	1.3	0.9	0.7	0.9	0.6	0.16
PTH (pg/ml)	16.3	30.6	19.5	37.1	23.2	19.1	34.6	33.8	0.40
VitD ₃ (nmol/l)	170.1	101.8	157.5	106.6	212.9	19.3	123.7	92.9	0.47

Table 3 Bone mineral content at 10 months post OVX

	Group 1 (OVX)		Group 2 (OVX, low Ca)		Group 3 (Sham)		Group 4 (Sham, low Ca)		p-Value
	Mean	SD	Mean	SD	Mean	SD	Mean	SD	
DXA (g/cm²)									
Neck	1.47	0.08	1.56	0.16	1.48	0.11	1.43	0.15	0.34
Trochanter	1.59	0.14	1.58	0.09	1.76	0.2	1.6	0.34	0.51
Total	1.67	0.09	1.67	0.05	1.76	0.09	1.62	0.26	0.54
Bone ash values (mg/cm³)									
Radius	380.0	61.0	393.7	108.9	393.7	49.1	346.2	45.5	0.65
Tibia	344.3	95.4	339.3	113.8	312.0	94.5	282.7	78.9	0.65
Lumbar vertebra	394.7	55.3	374.9	60.7	394.3	65.3	385.2	39.4	0.91

Table 4 μ CT data of respective bone sites at 10 months post OVX

Bone site	Group 1 (OVX)		Group 2 (OVX, low Ca)		Group 3 (Sham)		Group 4 (Sham, low Ca)		p-Value
	Mean	SD	Mean	SD	Mean	SD	Mean	SD	
Radius									
BMD apparent (mg/cm ³)	357.5	68.4	404.3	111.0	376.0	58.2	341.2	43.1	0.50
BMD tissue (mg/cm ³)	1048.1	40.9	1093.3	120.6	1047.8	19.9	1075.6	24.6	0.58
BV/TV (%)	0.327	0.058	0.349	0.090	0.339	0.048	0.304	0.038	0.63
BS/BV (%)	12.594	2.160	12.062	2.374	12.504	1.932	12.725	1.050	0.93
Tb.N (/mm)	2.019	0.212	2.031	0.234	2.087	0.104	1.922	0.123	0.52
Tb. Th (mm)	0.162	0.024	0.171	0.032	0.162	0.024	0.158	0.013	0.81
Tb. Sp (mm)	0.337	0.061	0.327	0.079	0.317	0.028	0.363	0.040	0.57
Degree of anisotropy	2.303	0.286	2.184	0.439	2.142	0.301	2.429	0.198	0.44
ConnDens (/mm ³)	5.818	1.547	7.635	3.236	6.118	1.021	5.837	1.254	0.33
Structure model index	-0.439	0.276	-0.448	0.693	-0.472	0.289	-0.053	0.351	0.34
Tibia									
BMD apparent (mg/cm ³)	320.2	98.9	298.1	108.3	294.3	101.5	255.9	85.1	0.69
BMD tissue (mg/cm ³)	1073.9	44.2	1044.3	55.6	1086.0	25.8	1094.5	54.5	0.24
BV/TV (%)	0.291	0.086	0.282	0.096	0.264	0.090	0.238	0.068	0.70
BS/BV (%)	15.078	4.272	15.877	4.196	15.872	4.047	16.874	3.638	0.88
Tb. N (/mm)	2.040	0.175	2.074	0.235	1.969	0.200	1.920	0.228	0.54
Tb. Th (mm)	0.141	0.035	0.134	0.039	0.133	0.035	0.123	0.030	0.84
Tb. Sp (mm)	0.351	0.067	0.352	0.079	0.379	0.072	0.403	0.081	0.55
Degree of anisotropy	1.820	0.376	1.880	0.307	1.917	0.315	1.841	0.424	0.96
ConnDens (/mm ³)	8.610	2.261	9.138	3.024	8.103	0.698	8.287	2.700	0.87
Structure model index	0.144	0.530	0.132	0.727	0.299	0.641	0.450	0.622	0.77
L6									
BMD apparent (mg/cm ³)	367.3	65.1	347.9	80.2	399.8	53.2	383.6	38.2	0.53
BMD tissue (mg/cm ³)	1014.5	49.7	994.8	64.0	1036.4	12.7	1006.0	46.6	0.54
BV/TV (%)	0.339	0.046	0.327	0.053	0.352	0.047	0.348	0.033	0.77
BS/BV (%)	13.706	2.656	15.014	1.813	13.159	2.135	14.300	1.270	0.44
Tb. N (/mm)	2.289	0.265	2.421	0.232	2.286	0.265	2.477	0.144	0.39
Tb. Th (mm)	0.149	0.024	0.134	0.016	0.155	0.029	0.140	0.014	0.36
Tb.Sp (mm)	0.292	0.045	0.281	0.049	0.286	0.041	0.263	0.023	0.66
Degree of anisotropy	2.196	0.241	2.065	0.169	2.118	0.170	2.041	0.135	0.33
ConnDens (/mm ³)	3.902	1.354	4.332	1.358	4.027	1.247	3.989	1.297	0.93
Structure model index	-1.258	0.385	-1.259	0.364	-1.405	0.452	-1.418	0.365	0.80

been expected already within 6 months according to the available literature dealing with growing minipigs [4]. Interestingly, OVX and calcium shortage alone as well as in combination did not influence bone metabolism and microstructure. This was unexpected, as other authors observed increasing PTH, calcitriol, and BAP plasma levels, albeit transient and moderate, in sows of the same age group as the ones investigated here, although those were ovariectomized but fed a standard diet [5]. Nevertheless, these authors also did not find any significant changes concerning bone chemistry and histomorphometry. One reason for the observed dynamic rigidity, meaning the inability to respond to estrogen and calcium shortage, might be the notably high BMD of adult pigs, making efforts to artificially reduce bone mass and weakening bone a hardly achievable task. Lactating sows have to nourish litters of up to 13 piglets and

thus produce large amounts of milk with 10–11 kg per day containing approximately 50 mmol/l calcium, which corresponds to a 14 times higher calcium concentration than human milk, consequently giving evidence of the intense need for huge calcium resources [13]. Despite these tremendous physiologic needs for calcium the skeletal apparatus has to function properly. This may provide some explanation for the peculiarities of bone physiology of adult sows.

In other ungulates investigated so far, that is, sheep, OVX alone reduces bone mass, which can be enforced by an additional nutritive calcium shortage. BMD of L5 and distal radius as evaluated by DXA was significantly changed after 6 months and the one of L4 1 year after OVX, whereas the proximal parts of femur, humerus, and tibia did not exhibit alterations to that extent [14]. However, MacLeay and colleagues [15] were not able to detect

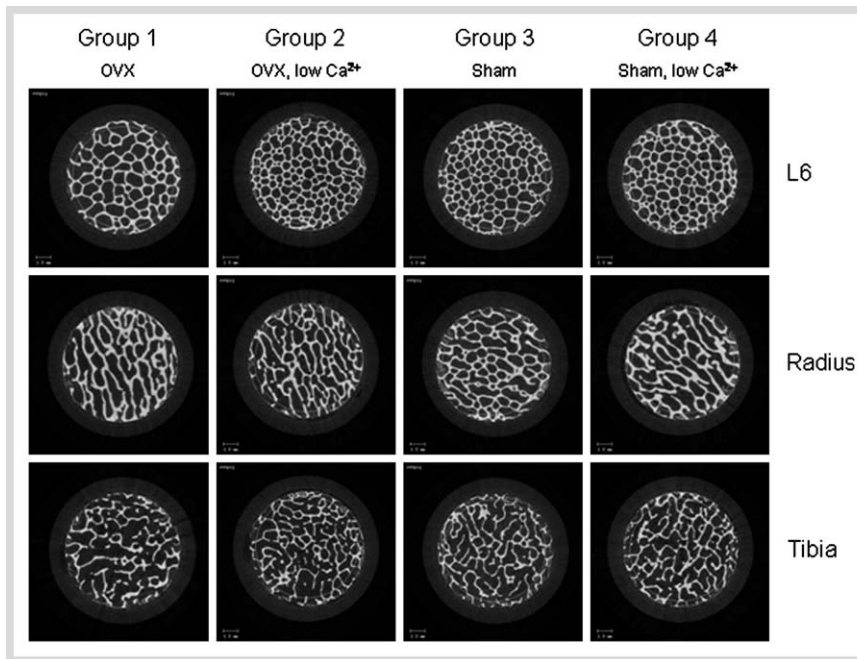


Fig. 1 2-dimensional μ CT images of trabecular bone of distal radius, proximal tibia, and sixth lumbar vertebra of representative sows of each group. Measurements were started at a predefined distance from the end of the samples to avoid inclusion of growth plates. Different anatomical sites exhibited characteristic micromorphological features but were uniform between groups.

areal BMD changes in lumbar vertebrae in ovariectomized sheep 3 months after surgery. Another study showed a significantly decreased femoral but not lumbar vertebral BMD as well as significant effects on cortical bone parameters by 6 months after OVX [16]. The lesser effects of single OVX treatment on bone mass in sheep are discussed by their probable ability of extragonadal estrogen production, which has not been reported for pigs so far. Interesting and seemingly inconsistent with the hypothesis of extragonadal estrogen synthesis is the fact that sheep experience significant microarchitectural changes in vertebral cancellous bone (decreased BV/TV by approximately 30%, trabecular thickness by 13%, and increased trabecular separation by 46%) 2 years after OVX and show significantly increased osteoclast numbers already 3 months after surgery [17,18]. Another study reported the advantages of combining OVX and glucocorticoid administration over combining OVX and calcium restriction with a higher decrease of BMD of distal radius, distal tibia, and calcaneus (spongiosa by 25% and corticalis by 17% in the former group and 10 and 5%, respectively, in the latter) [19]. Combining all 3 measures led to the most pronounced reductions (60 and 25%). When using corticosteroids, the disadvantage of immunosuppressive side effects causing local and/or systemic opportunistic infections [20] and of hampering osteoimmunological analyses has to be kept in mind.

Sows seem to differently handle loss of ovarian function and nutritive calcium shortage with respect to bone morphology, but also their cellular immune system gives no evidence of being affected by these measures. This is another difference to postmenopausal women, whose T cells have been shown to be activated as a consequence of increased IFN- γ , TNF- α , and RANKL synthesis [21]. Also, postmenopausal women have been shown to harbor higher levels of CD8⁺CD57⁺ cells and to suffer from a proinflammatory state [22,23], none of which has been observed in our model.

Another way to possibly achieve an osteoporosis-like phenotype in adult sows could be glucocorticoid treatment with all the disadvantages discussed above. Glucocorticoid-induced osteoporosis has already been shown as an option in the porcine model, albeit again minipigs have been used for these experiments.

Scholz-Ahrens and colleagues [24] induced an osteoporotic phenotype in adult (30 months old) primiparous Göttingen minipigs by daily oral prednisolone treatment at a dose of 1 mg/kg for 2 months with a reduction of this dose to 0.5 mg/kg thereafter until the end of the experiment, which was after 8 months in the short-term group and 15 months in the long-term group. In the short term, glucocorticoids reduced BMD at the lumbar spine by 48 mg/cm³ from baseline, whereas in the control group reduction was 12 mg/cm³. These changes were also evidenced by plasma BAP levels, which decreased significantly in the glucocorticoid group. In the long term, the loss of BMD became more pronounced, and bone mineral content, trabecular thickness, and mechanical stability tended to be lower compared with the control group. There was a negative association between the cumulative dose of glucocorticoids and BMD, which could be traced back to impaired osteoblastogenesis.

The present study has 2 major limitations. First, we analyzed only 8 sows per group. This small number is a consequence of the animals' dimension with each sow weighing over 300 kg. On the other hand, comparable large animal studies included even smaller numbers of pigs [4,25]. The second limitation is the cross-sectional design instead of a longitudinal one, which was due to logistic reasons. One additional limitation is the impropriety of serum estrogen measurement for validation of successful OVX in the pig model, which in the case of large animal surgery might not be considered a serious drawback as the ovaries are clearly visible and of a comparably large size. The aforementioned ineligibility is due to the fact that on the one hand serum estrogens are below the detection limit in non-ovariectomized sows during diestrus and on the other hand this study gives evidence of extragonadal sexual steroid synthesis in ovariectomized sows. Nevertheless, the percentage of animals with a measurable systemic estrogen titer was significantly lower in the group of ovariectomized sows when compared to non-ovariectomized ones.

In conclusion, the skeleton of adult conventional sows is seemingly protected from effects of OVX and calcium shortage, which is a very interesting finding. Hence, these animals do not appear to be a suitable model for investigations concerning postmenopausal osteoporosis but could be used to identify factors that

protect bone from calcium or sex hormone deficiency. Moreover, this study additionally provides valuable information about yet unknown physiological data on bone metabolism parameters in adult sows. At the moment we have to accept that bone of some species such as laboratory rodents reacts to generally accepted osteoporosis-inducing stimuli such as OVX and calcium shortage as would human bone, whereas the skeleton of other species, such as the adult pig, seems to be resistant to the development of osteopenia or osteoporosis. The skeleton of bears may serve as another contradictory example, as it remains unaltered in the context of immobilization during hibernation [26]. Future studies should focus on unraveling the endocrinological and perhaps immunological mechanisms, which function as protectors of bone mass and structure in these species.

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