Melatonin prevents bone destruction in mice with retinoic acid–induced osteoporosis

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Abstract

Background: The protective effect of melatonin against bone metabolism imbalance in osteoporosis (OP) induced by drugs such as retinoic acid (RA) is unclear. The aim of this study was to explore the role of melatonin in bone destruction based on a mouse model.

Methods: RA-induced OP model mice were established. To assess the effect of melatonin on these mice, micro-CT was used to characterize the trabecular structure of normal mice and those treated with RA (model), RA + low-dose melatonin (Mlt-L), RA + high-dose melatonin (Mlt-H), and RA + alendronate sodium (positive control). The shape of the trabecular bone, the length and diameter of the femoral head and the height and diameter of vertebra (L1) of each group were also measured and the number of osteoclasts was determined by Tartrate-resistant acid phosphatase (TRACP) staining. Meanwhile, the expression of alkaline phosphatase (ALP) was evaluated by immunohistochemistry assays. The differences between groups in terms of liver and kidney oxidation–related indexes and serum and urinary indicators related to bone metabolism were also analyzed. Furthermore, qRT-PCR and western blotting were used to evaluate the effect of melatonin on osteogenic and osteoclastic differentiation in MC3T3-E1 and RAW264.7 cells, respectively.

Results: RA induction led to a decrease in the amount and density of trabecular bone, a decrease in the length and diameter of the femur and height, diameter of the vertebra (L1), a decrease in bone mass and density and the expression of ALP, and an increase in the number of osteoclasts. Melatonin treatment alleviated these effects induced by RA, increasing the amount of trabecular bone in OP mice, improving the microstructure of the femur and vertebra (L1) and increasing bone mass bone density and the expression of ALP, as well as decreasing the number of osteoclasts. Additionally, blood and urinary bone metabolism-related indicators showed that melatonin promoted bone formation and inhibited bone resorption. Determination of oxidant and antioxidant biomarkers in the livers and kidneys of the mice revealed that melatonin promoted the antioxidant level and suppressed the level of oxidant molecules in these organs. In vitro, RA promoted osteoclasts and inhibited osteogenesis by increasing oxidative stress levels in the RAW264.7 and MC3T3-E1 cells, but melatonin reversed this effect. Melatonin may, therefore, play a role in the ERK/SMAD and NF-κB pathways.

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Background
Osteoporosis (OP) is a systemic disease characterized by low bone mass and micro-architectural deterioration of bone tissue, which increases the risk of bone fracture (Sanchez-Barcelo et al. 2010). It is common in the elderly. Worldwide, approximately 89 million people are diagnosed with OP fracture every year, which is equivalent to one case every 3 s (Johnell and Kanis 2006). OP is divided into two major categories: primary (including mainly postmenopausal and senile OP) and secondary (including mainly endocrine and metabolic OP, disuse osteoporosis, and drug-induced OP). Treatment and care for OP requires enormous human and material resources, resulting in heavy burdens on affected families and society in general. It is believed that the main pathogenesis of OP is an imbalance of bone metabolism, in which osteogenic capacity is decreased and the likelihood of breaking bones is increased (Bonucci and Ballanti 2014). Anti-OP drugs (e.g., bisphosphonate and parathyroid hormone) currently on the market have many disadvantages, such as side effects, high price, poor patient compliance, tumorigenic risks, and an inability to fully restore the balance of bone metabolism.

Melatonin is a neuroendocrine hormone secreted by the pineal gland that has a wide range of effects, including sleep regulation, anti-aging, anti-inflammatory, and anti-oxidation activities (Jarzynka et al. 2009; Zawilska et al. 2009; Witt-Enderby et al. 2006). In the field of orthopedics, it is considered to be a promoter of bone and cartilage de-

Methods
Animal model construction and experimental design
A total of 42 female specific-pathogen-free (SPF) C57 mice aged 8 to 9 months were purchased from Charles River Laboratories (Beijing, China) and housed in a metabolic cage under 12 h light/12 h dark conditions. Mice were randomly divided into two groups: control group (n = 12) and model group (RA, 70 mg/kg/d, vegetable oil solvent, n = 30). After 15 d of administration of RA, six mice in the control group and six in the model group were tested to confirm the success of modeling. Then, mice in the model groups were randomly assigned to four groups (n = 6 per group): model group, model + low-melatonin group (5 mg/kg/d, solvent: 5% ethanol in saline), model + high-melatonin group (50 mg/kg/d, solvent: 5% ethanol in saline), and model + alendronate sodium group (10 mg/kg/week, solvent: deionized water), which were then subjected to the corresponding treatment. RA was administered by gavage, and melatonin and alendronate sodium were administered by intraperitoneal injection. The body weight of the mice was recorded weekly for 8 weeks. Two months after modeling, liver and kidney tissues of the mice in each group as well as the left and right tibia and

Conclusions: Melatonin can alleviate bone loss in RA-induced OP model mice, repair the trabecular microstructure, and promote bone formation. These effects may be related to reducing oxidation levels in vivo and vitro through the ERK/SMAD and NF-κB pathways.

Keywords: Osteoporosis, Melatonin, Bone metabolism, Bone remodeling, Oxidant stress
femur were dissected. In addition, blood samples were collected by removing the eyeball. Urine was collected using metabolic cages. The Committee of Institute Research Medical Ethics of Sun Yat-sen University granted approval for this study.

**Measurement of BMD**
For the measurement of BMD, mouse femur samples were immersed in PBS at room temperature. X-ray imaging was performed using an X-ray apparatus (SRO-M50; SOFRON, Tokyo, Japan) 15 d after gavage. The BMD was measured by a dual-energy X-ray absorptiometry apparatus (DCS-600R; Aloka, Tokyo, Japan) and expressed as the bone mineral content per unit area.

**Bone microstructure observation**
To observe the bone microstructure, mouse femur and vertebra (L1) samples were dissolved in PBS at room temperature. Micro-CT was used for two-dimensional (2D) and three-dimensional (3D) quantitative determination before sacrificing as previously described (Zhang et al. 2010b), and the microstructure parameters of cancellous bone, such as bone volume/total volume (BV/TV), trabecular number (Tb.N), trabecular thickness (Tb.Th), trabecular separation (Tb.Sp), trabecular pattern factor, and cortical bone thickness (Ct.Th), were measured automatically from the Siemens Preclinical Imaging System in a multislice in standard resolution mode (Gong et al. 2005; Hildebrand et al. 1999).

**Organ index, femur length, diameter and vertebra(L1) height, diameter determination**
Weights of the liver, spleen, left kidney, right kidney, left tibia, left femur, right tibia, right femur, and the whole body of the mice were determined and used to calculate the organ indexes using the following formula: organ index = mean organ weight (mg)/mean body weight (g) (Yang et al. 2015). In addition, the length diameter of the femur and height, diameter of the vertebra(L1) were measured and recorded.

**Hematoxylin and eosin (HE) staining and immunohistochemical (IHC) staining**
Bone samples were fixed with 4% paraformaldehyde, decalcified, dehydrated, embedded in paraffin, and cut into 5 μm slices, after which they were dewaxed by xylene and alcohol (in the following order: 100, 95, 75, and 50%). Slices were immersed in distilled water and HE staining was performed in accordance with standard protocols. Fields with positive staining were captured by microscopy. Additionally, primary antibody anti-ALP (abcam, UK) was incubated overnight at 4°C and, subsequently, incubated with anti-mouse IgG-HRP (SV-0001; Boster) for 1 h. Then, sections were observed under the microscope (OPTEC CCD TP510) with a 100× objective and pictures were collected. The staining intensity was scored according to previous research (Lian et al. 2019) as follows: 0 (negative), 1 (weakly positive), 2 (moderately positive), and 3 (strongly positive). The percent of positivity was scored according to five categories: 0 (<5%), 1 (5–25%), 2 (25–50%), 3 (50–75%), and 4 (>75%). Finally, the percent positivity score was multiplied by the staining intensity score to generate final expression scores, which ranged from 0 to 12.

**Tartrate-resistant acid phosphatase (TRACP) staining**
After treatment with melatonin, osteoclasts were fixed in 4% paraformaldehyde and then incubated in TRACP solution including tartaric acid, hexa-azo deoxymethylene red, and napthol AS-BI phosphate, among others, at 37°C for 1 h for staining. The staining procedure was carried out in accordance with the manufacturer’s instructions (Jinqiao, Xuzhou, China); TRACP-positive signals were observed and photographed using a microscope. At least four images of 100× magnification were taken at random for each group. The number of TRACP-positive cells in each image was quantified using Image-Pro Plus 6.0 (Media Cybernetics, Inc., MD, USA).

**Oxidation level detection**
Liver and kidney tissues were dissected and ground with liquid nitrogen. The resultant supernatant was collected and centrifuged. The levels of malondialdehyde (MDA), superoxide dismutase (SOD), and glutathione peroxidase (GPX) in the tissues of each group were determined using an MDA assay kit (TBA method), a T-SOD assay kit (hydroxyamine method), and a GSH-PX assay kit in accordance with standard instructions (liancheng, Nanjing, China).

**Biochemical testing**
Blood and urine samples were collected 2 months after establishment of the RA-induced model. Samples were centrifuged at 3000–4000 rpm for 10 min and the supernatant was collected for subsequent analysis. The concentrations of serum calcium, phosphorus, and ALP, as well as urinary phosphorus and urinary calcium, were tested in accordance with kit instructions (Huili, Changchun, China). Serum receptor activator of nuclear factor-κB ligand (RANKL), osteoprotegerin (OPG), and TRACP assays were performed in accordance with the mouse soluble receptor activator of nuclear factor-κB ligand (sRANKL) ELISA kit, mouse osteoprotegerin (OPG) ELISA kit, and mouse TRACP-5b ELISA kit instructions (Cusabio, Changchun, China).

**Cell culture and melatonin application**
The RAW264.7 murine monocyte/macrophage cell line and mouse osteoblast cells, MC3T3-E1, were purchased.
from the Shanghai Cell Bank, Chinese Academy of Sciences (Shanghai, China). Then, 30 ng/ml RANKL (R&D Systems, Minnesota, USA) was added to RAW264.7 cells to initiate differentiation into osteoclast-like cells, which were then cultured with 1 μM RA with or without 100 nM melatonin. To induce osteoblast differentiation, 1 μM RA with or without 100 nM melatonin in α-MEM supplemented with 10% fetal bovine serum (FBS), 50 μg/ml ascorbic acid, and 10 mmol/L β-glycerol phosphate (zqxbio, Shanghai, China) were added to the MC3T3 cells.

**Quantitative real-time polymerase chain reaction (qRT-PCR)**

Total RNA was isolated from cells using Trizol according to the manufacturer’s instructions. The cDNA was synthesized using the PrimeScript RT Master Mix (TaKaRa) according to the manufacturer’s protocols. Real-time qPCR was conducted using a real-time RT-PCR assay (TsongKe, Guangzhou, China) in a final volume of 10 μl according to the manufacturer’s protocol, and the results were analyzed using Applied Biosystems 7500 Fast System SDS software. The expression levels of mRNAs were calculated by the standard 2-△△Ct method with Gapdh as internal controls. The primer pairs used are shown in Table 1.

**Western blot analysis**

The treated cells lysates were prepared using RIPA lysis buffer containing proteinase and phosphatase inhibitors (Sigma-Aldrich, MO, USA). Protein concentration was quantified with a Pierce™ BCA Protein Assay kit (ThermoFisher, MA, USA), and then proteins were separated by 10% SDS-polyacrylamide gel before transfer to a PVDF membrane. The blotting membrane was blocked with TBST containing 5% BSA for 1 h, and then incubated with anti-Rank, anti-Runx2, anti-Sod1, anti-Sod2, anti-p-Erk1/2 (Thr202/Tyr204), anti-Erk1/2, anti-p65, Cu/Zn superoxide dismutase (zqxbio, Shanghai, China) were added to the MC3T3 cells.

| Table 1: All primers for reverse transcription-quantitative polymerase chain reaction |
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| **Gene** | **Primer sequence(5′→3′)** |
| Gapdh | Forward:AGGTCCGGTGTGAAACGGATTTG; Reverse: TGGACACCACTGATTGGAGTCA; |
| Tnfrsf11a | Forward: GGACGGTGTGGTCACAGAAT; Reverse: GCAGTGTACCTGACGAGTTT |
| Ctsk | Forward: GAAGAAGACTCACCAGAAGC; Reverse: GCAGTCTGAGTATTGGACAGG; |
| Calcr | Forward: AACACGGAATGAGGATCAGC; Reverse: CTGGCTCACTAGAGGAGCAG; |
| Ocn | Forward: CTCGGTGGCGTTGAGATTGTT; Reverse: CCACCATGCTTGACATCCAGT; |
| Opn | Forward: ACCCAGAAGTGCGTACTAGC; Reverse: CTGCTGTACAAATACACTACACT; |
| Runx2 | Forward: TCTCCAGGTCTATCCATCTGCA; Reverse: GCCTGACATACGTCTGATGTT; |
| Nox1 | Forward: TTGTTGTTGGGCTTGAATGTC; Reverse: CTGAGAAAGAGGAGGACAGTG; |
| Nox2 | Forward: AACCAGTGGTTGTGTAGGAC; Reverse: CACCACCATGCTTTGAGATGAGG; |
| Sod1 | Forward: GAAGAAGACTCACCAGAAGC; Reverse: GCAGTCTGAGTATTGGACAGG; |
| Sod2 | Forward: CAGACCTGGCTTCATAGCTACTG; Reverse: CTCCGGTGGCTGTTACAGG; |

**Results**

**Construction of OP model induced by RA**

The OP model was constructed using RA and subjected to femoral micro-CT testing 15 d after gavage. The 2D and 3D reconstruction micro-CT results showed that the amount of trabecular bone in the model group was reduced and its density was low (Fig. 1a). In addition, BV/TV, Tb. Th, and Tb. N showed significant decrease in the OP model mice, while the value of Tb. Sp was slightly increased (Fig. 1b-e).

HE staining showed that in the model group, the number of femoral trabeculae was reduced and the bone structure had a low density (Fig. 2a). Compared to the normal control group, the bone mass density also significantly decreased in the model group (Fig. 2b). TRACP staining showed a significant increase in the number of TRACP-positive cells in the model group compared with that in the control (Fig. 2c and d), indicating that the OP mouse model had been successfully constructed.
Melatonin improves femoral and vertebral (L1) changes in OP mice
To study the effect of melatonin on the femurs and vertebra (L1) of OP mice, the control group, a model group, a low-dose melatonin model group (Mlt-L), and a high-dose melatonin model group (Mlt-H) were established, while OP mice treated with alendronate were used as a positive control. During the modeling period, the weight of the mice generally decreased, then gradually increased again after the modeling stopped (Fig. 3a). Analysis of the organ weight index showed that melatonin can reverse changes induced by RA in the weight indexes of the liver, both kidneys, both tibia, and both femurs of mice (Fig. 3b). Measurements showed that the length, and height, diameter of the vertebral (L1) were significantly reduced in the model group, and the melatonin treatment (both Mlt-L and Mlt-H) restored the length, diameter of the femur and height, diameter of the vertebra to the same values as the control group (Fig. 3c-f).

In addition, the trabecular microstructure of the femur and the vertebra (L1) were characterized by micro-CT after the administration of melatonin or alendronate. Micro-CT 2D imaging and 3D reconstructions showed a reduction in the number of femoral and vertebral (L1) trabeculae and the thickness of femur diaphysis in the model group (Figs. 4a, b, and 5a), whereas mice treated with melatonin (both Mlt-L and Mlt-H) had significantly higher BV/TV values than OP model mice (Figs. 4c and 5b). In addition, Tb. Th, Tb. N, and Ct. Th in the Mlt-L and Mlt-H mice were found to be significantly increased (Figs. 4d, e, h and 5c, d) while Tb. Sp and the trabecular pattern factor were significantly decreased (Figs. 4f, g, and 5e) compared to the OP model mice.

Melatonin improves OP in mice by reducing the number of osteoclasts and increasing the expression of ALP
To investigate the effect of melatonin on the number of femoral trabecular cells, HE staining, TRACP staining and ALP IHC staining were used to observe pathological changes in the distal femur and trabecular bone. HE staining revealed that the amount of trabecular bone decreased and the bone structure became sparse in OP, and that this condition was improved in the Mlt-L, Mlt-H, and positive control groups, in that the trabecular bone became more abundant (Fig. 6a). TRACP staining showed an increase in the number of osteoclasts in OP mice, which decreased after the administration of melatonin; TRACP-positive signals in the Mlt-L and Mlt-H groups were significantly lower than in the model group and the positive control group, and were similar to that in the control group (Fig. 6b and e). In addition, BMD measurements in each group showed a significant reduction in bone density in OP mice, but melatonin and alendronate treatments restored the bone density of the model mice to the normal control level (Fig. 6d). Further confirming that melatonin could improve OP, the expression of ALP measured by IHC was inhibited in OP mice, but upregulated in the melatonin and alendronate groups (Fig. 6c and f). Additionally, HE staining, TRACP staining, and ALP IHC staining were performed on vertebra, which more comprehensively confirmed that melatonin can improve OP (Additional file 1: Figure S1).

Melatonin improves bone metabolism and oxidation in OP mice
After treatment with RA for 15 d, serum and urinary calcium and phosphorus concentration as well as the expression of ALP, RANKL, TRACP and OPG in the serum of mice were tested (Additional file 2: Figure S2). There was no difference in the expression of detection indicators in all mice in the model group. And after randomized treatment with melatonin and alendronate, the corresponding expression was different. The results confirmed that melatonin had a therapeutic effect on RA-induced osteoporosis. In the model group, the concentrations of serum RANKL, TRACP, ALP,
phosphorus, and calcium were increased (Fig. 7a, b, and d-f), while OPG (Fig. 7c) was decreased significantly compared to control mice. In both Mlt-L and Mlt-H, these changes were repaired. Urinary sample testing also showed increases in urinary calcium and phosphorus concentrations in the model mice, and melatonin and alendronate treatments reversed this increase in the model group (Fig. 7g and h).

In addition, the oxidation level of mouse livers and kidneys was measured. The level of MDA was increased

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**Fig. 2** Osteoporosis (OP) was induced in mice with retinoic acid (RA). HE staining (40X) (a), femur bone mineral density determination (b), and TRACP staining (100X) (c) were performed to investigate the changes in femoral trabecular structure, bone density, and number of osteoclasts (d) in the OP model. Control: normal mice, Model: RA-induced OP model mice. *, P < 0.05 vs control; n = 6 per group

**Fig. 3** Melatonin improves femoral and vertebral (L1) changes, body weight, and organ index in osteoporosis (OP) mice. Effects of treatment with melatonin at different doses on body weight (a), organ index (b), femur diameter (c), femur length (d), vertebra diameter (e), and vertebra height (f). Control: normal mice, Model: retinoic acid (RA)-induced OP model mice, Mlt-L: low-dose melatonin-treated OP model mice, Mlt-H: high-dose melatonin-treated OP model mice. Positive control: alendronate-treated OP mice. *, P < 0.05 vs control. #, P < 0.05 vs model; n = 6 per group
and the levels of GPX and SOD were significantly decreased in model mice. Meanwhile, in both Mlt-L and Mlt-H as well as alendronate-treated mice, the level of MDA was suppressed and the levels of GPX and SOD were restored to those of the control group (Fig. 8).

Melatonin can inhibit osteoclast differentiation and promote osteogenic differentiation in vitro

To investigate the effect of melatonin on osteogenic and osteoclastic differentiation in vitro in RAW264.7 and MC3T3-E1 cells, respectively, were induced. TRACP staining showed an increase in the number of osteoclasts in the RA-induced RAW264.7 cells with RANKL, but the number of osteoclasts was decreased in the RA + Mlt groups (Fig. 9a). Further, markers of osteoclast differentiation (Calcr, Ctsk, and Rank) and oxidation indexes (Nox1 and Nox2) were upregulated, as detected by qPCR and western blot, and antioxidant enzymes (Sod1 and Sod2) were downregulated in RA-induced RAW264.7 cells. However, melatonin could reverse these changes (Fig. 9b, c, f, and g).

Meanwhile, we induced osteogenic differentiation in MC3T3-E1 cells and studied the effect of melatonin on osteogenic differentiation. QPCR and western blot results showed that melatonin could increase the markers of osteogenic differentiation (Runx2, Opn, and Ocn) and antioxidant enzymes (Sod1 and Sod2) and inhibit the expression of oxidation enzymes (Nox1 and Nox2) in RA-induced MC3T3-E1 cells (Fig. 9d, e, i, and j). Finally, we studied the signaling pathways involved and found that RA could upregulate the expression of p-p65 and p-1κBα in RAW264.7 induced by RANKL, whereas melatonin downregulated this expression (Fig. 9h). When osteogenic differentiation was induced in MC3T3-E1 cells with RA, the expression of p-ERK and p-SMAD1 were decreased, but melatonin reversed this (Fig. 9k).

Discussion

Osteoporosis is a systemic bone disease that results in thinned bone mass, altered microstructure of bone tissues, and higher risk of fracture. RA and ovariectomy (OVX) treatments are frequently used to induce osteoporosis. Compared with the long and complicated experimental process of the OVX model, the RA-induced osteoporosis model is simple, quick, and...
highly successful, and shows the typical symptoms of osteoporosis (Cowan et al. 2005; Wei et al. 2007; Oršolić et al. 2018). As a derivative of vitamin A and an anti-tumor drug, RA can activate osteoclasts and promote bone absorption (Yang et al. 2016). Meanwhile, it has been reported that RA can damage the ovarian function, decrease the estrogen level, and also affect the occurrence of osteoporosis (Zhao et al. 2016).

In recent years, the RA-induced OP model has been widely used in drug-induced secondary OP studies (Yang et al. 2016), especially in the research and development of Chinese traditional medicine. Using RA-induced animal models, previous studies found that nylestriol/levonorgestrel (Liao et al. 2003), naringin, and quercetin play beneficial roles in protecting against RA-induced OP in rats (Wei et al. 2007; Oršolić et al. 2018). In addition, deer tendon collagen and isoflavone-enriched soy protein have also been found to exert protective effects in an RA-induced mouse model (Zhang et al. 2010c; Yang et al. 2016).

Accumulating evidence shows that melatonin has protective effects against OP in menopausal women. However, as previous studies mainly focused on the role of melatonin in diabetes-induced OP and ovariectomy-induced OP, its role in drug-induced OP models remained unclear. In the current study, by constructing an RA-induced OP mouse model developed to investigate drug-induced OP (Yang et al. 2016), we analyzed differences in trabecular bone, bone metabolism markers, and liver and kidney oxidative stress between groups of mice (normal mice, RA-induced model group, low-dose and high-dose melatonin-treated model mice, and a sodium alendronate-treated group as a positive control) and found that RA induction can lead to changes in trabecular morphology and microstructure, decreased bone density and the expression of ALP, and increased osteoclasts. Meanwhile, melatonin administration can alleviate the effects of RA, rebuilding trabecular bone structure, increasing bone mass, bone density and the expression of ALP, and decreasing the number of osteoclasts. Our results suggest that melatonin has a treatment effect against drug-induced OP. Additionally, our study established a positive control in which mice were treated with alendronate, a drug thought to protect against bone loss and reduce the incidence of fractures.
in menopausal women (Saag et al. 2017; Liberman et al. 1995). According to a number of indicators, melatonin worked similarly to alendronate. This suggests the therapeutic function and clinical value of melatonin in drug-induced OP bone metabolism imbalance.

Micro-architectural deterioration of bone tissues and low bone mass are two characteristics of OP. Another study has shown that the RA-induced OP model exhibits bone loss and bone structural changes within 1–3 weeks (Fahmy and Soliman 2009). Melatonin treatment of ovariectomized model OP mice has been reported to improve bone microstructure (Uslu et al. 2007). Our current IHC assay and micro-CT data provide evidence that bone microstructure and bone density changed after 15 d of RA induction; moreover, we found that bone structure and bone metabolism imbalance were improved after melatonin treatment. These observations suggest that melatonin has beneficial effects on bone microstructure and enhances bone density, which is consistent with the research results of Sharan et al. (2017). Their results revealed that oral administration of melatonin can increase bone mass and reverse the OVX-induced microstructural and functional degeneration of bone through increasing bone formation.

Various types of cells, particularly osteoclasts and osteoblasts, are involved in the bone remodeling process. Factors such as RANKL and its decoy receptor, OPG, which is secreted from osteoblasts (Martin and Sims 2015), are essential for promoting osteoclast differentiation and inhibiting bone formation. OP is one of the results of an imbalance in bone remodeling. As reviewed by Sanchez-Barcelo et al. (2010), the mechanism of action of melatonin in protecting against OP mainly involves the following aspects: enhanced osteoblast differentiation and activity;

**Fig. 6** Melatonin improves the femur trabecular bone and reduces the number of osteoclasts of osteoporosis (OP) model mice. After the administration of melatonin or alendronate to trabecular bone for 2 months, distal femur was detected by HE staining (40X) (a), TRACP staining (100X) (b), and ALP IHC staining (100X and 400X) (c). Bone density was assessed (d), and the number of TRACP-stained osteoclasts was quantified (e). The IHC staining intensity was scored (f). Control: normal mice, Model: retinoic acid (RA)-induced OP model mice, Mit-L: low-dose melatonin-treated OP model mice, Mit-H: high-dose melatonin-treated OP model mice. Positive control: alendronate-treated OP mice. *, P < 0.05 vs control. #, P < 0.05 vs model; n = 6 per group.
suppressed differentiation of osteoclasts; and the clearing of free radicals produced by osteoclasts and responsible for bone resorption. When bone resorption occurs more rapidly than bone formation, bone loss occurs (Isomura et al. 2004). Bone resorptive markers secreted by osteoclasts such as TRACP-5b (Kuo and Chen 2017; Liu et al. 2015), a commonly used biomarker reflecting bone formation, and ALP are upregulated in OP patients and RA-induced animal models (Oršolić et al. 2018; Fink et al. 2016). Consistent with previous studies, the current study revealed that the expression levels of RANKL, which promotes osteoclast differentiation; TRACP-5b, which promotes osteoclast-mediated bone resorption; and ALP, which inhibits osteoclast differentiation, were upregulated, and that these changes were reversed in OP mice treated with melatonin. These findings indicate that melatonin inhibits bone resorption and osteoclastic differentiation in RA-induced OP mice. Additionally, TRACP staining showed that melatonin reduced the number of osteoclasts in the femurs of OP mice. Meanwhile, the effect of melatonin on the osteoclast differentiation of RAW264.7 cells and osteogenic differentiation of MC3T3-E1 cells were performed. We found that melatonin could inhibit the expression of the markers of osteoclast differentiation (Calcr, Ctsk, and Rank) in the RA-induced RAW264.7 cells, but increase the markers of osteogenic differentiation (Runx2, Opn, and Ocn) in the RA-induced MC3T3-E1 cells. Previous study has showed that melatonin could promote human adult mesenchymal stem cells (hAMSCs) differentiation into osteoblasts through an increase in mRNA expression of osteogenic genes Runx2, Bmp-2, and Osteocalcin and this may promote bone formation (Sethi et al. 2010). Therefore, it is possible that melatonin promotes bone remodeling in OP mice by inhibiting osteoclast differentiation and promoting osteogenic differentiation.

Calcium and phosphorus are important bone minerals. Evidence has shown that patients with OP have a high incidence of hypercalcuria (El-Husseini et al. 2017). RA-induced OP animal models also exhibited elevated levels of serum calcium and phosphorus in the present study, which may be related to the increase in the number of osteoclasts in OP; the
increase in osteoclasts promotes bone resorption and bone mineral dissolution, decreases bone calcium and phosphorus levels, and increases serum and urinary levels of calcium and phosphorus. Increased urinary calcium excretion suggests increased bone resorption. The application of melatonin reduces the levels of serum and urinary calcium and phosphorus, indicating that melatonin alleviates OP.

Oxidative stress plays an important role in the formation of bone. Excessive free radical production and aggregation can induce oxidative stress, cause lipid peroxidation, inhibit osteogenic differentiation, and promote bone resorption (Bai et al. 2004; Wauquier et al. 2009; Callaway and Jiang 2015). MDA, the end-product of important oxygen free lipid peroxidation, is an important marker of the degree of damage of reactive oxygen free radicals (Spirlandeli et al. 2014). Studies have found that it is significantly elevated in the serum of women with OP after menopause (Ozgocmen et al. 2007; Sendur et al. 2009). The antioxidant enzymes SOD and GPX scavenge free radicals, and their levels have been found to be significantly lower in the serum of women with OP after menopause than in healthy postmenopausal women (Sharma et al. 2015). If excessive free radical production exceeds the coping ability of the natural antioxidant defense mechanism, bone resorption may occur (Sheweita and Khoshhal 2007). In the present study, we found that MDA increased in the model group, suggesting an increase in oxidative stress and decreases in SOD and GPX, in turn suggesting a decrease in antioxidant levels. In contrast, SOD and GPX in the melatonin-treated group were significantly improved, suggesting that melatonin has an antioxidative effect, possibly due to scavenging free radicals, in the RA-induced OP model (Orsolic et al. 2014).

![Fig. 8 Melatonin improves oxidation in osteoporosis (OP) mice. Oxidative stress-related liver superoxide dismutase (SOD) (a), malondialdehyde (MDA) (b), and glutathione peroxidase (GPX) (c); left kidney SOD (d), MDA (e), and GPX (f); and right kidney SOD (g), MDA (h), and GPX (i) were determined after the administration of melatonin or alendronate for 2 months. Control: normal mice, Model: retinoic acid (RA)-induced OP model mice, Mlt-L: low-dose melatonin-treated OP model mice, Mlt-H: high-dose melatonin-treated OP model mice. Positive control: alendronate-treated OP mice. *, P < 0.05 vs control. #, P < 0.05 vs model; n = 6 per group.](attachment:image.png)
Conclusions

In summary, melatonin can improve the RA-induced trabecular microstructure of OP model mice, inhibit osteoclast differentiation, and promote bone formation. These effects may be related to the reduction of oxidative stress in vivo and vitro through the ERK/SMAD and NF-κB pathways.

Additional files

**Additional file 1:** Figure S1. Melatonin improves the vertebral (L1) trabecular bone and reduces the number of osteoclasts of osteoporosis (OP) model mice. After the administration of melatonin or alendronate for 2 months, vertebra were examined by HE staining (40X) (**A**), TRACP staining (100X) (**B**), and ALP IHC staining (100X and 400X) (**C**). The number of TRACP-stained osteoclasts was quantified (**D**). The expression levels of Calcr, Ctsk, Rank, Nox1, Nox2, Sod1, and Sod2 were examined by RT-PCR and western blot (**E, F, G, H, I, J, K**). Signaling pathway related proteins (p-p65/p65, p-IκBα/IκBα, p-Erk/Erk, and p-Smad1/Smad1) were examined by western blot (**L, M, N, O, P, Q**). Control: normal mice, Model: retinoic acid (RA)-induced OP model mice, Mlt-L: low-dose melatonin-treated OP model mice, Mlt-H: high-dose melatonin-treated OP model mice. Positive control: alendronate-treated OP mice. *, P < 0.05 vs control. #, P < 0.05 vs model; n = 6 per group. (TIF 8735 kb)

**Additional file 2:** Figure S2. Biochemical markers bone metabolism in osteoporosis (OP) mice. Laboratory tests for biochemical markers of bone metabolism. Serum bone metabolism-related indicators RANKL (**A**), TRACP (**B**), OPG (**C**), ALP (**D**), phosphorus (**E**), calcium (**F**), and urinary phosphorus (**G**), and calcium (**H**) were determined after the administration of retinoic acid (RA) for 15 d. Control: normal mice, Model: RA-induced OP model mice, Mlt-L: low-dose melatonin-treated OP model mice, Mlt-H: high-dose melatonin-treated OP model mice. Positive control: alendronate-treated OP mice. *, P < 0.05 vs control. #, P < 0.05 vs model; n = 6 per group. (TIF 1247 kb)

**Abbreviations**

ALP: Alkaline phosphatase; BV/TV: Bone volume/Total volume; Calcr: Calcitonin receptor; Ctsk: Cathepsin K; GPX: Glutathione peroxidase; HE: Hematoxylin and eosin; MDA: Malondialdehyde; Mlt-H: High-dose melatonin; Mlt-L: Low-dose melatonin; Nox1: NADPH oxidase 1; Nox2: NADPH oxidase 2; Ocn: Osteocalcin; OP: Osteoporosis; OPG: Osteoprotegerin; Opn: Osteopontin;
RA: Retinoic acid; RANKL: Receptor activator of nuclear factor-κB ligand; Runt2: Runt-related transcription factor-2; SOD: Superoxide dismutase; Sod1: Cu/Zn superoxide dismutase; Sod2: Manganese superoxide dismutase; SPP: Specific-pathogen-free; sRANKL: Soluble receptor activator of nuclear factor-κB ligand; Tb:N: Trabecular number; Tb:Sp: Trabecular separation; Tb:Th: Trabecular thickness; Tnfrsf11a: Tumor necrosis factor receptor superfamily member 11a; TRACP: Tartrate-resistant acid phosphatase; TRACP-Sb: Tartrate-resistant acid phosphatase-Sb

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Authors’ contributions
DH and PS designed the experiments. XW, TL, YZ, XQ, CL, BG, YP, AL, HZ, XY, ZL, YL, and CX conducted the experiments. XW, TL, and YZ analyzed the data. XW, TL, YZ, DH, and PS wrote the manuscript. All authors read and approved the final manuscript.

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Availability of data and materials
All relevant data supporting the conclusions of this article is included within the manuscript.

Ethics approval and consent to participate
The Committee of Institute Research Medical Ethics Committee of Sun Yat-sen University granted approval for this study.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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