Obesity does not aggravate osteoporosis or osteoblastic insulin resistance in orchietomized rats

Saranyapin Potikanond^{1,2}, Pinyada Rattanachote¹, Hiranya Pintana², Panan Suntornsaratoon^{3,4}, Narattaphol Charoenphandhu^{3,4}, Nipon Chattipakorn^{2,5,6} and Siriporn Chattipakorn^{2,7}

¹Department of Pharmacology, Faculty of Medicine, Chiang Mai University, Chiang Mai, Thailand

²Neurophysiology Unit, Cardiac Electrophysiology Research and Training Center, Faculty of Medicine, Chiang Mai University, Chiang Mai, Thailand

³Center of Calcium and Bone Research (COCAB), Faculty of Science, Mahidol University, Bangkok, Thailand

⁴Department of Physiology, Faculty of Science, Mahidol University, Bangkok, Thailand

⁵Cardiac Electrophysiology Unit, Department of Physiology, Faculty of Medicine, Chiang Mai University, Chiang Mai, Thailand

⁶Center of Excellence in Cardiac Electrophysiology Research, Chiang Mai University, Chiang Mai, Thailand

⁷Department of Oral Biology and Diagnostic Sciences, Faculty of Dentistry, Chiang Mai University, Chiang Mai, Thailand

Correspondence should be addressed to S C Chattipakorn
Email
scchattipakorn@gmail.com or siriporn.c@cmu.ac.th

Abstract

The present study aimed to test the hypothesis that testosterone deprivation impairs osteoblastic insulin signaling, decreases osteoblast survival, reduces bone density, and that obesity aggravates those deleterious effects in testosterone-deprived rats. Twenty four male Wistar rats underwent either a bilateral orchietomy (O, $n = 12$) or a sham operation (S, $n = 12$). Then the rats in each group were further divided into two subgroups fed with either a normal diet (ND) or a high-fat diet (HF) for 12 weeks. At the end of the protocol, blood samples were collected to determine metabolic parameters and osteocalcin ratios. The tibiae were collected to determine bone mass using microcomputed tomography and for osteoblast isolation. The results showed that rats fed with HF (sham-operated HF-fed rats (HFS) and ORX HF-fed rats (HFO)) developed peripheral insulin resistance and had decreased trabecular bone density. In ND-fed rats, only the ORX ND-fed rats (NDO) group had decreased trabecular bone density. In addition, osteoblastic insulin resistance, as indicated by a decrease in tyrosine phosphorylation of the insulin receptor and Akt, were observed in all groups except the sham-operated ND-fed rats (NDS) rats. Those groups, again with the exception of the NDS rats, also had decreased osteoblastic survival. No differences in the levels of osteoblastic insulin resistance and osteoblastic survival were found among the NDO, HFS, and HFO groups. These findings suggest that either testosterone deprivation or obesity alone can impair osteoblastic insulin signaling and decrease osteoblastic survival leading to the development of osteoporosis. However, obesity does not aggravate those deleterious effects in the bone of testosterone-deprived rats.

Key Words

- ▶ osteoblastic insulin signaling
- ▶ testosterone deprivation
- ▶ high-fat diet induced obesity

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Introduction

Insulin and insulin receptors play important roles in bone development and in promoting bone formation (Pun *et al.* 1989, Fulzele *et al.* 2010). Several studies have demonstrated that the downstream signaling molecules of insulin receptors, including IR, IRS1 and Akt, are required for osteoblastic proliferation and differentiation (Ogata *et al.* 2000, Kawamura *et al.* 2007). Our previous study demonstrated that obese-insulin resistant rats induced by a 12-week high-fat diet (HF) resulted in impairment of osteoblastic insulin signaling, decreased osteoblastic proliferation, increased osteoblastic apoptosis, and can lead to osteoporosis in the jaw bone (Pramojanee *et al.* 2013).

Preclinical and clinical studies have reported that obesity is associated with low serum testosterone levels or testosterone deficiency (Wang *et al.* 2011, Saboor Aftab *et al.* 2013). In addition, studies indicate that a state of testosterone deficiency could impair glucose homeostasis and insulin signaling, leading to the development of obesity, insulin resistance, and type 2 diabetes mellitus (Grossmann *et al.* 2009). Several studies have also demonstrated that a testosterone-deficient condition causes osteopenia and osteoporosis (Dupree & Dobs 2004, Tsujimura 2013). However, the effects of combined testosterone deprivation and obese insulin resistance on bone morphology, osteoblastic insulin signaling and cell survival have not yet been investigated. The present study aimed to determine the effect of combined testosterone deprivation plus an obese-insulin resistant condition on bone morphology, osteoblastic insulin signaling, and osteoblastic survival. The hypotheses of the present study are that i) either testosterone deprivation alone or an obese-insulin resistant condition alone causes impairment of osteoblastic insulin signaling, a decrease in osteoblastic survival and leads to osteoporosis, or ii) an obesity-induced insulin resistant condition aggravates those deleterious effects on bone under testosterone-deprived conditions.

Materials and methods

Animal preparation

All experiments were conducted in accordance with the approved protocol of the Faculty of Medicine, Chiang Mai University and the Institutional Animal Care Committee. Twenty-four male Wistar rats weighing 180–200 g were purchased from the National Animal Center, Salaya Campus, Mahidol University, Bangkok, Thailand.

Experimental protocol

The rats were divided into two groups ($n=12/\text{group}$), either the sham-operation (S) group or the bilateral orchiectomy (O) group. After 1 week following the orchiectomy, the rats in each group were further divided into two subgroups to receive either a normal diet (ND) or a HF for an additional 12 weeks ($n=6/\text{sub-group}$), following the procedure described in a study by Pramojanee *et al.* (2013). Blood samples were collected from the rats' tails for determination of glucose, cholesterol, insulin, testosterone, and osteocalcin levels at week 0 and week 12. At the end of the experiment, animals were deeply anesthetized and sacrificed. The tibiae were removed, cleaned, and kept with in normal saline at -80°C before micro-computed tomography (micro-CT) and histomorphometry analysis. Also, total visceral fat, including peritoneal, periovarian, and perirenal fat pads were removed and weighed (Prachayasakul *et al.* 2014). A summary of the experimental protocol is provided in Fig. 1.

Orchiectomy procedure

Orchiectomy was performed using the method described in previous studies (Dulisch 1976, Pongkan *et al.* 2015).

Plasma analysis

Plasma glucose and cholesterol levels were determined by a colorimetric assay (Biotech, Bangkok, Thailand). Plasma HDL and LDL levels were determined using a commercial colorimetric assay kit (Biovision, CA, USA). Plasma insulin levels were determined using a Sandwich ELISA kit (LINCO Research, St. Claire, MO, USA). Plasma testosterone levels were measured using the electrochemiluminescence immunoassay 'ECLIA' technique (Roche Diagnostic). Peripheral insulin resistance was assessed by the homeostasis model assessment (HOMA) (Haffner *et al.* 1997,

Figure 1

Experimental schematic. At the start of the first week, rats were divided into two groups: a sham-operation (S) group and a bilateral orchiectomy (ORX) group. One week after surgery, rats in each group were divided into two subgroups, receiving either a ND or a HF for 12 weeks.

Appleton *et al.* 2005), a mathematical model describing the degree of insulin resistance, calculated from fasting plasma insulin and fasting plasma glucose concentration. A higher HOMA index indicates a higher degree of insulin resistance. The HOMA index was determined by the following equation: (fasting plasma insulin ($\mu\text{U/ml}$)) $\times$ (fasting plasma glucose (mmol/l))/22.5 (Haffner *et al.* 1997, Appleton *et al.* 2005). The plasma Gla and Glu osteocalcin levels were measured by a rat Glu-Osteocalcin high sensitive EIA Kit (Takara Bio, Inc., Shiga, Japan) with 0.125 ng/ml minimum sensitivity of detection, 4.8% inter assay, and 6.3% intra assay.

Primary osteoblastic culture

Left tibiae were prepared in pieces for osteoblastic isolation using the serial enzymatic method (Pramojane *et al.* 2013). Osteoblastic cells were identified with Alizarin red (Sigma–Aldrich) and alkaline phosphatase (Wako, Osaka, Japan), as described in a previous study (Pramojane *et al.* 2013).

Cell proliferation and apoptosis assay

The osteoblastic cells were plated on glass coverslips in 96-well plates at a density of 1×10^4 cells/well and cultured for 24 h in DMEM which was supplemented with 10% FBS as previously described (Pramojane *et al.* 2013). Cell proliferation and apoptosis assay was carried out by the following method: the osteoblasts isolated from all groups were plated in 96-well plates at the density of 1×10^4 cells/well and cultured in DMEM supplemented with 10% FBS for 24 h. Cells were starved by DMEM supplemented with 1% FBS for 24 h before stimulating with 10 nM insulin for 24 h. Cell proliferation levels were measured by a Alamarblue™ fluorometric cell viability assay (Biosource, California, USA). A TUNEL assay (Click-iT TUNEL Alexa Fluor imaging assays; Invitrogen) was used to determine cellular apoptosis levels, as described in a previous study: (Pramojane *et al.* 2013). Cell proliferation/apoptosis was given as a percentage of the control (non-insulin stimulated cells). Three sections per sample were used in the TUNEL analysis.

Western blot analysis

The osteoblastic cells were lysed in a lysis buffer (20 mmol/l Tris–HCl, 1 mmol/l Na_3VO_4 , 5 mmol/l NaF and phosphatase inhibitor) containing protease inhibitors. Protein analysis using western blot analysis was performed as previously described (Pramojane *et al.*

2013). The protein preparations were separated by 10% w/v SDS–PAGE and transferred to PVDF membranes (Millipore, MA, USA). Primary antibodies at 1:1000 were used for phosphorylation of IR (Tyr 1162/1163), IRB-subunit and Cyclin D1 (Santa Cruz Biotechnology, Inc.). β -actin at 1:2000 (Santa Cruz Biotechnology, Inc.) was used as a loading control. pAkt (Ser 473), Akt and secondary at 1: 1000, anti-rabbit or anti-mouse HRP conjugated antibodies at 1:2000 were obtained from Cell Signaling Technology, Inc. (Danvers, MA, USA). The protein bands were visualized by ECL western blotting detection reagent and quantitated using ImageJ Software (NIH, Bethesda, MA, USA; <http://imagej.nih.gov/ij/>).

Micro-CT and trabecular microstructural analysis

Right tibiae were analyzed by a micro-CT system (model SkyScan 1178; SkyScan, Kontich, Belgium) followed by hisomorphometry for trabecular microstructural analysis as previously described (Suntornsaratoon *et al.* 2014). We analyzed the results by using a section in the mid-sagittal plane, and examined 25 areas within the tibial secondary spongiosa.

Statistical analysis

All data were expressed as mean $\pm$ S.E.M. One-way ANOVA with Dunnett's multiple comparison test was used for multiple sets of data. $P < 0.05$ was considered statistically significant.

Results

HF consumption, but not testosterone deprivation, caused peripheral insulin resistance

At baseline levels (before orchietomy), no metabolic parameters were significantly different between rats in the sham-operation (S) group and rats in the orchietomized group (Table 1). The success of the orichectomy was

Table 1 Metabolic parameters at baseline (week 0) before operating on the sham-operated group (SHAM) and the orchietomized group (ORX)

Metabolic parameters	SHAM	ORX
Body weight (g)	191.67 $\pm$ 3.33	197.50 $\pm$ 4.03
Plasma glucose (mg/dl)	105.64 $\pm$ 6.49	109.18 $\pm$ 4.54
Plasma insulin (ng/ml)	2.31 $\pm$ 0.33	2.33 $\pm$ 0.31
HOMA index	14.02 $\pm$ 1.19	14.98 $\pm$ 1.98
Plasma total cholesterol (mg/dl)	48.33 $\pm$ 3.69	47.72 $\pm$ 2.53

Table 2 Metabolic parameters and plasma osteoclastin levels following either orchietomy or sham operation

Metabolic parameters	NDS	NDO	HFS	HFO
Body weight (g)	475 ± 8.5	425.8 ± 10.8 ^{*,†}	550 ± 16.9 [*]	510 ± 14.1 [†]
Visceral fat (g)	31.8 ± 2.8	18.8 ± 3.7 ^{*,†}	46.0 ± 4.5 [*]	30.7 ± 2.8 [†]
Plasma insulin (ng/ml)	1.7 ± 0.2	2.0 ± 0.3	3.1 ± 0.4 [*]	2.9 ± 0.3 [*]
Plasma glucose (mg/dl)	145.2 ± 6.0	143.2 ± 12.5	147.4 ± 6.5	146.3 ± 7.3
HOMA index	14.6 ± 1.7	17.2 ± 3.2	26.6 ± 3.0 [*]	25.2 ± 4.1 [*]
Plasma total cholesterol (mg/dl)	49.4 ± 3.7	50.0 ± 5.7	70.7 ± 3.3 [*]	65.3 ± 7.0 [*]
Plasma testosterone (ng/dl)	0.75 ± 0.2	0.07 ± 0.01 ^{*,†}	0.46 ± 0.01 [*]	0.15 ± 0.06 ^{*,†}
Gla osteoclastin (ng/ml)	150.3 ± 11.6	171.9 ± 13.1	171.7 ± 14.0	163.8 ± 30.0
Glu osteoclastin (ng/ml)	39.2 ± 3.5	51.8 ± 3.9 [*]	55.4 ± 5.6 [*]	55.2 ± 8.6 [*]
Osteocalcin Gla/Glu ratio	4.5 ± 0.4	3.3 ± 0.2 [*]	3.3 ± 0.3 [*]	3.8 ± 0.3 [*]

* $P < 0.05$ vs NDS, $^{\dagger}P < 0.05$ vs HFS.

confirmed by decreased plasma testosterone levels (Table 2). Interestingly, the testosterone levels of the sham-operated HF-fed rats (HFS) group also decreased significantly compared with that of the sham-operated ND-fed rats (NDS) group. After 12 weeks of the dietary protocol, both the HFS and the ORX HF-fed rats (HFO) rats presented characteristics of impaired peripheral insulin sensitivity as indicated by increased body weight, increased visceral fat, hyperinsulinemia with euglycemia, an increased HOMA index and increased plasma cholesterol levels when compared with the NDS group (Table 2). Although the body weight and quantity of visceral fat in the HFO group were significantly greater than in the NDS group, the HFO group had a lower body weight and less visceral fat than the HFS group (Table 2). In addition, the ORX ND-fed rats (NDO) rats had significantly decreased body weight and visceral fat compared with the NDS group. These findings suggest that a HF, but not testosterone deprivation, caused the obesity and the peripheral insulin resistance.

Decreased Gla/Glu osteocalcin ratio observed in both HF-fed rats and testosterone-deprived rats

The plasma Gla/Glu-osteocalcin ratio was used as a marker of bone formation. It was found that the Gla/Glu osteocalcin ratio was significantly decreased in the NDO, HFS, and HFO groups compared with that of the NDS group (Table 2). However, the ratios were not different among the NDO, HFS, and HFO rats. These findings suggest that either HF consumption or testosterone deprivation can cause a reduction in bone formation. Obesity, however, did not aggravate this deleterious effect in testosterone-deprived rats.

Trabecular bone microstructure decreased in HF-fed rats and in testosterone-deprived rats

The morphology of the tibial bones was analysed using a micro-CT. The cortical parameters, including x -axis moment of inertia, y -axis moment of inertia, cortical thickness, cortical bone area, cortical periosteal perimeter, and cortical endosteal perimeter were not different among the groups (Fig. 2A, B, C, D, and F). Interestingly, trabecular bone mineral density was significantly decreased in the NDO, HFS, and HFO groups (Fig. 2G). This indicated that osteoporosis occurred in this area. Bone histomorphometry was used to further investigate the bone microstructure. Goldner's trichrome-stained tibial sections indicated that testosterone deprivation and HF consumption led to a decrease in mineralized tissue (green) and bone volume (Fig. 3A and B). Testosterone deprivation in both ND and HF rats increased trabecular separation (Fig. 3D) and decreased trabecular number (Fig. 3E). The decrease in trabecular bone volume in HFS was probably the result of the decreased trabecular number rather than the decreased trabecular thickness (Fig. 3E).

Osteoblastic proliferation significantly decreased in HF-fed rats and testosterone-deprived rats

Cells isolated from tibiae and cultured in osteogenic medium were positively stained with Alizarin red (Fig. 4A, C, E, and G) and alkaline phosphatase (Fig. 4B, D, F, and H), to identify osteoblastic characteristics. AlamarBlue fluorescent cell viability assay was performed at 1, 2, 4, 6, 24, 48, and 72 h after initial culturing. The number of cells was significantly decreased in the NDO, HFS, and HFO groups at 48 and 72 h when compared with the NDS group (Fig. 4J).

Figure 2

The tibial diaphysis of rats at 12 weeks post surgery was determined by micro-CT, x-axis moment of inertia (MMIx; A), y-axis moment of inertia (MMIy; B), cortical thickness (Ct.Th; C), cortical bone area (Ct.B.Ar; D),

cortical periosteal perimeter (Ct.Pe.Pm; E), cortical endosteal perimeter (Ct.En.Pm; F), and tibial bone mineral density (T.BMD; G). * $P < 0.05$ vs NDS group.

A proliferation signal, Cyclin D1 protein, was decreased in the HF-fed rats and testosterone-deprived rats (Fig. 4L). No differences in these factors were found among the NDO, HFS, and HFO groups. These findings suggest that either

testosterone deprivation alone or HF consumption alone can cause a reduction in osteoblastic proliferation, but that obesity induced by HF consumption does not aggravate this effect in testosterone-deprived rats.

Figure 3

Micrographs of proximal tibial metaphyses (A). Scale bars = 1000 μ m. Bar graphs represent bone volume fraction (BV/TV) (B), trabecular thickness (Tb.Th) (C), trabecular separation (Tb.Sp) (D), and trabecular number

(Tb.N) (E). * $P < 0.05$ vs NDS group; ** $P < 0.01$ vs NDS group; *** $P < 0.001$ vs the NDS group; † $P < 0.05$ vs the HFS group; †† $P < 0.01$ vs the HFS group. A full colour version of this figure is available at <http://dx.doi.org/10.1530/JOE-15-0333>.

Figure 4

Alizarin red (A, C, E, and G) and alkaline phosphatase staining (B, D, F, and H) were performed to characterize osteoblasts. (I) Magnified image of cultured osteoblasts (stellate cell with nucleus; arrow) from NDS group. Alizarin-stained nodules (red) are scattered in matrix. The percentage change of cell viability using alamarBlue fluoremetric cell viability assay is shown in J. The representation of cyclinD protein expression with insulin

stimulation (+) and without insulin stimulation (–) are shown in K. Bar graphs represent densitometric values of cyclin D calculating by the level of cyclin D without insulin stimulation normalizing with level of actin (L). * $P < 0.05$ vs NDS group. A full colour version of this figure is available at <http://dx.doi.org/10.1530/JOE-15-0333>.

Osteoblastic apoptosis significantly increased in both HF-fed rats and testosterone-deprived rats

Apoptotic cell death, measured with TUNEL assay, significantly increased in the NDO, HFS, and HFO groups when compared with the NDS group (Fig. 5). No differences in apoptotic cell death were found among the NDO, HFS, and HFO groups. These findings suggest that either obesity alone or testosterone deprivation alone results in increased osteoblastic apoptosis. However, obesity induced by HF consumption did not aggravate apoptotic cell death in testosterone-deprived rats.

Osteoblastic insulin signaling impaired in both HF-fed rats and testosterone-deprived rats

The osteoblastic insulin signaling activity was determined by tyrosine phosphorylation of insulin receptors after stimulation with 10 nM of insulin for 10 min. The phosphorylation of the insulin receptor was decreased in the NDO, HFS, and HFO groups compared with that of the NDS group (Fig. 6A and B). In addition, the phosphorylation of the downstream insulin signaling molecule, Akt, was significantly decreased in the NDO,

HFS, and HFO groups when compared with that in the NDS group (Fig. 6A and C).

Discussion

Based on the findings of this study, it appears that HF consumption, not testosterone deprivation, causes obesity and peripheral insulin resistance. Furthermore, testosterone deprivation alone and HF-induced obesity alone lead to a decrease in serum osteocalcin, decreased osteoblastic proliferation, increased osteoblastic apoptosis, osteoblastic insulin resistance, and the impairment of bone microarchitectures. In addition, an obesity-induced insulin resistant condition does not aggravate those deleterious effects in the bones of testosterone-deprived rats.

The present findings and those of previous reports (Winzell & Ahren 2004, Pramojanee *et al.* 2013) demonstrate that obesity induced by 12-week HF consumption causes peripheral insulin resistance as indicated by increased body weight, increased visceral fat, hyperinsulinemia with euglycemia, and an increased HOMA index. The development of peripheral insulin resistance in the present study was dependent on HF-induced obesity, but not testosterone deprivation, as confirmed by the fact

Figure 5

Measurement of apoptotic cell death was performed by TUNEL assay. Apoptotic cells were labeled with green fluorescent (A, C, E, and G) and are indicated by arrows. Nucleus labeling with Hoechst stain is shown in B, D,

F, and H. Scale bars are 100 µm. The bar graph represents the percentage of positive TUNEL cells (I). * $P < 0.05$ vs NDS group. A full colour version of this figure is available at <http://dx.doi.org/10.1530/JOE-15-0333>.

that there was no change in the HOMA index in the NDO rats. These findings are consistent with a report by Christoffersen *et al.* (2006). However, several previous studies have reported that testosterone deprivation alone via orchiectomy (ORX) in male rodents causes impairment of insulin sensitivity (Xia *et al.* 2013, Jayaraman *et al.* 2014a). For example, Xia *et al.* (2013) found that ORX-induced testosterone deficiency increased fasting blood glucose in mature adult (10 week old) male Sprague–Dawley rats. Jayaraman *et al.* (2014a b) demonstrated that ORX adult mice (3 months of age) had impaired insulin sensitivity. Differences in the breed and age at the start of the ORX procedure (this study used young adult male Wistar rats orchiectomized at 5–6 weeks of age rather than Sprague–Dawley rats) might be the cause of the differences in the results.

Previous studies reported the role of osteocalcin in increased insulin sensitivity in the target organs (Saleem

et al. 2010, Xiao *et al.* 2014). The present study showed that the ratio of Gla/Glu ratio and Glu osteocalcin levels in NDO, HFS, and HFO groups significantly reduced, when compared with that of the NDS group. However, Gla osteocalcin levels among all four groups were not significantly different. These findings suggested that the reduction of Gla/Glu osteocalcin ratio in NDO, HFS, and HFO groups was due to increased Glu levels, implying that decreased insulin sensitivity could occur in NDO, HFS, and HFO groups. The present study, however, also demonstrated a reduction in the Gla/Glu osteocalcin ratio as well as the impairment of osteoblastic insulin signaling as indicated by decreased phosphorylation of the insulin receptor and its downstream signaling molecule, Akt protein, in either orchiectomy alone or HF-induced obesity alone. Our findings suggested that the osteoblastic insulin resistance in both testosterone-deprived rats and obese-insulin resistant rats could possibly occur via the

Figure 6

Effects of HFD and testosterone deprivation on insulin signaling. Cells were stimulated either with or without 10 nM of insulin for 10 min. Western blotting bands of pIR, total IR, pAkt, total Akt, and actin were shown in A. The bar graphs represent densitometric values of pIR (B) and pAkt (C) which

are calculating by divided the insulin stimulation by without insulin stimulation (insulin+/insulin-), then normalizing with total protein expression. * $P < 0.05$ vs NDS group.

effects of low osteocalcin levels as previously observed in pancreatic β cells and adipocytes (Ferron *et al.* 2008, Hwang *et al.* 2012).

The testosterone deprivation following orchiectomy was confirmed by the decreased testosterone level in the orchiectomized rats. However, the HFS rats also showed a decrease in plasma testosterone levels, a finding consistent with previous clinical reports which demonstrated that obesity in men is associated with low testosterone levels (Barrett-Connor 1992, Andersson *et al.* 1994, Grossmann 2013, Mogri *et al.* 2013). The reduction of testosterone levels in the HFS rats could be a factor in the decrease in the hypothalamo-pituitary-gonadal axis due to inflammation following obesity as described in previous studies (Cohen 1999, Gregor & Hotamisligil 2011, Mammi *et al.* 2012, Jayaraman *et al.* 2014b). Although it has been shown that obesity could increase the estrogen levels in female rats (Pratchayasakul *et al.* 2011), our previous study demonstrated that 12 weeks of high-fat diet feeding did not affect the circulating estrogen levels in male rats (Pratchayasakul *et al.* 2011). Moreover, a recent report also demonstrated that estradiol administration did not affect cardiac function in orchiectomized male rats (Ayaz *et al.* 2015). All of these findings indicated the important role of decreased testosterone levels rather than increased estrogen levels in obese male rats.

A previous study demonstrated that orchiectomy-induced trabecular bone degeneration was possibly caused by a decrease in the expressions of osteogenic genes (Chin & Ima-Nirwana 2015). Several studies have demonstrated that testosterone played an important role in osteoblastic proliferation by enhancing mitogenesis and differentiation in bone cells (Sömjen *et al.* 1989, Kasperk *et al.* 1990, Gray *et al.* 1992). In addition, testosterone has been

shown to play a role in osteoblast apoptosis in which a lack of testosterone enhanced osteoblast apoptosis through an increased Bax/Bcl-2 ratio (Wiren *et al.* 2006, Chen *et al.* 2010). Our results showed that a reduction of the Gla/Glu osteocalcin ratio, decreased osteoblastic proliferation and increased osteoblastic apoptosis in testosterone-deprived rats was associated with microstructural changes in the tibial trabecular bones and a significant overall reduction in mineral density of those bones. Our findings and a previous report (Fulzele *et al.* 2010) demonstrated that the impairment of osteoblastic insulin receptors results in a decrease in osteoblast proliferation. All of these findings suggest that testosterone deprivation decreases osteoblastic activity and osteoblastic proliferation while increasing osteoblastic apoptosis, leading to osteoporosis.

The effects of obesity on bone density is still unclear. Several studies have shown that obesity could increase bone density (Harris *et al.* 1992, Albala *et al.* 1996, Vandewalle *et al.* 2013); however, other studies have found that obesity has either no effect on bone density (Greco *et al.* 2010) or increases bone porosity (Rosen & Buxsein 2006, Zhao *et al.* 2008, Gómez *et al.* 2009). The present study, also, found that obese and insulin resistant rats had a decreased Gla/Glu-osteocalcin ratio, decreased osteoblastic proliferation, increased osteoblastic apoptosis, osteoblastic insulin resistance and increased bone porosity. These findings indicate the deleterious effects of obesity on bone quality. Interestingly, obesity did not aggravate those forms of deterioration in testosterone-deprived rats as indicated by the lack of differences in trabecular bone quality between the HFO and the NDO groups (Fig. 3B, D, and E). There are several possible explanations for these findings. First, the deleterious effects found in osteoblasts of testosterone-deprived rats

could have been at their maximum level, thus obesity could not have had further additive effects on those osteoblastic cells. Secondly, it could be that the lack of testosterone, rather than obesity, that was the major aggressive factor causing the impairment of trabecular bone structures as this impairment was also observed in NDO rats. However, in the present study no significant differences in cortical bones were observed among the experimental groups. Two previous studies found that the cortical bone markers had markedly decreased 6 months after castration (Danielsen *et al.* 1992, Reim *et al.* 2008). It is possible that the 3-month period of castration in the present study might not have been long enough to observe cortical bone change. However, the hypothesis of this study is supported by the fact that the response of cortical bone mass to HF-induced obesity differs by age (Ionova-Martin *et al.* 2011). The Ionova-Martin group demonstrated that femur thickness does not change in young mice after 16 weeks of HF consumption although those mice, as in the mice in the present study, had poor bone quality.

The limitation of the present study was that testosterone deprivation and obesity were not independently assessed as the cause of osteoporosis due to the fact that decreased testosterone levels was also observed in the obese-insulin resistant rats. However, it has been shown previously that testosterone deprivation alone could cause osteoporosis (Tuck & Francis 2009). Moreover, the recent study by Donner *et al.* (2015) demonstrated that diet-induced visceral obesity resulted in decreased bone mineral area and content and impaired femoral stiffness and strength without changes in testosterone levels. All of these findings indicated that either obesity or testosterone deprivation could lead to osteoporosis. In addition, the aim of the present study was to mainly investigate the synergistic effects of obesity and testosterone deprivation on osteoporosis. Findings from the present study clearly demonstrated that either obesity alone or testosterone deprivation alone led to osteoporosis, however no additive or synergistic effect of obesity and testosterone deprivation on osteoporosis could be found.

Lack of results pertaining to bone cellular activities is one of the limitations of the present study. However, neither serum bone biomarkers nor tissue samples for dynamic bone histomorphometry were collected from those rats. Therefore, it is not possible for us to quantify bone cellular activities in the present study. However, we believe that the available bone parameters provided in the present study provided sufficient evidence to confirm the detrimental effects of testosterone deprivation and insulin

resistance on bone. The overall adiposity was not measured. However, the visceral fat was determined and used to represent the level of metabolic complications in this obese-insulin resistant model (Hansen *et al.* 1997, Morakinyo *et al.* 2015).

Conclusions

Either testosterone deprivation alone or HF-induced obesity alone can cause impairment of osteoblastic insulin signaling and reduction in osteoblastic survival, leading to osteoporosis. However, obesity does not augment the adverse effects on bone density under testosterone-deprived conditions.

Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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Author contribution statement

S P performed the experiments, analyzed the data, and wrote the manuscript. P R performed the experiments and analyzed the data. H P performed the animal experiments and analyzed the metabolic data. P S and Na C performed and analyzed the bone histomorphometry. N C designed the study and wrote the manuscript. S C designed the study, analyzed the data, and wrote the manuscript.

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