CGRP Regulates the Age-Related Switch Between Osteoblast and Adipocyte Differentiation

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Osteoporosis is a chronic age-related disease. During aging, bone marrow-derived mesenchymal stem cells (BMSCs) display increased adipogenic, along with decreased osteogenic, differentiation capacity. The aim of the present study was to investigate the effect of calcitonin gene-related peptide (CGRP) on the osteogenic and adipogenic differentiation potential of BMSC-derived osteoblasts. Here, we found that the level of CGRP was markedly lower in bone marrow supernatant from aged mice compared with that in young mice.

In vitro experiments indicated that CGRP promoted the osteogenic differentiation of BMSCs while inhibiting their adipogenic differentiation. Compared with vehicle-treated controls, aged mice treated with CGRP showed a substantial promotion of bone formation and a reduction in fat accumulation in the bone marrow. Similarly, we found that CGRP could significantly enhance bone formation in ovariectomized (OVX) mice in vivo. Together, our results suggested that CGRP may be a key regulator of the age-related switch between osteogenesis and adipogenesis in BMSCs and may represent a potential therapeutic strategy for the treatment of age-related bone loss.

Keywords: CGPR, osteoporosis, BMSCs, osteogenic, adipogenic, age-related

INTRODUCTION

Osteoporosis is a chronic disease caused by the breakdown of bone homeostasis, with elderly and postmenopausal women being the populations most at risk of developing this condition (NIH Consensus Development Panel on Osteoporosis Prevention, Diagnosis, and Therapy, 2001; Rachner et al., 2011). Age-related bone loss and osteoporosis have been associated with reduced numbers of osteoblasts and increased numbers of adipocytes (Idris et al., 2009; Yu and Wang, 2016). It is known that bone marrow mesenchymal stem cells (BMSCs) have the potential to differentiate into osteoblasts, adipocytes, and osteoclasts, thereby playing an important role in bone formation (Li et al., 2016; Chen et al., 2018; Peng et al., 2019). Overall, bone homeostasis depends on the balance between the osteogenic and adipogenic differentiation of BMSCs (Li et al., 2015;
LI et al., 2018). Age-related osteoporosis results from reduced osteogenic differentiation and increased adipogenic differentiation of BMSCs in elderly patients (Shen et al., 2012; Childs et al., 2015; Li et al., 2017).

Calcitonin gene-related peptide (CGRP), a member of the calcitonin protein family, is a 37-amino acid peptide generated through the alternative splicing of primary transcripts of the calcitonin gene (Amara et al., 1982; Rosenfeld et al., 1983; Naot and Cornish, 2008). In humans and mice, CGRP exists as two isoforms—α-CGRP and β-CGRP—which are encoded by the CALCA and CALCB genes, respectively. In mice, α-CGRP is highly expressed in the central nervous system and peripheral nervous system, while β-CGRP is expressed in the enteric nervous system (Sternini, 1992; Russell et al., 2014). In addition, in mice, the Calca gene has been shown to affect bone remodeling; however, the underlying mechanism remains unclear (Schinke et al., 2004).

CGRP is a neuropeptide that is released from sensory nerve endings and can also be found in bone cells and endothelial cells (Russell et al., 2014). There is evidence that CGRP is involved in the regulation of cell proliferation and differentiation and may also be important for connecting the systems involved in bone metabolism (Bjurholm et al., 1988; Chattergoon et al., 2005; Thievent et al., 2005; Zhang et al., 2016; Xu et al., 2019). Additionally, studies have confirmed that CGRP can promote osteogenesis and inhibit osteoclast formation (Imai et al., 1997; Villa et al., 2003; Yoo et al., 2014). However, few studies have investigated the roles of CGRP in the age-related switch between osteoblast and adipocyte differentiation in bone marrow. Consequently, the aim of this study was to investigate whether CGRP is associated with increased bone formation and inhibition of adipocyte accumulation in age-related bone loss.

In the current study, we show that the level of CGRP in mouse bone marrow-derived supernatant decreases with aging. Furthermore, we found that CGRP not only promotes the osteogenic differentiation of BMSCs but also inhibits their adipogenic differentiation and senescence. Importantly, we demonstrate that exogenous application of CGRP can accelerate bone formation in aged and OVX mice in vivo, implying that CGRP may be a potential therapeutic target for the prevention of osteoporosis.

### MATERIALS AND METHODS

#### Mice

C57BL/6J mice were purchased from Hunan Slaccas Jingda (Changsha, China). Two-month-old female C57BL/6J mice underwent bilateral ovarian resection to establish the OVX model. For *in vitro* CGRP treatment experiments, 12-month-old OVX mice and sham-operated mice were injected with CGRP (10 mg/kg) via the tail vein three times weekly for 1 month. Mice treated with 1 × PBS were used as controls. All mice were maintained in the specific-pathogen-free facility of the Laboratory Animal Research Center of Central South University. The mice were kept under a 12 h/12 h light: dark cycle.
Briefly, the femora and tibiae of four 7-day-old mice were cut and digested with Liberase DL (26 U/mL) (Roche) in a water bath at 37°C and digested with 0.25% trypsin and diluted with osteogenic induction medium containing 10% FBS, 1% penicillin/streptomycin, 0.1 mM dexamethasone, and 5 µg/mL insulin.

### Isolation and Culture of BMSCs

BMSCs were isolated as previously described (Picke et al., 2018). Bone marrow was separated, centrifuged at 1,000 rpm for 10 min, and then the supernatant was transferred to a new 10-cm culture dish. The isolated BMSCs were plated in 6-well plates at 2.5 × 10⁶ cells/well and cultured in α-MEM containing 10% FBS, 1% penicillin/streptomycin, and 0.5 mM 3-isobutyl-1-methylxanthine, 1 mM dexamethasone, and 5 µg/mL insulin. The adipogenic induction medium was renewed every 3 days for 14 days. The cells were then fixed in 4% paraformaldehyde and stained with Oil Red solution (Sigma-Aldrich). The dye was solubilized in isopropanol and the absorption at 540 nm was measured using a BioTek Epoch microplate spectrophotometer (BioTek Instruments).

### Osteogenic Differentiation and Mineralization Assay

Isolated BMSCs were cultured in a 10-cm culture dish. The medium was replaced containing 1% penicillin/streptomycin and 10% FBS, and then cultured in a 10-cm culture dish. The medium was replaced every other day and CD11b⁺ cells were cleared using anti-CD11b antibody-coated magnetic beads.

### Enzyme-Linked Immunosorbent Assay (ELISA)

Bone marrow was separated, centrifuged at 1,000 rpm for 10 min, and then the supernatant was transferred to a new sterile centrifuge tube. ELISA was performed using a CGPR (rat, mouse) EIA-Kit (K-01509, Phoenix Pharmaceuticals) according to the manufacturer’s instructions.

## TABLE 1 | Continued

| Name     | Sequences (5'-3') | Amplicon size (bp) |
|----------|------------------|--------------------|
| Hmg2-R   | TGGGCCCTTTTCTCCAAATG  | 233                |
| Ptgds-2F | TGTGAGCTTGACCCGGACTG  | 231                |
| Ptgds-2R | TGCCACATTGGAATGATGTC  | 106                |
| Vgf-F    | AGGGTGGACGCGTGACCAAG  | 114                |
| Vgf-R    | TGCCCTCAACAGTACCGAG  | 101                |
| Per2-F   | GAAAGTGTCACACCACATGAA | 86                |
| Apmap-R  | CCTTGCCATTCCCCTACTTGCGA | 109            |
| Dgat1-R  | TGAGCCCGGGTGGTAGTG  | 199                |
| Prkab2-F | AAGGTGAAGAGCATCATAACCCT  | 114            |
| Dgat1-F  | TCCGTCCAGGGTGGTAGTG  | 199                |
| Prkab2-R | CCATTCGACACTTCTTTTTT  | 103                |
| Dgat1-F  | TCCGTCCAGGGTGGTAGTG  | 199                |
| Hdac6-F  | ACGCCTGATGACACTTCTTT  | 103                |
| Prkab2-R | CCATTCGACACTTCTTTTTT  | 103                |
| Dgat1-F  | TCCGTCCAGGGTGGTAGTG  | 199                |
| Hdac6-R  | CACACACACTTCTTCACCAC | 109                |
| Egr2-F   | GGAAGGCGCGTAGACAAATAC | 154            |
| Egr2-R   | CCACTGGCTTCATGCTGCA  | 116                |
| Prkab2-R | CCCGCTTGCTCCGACTTGCA  | 116                |
| Apmap-R  | ACTTGATTTGAGTACGAAACA | 116                |
| FABP4-F  | AAGGTGAAGAGCATCATAACCCT  | 133            |
| FABP4-R  | TCAGCGCTTCCATACGAGCTCC | 133            |
| PPARγ-F  | CGCGCTGATGACACTTCTTT  | 103                |
| PPARγ-R  | CCATTCGACACTTCTTTTTT  | 103                |
| β-actin-F | GGCGTGTATCTCCGCTCATCG  | 154                |
| β-actin-R | CACAGTTGTAACAAATGCGATGT  | 154                |

β-glycerol phosphate, and 50 mM ascorbate. The osteogenic induction medium was renewed every 3 days for 21 days. The cells were subsequently fixed in 4% paraformaldehyde and stained with 2% Alizarin Red S (Sigma-Aldrich). The Alizarin Red S was dissolved in cetylpyridinium chloride solution and quantified by spectrophotometry at 540 nm.

### Adipogenic Differentiation Assay

Isolated BMSCs were plated in 6-well plates at 2.5 × 10⁶ cells/well in adipogenic differentiation medium containing 10% FBS, 1% penicillin/streptomycin, 0.5 mM 3-isobutyl-1-methylxanthine, 1 mM dexamethasone, and 5 µg/mL insulin. The adipogenic induction medium was renewed every 3 days for 14 days. The cells were then fixed in 4% paraformaldehyde and stained with Oil Red solution (Sigma-Aldrich). The dye was solubilized in isopropanol and the absorption at 540 nm was measured using a BioTek Epoch microplate spectrophotometer (BioTek Instruments).

### Quantitative Real-Time PCR Analysis

Total RNA (1 µg) isolated from CGRP-treated BMSCs was treated with gDNA Eraser to remove residual genomic DNA and then reverse-transcribed into cDNA using the PrimeScript RT reagent Kit (Takara, Japan). Real-time PCR analysis was performed with SYBR Green (Takara, Japan) in an Applied Biosystems QuantStudio 3 Real-Time PCR System (Applied Biosystems). The amplification reactions were performed in a 96-well plate and consisted of 40 cycles of 95°C for 5 s and 60°C for 30 s. The relative transcript levels of target genes were quantified using the 2−△△Ct method with beta-actin serving as the internal control. The primer sequences are listed in Table 1.

### Micro-CT Analysis

Micro-computed tomography (micro-CT) analysis was performed as previously reported (Li et al., 2015; Yang et al., 2019). The femora of mice from both the CGRP treatment group and the control group were isolated, fixed in 4% paraformaldehyde for 24 h, and then scanned by X-ray microtomography (Skyscan 1172, Bruker) at a pixel size of 13.98 µm. For the distal femur, the region-of-interest (ROI) was defined from 0.215 to 1.72 mm below the growth plate. The bone volume as a fraction of total bone volume (BV/TV), trabecular thickness (Tb. Th), trabecular number (Tb. N), and trabecular separation (Tb. Sp) were measured.

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**Isolation and Culture of BMSCs**

BMSCs were isolated as previously described (Picke et al., 2018). Briefly, the femora and tibiae of four 7-day-old mice were cut and digested with Liberase DL (26 U/mL) (Roche) in a water bath at 37°C for 2 h. The cell suspension was centrifuged at 1,000 rpm for 5 min at 4°C, resuspended in 1 mL of α-MEM containing 1% penicillin/streptomycin and 10% FBS, and then cultured in a 10-cm culture dish. The medium was replaced every other day and CD11b⁺ cells were cleared using anti-CD11b antibody-coated magnetic beads.

**Osteogenic Differentiation and Mineralization Assay**

Isolated BMSCs were digested with 0.25% trypsin and diluted to 1 × 10⁷ cells/mL. Then, 200 µL of the cell suspension was plated in 6-well plates at a density of 2 × 10⁶ cells/well. At 80% confluence, the medium was replaced with osteogenic induction medium containing 10% FBS, 1% penicillin/streptomycin, 0.1 mM dexamethasone, 10 mM β-glycerol phosphate, and 50 mM ascorbate. The osteogenic induction medium was renewed every 3 days for 21 days. The cells were subsequently fixed in 4% paraformaldehyde and stained with 2% Alizarin Red S (Sigma-Aldrich). The Alizarin Red S was dissolved in cetylpyridinium chloride solution and quantified by spectrophotometry at 540 nm.
CGRP Regulates Osteoblast and Adipocyte

Immunohistochemistry and Tartrate-Resistant Acid Phosphatase (TRAP) Staining

The femora were decalcified with 0.5 M EDTA for 1–2 weeks and then embedded in paraffin. The paraffin-embedded femora were sectioned (4 µm) using a RM2135 rotary microtome (Leica Geosystems). Sections were roasted at 60°C in an oven for 2 h, dewaxed with dimethyl benzene, dehydrated with alcohol, and treated with an antigen retrieval solution. The sections were subsequently washed with 1 × TTBS, blocked with 5% goat serum, and incubated with an anti-osteocalcin primary antibody (diluted 1:500; Cat# M041, Takara) overnight at 4°C. The next day, the sections were washed with 1 × TTBS, incubated with biotinylated secondary antibody (anti-mouse,1:200; Cat# PV9000, Beijing Zhongshan Jinqiao Biotechnology Co. Ltd.) for 1 h, washed with 1 × TTBS, and counterstained with hematoxylin (Sigma–Aldrich) for immunohistochemical analysis. An inverted fluorescence microscope was used for imaging. For TRAP staining, an Osteoclast Staining Kit (Sigma–Aldrich) was used according to the manufacturer's instructions. The number of osteoblasts on the bone surface and osteoclast number per bone perimeter were measured in the femora.

β-Galactosidase Staining

BMSCs were washed with 1 × PBS and fixed in 4% formaldehyde for 30 min. Cell senescence was assessed using a β-Galactosidase Staining Kit (Cell Signaling Technology, 9860) according to the manufacturer's instructions. The percentage of senescent cells was determined using ImageJ software.

Statistical Analysis

Data were analyzed by unpaired, two-tailed, Student's t-tests or one-way or two-way analysis of variance (ANOVA) followed by Bonferroni’s post-test using GraphPad Prism 7.0 software. All data are presented as means ± SEM. A p-value < 0.05 was considered statistically significant.

RESULTS

CGRP Levels Were Decreased in the Bone Marrow Supernatant of Mice During Aging

CGRP is a 37-residue neuropeptide primarily expressed in the central and peripheral nervous systems (Emeson et al., 1989, 1992). That CGRP plays a vital role in bone metabolism is supported by evidence showing that the lack of CGRP results in reduced bone formation and impaired bone regeneration in mice (Schinke et al., 2004; Appelt et al., 2020). To assess the expression of Calca (encoding α-CGRP) in bone tissue during aging, we conducted RT-qPCR analysis on total RNA extracted from the bone tissue of mice at 3, 6, 9, 12, 15, 18, 21, and 24 months of age (n = 5 per age group). We found that the Calca level was significantly decreased during aging (Figure 1A). CGRP is thought to be a secreted neuropeptide that is released from sensory nerve endings (Tsujikawa et al., 2007). To assess the CGRP levels in bone marrow, we collected bone marrow supernatant from male C57BL/6J mice aged 3, 12, and 24 months (n = 5 per age group) and measured CGRP levels by ELISA. As shown in Figure 1B, CGRP levels were lower in aged mice than in young mice. Given the vital role of CGRP in bone metabolism (Irie et al., 2002; Schinke et al., 2004; Appelt et al., 2020) and that CGRP has been reported to stimulate the proliferation and osteogenic differentiation of rat-derived BMSCs (Liang et al., 2015), we subsequently hypothesized that CGRP may be involved in regulating BMSC functions during the aging process in mice.

CGRP Treatment Promoted BMSC Osteogenic Differentiation and Reduced BMSC Senescence

To evaluate the effect of CGRP on the osteogenic differentiation potential of BMSCs, we isolated BMSCs from femoral and
FIGURE 2 | CGRP induced the osteogenic differentiation of bone marrow-derived stem cells (BMSCs) and reduced BMSC senescence. (A) Representative images of Alizarin Red staining and (B) quantitative analysis of matrix mineralization in BMSCs treated with or without CGRP. (C) Representative images of beta-galactosidase staining and (D) quantitative analysis of the percentage of senescent cells in BMSCs treated with or without CGRP. (E) Volcano map and (F) histogram of differentially expressed genes between the CGRP treatment and control groups. Red spots represent downregulated genes and green spots represent upregulated genes. (G) Enrichment of differentially expressed genes was tested using Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway analysis and (H) Gene Ontology (GO) analysis. (I) A heatmap of the mRNA-seq profile of BMSCs treated with or without CGRP. Fold-change ≥ 2.0. (J) RT-qPCR analysis of gene expression levels in BMSCs treated with or without CGRP. Scale bar, 50 µm. Data are presented as means ± SEM. *P < 0.05, **P < 0.01, ***P < 0.001.
tibial bone marrow of mice and cultured them first in a complete medium, and then in an osteogenesis induction medium containing CGRP at 0, 50, or 100 ng/mL. Alizarin Red staining and quantitative analysis of calcium content indicated that the ability of BMSCs to form mineralized nodules was enhanced in the CGRP treatment group compared with that in the control group (Figures 2A,B). A similar result was obtained for BMSCs transfected with a CGRP expression plasmid (Supplementary Figures 1A,B). In mice, the capacity of BMSCs to differentiate into osteoblasts is known to decrease with age. Consequently, we next measured the effect of CGRP on BMSC senescence by β-galactosidase staining. As expected, the percentage of senescent cells (β-galactosidase+) was significantly lower in the CGRP treatment group than in the control group (Figures 2C,D). To determine the mechanism underlying the effect of CGRP on osteogenic differentiation of BMSCs and BMSC senescence, we performed RNA-seq to identify differences in mRNA expression levels between the CGRP treatment group and the control group. A total of 1,020 differentially expressed mRNAs (log2 fold-change ≥ 2) were identified (Figures 2E,F). Using Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway and Gene Ontology (GO) enrichment analyses of the differentially expressed genes (Figures 2G,H), we identified several biological processes involved in osteogenic and fat cell differentiation (Figure 2H). Analysis of a heatmap depicting the differentially expressed genes involved in the regulation of osteoblast differentiation demonstrated that almost all of these genes, including Gdpd2, Igf1, Fgf9, Wnt110b, Gli2, Rspo2, Jag1, Sox2, Igfbp3, Igf2, Bmp4, Bmp6, Bmp8a, Id4, Acvr2a, Pth1r, Lrp5, Hdac5, Hey1, Dlk1, Tmem119, Fam20c, Gja1, Tgfbr3, Colla1, and Ifi80, were markedly upregulated in the CGRP treatment group compared with that in the control group (Figure 2I). The result of the RNA-seq was further confirmed by RT-qPCR analysis (Figure 2J). Combined, the results showed that CGRP treatment dose-dependently increased the osteogenic differentiation potential of BMSCs and reduced BMSC senescence.

**CGRP Inhibited the Adipogenic Differentiation of BMSCs**

BMSCs are multipotent stem cells capable of differentiating into multiple lineages, including osteoblasts and adipocytes (Pittenger et al., 1999; James, 2013). The differentiation potential of BMSCs is influenced by multiple factors, such as injury. Given our above results showing that CGRP can induce the osteogenic differentiation of BMSCs and reduce BMSC senescence, we then tested whether CGRP treatment could also affect the adipogenic differentiation capacity of BMSCs. To evaluate this possibility in vitro, BMSCs were cultured in an adipogenesis induction medium containing CGRP at 0, 50, or 100 ng/mL. Oil Red O staining results showed that there were fewer lipid droplets and their size was reduced in CGRP-treated BMSCs compared with that in control BMSCs (Figures 3A,B). We obtained a similar result in BMSCs transfected with a CGRP expression plasmid (Supplementary Figures 1C,D). Next, we
identified genes involved in promoting adipogenesis that were differentially expressed between the CGRP treatment group and the control group. Heatmap analysis showed that, compared with control BMSCs, the expression of Hmga2, Ptgs2, Vgf, Per2, Acads, Fabp3, Prkab2, Dgat1, Hdac6, Egr2, and Apmap was markedly downregulated in those treated with CGRP (Figure 3C). RT-qPCR analysis further confirmed the RNA-seq results (Figure 3D). These data indicated that CGRP treatment downregulated the expression of adipogenesis-related genes and inhibited the adipogenic differentiation capacity of BMSCs.

### CGRP Treatment Promoted Bone Formation in Aged Mice

To evaluate whether CGRP could promote bone formation in aged mice in vivo, 12-month-old male C57BL/6J mice (n = 5) were administered CGRP (10 mg/kg) or 1 × PBS via tail vein injection three times weekly (Figure 4A). After 1 month, the bone phenotype of the mice in both groups was analyzed. The results showed that bone mass, BV, Tb. Th, and Tb. N were markedly higher, and the Tb. Sp lower, in mice treated with CGRP when compared with those treated with PBS (Figures 4B–F). Hematoxylin and eosin (H&E) staining results showed that, compared with PBS-treated mice, those treated with CGRP had fewer adipocytes, and the area occupied by them was smaller (Figures 4G,H). Moreover, the number of osteocalcin⁺ osteoblasts on the trabecular surface was also higher in mice treated with CGRP compared with that of the control group (Figures 4I,J). These results suggested that CGRP treatment induced osteoblast differentiation and promoted bone formation in aged mice.

### CGRP Treatment Promoted Bone Formation in OVX Mice

The OVX rodent model is well-established as a means for investigating osteoporosis and osteoporotic therapies (Mathavan et al., 2015). Here, to test the effect of CGRP on bone formation in OVX mice, we generated a model of postmenopausal osteoporosis via bilateral ovarian resection in 2-month-old female C57BL/6J mice (n = 12/group). One month after the operation, CGRP (10 mg/kg) or 1 × PBS (10 mg/kg) was administered into mice of the OVX and sham operation groups via tail vein injection and ovarian injections three times weekly (Figure 4A). RT-qPCR analysis further confirmed the RNA-seq results (Figure 3D). These data indicated that CGRP treatment downregulated the expression of adipogenesis-related genes and inhibited the adipogenic differentiation capacity of BMSCs.

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FIGURE 5 | CGRP treatment promoted bone formation in ovariectomized (OVX) mice. (A) A schematic representation of the injection protocol (CGRP or 1 × PBS via the tail vein) for OVX and control (sham-operated) mice. (B) Representative images of the ovaries of OVX and control (sham-operated) mice treated with or without CGRP. (C) Representative micro-computed tomographs (n = 6/group) and (D–G) quantitative analysis of bone volume as a fraction of total bone volume (BV/TV), trabecular thickness (Tb. Th), trabecular number (Tb. N), and trabecular separation (Tb. Sp) of femora from OVX and control (sham-operated) mice treated with or without CGRP. (H) Representative images of hematoxylin and eosin staining and (I) quantitative analysis of the number and area of adipocytes in the bone marrow of OVX and control (sham-operated) mice treated with or without CGRP. (J) Representative images of osteocalcin (OCN) staining and (K) quantification of osteoblast bone surface density (N.Ob/B.Pm) in the femora of OVX and control (sham-operated) mice treated with or without CGRP. (L) Representative images of tartrate-resistant acid phosphatase (TRAP) staining and (M) quantification of osteoclast bone surface density (N.Oc/B.Pm) in the femora of OVX and control (sham-operated) mice treated with or without CGRP. Scale bar, 300 µm. Data are presented as means ± SEM. *P < 0.05, **P < 0.01, ***P < 0.001. # means no statistical significance.
times weekly (Figures 5A,B). We found that the BV and Tb. Th were markedly reduced in the OVX group compared with those in the sham operation group, indicating that the model of postmenopausal osteoporosis had been successfully generated (Figures 5C–G). Moreover, bone mass, BV, Tb. Th, and Tb. N were markedly increased, while the Tb. Sp. was reduced, in CGRP-treated mice from both the OVX and sham operation groups compared with that in PBS-treated mice from both groups (Figure 5C–G). Additionally, when compared with control mice, CGRP-treated mice from the OVX and sham operation groups displayed fewer adipocytes and smaller adipocyte-containing areas in the bone marrow (Figures 5H,I), as well as greater numbers of osteocalcin+ osteoblasts and alkaline phosphatase+ osteoprogenitors on the bone surfaces (Figures 5J,K). In contrast, CGRP-treated mice from the OVX and sham operation groups had fewer TRAP+ osteoclasts on the bone surfaces (Figures 5L,M), which was consistent with the results of previous in vitro studies (Wang et al., 2010). Taken together, these results suggested that CGRP can promote bone formation in both aged and osteoporotic (OVX) mice, and suggest a potential approach for the treatment of age-related osteoporosis.

**DISCUSSION**

Osteoporosis is a chronic, age-related disease that seriously affects the quality of life of both the elderly and postmenopausal women (Rachner et al., 2011). Owing to the aging of the world’s population, it is increasingly important to understand the role of neuropeptides and their receptors in the aging process (Naot and Cornish, 2008). In the elderly, BMSCs have a greater capacity to differentiate along adipocytic lineages than along osteoblastic lineages, leading to the gradual accumulation of fat and the loss of bone (Bartel, 2004). Here, we demonstrated that CGRP is involved in the shift in BMSC cell lineage commitment, which leads to increased osteogenic differentiation and reduced adipocytic differentiation of BMSCs. Furthermore, we demonstrated for the first time that systemic CGRP administration can reduce fat accumulation, as well as promote bone formation, in both aged and OVX mice, likely by acting on BMSCs.

CGRP can reportedly regulate BMSC lineage commitment (Villa et al., 2003; Wang et al., 2010; Naot et al., 2019). In addition, several studies have shown that CGRP levels are influenced by age, and that these age-related changes can affect osteogenesis (Imai and Matsusue, 2002; Villa et al., 2003; Niedermair et al., 2020). These results are consistent with those reported here. We found that the level of CGRP in bone marrow supernatants was inversely proportional to age. Furthermore, CGRP treatment enhanced the ability of BMSCs to form mineralized nodules and significantly reduced the percentage of senescent BMSCs. Given the interaction between osteogenesis and adipogenesis, we further examined the role of CGRP in adipocytic differentiation, and report for the first time that CGRP can inhibit the adipogenic differentiation of BMSCs. These results indicated that CGRP regulates BMSC lineage commitment during aging and contributes to age-related bone formation.

It has recently been suggested that CGRP is a key neuropeptide in bone metabolism. Niedermair et al. (2020) demonstrated that CGRP regulates the bone remodeling properties of osteoblasts and osteoclasts in an age-dependent manner. Further, Mi et al. (2021) reported that CGRP can increase the endothelial progenitor cell population in the endothelial differentiation of BMSCs in vitro, thereby promoting bone regeneration in a rat model of distraction osteogenesis. In addition, it has been shown that CGRP can enhance BMP2 signal transduction, the expression of related osteogenic genes, and mineralization in vitro (Tuzmen and Campbell, 2018). He et al. (2016) reported that CGRP can maintain bone mass by stimulating osteoblast differentiation and inhibiting RANKL-induced osteoclastogenesis and bone resorption. CGRP has also been indicated to serve as a protective mechanism against particle-induced osteolysis (Kauther et al., 2011). However, to date, no work has reported on the role of CGRP in the regulation of BMSC function. In the present study, we defined a new mechanism through which CGRP regulates the BMSC switch in the bone. We further identified biological processes involved in both osteogenic and adipocyte differentiation and found that CGRP treatment led to a significant upregulation of the expression of almost all osteogenesis-related genes. Furthermore, based on the decrease in the expression level of adipogenesis-related genes, we reasoned that CGRP might suppress adipogenesis.

These results suggested that CGRP influences the direction of BMSC differentiation by regulating the expression of related genes, thereby providing a neuropeptide-mediated link to the age-related transition between osteoblast and adipocyte differentiation. During the aging process, the level of CGRP and the expression of osteogenic-related genes in the bone marrow decreases, whereas that of lipid-related genes increases. Consequently, BMSCs tend to differentiate into adipocytes, leading to a reduction in the number of osteoblasts and an increase in that of adipocytes, resulting in age-related bone loss. Here, we identified a new mechanism through which CGRP regulates the differentiation of BMSCs during aging. Relatively few studies have investigated the role of CGRP in bone metabolism in vivo. Here, via the delivery of CGRP to BMSCs by tail vein injection, we confirmed in vivo that CGRP can promote the osteogenic differentiation of BMSCs and inhibit fat accumulation in the bone marrow. The results of bone histomorphometry in the CGRP-treated group were significantly better than those in the control group in both aged and OVX mice. In vitro and in vivo experiments both confirmed that CGRP can promote the osteogenic differentiation and inhibit the adipogenic differentiation of BMSCs.

With the aging of the world’s population, age-related osteoporosis has become a serious public health issue (Curtis and Safford, 2012). However, most drugs currently used to treat osteoporosis are aimed at inhibiting bone resorption without promoting bone formation, and most are accompanied by severe side effects (Hu et al., 2020). In this study, we identified a new neuropeptide that may serve as potential therapeutic target for
the treatment of osteoporosis. To date, CGRP has been applied in the treatment of several diseases, such as migraine, diabetes, and liver damage, as well as for cardioprotection (Kroeger et al., 2009; Yuan et al., 2017; Guo et al., 2018, 2020), suggesting that CGRP can be used safely and effectively in the treatment of age-related osteoporosis.

CONCLUSION

In conclusion, our findings support that the age-related changes in CGRP levels regulate BMSC differentiation. Our results revealed a novel mechanism underlying age-related bone loss and provide a potential therapeutic strategy to treat age-related osteoporosis.

DATA AVAILABILITY STATEMENT

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

ETHICS STATEMENT

All animal care protocols and experiments were reviewed and approved by the Animal Care and Use Committee of the Laboratory Animal Research Center at the Second Xiangya Hospital of Central South University. All mice were maintained in the specific pathogen–free facility of the Laboratory Animal Research Center at Central South University.

AUTHOR CONTRIBUTIONS

QL conceived the study. HL and RW drafted the manuscript. HZ and JW designed the figures. XX and JQ designed the tables. QL and RW revised the manuscript. All the authors were involved in the critical revision of the manuscript and approved the final version.

FUNDING

This work was supported by the Natural Science Foundation of Hunan Province (Nos. 2020YFC2009000 and 2020YFC2009001), and the Natural Science Foundation of Changsha (KQ2007051).

SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fcell.2021.675503/full#supplementary-material

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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.