GGCX and VKORC1 inhibit osteocalcin endocrine functions

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Osteocalcin (OCN) is an osteoblast-derived hormone favoring glucose homeostasis, energy expenditure, male fertility, brain development, and cognition. Before being secreted by osteoblasts in the bone extracellular matrix, OCN is γ-carboxylated by the γ-carboxylase (GGCX) on three glutamic acid residues, a cellular process requiring reduction of vitamin K (VK) by a second enzyme, a reductase called VKORC1. Although circumstantial evidence suggests that γ-carboxylation may inhibit OCN endocrine functions, genetic evidence that it is the case is still lacking. Here we show using cell-specific gene inactivation models that γ-carboxylation of OCN by GGCX inhibits its endocrine function. We further show that VKORC1 is required for OCN γ-carboxylation in osteoblasts, whereas its paralogue, VKORC1L1, is dispensable for this function and cannot compensate for the absence of VKORC1 in osteoblasts. This study genetically and biochemically delineates the functions of the enzymes required for OCN modification and demonstrates that it is the uncarboxylated form of OCN that acts as a hormone.

Introduction

It was recently established that the osteoblast, the bone-forming cell, produces a hormone, osteocalcin (OCN), that promotes insulin secretion by pancreatic β cells and glucose utilization by peripheral tissues (Ferron and Lacombe, 2014). Accordingly, when fed a normal chow diet (ND), OCN-deficient mice (Ocn−/−) display glucose intolerance, low circulating levels of insulin, and insulin insensitivity (Lee et al., 2007). Cell-based and genetic studies have further shown that OCN stimulates insulin secretion by acting directly on pancreatic β cells (Hinoi et al., 2008; Pi et al., 2011; Wei et al., 2014b) and indirectly through stimulation of glucagon-like peptide-1 (GLP-1) secretion by the gut (Mizokami et al., 2013). Moreover, OCN promotes β cell proliferation by increasing Ccnd1, Ccnd2, and Cdk4 expression in these cells (Ferron et al., 2008) and favors insulin sensitivity in liver, muscle, and adipose tissue (Lee et al., 2007; Wei et al., 2014a). In addition, OCN increases energy expenditure and prevents high-fat diet (HFD)–induced obesity by stimulating Ucp1 expression in brown adipose tissue (Ferron et al., 2008, 2012). Independently of its metabolic actions, OCN promotes male fertility by stimulating testosterone production by Leydig cells in the testis (Oury et al., 2011). OCN signaling in β cells and in Leydig cells occurs through GPRC6A, a G protein–coupled receptor (GPCR) of the class C (Pi et al., 2008, 2011; Oury et al., 2011, 2013a; Wei et al., 2014b). Lastly, OCN is necessary for hippocampal development and favors spatial learning and memory in part by enhancing the synthesis of monoamine neurotransmitters in the brain (Oury et al., 2013b).

To become a hormone, OCN undergoes two posttranslational modifications. First, inside the osteoblast, three glutamic acid residues of OCN are converted to γ-carboxyglutamic acid (GLA) residues through γ-carboxylation (Hauschka et al., 1989). Extensive study of the γ-carboxylation of coagulation factors in the liver has demonstrated that this posttranslational modification generally involves two enzymes, γ-glutamyl carboxylase

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Abbreviations used in this paper: GTT, glucose tolerance test; HFD, high-fat diet; ITT, insulin tolerance test; ND, normal chow diet; OCN, osteocalcin; GPCR, quantitative PCR; VK, vitamin K; VKO, VK epoxide; WT, wild type.
(γ-carboxylase or GGCX) and vitamin K (VK) epoxide (VKO) reductase (VKOR or VKORC1), which together constitute the VK cycle (Stafford, 2005). Indeed, γ-carboxylase requires reduced VK (VKH2) as an obligate cofactor, which upon γ-carboxylation is oxidized to VKO by GG CX. VKO is next converted back to VKH2 by VKORC1, allowing another γ-carboxylation reaction to take place. Because the γ-carboxylation of OCN increases its affinity for hydroxyapatite, the mineral component of the bone ECM (Hoang et al., 2003), the vast majority of OCN secreted by osteoblasts gets trapped in bone ECM, where it is the most abundant bone noncollagenous protein (Hauschka et al., 1989).

The second posttranslational modification occurs during the bone resorption phase of bone remodeling when the low pH of the resorption lacuna allows a partial decarboxylation of the OCN molecules present in the bone ECM before reaching the bloodstream (Ferron et al., 2010a; Lacombe et al., 2013). Indeed, both the γ-carboxylated (GLA) and the undercarboxylated (GLU and GLU13) forms of OCN can be detected in the general circulation. This raised the question of which forms of OCN are endowed with the hormonal functions.

Several lines of evidence of purely correlatively nature have suggested that the endocrine functions of OCN may be fulfilled by its undercarboxylated form (Fulzele et al., 2010; Rached et al., 2010; Pi et al., 2011; Ferron et al., 2012; Mizokami et al., 2013; Zhou et al., 2013), and many but not all studies conducted in humans have shown a negative correlation between the serum levels of GLU-OCN and blood glucose levels, insulin resistance, obesity, diabetes, or markers of the metabolic syndrome (Hwang et al., 2009; Kanazawa et al., 2011; Levinger et al., 2011; Pollock et al., 2011; Bulló et al., 2012; Furu syo et al., 2013; Wang et al., 2013; Chen et al., 2014). In that model, OCN produced by osteoblasts would be stored as a γ-carboxylated and inactive protein in the bone ECM, before being activated by decarboxylation during bone resorption. If this model were correct, then blocking OCN γ-carboxylation would prevent OCN accumulation in the bone ECM and improve glucose homeostasis. This assumption has never been tested, and as a result, direct evidence demonstrating that γ-carboxylation is detrimental for the endocrine functions of OCN is still lacking. This is a critical question to address because several other human studies have questioned the existence of a correlation between serum levels of GLU-OCN and insulin resistance or insulin secretion (Shea et al., 2009; Diamanti-Kandarakis et al., 2011; Absey et al., 2012; Knapen et al., 2012; Lu et al., 2012; Mori et al., 2012; Polgreen et al., 2012; Diaz-López et al., 2013).

All vertebrates possess a second VKORC1-like encoding gene named VKORCIL1 (Robertson, 2004). Despite a relatively high degree of similarity (45%) with VKORC1, the physiologic al functions of VKORCIL1 in any cell type remain unknown. In this study, using cell-specific gene deletion mouse models for Ggcx, Vkorcl, or Vkorcll, we directly assayed what was the influence of γ-carboxylation on OCN endocrine function in vivo. We also tested whether VKORC1 and VKORCIL1 have redundant functions in terms of regulating OCN γ-carboxylation in vivo or in cell culture. To that end, we used mainly the regulation of energy metabolism by OCN as a readout of its endocrine functions. We show here that even a partial Ggcx inactivation in osteoblasts is sufficient to improve glucose tolerance. Furthermore, the osteoblast-specific inactivation of Vkorcl decreases the γ-carboxylation of OCN, whereas whole body or cell-specific inactivation of Vkorclll does not. These results establish genetically that it is the undercarboxylated form of OCN that fulfills its endocrine functions.

**Results**

**Partial osteoblast-specific inactivation of Ggcx improves glucose metabolism**

We first assessed ex vivo the respective importance of the γ-carboxylase (GGCX) and VK for the γ-carboxylation of OCN. For that purpose, Ggcxfl/fl osteoblasts were infected with adenovirus expressing either GFP or the Cre recombinase to generate control or Ggcx-deficient osteoblasts, respectively (see Materials and methods and Fig. S1, A–C). Ggcx expression was decreased by >90%, and GGCX protein levels were nearly unde tectable in Cre-expressing compared with controls osteoblasts, demonstrating that an efficient inactivation has occurred (Fig. 1 A). These osteoblasts were then cultured either in the absence of VK or in the presence of VKO or VK1 for 18 h, before assessing the γ-carboxylation status of OCN in the culture media. As shown in Fig. 1 B and in Fig. S1 D, even in the absence of VK, deletion of Ggcx in osteoblasts reduced the amount and the percentage of γ-carboxylated OCN secreted. Importantly, although the addition of VKO and VK1 increased, albeit modestly, γ-carboxylation of OCN found in the culture media of control cells, it did not affect OCN γ-carboxylation in Ggcx-deficient osteoblasts (Fig. 1 B and Fig. S1 D). This result confirms that GGCX is required for OCN γ-carboxylation ex vivo.

To determine whether GGCX affects OCN endocrine functions, we generated osteoblast-specific, Ggcx-deficient mice by crossing α1(I)collagen-Cre (Col1a1-Cre) transgenic mice, which delete floxed genes in osteoblasts only (Dacquin et al., 2002), with mice harboring a floxed allele of Ggcx (Fig. S1, A–C). Recombination at the Ggcx locus in the Ggcxfl/fl;Col1a1-Cre mice was detected in bone, but in no other tissues (Fig. 1 C). The efficiency of recombination at the genomic level did not exceed 50% in osteoblast cultures derived from Ggcxfl/fl;Col1a1-Cre bone marrow (Fig. S1 E). Measurement of the different forms of OCN in the serum revealed that the circulating levels of GLA-OCN and total OCN were not significantly altered and that the circulating levels of GLU-OCN were increased three to four times in Ggcxfl/fl;Col1a1-Cre compared with control Ggcxfl/fl mice (Fig. 1, D and E). Together, these results indicate that we had achieved a specific, but partial inactivation of Ggcx in osteoblasts in vivo.

As a way to characterize the impact of reducing GGCX activity in osteoblasts on the endocrine functions of OCN, we tested, in adult Ggcxfl/fl;Col1a1-Cre and control mice fed an ND, an array of physiological functions known to be regulated by OCN (Lee et al., 2007; Ferron et al., 2008, 2010a; Fulzele et al., 2010). Intraperitoneal glucose tolerance tests (GTTs) revealed that Ggcxfl/fl;Col1a1-Cre male mice handled a glucose load significantly better than control animals (Fig. 2 A). This was explained, at least in part, by an enhanced insulin sensitivity in
**Partial osteoblast-specific inactivation of Ggcx prevents diet-induced glucose intolerance**

In view of the results presented above, we next asked whether the chronic elevation of the circulating GLU-OCN levels in Ggcx\textsuperscript{fl/fl},Col1a1-Cre mice could weaken if not prevent the deleterious metabolic effects associated with consuming an HFD. To test this contention, 4-wk-old control and Ggcx\textsuperscript{fl/fl},Col1a1-Cre mice were fed an HFD for 14 wk, and body weight gain, glucose tolerance, and insulin sensitivity were then assessed at the end of this period. HFD-fed Ggcx\textsuperscript{fl/fl},Col1a1-Cre mice gained significantly less weight than age-matched control mice fed the same HFD, and this difference remained significant from 6 wk onward (Fig. 3 A). HFD-fed Ggcx\textsuperscript{fl/fl},Col1a1-Cre mice also had a better insulin sensitivity (Fig. 2 C), suggesting that the reduction in body weight was secondary to a decrease in the size of the fat depots in these mice. Because GLU-OCN injections or infusions can prevent fat accumulation by increasing whole body energy expenditure in mice (Ferron et al., 2008, 2012; Zhou et al., 2013), we also assessed energy expenditure of the Ggcx\textsuperscript{fl/fl},Col1a1-Cre mice through indirect calorimetry. Ggcx\textsuperscript{fl/fl},Col1a1-Cre mice had increased O\textsubscript{2} consumption, CO\textsubscript{2} production, and overall energy expenditure when compared with control littermates (Fig. 2 D).

As hypothesized, fasting blood glucose was increased in control mice fed an HFD compared with those fed an ND. In contrast, Ggcx\textsuperscript{fl/fl},Col1a1-Cre mice fed an HFD displayed fasting glycemia comparable with the ones of control mice fed an ND (Fig. 3 B). Strikingly, glucose tolerance determined by GTT was indistinguishable in Ggcx\textsuperscript{fl/fl},Col1a1-Cre mice fed an HFD and control mice fed an ND, whereas Ggcx\textsuperscript{fl/fl} fed an HFD displayed the expected glucose intolerance (Fig. 3 C). Ggcx\textsuperscript{fl/fl},Col1a1-Cre mice fed an HFD also had a better insulin sensitivity compared with control mice fed an HFD (Fig. 3 D). Although control and Ggcx\textsuperscript{fl/fl},Col1a1-Cre mice fed an HFD were equally hyperinsulinemic when compared with mice fed an ND, HOMA-IR, an index of insulin sensitivity, was consistently...
reduced, although not significantly, in $Ggcx^{fl/fl}; Col1a1$-Cre mice (Fig. S1, G and H). These data indicate that a chronic elevation of GLU-OCN can prevent the obesity and glucose intolerance induced by HFD in wild-type (WT) mice.

**Consequences of a more complete inactivation of Ggcx on OCN endocrine functions**

Even though there was a significant increase in circulating GLU-OCN levels and an improvement of their glucose tolerance in the $Ggcx^{fl/fl}; Col1a1$-Cre mice, the inactivation of GGCX in osteoblasts was rather partial in these mice because GLA-OCN serum levels were marginally decreased (Fig. 1 D), further suggesting that a much more efficient inactivation of GGCX occurred in osteoblasts derived from $Ggcx^{fl/fl}; Col1a1$-Cre mice compared with control mice (Fig. 4 E), establishing in vivo that the deletion of $Ggcx$ had been achieved when using the $OC$-Cre deleter mice.

GLA residues allow OCN to bind to hydroxyapatite, the mineral component of the bone ECM (Hoang et al., 2003), and in vitro GLU-OCN has a reduced binding capacity for hydroxyapatite compared with control $Ggcx^{fl/fl}$ mice (Fig. 4 C). In addition, serum levels of GLA-OCN were increased >15 times (from 10 to over 150 ng/ml) in $Ggcx^{fl/fl}; OC$-Cre mice compared with control $Ggcx^{fl/fl}$ mice (Fig. 1 D), further suggesting that a much more efficient inactivation of $Ggcx$ had been achieved when using the $OC$-Cre deleter mice.
3-mo-old mice did not affect trabecular or cortical bone density as assessed by µCT or histologically; the same was true for osteoblast and osteoclast numbers, for bone formation rate, and for osteoclastic activity, as estimated by serum CTx levels (Fig. S2, A and B). Moreover, in cell culture differentiation markers of osteoblasts were normally expressed in Ggcx-deficient primary osteoblasts, and the deletion of Ggcx did not overly impact the ability of osteoblasts to mineralize the ECM (Fig. S2, C and D).

As observed in Ggcx<sup>fl/fl</sup>;Col1a1-Cre, 3-mo-old Ggcx<sup>fl/fl</sup>, OC-Cre mice displayed an improved glucose tolerance compared with control (Fig. 4 F) without hypergonadism (Fig. S1 F). This metabolic phenotype was less severe in Ggcx<sup>fl/fl</sup>;Col1a1-Cre;OC-Cre mice. This is consistent with the bell-shaped dose-response curve of target cells to OCN in cell culture (Ferron et al., 2008; Oury et al., 2011) and with the notion previously advanced that desensitization can occur when circulating OCN levels are too high (Riddle et al., 2014). To test whether VKORC1 contributes to OCN γ-carboxylation (Fig. 5 A), mice harboring a floxed allele of Vkorc1 (Fig. S3, A–C) were first bred with Cmv-Cre mice (Schwenk et al., 1995) to inactivate Vkorc1 in germ cells and to obtain mice carrying a null allele of this gene (Vkorc1<sup>−/−</sup>; Fig. S3 D). When Vkorc1<sup>−/−</sup> mice were intercrossed, no Vkorc1<sup>−/−</sup> pups were retrieved at postnatal day 14 (P14; Fig. S3 E). A careful monitoring revealed that Vkorc1<sup>−/−</sup> mice were born at the expected Mendelian ratio but died within 2–5 d of intra-abdominal hemorrhages (Fig. S3, E and F). These results, which confirm a previous study on global Vkorc1 deletion (Spohn et al., 2009), established that an efficient ablation of VKORC1 activity occurred in Vkorc1<sup>−/−</sup> mice in vivo. Western blot analysis also confirmed the absence of Vkorc1 protein in Vkorc1<sup>−/−</sup> liver (Fig. S3 G).

Vkorc1<sup>−/−</sup> mice were next bred with OC-Cre mice to inactivate Vkorc1 only in osteoblasts. As shown in Fig. 5 B, VKORC1 was deleted in bone but not in any of the other tested tissues in Vkorc1<sup>−/−</sup>;OC-Cre mice. Unlike Vkorc1<sup>−/−</sup>/mice, Vkorc1<sup>−/−</sup>;OC-Cre mice were viable and did not present any obvious hemostatic defects. Western blotting analyses demonstrated that VKORC1 protein accumulation was considerably reduced in Vkorc1<sup>−/−</sup>;OC-Cre osteoblasts compared with controls (Fig. 5 C). Serum measurement of the various forms of OCN revealed that circulating levels of GLA-OCN were decreased by ∼40%, whereas those of GLU-OCN were increased by ∼10-fold in Vkorc1<sup>−/−</sup>;OC-Cre mice (Fig. 5, D and E). Circulating levels of total OCN were unaffected in the same animals (Fig. 5 D). Of note, very similar changes in OCN γ-carboxylation status were observed in the serum of newborn Vkorc1<sup>−/−</sup> pups (Fig. S3 H), indicating that VKORC1 expression in osteoblasts is sufficient to mediate OCN γ-carboxylation. Similarly to what was observed in Ggcx<sup>fl/fl</sup>;Col1a1-Cre mice, the

VKORC1 is necessary for OCN γ-carboxylation in vivo and in cell culture

VKORC1 is the enzyme responsible for reducing VKO in hepatocytes, thereby allowing the γ-carboxylation of various coagulation factors (Li et al., 2004; Rost et al., 2004). To test whether VKORC1 contributes to OCN γ-carboxylation (Fig. 5 A), mice harboring a floxed allele of Vkorc1 (Fig. S3, A–C) were first bred with Cmv-Cre mice (Schwenk et al., 1995) to inactivate Vkorc1 in germ cells and to obtain mice carrying a null allele of this gene (Vkorc1<sup>−/−</sup>; Fig. S3 D). When Vkorc1<sup>−/−</sup> mice were intercrossed, no Vkorc1<sup>−/−</sup> pups were retrieved at postnatal day 14 (P14; Fig. S3 E). A careful monitoring revealed that Vkorc1<sup>−/−</sup> mice were born at the expected Mendelian ratio but died within 2–5 d of intra-abdominal hemorrhages (Fig. S3, E and F). These results, which confirm a previous study on global Vkorc1 deletion (Spohn et al., 2009), established that an efficient ablation of VKORC1 activity occurred in Vkorc1<sup>−/−</sup> mice in vivo. Western blot analysis also confirmed the absence of Vkorc1 protein in Vkorc1<sup>−/−</sup> liver (Fig. S3 G).

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Although GGXC and VKORC1 proteins were decreased to similar levels by the OC-Cre–mediated inactivation, circulating GLA-OCN levels were only 40% reduced in Vkorc1fl/fl;OC-Cre mice, whereas they were reduced by >80% in Ggcxfl/fl;OC-Cre mice. Contents of GLA-OCN and total OCN in bone were reduced by 22 and 8 times, respectively, in Vkorc1fl/fl;OC-Cre mice (Fig. 5F). Altogether, these results suggest that Vkorc1 in osteoblasts is required for efficient OCN γ-carboxylation in vivo.

Figure 4. Efficient reduction of OCN γ-carboxylation in Ggcxfl/fl;OC-Cre mice. (A) Detection of the deletion allele (∆PCR) of Ggcx by PCR on genomic DNA isolated from tissues of Ggcxfl/fl;OC-Cre and Ggcxfl/fl mice. PCR for the floxed allele (3’lox PCR) was used as a loading control. (B) Western blotting analyses of GGXC expression in Ggcxfl/fl and Ggcxfl/fl;OC-Cre bone marrow–derived osteoblasts. VAT and BAT, white and brown adipose tissue, respectively. (C and D) Serum levels of GLA- and total OCN (C) and of GLU-OCN (D) in Ggcxfl/fl (n = 4) and Ggcxfl/fl;OC-Cre (n = 4) 2-mo-old mice. (E) Bone OCN content in Ggcxfl/fl (n = 6) and Ggcxfl/fl;OC-Cre (n = 6) 3-mo-old mice was assessed from whole bone extracts by Western blotting (left) or ELISA and normalized to the total protein content (right). (F) GTTs in Ggcxfl/fl (n = 13) and Ggcxfl/fl;OC-Cre (n = 16) 3-mo-old mice fed an ND. Mice were fasted for 16 h and injected i.p. with 2 g/kg glucose. (G) GTTs in Ggcxfl/fl (n = 7), Ggcxfl/fl;Ocn+/- (n = 9), Ggcxfl/fl;OC-Cre;Ocn+/- (n = 9), and Ggcxfl/fl;OC-Cre (n = 9) mice fed an ND. Mice were fasted for 16 h and injected i.p. with 1.3 g/kg glucose. Results are given as means ± SEM. **, P < 0.01; ***, P < 0.001. (G) #, P < 0.05 when comparing with Ggcxfl/fl, Ggcxfl/fl;Ocn+/-, and Ggcxfl/fl;OC-Cre;Ocn+/- mice.
mice (Figs. 4 C and 5 D). This observation implied that the absence of VKORC1 in osteoblasts might be partially compensated in vivo. It has previously been shown that exogenous VK1 could rescue bleeding defects in Vkorc1<sup>−/−</sup> mice or in humans carrying a mutation in VKORC1 (Rost et al., 2004; Spohn et al., 2009). We therefore hypothesized that the remaining GLA-OCN detected in Vkorc1<sup>−/−</sup>; OC-Cre could be explained, in part, by the absorption of VK1 through food, milk, and/or more likely feces because mice are coprophagic (Fig. 5 A). In support of this hypothesis, circulating GLA-OCN levels were further reduced in Vkorc1<sup>−/−</sup>; OC-Cre mice when coprophagy was prevented (Fig. S3 I). In addition, Vkorc1<sup>−/−</sup> osteoblasts displayed a...
That the improvement in glucose tolerance is consistently more pronounced in Ggcxfl/fl;Col1a1-Cre than in Vkorc1fl/fl;OC-Cre mice, even though they had a more modest increase in GLU-OCN (i.e., 4 vs. 8 times), further suggests that chronically high levels of circulating GLU-OCN may result in OCN receptor desensitization.

Another possible explanation for the results presented could be that a second enzyme harboring VK1 reductase activity and compensating for the loss of VCORC1 may exist in osteoblasts (Fig. 6 A). This hypothesis is indirectly supported by the fact that a paralogue of VKORC1, VKORC1L1, is present in the defect in OCN γ-carboxylation in the absence of VK or in the presence of VKO, similar to the one observed in Ggcx−/− osteoblasts (Fig. 5, G and H; and Fig. S3 J). However, unlike what was observed in Ggcx−/− osteoblasts, the OCN γ-carboxylation defect in Vkorc1−/− osteoblasts was rescued by addition of large (micromolar) amounts of VK1 in the media (compare Fig. 5 H with Fig. 1 B). Deletion of Vkorc1 did not alter the capacity of osteoblasts to mineralize the ECM in culture (Fig. S3 K). These results indicate that although VKORC1 is required for OCN γ-carboxylation in osteoblasts, its absence can be compensated by exogenous high concentrations of VK1 in the mouse.

Inactivation of Vkorc1 in osteoblasts caused a slight and not significant improvement in glucose tolerance when fed a chow diet (Fig. 5 I). That the improvement in glucose tolerance is consistently more pronounced in GgcxloxP;Coll1−/Cre than in Vkorc1loxP;OC-Cre mice, even though they had a more modest increase in GLU-OCN (i.e., 4 vs. 8 times), further suggests that chronically high levels of circulating GLU-OCN may result in OCN receptor desensitization.

**VKORC1L1 is dispensable for OCN γ-carboxylation and for hemostasis in vivo**

Another possible explanation for the results presented could be that a second enzyme harboring VK1 reductase activity and compensating for the loss of VCORC1 may exist in osteoblasts (Fig. 6 A). This hypothesis is indirectly supported by the fact that a paralogue of VKORC1, VKORC1L1, is present in the
genome of all vertebrates and that in vitro and cell-based assays have indicated that VKORC1L1 can have VKO and VK1 reductase activity (Westhofen et al., 2011; Hammed et al., 2013; Tie et al., 2013). Thus, we tested the importance of VKORC1L1 for OCN \(\gamma\)-carboxylation in vivo by generating mice lacking Vkorcl1 globally or only in osteoblasts.

Mice harboring a floxed allele of Vkorcl1 were bred with OCN-Cre mice to generate animals in which VKORC1L1 was deleted in osteoblasts only (Fig. S4, A–C). In these Vkorcl1\(^{fl/fl}\), OCN-Cre mice, the deletion of Vkorcl1 could be detected in bone tissue, but in no other tissues (Fig. 6 B). Quantitative PCR (QPCR) analysis demonstrated that Vkorcl1 expression was efficiently reduced in osteoblasts derived from Vkorcl1\(^{fl/fl}\), OCN-Cre mice compared with control cells (Fig. S4 D). However, measurement of the circulating levels of the various forms of OCN revealed that deleting Vkorcl1 in osteoblasts had no detectable impact on the circulating levels of GLA-, GLU-, or OCN levels in those double mutant mice with those of either VKORC1, VKORC1L1, or both, these data further support the notion that in this cell type GLA proteins become \(\gamma\)-carboxylated while they are transiting through the ER. Although it is assumed that \(\gamma\)-carboxylation also takes place in the ER of osteoblasts, the intracellular localization of the VK cycle components in these cells has never been tested. Our cell biology analysis of this aspect of VK biology showed that immunostaining for GGCX and VKORC1 both overlapped with the signal obtained with an antibody raised against the KDEL sequence, a marker of ER (Fig. 7 E, top and middle). In contrast, the intracellular localization pattern of VKORC1L1 differed greatly and did not show such overlap with the ER (Fig. 7 E, bottom). These results suggest that in osteoblasts GGCX and VKORC1 are enriched in the ER, whereas \(\gamma\)-carboxylation was shown to take place, whereas VKORC1L1 is not localized to this compartment. Combined with the data obtained in mice lacking VKORC1, VKORC1L1, or both, these data further support the conclusion that VKORC1L1 is not involved in the \(\gamma\)-carboxylation of OCN.

**Discussion**

### Role of GLU-OCN in glucose metabolism

Several lines of evidence have established that bone is an endocrine organ (Karsenty and Ferron, 2012). In vitro and in vivo evidence indicates that OCN, an osteoblast-specific protein, affects glucose metabolism in several ways (Lee et al., 2007; Ferron et al., 2008). In addition, ablation of osteoblasts in adult mice or the suppression of OCN production by glucocorticoids results in metabolic abnormalities that are mostly rescued by injections of GLU-OCN (Yoshikawa et al., 2011; Brennan-Speranza et al., 2012). These data further suggest that osteoblasts and GLU-OCN play a central role in the control of energy metabolism by bone.

Since the identification of the metabolic functions of OCN in mice, several cross-sectional and observational studies have addressed the role of OCN in glucose and energy metabolism in humans (reviewed by Ferron and Lacombe [2014]). The majority of those studies reported that serum levels of GLU-OCN negatively correlate with blood glucose levels, insulin resistance, diabetes, obesity, or markers of the metabolic syndrome. Moreover, some data obtained in humans also support a role for...
Figure 7. **VKORC1L1 does not compensate for VKORC1 absence in osteoblasts.** (A) Serum levels of GLU-OCN in control (n = 12), Vkorc1fl/fl;OC-Cre (n = 4), Vkorc1l1fl/fl;OC-Cre (n = 5), Vkorc1fl/fl;Vkorc1l1+/fl;OC-Cre (n = 5), and Vkorc1fl/fl;Vkorc1l1fl/fl;OC-Cre (n = 6) 2-mo-old mice. (B) Bone OCN content in 2-mo-old mice was assessed from whole bone extracts by Western blotting. Each lane represents an individual animal. (C) Cre-mediated inactivation of VKORC1 and VKORC1L1 in Vkorc1fl/fl;Vkorc1l1fl/fl osteoblasts infected with either Ad-GFP or Ad-Cre was assessed by Western blotting. (D) Percentage of GLA over total OCN measured in the supernatant of osteoblasts of the indicated genotype cultured in the absence of VK (−VK) or in the presence of VKO (+VKO) or VK1 (+VK1; n = 4 for each condition). (E) Immunofluorescence analyses of mouse osteoblasts. Cells were stained with rabbit antibodies against GGCX (top), VKORC1 (middle), or VKORC1L1 (bottom) and with a mouse anti-KDEL antibody to label the ER. The areas boxed on the merge panels are zoomed in the panels labeled “merge (zoom).” The graphs displayed on the right show the intensity of each of the fluorescent signals (green, red, and blue) in a 50-µm cross-section of each cell (dashed lines). At least 20 individual cells were analyzed for each staining condition, and comparable results were obtained in all cases. Results are given as means ± SEM. ***, P < 0.001.
GLU-OCN in insulin secretion. However, the notion that GLU-OCN is the form of OCN responsible for the endocrine functions fulfilled by this molecule has been questioned in several studies that failed to detect a correlation between GLU-OCN and insulin sensitivity (Shea et al., 2009; Lu et al., 2012; Mori et al., 2012) or reported a negative correlation between GLA-OCN and insulin sensitivity or obesity (Shea et al., 2009; Díaz-López et al., 2013). A possible explanation for these discrepant results could be that the association strength between undercarboxylated OCN and various parameters of energy metabolism varies from one population to another or that the methods of OCN measurement were not similar across these multiple studies.

To go beyond these experiments, we relied on cell-specific gene deletion approaches in the mouse. This effort started with the study of mice lacking γ-carboxylase in osteoblasts. Analysis of these mice provides direct evidence that increasing circulating levels of GLU-OCN improves glucose metabolism in mice fed an ND. These Ggcxfl/fl;Col1a1-Cre mice were also protected from the deleterious effect of an HFD on glucose metabolism. Ggcxfl/fl;OC-Cre mice had increased GLU-OCN serum levels, whereas GLA-OCN circulating levels were concurrently decreased. Importantly, the levels of GLA-OCN and of total OCN were not changed in the serum of Ggcxfl/fl;Collal-Cre mice, yet the two lines of mice exhibited similar improvements in glucose tolerance. These data add further support to the notion that GLA-OCN is not involved in the endocrine function of OCN in the mouse (Fig. 8). This is also supported by the fact that GLA-OCN does not cross the placenta or the blood–brain barrier (Oury et al., 2013b). That the deletion of one allele of Ocn in the background of Ggcxfl/fl;OC-Cre mice normalized their glucose tolerance provides direct genetic support to the notion that deletion of Ggcx in osteoblasts results in a gain-of-function of OCN, although it does not rule out in any way the possibility that additional γ-carboxylated proteins produced by osteoblasts could influence important physiological functions.

Even though the increase in serum GLU-OCN in Ggcxfl/fl;Collal-Cre mice was associated with improved glucose tolerance, better insulin sensitivity, and a reduction in fat mass, it did not overtly affect insulin secretion or β cell mass (unpublished data). We have previously shown that continuous infusions of low doses of GLU-OCN (0.3–3 ng/h) efficiently stimulate insulin secretion, whereas high doses of the same protein (10–30 ng/h) increase insulin sensitivity without a noticeable impact on insulin secretion or β cell proliferation (Ferron et al., 2008). Therefore, we speculate that the levels of GLU-OCN observed in Ggcxfl/fl;Collal-Cre mice (~40 ng/ml) were too high to affect insulin secretion, but efficiently affected insulin sensitivity. Along these lines, we also noticed that the improvement in glucose tolerance was more marked in Ggcxfl/fl;Collal-Cre than in Ggcxfl/fl;OC-Cre and Vkorclfl/fl;OC-Cre mice, despite a more modest increase in circulating GLU-OCN levels (i.e., 4 vs. 8–15 times). These results suggest that chronically high levels of circulating GLU-OCN may cause a desensitization of the OCN receptor. Although we do not have direct biochemical evidence supporting this hypothesis here, this possibility is consistent with the bell-shaped dose–response curve observed when target cells of OCN are treated with increasing concentration of this hormone (Ferron et al., 2008; Oury et al., 2011). It was also proposed to explain the absence of improvement in glucose tolerance in Tsc2 osteoblast-specific deficient mice, which displayed a 10-fold increase in undercarboxylated OCN (Riddle et al., 2014).

Regulation of OCN structure and function by γ-carboxylation

The structure of GLA-OCN was determined using x-ray crystallography more than 10 years ago (Hoang et al., 2003), whereas the one for GLU-OCN was solved just recently (Malashkevich et al., 2013). GLU and GLA-OCN structures bear many similarities. For instance, they both consist of three α helices surrounding a hydrophobic core and a disulfide bond between two of the helices. However, unlike what is the case for GLA-OCN, the GLU form of this protein does not bind Ca²⁺ and does not require elevated Ca²⁺ concentration to fold into a helical structure (Hauschka and Carr, 1982; Dowd et al., 2001; Malashkevich et al., 2013). These observations suggest that at Ca²⁺ concentrations around 1 mM (normally found in cell culture media or serum in vivo), only GLU-OCN will exhibit a helical conformation and presumably be able to bind to its receptor or receptors. This provides a structural explanation for the lack of biological activity of GLA-OCN in glucose metabolism.

The notion that GLA residues of OCN allow binding of this molecule to hydroxyapatite, the mineral component of bone ECM, has never been established in vivo. Here we show that mice lacking Ggcx or Vkorc1 in osteoblasts are characterized by a nearly complete absence of OCN in their bones. These results provide the first genetic in vivo evidence that γ-carboxylation of OCN is absolutely necessary for its accumulation in the bone ECM. Surprisingly, however, despite an absence of OCN in bone, the total serum levels of OCN were not increased in Ggcxfl/fl;OC-Cre or in Vkorclfl/fl;OC-Cre mice. One possible explanation for this apparent discrepancy could be that GLU-OCN has a shorter half-life than GLA-OCN in the serum or that other compensatory mechanisms exist to reduce circulating levels of GLU-OCN.

The importance of VKORC1 for OCN γ-carboxylation

Because it requires VKH₂ as a cofactor, γ-carboxylation is dependent on the reduction of VKO by a VKO reductase. Genetic and biochemical evidence supports the notion that VKORC1 is the main enzyme accomplishing this function in hepatocytes (Li et al., 2004; Chu et al., 2006). We demonstrate here that VKORC1 plays the same role in osteoblasts in vivo. Importantly, because the decrease in serum GLA-OCN was more pronounced in Ggcxfl/fl;OC-Cre than in Vkorclfl/fl;OC-Cre mice (80% vs. 40%), we hypothesized that this difference could be caused by the exogenous intake of VK in Vkorclfl/fl;OC-Cre mice, which would then be reduced to VKH₂ in osteoblasts. It is well known that rodents are generally resistant to VK deficiency because they can obtain sufficient VK through coprophagy (Groenen-Van Dooren et al., 1995). In support of this notion,
we found that ex vivo, Ggcx- and Vkorc1-null osteoblasts are equally ineffective at γ-carboxylating OCN in the absence of VK. In contrast, the addition of VK1 could rescue the defect in Vkorc1<sup>−/−</sup> osteoblasts only. Moreover, we provide evidence that preventing coprophagy even for a short period of 5 d reduced even further OCN γ-carboxylation in Vkorc1<sup>−/−</sup>;OC-Cre mice. In line with this result, injections of massive doses of VK1 correct the bleeding defect in mice or humans lacking a functional VKORC1 (Rost et al., 2004; Spohn et al., 2009) or rats treated with the VKORC1 inhibitor warfarin (Price et al., 1982). These observations suggest the existence of a warfarin-resistant “antidotal enzyme” reducing VK1 to VKH<sub>2</sub> in hepatocytes. Studies by Wallin and Martin (1985) and Ingram et al. (2013) provide evidence that a yet to be identified NAD(P)H-dependent VK reductase present in liver microsomes is involved in this antidotal response. Although our data do suggest that an antidotal VK reductase enzyme is also present in osteoblasts, we do not yet know the identity of this enzyme.

**VKORC1L1 is not implicated in γ-carboxylation in osteoblasts**

VKORC1L1 is a vertebrate parologue of VKORC1, which has been suspected to possess VKO and/or VK reductase activity (Westhofen et al., 2011; Tie et al., 2014), and therefore could compensate for the absence of VKORC1 in cells and some mouse tissues (Hammed et al., 2013; Tie et al., 2013). We tested in vivo whether VKORC1L1 exerts a redundant function with VKORC1 by generating global and cell-specific deletion of these genes and by measuring the γ-carboxylation of OCN. Using mice lacking either VKORC1 or VKORC1L1 in osteoblasts, we found that only VKORC1 is required for the γ-carboxylation of OCN in these cells. Moreover, the combined
deletion of Vkorc1 and Vkorc1l1 did not further increase GLU-OCN in vivo or in cell culture. Finally, that VKORC1 and GGCX but not VKORC1L1 localized in the ER in osteoblasts further indicates that, at least in osteoblasts, VKORC1 and VKORC1L1 have different cellular functions.

Our data also show that in hepatocytes and in osteoblasts, VKORC1L1 is not involved in γ-carboxylation and is not able to compensate for the absence of VKORC1. The notion that VKORC1 and VKORC1L1 may have different cellular and physiological functions is further supported by a recent study showing that, although VKORC1 is a three-transmembrane domain (TMD) membrane protein with its N terminus in the ER lumen and its C terminus in the cytoplasm, VKORC1L1 is a four-TMD membrane protein with both termini located in the cytoplasm (Tie et al., 2014). Moreover, it was shown recently that an Arg98Trrp mutation in VKORC1 that causes VK-dependent clotting factor type 2 deficiency disrupts a di-arginine-based ER retention motif responsible for ~80% of the ER localization of this protein (Czogalla et al., 2014). Interestingly, alignment of VKORC1 and VKORC1L1 failed to identify this di-arginine motif in VKORC1L1 (unpublished data), thus supporting the notion that these two proteins may have different subcellular localizations. Importantly, our results do not exclude in any way the possibility that VKORC1L1 may possess VKO or VK1 reductase activity or that in other yet-undefined cell types VKORC1L1 can support γ-carboxylation. It is also possible that in osteoblasts VKORC1L1 acts as a VK-binding protein that would not, however, be involved in γ-carboxylation of OCN. Further investigations will address these questions.

In summary, this work provides in vivo and direct evidence that γ-carboxylation of OCN by GGCX is a negative regulator of its endocrine function (Fig. 8). Moreover, using genetic approaches, we delineated the importance of VKORC1 in the process of γ-carboxylation in osteoblasts and suggest that its paralogue, VKORC1L1, may have a distinct physiological function.

Materials and methods

Mouse models

Ocn−/− mice were generated by simultaneously deleting Ocn1 and Ocn2 genes in the mouse OCN cluster and by replacing them with a Neomycin resistance cassette by homologous recombination (Ducy et al., 1996). The Col1a1-Cre transgenic mice express the Cre recombinase under the control of the 2.3-kb proximal fragment of the α1(I)-collagen promoter, which is expressed at a high level in osteoblasts (Daquin et al., 2002). The OCC-Cre transgenic mice express Cre recombinase under the control of a fragment of the human OCN promoter (Zhang et al., 2002). To obtain mice harboring conditional alleles of Ggcx, Vkorc1, and Vkorc1l1, we constructed three DNA targeting vectors to allow the introduction by homologous recombination of two loxP sites in the Ggcx, Vkorc1, and Vkorc1l1 locus, respectively (Figs. S1 A, S2 A, and S4 A). A Neomycin resistance cassette (NEO) flanked by two loxP sites was also included in each construct to allow positive selection, whereas a thymidine kinase selection cassette served as a negative selection for nonspecific recombination events in the Ggcx and the Vkorc1 constructs. These vectors were individually electroporated into embryonic stem cells that were subsequently selected with G418 and ganciclovir, and correctly targeted clones were identified by Southern blot analysis (Figs. S1 B, S2 B, and S4 B). Positive clones were injected in blastocysts and implanted in pseudopregnant females to generate chimeric mice. The chimeras were bred with WT mice, and germline transmission of the targeted allele was confirmed by Southern blot hybridization. These mice were next crossed with Eta-Cre mice, which allow partial recombination of loxP sequence in early mouse embryo (Xu et al., 2001), thus resulting in the excision of the NEO cassette and the generation of the floxed (fl) alleles. PCR-based genotyping confirmed that we have obtained Ggcxfl/fl, Vkorc1fl/fl, and Vkorc1l1fl/fl mice (Figs. S1 C, S2 C, and S4 C). In a separated cross, the Vkorc1−/− and Vkorc1l1−/− mice were bred with CMV-Cre (Schwenk et al., 1995) to inactivate Vkorc1 and Vkorc1l1 in the germ cells and obtain mice carrying a null allele of these genes, i.e., Vkorc1−/− and Vkorc1l1−/− (Figs. S2 D and S4 F). Male mice maintained on ND were used in all experiments, except where indicated. Tissue DNA was extracted by standard procedure, and PCR was performed with the indicated genotyping PCR primers (Table S1). All of the animal protocols were reviewed and approved by an Institutional Animal Care and Use Committee.

Metabolic analysis

In adult mice, GTTs were performed after overnight fasting. A dose of glucose (1.3 or 2 g/kg, depending on the experiment) was administered i.p., and blood glucose levels were measured at the indicated time points.ITT was performed after 4 h of fasting: 0.75 U/kg insulin (Humulin; Lilly) was injected i.p., and blood glucose was measured at the indicated time points. In some experiments, mice were fed an HFD starting at 4 wk of age (Research Diets D12331). HOMA-IR was calculated using the following formula (Turner et al., 1979):

\[
\text{HOMA-IR} = \left( \frac{G_{0}}{l_{i}} \right) \times \frac{100}{\text{mU/mL}},
\]

where \(G_{0}\) is fasting blood glucose in mg/dl and \(l_{i}\) is fasting insulin in mU/ml.

The energy expenditure measurements were obtained using a six-chamber Oxymax system (Columbus Instruments). After 12-h acclimation to the apparatus, data for 24-h measurement were collected and analyzed as recommended by the manufacturer. Oxygen consumption was measured by calculating the difference between input oxygen flow and output oxygen flow. Similarly, carbon dioxide production was measured by calculating the difference between output and input carbon dioxide flows. Heat production was calculated by indirect calorimetry using the following formulas:

\[
\text{Heat} = CV \times V_{O_{2}} / BW, \\
CV = 3.815 \times 1.232 \times \text{RER}, \\
\text{RER} = \frac{V_{CO_{2}}}{V_{O_{2}}},
\]

Bone histomorphometry and μCT

For bone histomorphometry analysis, lumbar vertebrae were dissected, fixed in 10% formalin, dehydrated in graded ethanol series, and embedded in methyl methacrylate resin (Chappard et al., 1987). Van Kossa/Van Gieson staining was performed using 7-μm sections for bone volume over tissue volume (BV/TV) measurement. Calcine (Sigma-Aldrich) was dissolved in buffer [0.15 M NaCl and 2% NaHCO3] and injected twice at 0.125 mg/g body weight on days 1 and 4, and mice were sacrificed on day 6. 5-μm bone sections were cleared in xylene and used for bone formation rate (BFR) and mineralization apposition rate (MAR) measurements. For the analysis of osteoblast and osteoclast, sections were stained with toluidine blue and tartrate-resistant acid phosphatase (TRAP), respectively. Histomorphometric analyses were performed using the OsteoMeasure Analysis System [OsteoMetrics]. Microcomputed tomography (μCT) of fixed femur was performed at the Montreal Center for Bone and Periodontal Research on a SkyScan 1072 apparatus.

Cell culture

Calvaria osteoblasts were isolated from 3-d-old mice and cultured as previously described (Ducy and Karsenty, 1995). In brief, dissected calvarias were washed once with 1x PBS and digested at 37°C twice for 10 min and twice for 30 min in αMEM containing 0.1 mg/ml collagenase type 2 (Worthington Biochemical Corporation) and 0.25% trypsin. Cells of the first two digestions were discarded, whereas cells released from the third and fourth digestions were plated in αMEM supplemented with 10% FBS. Controls and Ggcx−/−, Vkorc1−/−, and Vkorc1l1−/− osteoblasts were generated ex vivo by infecting Ggcx−/−, Vkorc1−/−, and Vkorc1l1−/− with either GFP- or Cre-expressing adenoviruses (University of Iowa) at an MOI of 200. Thereafter, to induce osteoblast differentiation, αMEM was supplemented with 5 mM β-glycerophosphate and 100 μg/ml ascorbic acid and replaced every 2 d for 21 d. Von Kossa staining was used to assess mineralization of osteoblast culture. As indicated in some experiments, osteoblasts were derived in

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culture from bone marrow according to standard protocols (Owen and Pan, 2008). In this case, long bones (tibia and femur) from 2–4-wk-old animals were dissected, and bone marrows were flushed out using 1× PBS. Cells were seeded at 8 million/well in 6-well plates in αMEM supplemented with 10% FBS. Medium was changed 3 d later. After 7 d of culture and every third day thereafter, medium was changed for the differentiation medium: αMEM supplemented with 10% FBS, 100 µg/ml L-ascorbic acid, 5 mM β-glycerol phosphate, and 10 nM dexamethasone. After 2 wk of culture in the differentiation medium, mineralization nodules were observed. Cells were lysed, and the extracts were analyzed by Western blotting or by QPCR. To the differentiation medium, mineralization nodules were observed. Cells were seeded at 8 million/well in 6-well plates in osteoblast differentiation medium for 4 d. Cells were fixed in 4% formalin for 15 min. A second aliquot of 30% hydrogen peroxide (1:50 volume) was added to the reaction mixture, and the solution was incubated for an additional 45 min. The reaction was next diluted with distilled water and VKO extracted with hexane and concentrated under a stream of nitrogen. Bone extracts were prepared by homogenizing bone marrow–flushed tibia and femur in lysis buffer (20 mM Tris-HCl, pH 7.4, 150 mM NaCl, 1 mM EDTA, 1 mM EGTA, 1% Triton, 1 mM PMSF, and 1× protease inhibitors cocktail). To detect OCN, VKORC1 and VKORC1L1 proteins were resolved on 10% acrylamide Tris-Tricine gels, whereas 7.5% Tris-Glycine gels, were used for GGCX. Western blot analyses were performed according to standard protocols. Antibodies used were anti-GGCX (rabbit, 16209-1-AP, Proteintech), VKORC1L1 rabbit, ab116508; Abcam), β-actin (mouse, A5441; Sigma-Aldrich), KDEL (mouse clone 10C3, ADISA-827; Enzo Life Sciences), and OCN C terminus polyclonal goat antibody, which recognized an 26–46 corresponding to mature mouse OCN (Ferron et al., 2010b). Rabbit anti-VKORC1 antisera were obtained by immunizing rabbits with a GST fusion protein containing 30–80 corresponding to mouse VKORC1 protein. Anti-VKORC1 antibodies were next affinity purified using a peptide corresponding to VKORC1 aa 30–80. Antiserum and media levels of carboxylated, undercarboxylated, and total OCN were measured using EUSA assays as described previously (Ferron et al., 2010b). In brief, the GLA EUSA detects OCN only when it is carboxylated on the residue GLU13. The GLU EUSA quantifies the uncarboxylated form of OCN. The total EUSA detects OCN whether it is carboxylated or not.

Gene expression

Cells were collected, and RNA isolation, cDNA preparation, and real-time PCR analysis were performed according to standard protocols. In brief, total RNA was extracted (Chomczynski and Sacchi, 2006), DNase I treated, and media levels of carboxylated, undercarboxylated, and total OCN were measured using EUSA assays as described previously (Ferron et al., 2010b). In brief, the GLA EUSA detects OCN only when it is carboxylated on the residue GLU13. The GLU EUSA quantifies the uncarboxylated form of OCN. The total EUSA detects OCN whether it is carboxylated or not.

Microscopy

Mouse calvaria osteoblasts were plated on glass coverslips and cultured in osteoblast differentiation medium for 4 d. Cells were fixed in 4% formalin for 15 min, and immunofluorescence was performed according to a standard protocol (Cell Signaling Technology). The coverslips were next hybridized with primary rabbit [GGCX, VKORC1, or VKORC1L1; 1:50] and mouse antibody (KDEL; 1:100) overnight at 4°C. After three washes with 1× PBS, the coverslips were incubated with secondary anti-mouse Alexa Fluor 594 (1:1,000) and anti-rabbit Alexa Fluor 488 (1:1,000) for 1 h. Coverslips were then washed again, briefly washed with Hoechst solution, and mounted on slides with FluorSave reagent (EMD Millipore). Cells were imaged at RT on a confocal microscope (LSM 710; Carl Zeiss) using a Plan-Apochromat 63× objective (NA = 1.4) with oil Immersol 158F (Carl Zeiss) and the Zen 2009 software [Carl Zeiss]. The images were analyzed using Velocity software (PerkinElmer). Osteoblast cultures were imaged using a DeltaVision dissecting microscope (Discovery V12; Carl Zeiss) using a 0.63× objective at RT, a 0.8x motorized zoom, an AxioCam ERc5s camera, and the Zen 2012 software (Carl Zeiss). Histological sections of Von Kossa/Van Gieson staining were viewed under a light microscope (DM4000B LED; Leica) using a 2.5× objective (NA = 0.07) at RT, a DP72 camera (Olympus), and the CellSens Entry software (Olympus).

Results

Statistics

Results are given as means ± SEM. Statistical analyses were performed using unpaired, two-tailed Student’s t test for single time point measurements, F(1, 17) ANOVA for repeated measurements [MTT and FIT], two-way ANOVA tests for repeated measured followed by Bonferroni post tests were performed. In all figures, * P < 0.05; ** P < 0.01; *** P < 0.001. All experiments were repeated at least three times or performed on at least three independent animals.

Online supplemental material

Figs. S1 provides the targeting strategy used to generate Ggcxfl/fl mice, measured levels of GLA, and total OCN in Ggcxfl/fl osteoblast culture, recombination efficiency in the different mutant strains, data on testicular function in Ggcxfl/fl, Col1a1Cre and Ggcxfl/fl osteoblast mice, and fasting serum levels and HOMA-IR values for the Ggcxfl/fl, Col1a1Cre mice. Fig. S2 provides µCT and histological analyses of bones from Ggcxfl/fl, OCN-Cre mice and in vivo characterization of Ggcxfl/fl osteoblast function. Fig. S3 provides the targeting strategy used to generate Vkorclfl/fl mice, generation and characterization of Vkorclfl/fl mice, serum levels of OCN in Vkorclfl/fl pups, serum levels of GLA-OCN in Vkorclfl/fl, OCN-Cre mice in the absence of coprophagy, secreted levels of GLA and total OCN in Vkorcl−/− osteoblast culture, and mineralizing capacity of Vkorcl−/− osteoblasts. Fig. S4 provides the targeting strategy used to generate Vkorcl110/fl mice, generation and characterization of Vkorcl110/fl mice, expression levels of Vkorcl110 in Vkorcl110/fl, OCN-Cre and in Vkorcl110−/− osteoblasts, serum levels of OCN in Vkorcl110−/− mice, and secreted levels of GLA and total OCN in Vkorcl110−/− and in Vkorcl1−/− Vkorcl110−/− osteoblast cultures. Table S1 includes the sequence of the primers used in this study. Online supplemental material is available at http://www.jcb.org/cgi/content/full/jcb.201409111/DC1.

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References

Abseyi, N., Z. Sklkar, M. Berberoglu, B. Hachamidouglu, S. Savas Erdleve, and G. Gççl. 2012. Relationships between osteocalcin, glucose metabolism, and adiponectin in obese children: Is there crosstalk between bone tissue and glucose metabolism? J. Clin. Res. Pediatr. Endocrinol. 4:182–188. http://dx.doi.org/10.4274/prc.2012.271.831

Berkner, K.L., M. Harbeck, S. Lingenfelter, C. Bailey, C.M. Sanders-Hinck, and J.W. Suttie. 1992. Purification and identification of bovine liver gamma-carboxylase. Proc. Natl. Acad. Sci. USA. 89:6242–6246. http://dx.doi.org/10.1073/pnas.89.14.6242

Brennan-Speranza, T.C., H. Hennecke, S.J. Gasparini, K.I. Blankenstein, U. Hovhannessian, V.C. Oates, V. Svistounov, Y. Zhang, G.J. Cooney, F. Buttgeriet, et al. 2012. Osteoblasts mediate the adverse effects of glucocorticoids on fuel metabolism. J. Clin. Invest. 122:4172–4189. http://dx.doi.org/10.1172/JCI63377

Bullí, M., J.M. Moreno-Navarrete, J.M. Fernández-Real, and J. Salas-Salvadó. 2013. Total and undercarboxylated osteocalcin predict changes in insulin sensitivity and β cell function in elderly men at high cardiovascular risk. Am. J. Clin. Nutr. 95:249–255. http://dx.doi.org/10.3945/ajcn.111.016642

Chappard, D., P. Palle, C. Alexandre, L. Vico, and G. Riffat. 1987. Bone embedding in pure methyl methacrylate at low temperature preserves enzyme activities. Acta Histochem. 81:183–190. http://dx.doi.org/10.1016/S0001-2493(87)80012-9

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of male fertility by the skeleton. Cell. 144:796–809. http://dx.doi.org/10.
1016/j.cell.2011.02.004

Oury, F., M. Ferron, W. Haizhen, C. Conafreux, L. Xu, J. Lacombe, P. Srinivas,
A. Chamouni, F. Lugani, H. Lejeune, et al. 2013a. Osteocalcin regulates
murine and human fertility through a pancreas–bone-testis axis. J. Clin.
Invest. 123:2421–2433. http://dx.doi.org/10.1172/JCI126952

Oury, F., L. Khrimian, C.A. Denny, A. Gardin, A. Chamouni, N. Goeden, Y.Y.
Huang, H. Lee, P. Srinivas, X.B. Gao, et al. 2013b. Maternal and off-
spring pools of osteocalcin influence brain development and functions.
Cell. 155:228–241. http://dx.doi.org/10.1016/j.cell.2013.08.042

Owen, T.A., and L.C. Pan. 2008. Isolation and culture of rodent osteoprogeni-
tor cells. Methods Mol. Biol. 455:3–18. http://dx.doi.org/10.1007/978-
1-59745-104-8_1

Pi, M., L. Chen, M.Z. Huang, W. Zhu, B. Ringhofer, J. Luo, L. Christenson,
M. Pi, Y. Wu, and L.D. Quarles. 2011. GPRC6A mediates responses to os-
tin K epoxide reductase complex subunit 1-like 1 (VKORC1L1) are
involved in its active site regeneration.

Tishler, M., L.F. Fieser, and N.L. Wendler. 1940. Hydro, oxido and other deriva-
tives of vitamin K1 and related compounds. J. Am. Chem. Soc. 62:2866–
2871. http://dx.doi.org/10.1021/ja11867a066

Turner, R.C., R.R. Holman, D. Matthews, T.D. Hockaday, and J. Peto. 1979.
Insulin deficiency and insulin resistance interaction in diabetes: estima-
tion of their relative contribution by feedback analysis from basal plasma
insulin and glucose concentrations. Metabolism. 28:1086–1096. http://
dx.doi.org/10.1016/0026-0495(79)90146-X

Wallin, R., and L.F. Martin. 1985. Vitamin K-dependent carboxylase and vita-
mK metabolism in liver. Effects of warfarin. J. Clin. Invest. 76:1879–
1884. http://dx.doi.org/10.1172/JCI112182

Wang, Q., B. Zhang, Y. Xu, H. Xu, and N. Zhang. 2013. The relationship be-
tween serum osteocalcin concentration and glucose metabolism in pa-
tients with type 2 diabetes mellitus. Int. J. Endocrinol. 2013:842598.
http://dx.doi.org/10.1155/2013/842598

Wei, J., M. Ferron, C.J. Clarke, Y.A. Hamun, H. Jiang, W.S. Blaner, and G.
Karsenty. 2014a. Bone-specific insulin resistance disrupts whole-body
glucose homeostasis via decreased osteocalcin activation. J. Clin. Invest.
124:1–13.

Wei, J., T. Hanna, N. Suda, G. Karsenty, and P. Ducy. 2014b. Osteocalcin pro-
motes β-cell proliferation during development and adulthood through
Gperca. Diabetes. 63:1021–1031. http://dx.doi.org/10.2337/db13-0887

Westhoffen, P., M. Watzka, M. Marovina, M. Hass, G. Kirlf, J. Müller, C.G.
Bevans, C.R. Müller, and J. Oldenburg. 2011. Human vitamin K 2,3-
epoxide reductase complex subunit 1-like 1 (VKORC1L1) mediates vita-
mK-dependent intracellular antioxidant function. J. Biol. Chem. 286:
15085–15094. http://dx.doi.org/10.1074/jbc.M110.210971

Xu, X., C.L. Li, L. Garrett-Beal, D. Larson, A. Wynshaw-Boris, and C.X. Deng.
2001. Direct removal in the mouse of a floxed neo gene from a three-loxP
conditional knockout allele by two novel approaches. Genesis. 30:1–6.
http://dx.doi.org/10.1002/genesis.1025

Yoshikawa, Y., A. Kode, L. Xu, I. Mosialou, B.C. Silva, M. Ferron, T.L. Clemens,
A.N. Economides, and S. Kousteni. 2011. Genetic evidence points to an
osteocalcin-independent influence of osteoblasts on energy metabolism.
J. Bone Miner. Res. 26:2012–2025. http://dx.doi.org/10.1002/jbmr.417

Zhang, M., S. Xuan, M.L. Bouxsein, D. von Stechow, N. Akeno, M.C. Faugere,
H. Malluche, G. Zhao, C.J. Rosen, A. Ellstratadis, and T.L. Clemens.
2002. Osteoblast-specific knockout of the insulin-like growth factor
(IGF) receptor gene reveals an essential role of IGF signaling in bone
matrix mineralization. J. Biol. Chem. 277:44005–44012. http://dx.doi.
org/10.1074/jbc.M208265200

Zhou, B., H. Li, L. Xu, W. Zang, S. Wu, and H. Sun. 2013. Osteocalcin reverses
endoplasmic reticulum stress and improves impaired insulin sensitivity
secondary to diet-induced obesity through nuclear factor-κB signaling
pathway. Endocrinology. 154:1055–1068. http://dx.doi.org/10.1210/
end.2012-2144
