Melatonin Reverses the Loss of Stemness Induced by TNF-α in Human Bone Marrow Mesenchymal Stem Cells through Upregulation of YAP Expression

Xudong Wang,1 Tongzhou Liang,1 Jincheng Qiu,1 Xianjian Qiu,1 Bo Gao,1,1 Wenjie Gao,1 Chengjie Lian,2 Taiqiu Chen,1 Yuanxin Zhu,1 Anjing Liang,1 Peiqiang Su,2 Yan Peng,1 and Dongsheng Huang1

1Department of Orthopedics, Sun Yat-sen Memorial Hospital of Sun Yat-sen University, Guangzhou, Guangdong 510120, China
2Department of Orthopedics, The First Affiliated Hospital of Sun Yat-sen University, Guangzhou, Guangdong 510080, China

Correspondence should be addressed to Yan Peng; pygz258@126.com and Dongsheng Huang; hdongsh@mail.sysu.edu.cn

Received 12 August 2019; Revised 31 October 2019; Accepted 8 November 2019; Published 30 December 2019

Academic Editor: Federico Mussano

Copyright © 2019 Xudong Wang et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Mesenchymal stem cells (MSCs) are promising candidates for tissue regeneration and disease treatment. However, long-term in vitro culture results in loss of MSC stemness. The inflammation that occurs at stem cell transplant sites (such as that resulting from TNF-α) is a contributing factor for stem cell treatment failure. Currently, there is little evidence regarding the protective role of melatonin with regard to the negative effect of TNF-α on the stemness of MSCs. In this study, we report a melatonin-based method to reduce the inflammatory effects on the stemness of bone marrow mesenchymal stem cells (BMMSCs). The results of colony formation assays, Alizarin red staining, western blotting, and reverse transcription-polymerase chain reactions suggest that melatonin can reverse the inflammatory damage caused by TNF-α treatment in the third, seventh, and tenth generations of primary BMMSCs (vs. control and the TNF-α-treated group). Meanwhile, a detailed analysis of the molecular mechanisms showed that the melatonin receptor and YAP signaling pathway are closely related to the role that melatonin plays in negative inflammatory effects against BMMSCs. In addition, in vivo experiments showed that melatonin could reverse the damage caused by TNF-α on bone regeneration by BMMSCs in nude mice. Overall, our results suggest that melatonin can reverse the loss of stemness caused by inflammatory factor TNF-α in BMMSCs. Our results also provide a practical strategy for the application of BMMSCs in tissue engineering and cell therapy.

1. Introduction

Bone marrow mesenchymal stem cells (BMMSCs) are spindle-shaped adherent cells that were originally detected in bone marrow cultures. Subsequent studies have suggested that MSCs are mainly present in the bone marrow stroma [1]. Stemness is defined as the ability of stem cells to maintain the potential for proliferation and multiple routes of differentiation [2]. Under different induction conditions, MSCs can differentiate into a variety of mesodermal tissue cells, such as chondrocytes, osteoblasts, cardiomyocytes, and adipocytes. Meanwhile, MSCs can also differentiate into endoderm and ectoderm cells, such as hepatocyte-like cells and neuron-like cells [3]. In addition, multiple advantages of using MSCs in clinical applications have been reported, including low immunogenicity, multidirectional differentiation potential, induction of immune tolerance, immunosuppression, and lack of associated ethical issues [4]. Therefore, MSCs have become a primary candidate for cell therapy and tissue engineering. However, in clinical practice, MSCs need to be expanded in vitro to obtain sufficient quantities, but this subjects them to the deleterious effects of replicative aging. In addition, MSCs that experience oxidative stress may undergo premature aging, which can significantly affect their ability to differentiate into different types of cells [5, 6]. These factors limit the clinical application of MSCs [7].

Tumor necrosis factor-α (TNF-α) is a cytokine produced following the activation of macrophages/monocytes. This
factor plays an important role in the maintenance of cell stemness and in fat differentiation [8]. For example, TNF-α can inhibit the differentiation of stem cells into osteoblasts through multiple signaling pathways, including through wingless-type MMTV integration site family members (Wnt), bone morphogenetic protein- (BMP-) Smads, mitogen-activated protein kinase (MAPK), and nuclear transcription factor kapp a B (NF-xB) signaling [9, 10]. However, it is unclear what associated factors are required for TNF-α to maintain stemness and the potential for cell differentiation in BMMSCs [11].

Melatonin is a hormone secreted mainly from the pineal gland that has proven to have widespread effects [12]. Previous studies have shown that melatonin regulates various physiological functions such as sleep, circadian rhythms, and neuroendocrine activities [13]. Melatonin has well-known antioxidant properties and has been shown to eliminate excessive free radicals and increase synthesis of intracellular antioxidant enzymes [14, 15]. Melatonin protects cells from proinflammatory cytokines by reducing active oxygen production and increasing superoxide dismutase production [16]. In 1999, Dun et al. confirmed that high levels of melatonin were present in the bone marrow [17]. In recent years, melatonin has been shown to regulate pluripotent differentiation of MSCs [18]. Radio et al. found that melatonin enhanced alkaline phosphatase activity in human MSCs via the MAPK signaling pathway. Moreover, in vivo studies have confirmed that melatonin promotes bone formation in mice at pharmacological concentrations [19]. Therefore, the detailed roles of melatonin and TNF-α in the stemness of MSCs deserve further investigation.

In this study, we employed TNF-α to simulate inflammation in the environment of the third, seventh, and tenth generations of BMMSCs. Colony formation, Alizarin red staining, western blotting, and RT-PCR were used to assess the molecular effects of TNF-α and/or melatonin-treated human BMMSCs. In addition, an osteoporosis nude mouse model was used to study pathological changes with hematoxylin and eosin (H&E) staining and a micro-CT approach. The results obtained in this study provide detailed information regarding BMMSC stemness that will be useful in future clinical applications.

2.2. Flow Cytometry. BMMSCs were analyzed by flow cytometry (DxFLEX, Beckman Coulter, USA). Cells at P3 were incubated in PE anti-human CD73 (Ecto-5'-nucleotidase) (BioLegend, USA), PE anti-human CD73 (Ecto-5'-nucleotidase) (BioLegend, USA), APC anti-human CD90 (Thy1) (BioLegend, USA), APC anti-human CD34 (BioLegend, USA), Alexa Fluor® 488 anti-human CD45 (BioLegend, USA), and Alexa Fluor® 488 anti-human CD105 (Abcam, USA) at concentrations specified by the manufacturer. Corresponding isotype-identical antibodies served as controls.

2.3. BMMSC Treatment. BMMSCs at P3, P7, and P10 were induced toward osteogenic differentiation and adipogenic differentiation. For osteogenic induction, BMMSCs were plated in DMEM supplement with osteogenic differentiation assay (LiuheBio, China). After 2 weeks, differentiated BMMSCs were stained with Alizarin red S (ARS) solution (Sigma-Aldrich, USA). The detailed experimental processes of the tests were conducted as described previously [24]. For adipogenic induction, BMMSCs were seeded in mesenchymal stem cell-adipocyte differentiation medium (Sciencell, China). The experimental protocol was followed as provided by the manufacturer. The colony formation assays were conducted with untreated BMMSCs and BMMSCs treated with 20 ng/mL TNF-α, TNF-α +100 μM melatonin (Mel), TNF-α +Mel+10 μM luzindole (Luz), or TNF-α +Mel+5 μM verteporfin (VP, Yes-associated protein (YAP) inhibitor). About 500 cells per well were added to six-well culture plates. After a 2-week incubation, the cells were washed twice with PBS and stained with toluidine blue solution (BestBio, China). The clone formation efficiency was calculated as [number of colonies]/[number of cells inoculated]. Melatonin, TNF-α, Luz, and VP were obtained from Sigma-Aldrich (USA).

2.4. RT-PCR. RNAiso Plus reagent (Roche, Switzerland) was used to extract total RNA from BMMSCs. Reverse transcription-PCR (RT-PCR) analysis was carried out in 20 μL final volume from 400 ng total RNA using the TaKaRa PrimeScript II 1st Strand cDNA Synthesis Kit (D6210A, TaKaRa, Japan) according to manufacturer’s instructions. GAPDH, RUNX2, OPN, OCN, HTERT, SOX2, C-MYC, NANOG, PPARγ-2, LPL, ADIPONECTIN, and YAP mRNA levels were determined by RT-PCR, and primers were utilized as described in Table 1. Reactions were performed in 20 μL volumes containing 2x SYBR® Premix Ex Taq™ (TaKaRa, Japan), PCR Forward Primer (10 μM), PCR Reverse Primer (10 μM), cDNA template, and ddH₂O. RT-PCR was performed using a Bio-Rad CFX96 Real-Time PCR System. Thermal cycling conditions were as follows: 10 min at 95°C, followed by 40 cycles of 95°C for 10 s, 60°C for 20 s, and 72°C for 30 s. Data were collected and calculated using the Bio-Rad CFX Manager Software1.6. RNA expression was calculated based on a relative standard curve with the 2⁻ΔΔCt method.
2.5. Western Blotting. Total cellular protein from differently treated BMMSCs was isolated by the addition of 1% PMSF and RIPA lysis buffer (50 mM Tris-HCl (pH 7.4), 150 mM NaCl, 1% NP-40, 0.1% SDS). After boiling in SDS-PAGE sample buffer for 5 min, the samples underwent sodium dodecylsulfate-polyacrylamide gel electrophoresis (SDS-PAGE). The proteins were then transferred onto polyvinylidene difluoride (PVDF) membranes (Millipore, USA). After being blocked for 1 h at room temperature, the membranes were incubated with a 1:1000 dilution of the following antibodies (Abs) from Abcam (Cambridge, UK) overnight: Abs against HTERT, SOX2, C-MYC, NANOG, LPL, and ADIPONECTIN. After being blocked for 1 h at room temperature, the membranes were incubated with primary Abs against ADIPONECTIN, and GAPDH. Before detection with an ECL chemiluminescence detection kit (Beyotime Institute of Biotechnology, China), the membranes were washed three times with TBST to remove unbound Abs. Then, the membranes were reacted with HRP-conjugated goat anti-rabbit IgG (H&L) and HRP-conjugated goat ant-mouse IgG (H&L) secondary Abs (1:5000 dilution) for 1 h at room temperature. The bands were analyzed using Scanner (Scanco Medical AG, Switzerland) before sacrificing, multislice in standard resolution mode [27, 28].

2.6. Animal Treatment. Eight-week-old Balb/c nude mice were obtained from Charles River (Beijing, China). Sixty nude mice were randomly divided into six groups, including control (Con, n = 10), osteoporosis simulation group (OVX, n = 10), OVX+P3 BMSC treatment (OVX+P3, n = 10), OVX+P3 BMSCs+20 ng/mL TNF-α treatment (OVX+P3+TNF-α, n = 10), OVX+P3 BMSCs+TNF-α+100 µM Mel treatment (OVX+P3+TNF-α+Mel, n = 10), and OVX+P3 BMSCs+TNF-α+Mel+5 µM verteporfin (OVX+P3+TNF-α+Mel+VP, n = 10). After the model of osteoporosis was established (ovary removal surgery), BMMSCs treated with Mel, TNF-α, and VP were injected into the tail veins. Eight weeks after modeling, femur tissues were collected for further experimentation.

2.7. H&E Staining. H&E staining was conducted as previously reported [25]. Bone samples were fixed in 10% buffered formalin and embedded in paraffin. Three- to five-micrometer thick sections were stained with hematoxylin (Sigma H 3136) for 10 min and with eosin (Sigma E 4382) for 1 min to establish the diagnosis areas.

2.8. Bone Microstructure Observation. To observe the bone microstructure, micro-CT examination was carried out with SCANCO Medical AG (Switzerland) before sacrificing, as previously described [26]. The parameters were set as follows: voltage = 50 kV; current = 500 µA; resolution = 8.96 µm; exposure time = 5 ms; scanning layer = 2394. Subsequently, the microstructure parameters of bone, such as bone volume/total volume (BV/TV), trabecular number (Tb.N), trabecular thickness (Tb.Th), trabecular separation (Tb.Sp), and trabecular pattern factor, were measured automatically using the Siemens Preclinical Imaging System in a multislice in standard resolution mode [27, 28].

2.9. Statistical Analysis. Statistical differences in this study were assessed using SPSS 18.0 software. The mean values of the statistical results for the measured data are presented as the means ± standard error. The values in this study were analyzed by Student’s t-test. P values < 0.05 were considered to be a significant difference between two measurements.

3. Results

3.1. BMMSC Stemness Was Decreased along with Cell Passage. In this study, BMMSCs at P3 were subjected to flow cytometric analysis to estimate the presence of surface markers (Supplementary Figure 1). Most BMMSCs (>95%) were positive for CD73, CD90, and CD105, whereas <2% were positive for CD45 or CD34. The relationship between the stemness of BMMSCs and cell inheritance was examined by colony formation assays, western blotting, RT-PCR, and Alizarin red staining. For colony formation analysis, we selected the primary BMMSCs at P3, P7, and P10. The results showed that colony formation of P7 and P10 BMMSCs was significantly reduced compared with that of P3 and primary BMMSCs (P < 0.01) (Figure 1(a)). For western blot analysis, we evaluated biomarkers of BMMSC stemness (HTERT, SOX2, C-MYC, and NANOG), osteogenic differentiation (RUNX2, OPN, and OCN), and adipose differentiation (PPARγ-2, LPL, and ADIPONECTIN) in P3, P7, and P10 BMMSCs (Figures 1(b)–1(d)). The results revealed that abundance of proteins reflecting BMMSC stemness, osteogenic differentiation, and adipose differentiation was significantly downregulated in P7 and P10 cells compared with P3 cells. Meanwhile, RT-PCR analyses gave similar results for these
Figure 1: Continued.
biomarkers as western blot analyses. mRNA abundances of these biomarkers in P7 and P10 cells were all significantly reduced compared with those in P3 cells (P < 0.01) (Figures 1(e)–1(g)). In addition, we employed Alizarin red staining to investigate osteogenic differentiation in P3, P7, and P10 BMMSCs. Quantification analysis revealed that the optical density (OD) of staining in P7 and P10 cells was lower than that in P3 cells (Figure 1(h)). In summary, stemness was closely correlated with the generation of BMMSCs observed.

3.2. Melatonin Can Reverse the Loss of BMMS Stemness Caused by an Inflammatory Environment. In this study, we employed three different concentrations of TNF-α (10 ng/mL, 20 ng/mL, and 50 ng/mL) to simulate an inflammatory environment. RT-PCR analysis of biomarkers of BMMS stemness (SOX2, C-MYC, and NANOG) showed that mRNA abundances of these biomarkers were significantly lower in the three treated groups than in the control group (P < 0.01) (Figure 2(a)). Based on the relative mRNA abundances of these biomarkers in the treatment groups, 20 ng/mL was selected for further analysis. We also evaluated the effects of melatonin on TNF-α-treated BMMSCs. For colony formation assays, we selected the P3 generation of BMMSCs. The colony formation ratio in the TNF-α-treated group was significantly lower than that in the control (P < 0.01). However, colony formation in the TNF-α-treated group could be partially recovered with melatonin treatment (P < 0.01 vs. TNF-α-treated group) (Figure 2(b)). mRNA abundances of biomarkers of BMMS stemness (HTERT, SOX2, C-MYC, and NANOG), osteogenic differentiation (RUNX2, OPN, and OCN), and adipose differentiation (PPARγ-2, LPL, and ADIPONECTIN) in P3, P7, and P10 cells with different treatments were examined with RT-PCR (Figures 2(c)–2(e)). The results showed that the expression levels of HTERT, SOX2, C-MYC, NANOG, RUNX2, OPN, OCN, PPARγ-2, LPL, and ADIPONECTIN in TNF-α-treated P3, P7, and P10 cells were significantly lower than those in the control group (P < 0.01). However, the expression levels of these genes were recovered with additional melatonin treatment (P < 0.01 vs. TNF-α-treated group). Similar results were found in western blot analysis. Protein abundances of these biomarkers in TNF-α-treated P3, P7, and P10 cells were also significantly lower than those in the control group (P < 0.01). Meanwhile, melatonin treatment could partially recover the protein expression of these genes (P < 0.01) (Figures 2(f)–2(h)). Alizarin red staining was used to investigate osteogenic differentiation of BMMSCs in P3 cells. Quantification revealed that the OD in the TNF-α-treated group was significantly lower than that in the control group (P < 0.01). Melatonin treatment was able to increase mineral deposition in the TNF-α-treated group (P < 0.01 vs. TNF-α-treated group) (Figure 2(i)). Therefore, melatonin was effective at reversing the loss of BMMS stemness caused by inflammatory environment.

3.3. Melatonin Reverses the Inflammatory-Induced Loss of BMMS Stemness via Its Receptor. In this study, we have employed luzindole (a melatonin receptor inhibitor) to study the molecular mechanism of melatonin in BMMS P3 generation. BMMSCs were divided into four groups: control, TNF-α, TNF-α+Mel, and TNF-α+Mel+Luz. For colony formation analyses, colony formation ratio in the TNF-α group was significantly lower than that in the control (P < 0.01). Colony formation ratio in the TNF-α+Mel group was significantly higher than that in the TNF-α group (P < 0.01). However, this value in the TNF-α+Mel+Luz group was significantly lower than that in the TNF-α+Mel group (P < 0.01) (Figure 3(a)). RT-PCR and western blot analyses of BMMS stemness (HTERT, SOX2, C-MYC, and NANOG levels), osteogenic differentiation (RUNX2, OPN, and OCN levels), and adipose differentiation (PPARγ-2, LPL, and ADIPONECTIN levels) in P3 cells with different treatments were examined (Figures 3(b)–3(g)). The results show that the mRNA and protein expression levels in BMMS P3 cells of these molecular mechanisms in the TNF-α+Mel

![Figure 1: HMSC stemness assay. (a) Colony formation assays of HMSCs. Quantitative analysis of colony formation ratio of HMSCs. (b) Western blot analysis of biomarkers of osteogenic differentiation of HMSCs (OPN, OCN, and RUNX2). (c) Western blot analysis of biomarkers of adipose differentiation of HMSCs (PPARγ-2, LPL, and ADIPONECTIN). (e) RT-PCR analysis of biomarkers of HMSC stemness. (f) RT-PCR analysis of biomarkers of osteogenic differentiation of HMSCs. (g) RT-PCR analysis of biomarkers of adipose differentiation of HMSCs. (h) Alizarin red staining and quantification examination of osteogenic differentiation of HMSCs. P3: primary cells expanded to the third generation; P7: primary cells expanded to the seventh generation; P10: primary cells expanded to the tenth generation. *P < 0.05, **P < 0.01.](image)
Figure 2: Continued.
Figure 2: Continued.
primary cells expanded to the seventh generation; P10: primary cells expanded to the tenth generation.

quanti

fi

stemness, osteogenic di

diff

with di

treatments. (d) RT-PCR analysis of biomarkers of HMSC stemness, osteogenic differentiati

dation. BMMSCs were divided into four groups: control, TNF-α, TNF-α+Mel, and TNF-α+Mel+VP. RT-PCR and western blot analyses of YAP expression showed that TNF-α treatment could reverse YAP mRNA and protein expression in TNF-α-treated BMMSCs (P < 0.01) (Figures 4(a) and 4(b)). Moreover, Figure 4(c) shows that YAP protein expression in the TNF-α+Mel+VP group was significantly lower than that in the TNF-α+Mel group in P3, P7, and P10 BMMSCs. However, melatonin treatment could reverse YAP mRNA and protein expression in TNF-α-treated BMMSCs (P < 0.01) (Figures 4(a) and 4(b)). For colony formation analyses, colony formation ratio in the TNF-α+Mel+VP group was significantly lower than that in the TNF-α+Mel-treated group (P < 0.01) (Figure 4(d)).

In addition, a similar situation was observed in western blot and RT-PCR analyses of BMMSC stemness (HTERT, SOX2, C-MYC, and NANOG levels), osteogenic differentiation (RUNX2, OPN, and OCN levels), and adipose differentiation (PPARY-2, LPL, and ADIPONECTIN levels) in P3 cells with different treatments (Figures 4(e)–4(j)). The mRNA expression levels of these genes in the TNF-α+Mel+VP group was significantly lower than those in the TNF-α+Mel group (P < 0.01). In summary, YAP is a potential target of melatonin in reversing the impacts of inflammation on BMMSC stemness.

3.5. BMMSC Treatment in an Osteoporosis Nude Mouse Model. In this study, we constructed an osteoporosis mouse model to study the effects and potential molecular mechanisms of BMMSC treatment. Figure 5(a) shows the results of micro-CT of two-dimensional (2D) and three-dimensional (3D) map of trabecular bone in different groups. The results suggest that construction of the osteoporosis mouse model was successful (via OVX treatment). BMMSC treatment could partially recovery the damage caused by osteoporosis (OVX+P3 BMMSC treatment). TNF-α treatment aggravates osteoporosis. However, melatonin was able to reverse the damage of osteoporosis. VP, one of the YAP inhibitors, was shown to weaken the therapeutic effect of melatonin in osteoporosis. Similar results were also observed in H&E staining analysis (Figure 5(b)). Quantitative analysis of osteoporosis-related indices suggested that the values of bone density, BV/TV, Tb.N, and Tb.Th were significantly different in the six subgroups. For example, the above indices in the OVX group were significantly lower than those in the control group (P < 0.05). Meanwhile, the above indices in the OVX+P3 BMMSC+TNF-α+Mel group were significantly higher than those in the OVX+P3 BMMSC+TNF-α group (P < 0.05). However, VP could reverse the expression in the OVX+P3 BMMSC+TNF-α+Mel+VP group (Figures 5(c)–5(f)). In addition, Tb.Sp and trabecular pattern factor analysis indicated that both indices in different groups were also significantly different in the six subgroups. For example, both indices in the OVX group were significantly higher than those in the control group (P < 0.01).
OVX+P3 BMSC+TNF-α+Mel group were significantly lower than those in the OVX+P3 BMSC+TNF-α group (\( P < 0.01 \)). However, they were upregulated in the OVX+P3 BMSC+TNF-α+Mel+VP group compared with the OVX+P3 BMSC+TNF-α+Mel group (Figures 5(g) and 5(h)).

4. Discussion

Stemness is the ability of stem cells to maintain pluripotent differentiation potential and undergo no directional differentiation [29]. In the early stage of in vitro stem cell culture, the stemness of stem cells can be maintained. However, the stemness of stem cells decreases with an extension of culture time [30]. Meanwhile, when in vitro-cultured stem cells are injected into the body, the stemness of the MSCs is also reduced [31, 32]. Due to the reduction in stemness, MSCs reduce tissue damage mainly through anti-inflammatory effects rather than by differentiating into specific tissue types in tissue damage treatments following myocardial infarction and spinal cord injury [33, 34]. The maintenance of MSC stemness is related to multiple factors, including aging, in vitro generation, and origination. For aging, there are
Figure 4: Continued.
various features correlated with stemness reduction, including adipose tissue accumulation in the bone marrow and reduced ability of osteogenic differentiation [35]. Nicotinamide phosphoribosyltransferase (Nampt) has been shown to affect the balance between MSC adipogenesis and osteogenesis by regulating the activity of the silencing information regulator 1 (sirtl) gene. For in vitro generation and origination, the stemness of MSCs can be maintained in the early and middle generations of in vitro growth. However, stemness can experience a significant reduction once a critical time is exceeded [36, 37]. Meanwhile, stem cells from different tissue sources have significant differences in their abilities to maintain stemness during in vitro culture. For example, the stemness of BM-MSCs can be maintained until the 15th generation in vitro. However, the stemness then decreases with subsequent culture. Only a very small number of pluripotent cells can be maintained until the 20th generation in vitro [38]. Umbilical-derived MSCs showed stable stemness in the early and middle generations of culture; however, markers related to aging, such as CD105 and CD73, were significantly increased after the 21st generation [39]. In our study, with the passage of BM-MSCs, their stemness and osteogenic and adipogenic differentiation abilities gradually decreased with increasing generations. Therefore, it is important to achieve a detailed molecular understanding of the mechanisms underlying the maintenance of stemness in MSCs.

TNF-α was discovered in 1975 [40], and there are two main members of the TNF family, namely, TNF-α and TNF-β. TNF-α is mainly produced by activated mononuclear

**Figure 4:** Melatonin reverses loss of HMSC stemness via YAP gene expression. (a) Western blot analysis of YAP protein in P3, P7, and P10 HMSCs with different treatments. (b) RT-PCR analysis of YAP mRNA in P3, P7, and P10 HMSCs with different treatments. (c) Western blot analysis of YAP protein in P3, P7, and P10 HMSCs with different treatments. (d) Colony formation assays and quantitative analysis of P3 HMSCs with different treatments. (e) Western blot analysis of biomarkers of HMSC stemness of P3 HMSCs with different treatments. (f) Western blot analysis of biomarkers of osteogenic differentiation of P3 HMSCs with different treatments. (g) Western blot analysis of biomarkers of osteogenic differentiation of P3 HMSCs with different treatments. (h) RT-PCR analysis of biomarkers of HMSC stemness of P3 HMSCs with different treatments. (i) RT-PCR analysis of biomarkers of osteogenic differentiation of P3 HMSCs with different treatments. (j) RT-PCR analysis of biomarkers of adipogenic differentiation of P3 HMSCs with different treatments. P3: primary cells expanded to the third generation; P7: primary cells expanded to the seventh generation; P10: primary cells expanded to the tenth generation. *P < 0.05, **P < 0.01.
macrophages and is closely related to osteoporosis, as high TNF-α expression can be detected in cases of osteoporosis [41, 42]. This indicates that TNF-α is an important factor in diseases of bone metabolism. TNF-α is a trimeric inflammatory factor that binds to transmembrane receptors. TNF-α synergizes with a variety of cells to produce a concerted response to inflammatory products, bacterial toxins, and other aggressive stimuli [43]. BMMSCs undergo apoptosis in an inflammatory environment. TNF-α can promote apoptosis indirectly through NO regulation [44]. Meanwhile, TNF-α inhibits the expression of two transcription factors in HMSCs: Ostrix and Runx2. Therefore, TNF-α can reduce
the expression of specific markers of osteoblast differentiation, which ultimately inhibits the differentiation of terminal osteoblasts [45]. Moreover, TNF-α can regulate the expression of BMP-2 and POEM, thereby inhibiting the differentiation of osteoblasts [46]. In addition, TNF-α has a significant effect on the absorption of osteoclast bone [47]. Overall, TNF-α can inhibit bone remodeling and promote bone resorption, which results in a reduction of bone mass. In this study, 20 ng/mL TNF-α could effectively induce an inflammatory environment in BMMSCs.

Treatment with melatonin can significantly improve the viability of MSCs and stimulate angiogenesis, renal cell proliferation, and renal resuscitation [48]. In vitro cell assays have shown that melatonin can induce the production of antioxidant enzymes such as catalase and superoxide dismutase, which can reduce apoptosis of MSCs caused by oxidative stress [49]. Melatonin can also reduce the inflammation of MSCs induced by interleukin-1. Melatonin treatment can increase the mRNA expression of Cu/Zn superoxide dismutase (Cu/ZnSOD) and MnSOD and decrease the expression level of the apoptosis-related Bax gene. The melatonin receptor inhibitor luzindole could inhibit the effects caused by melatonin [50]. In this study, our results show that melatonin treatment in BMMSCs can effectively recover the loss of cell stemness caused by TNF-α treatment, which is in agreement with a previous study. Therefore, melatonin treatment was an effective means to rescue BMMSC stemness.

Maintenance of stemness in MSCs is regulated by a variety of cellular signals, such as BMP signaling and Wnt signaling, which can promote the differentiation of MSCs into osteoblasts. However, a lack of BMP signal or Wnt signal will inhibit the differentiation of MSCs into osteoblasts and promote the differentiation of MSCs into fat cells [51–53]. Platelet-derived growth factor BB (PDGF-BB), hepatocyte growth factor, transforming growth factor beta (TGF-β), and basic fibroblast growth factor (bFGF) signals can promote the differentiation of MSCs into muscle [54–56]. Recent studies have found that Hippo signaling also plays a key role in MSC fate determination. Hippo signaling is highly conserved in animals and mainly functions to regulate stem cell self-renewal, tissue regeneration, and organ size [57, 58]. YAP is a key transcriptional factor that is negatively regulated by the Hippo pathway, which is a conserved pathway that regulates organ size and tumorigenesis [59, 60]. Recent studies have found that YAP can regulate transcription factors that are crucial for bone homeostasis, such as Run-related transcription factor 2 (RUNX2) [61], signal transducer and activator of transcription factor 3 (STAT3) [62], and β-catenin [63]. RUNX2 is a basic transcription factor that stimulates osteogenesis, and YAP can be used as a coactivator of RUNX2 [61]. Pan et al. reported that YAP could stabilize β-catenin and thus increase nuclear β-catenin-mediated osteogenesis [63]. In this study, YAP expression was decreased as TNF-α inhibited BMMSC stemness. However, melatonin was able to restore YAP expression. Our results confirm that melatonin treatment can reverse the loss of BMMSC stemness caused by TNF-α via YAP signaling.

5. Conclusions

In summary, we demonstrated that HMSC stemness decreases with increased generations of in vitro culture. TNF-α could result in the inflammatory-based reduction in HMSC stemness. However, melatonin could reverse the inflammatory reduction of HMSC stemness via the melatonin receptor and YAP signaling. In osteoporotic nude mice, melatonin improved the effects of HMSC treatment. This work provides a detailed understanding of the effects of melatonin treatment of HMSCs.

Abbreviations

<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Full Form</th>
</tr>
</thead>
<tbody>
<tr>
<td>MSCs</td>
<td>Mesenchymal cell</td>
</tr>
<tr>
<td>BM-MSCs</td>
<td>Bone marrow mesenchymal stem cells</td>
</tr>
<tr>
<td>HMSCs</td>
<td>Human bone marrow mesenchymal stem cells</td>
</tr>
<tr>
<td>BMP</td>
<td>Bone morphogenetic protein</td>
</tr>
<tr>
<td>HTERT</td>
<td>Human telomerase reverse transcriptase</td>
</tr>
<tr>
<td>OPN</td>
<td>Osteopontin</td>
</tr>
<tr>
<td>OCN</td>
<td>Osteocalcin</td>
</tr>
<tr>
<td>RUNX2</td>
<td>Recombinant runt-related transcription factor 2</td>
</tr>
<tr>
<td>PPARγ-2</td>
<td>Peroxisome proliferator-activated receptor-γ</td>
</tr>
<tr>
<td>LPL</td>
<td>Lipoprotein lipase</td>
</tr>
<tr>
<td>H&amp;E</td>
<td>Hematoxylin and eosin staining</td>
</tr>
</tbody>
</table>

Data Availability

The datasets analyzed during the current study are available from the corresponding authors on reasonable request.

Ethical Approval

The Research Ethics Committee of the Sun Yat-sen University approved the collection of tissue samples for research.

Conflicts of Interest

All authors declare no conflict of interest.

Authors’ Contributions

DSH and YP designed the experiments. XDW, TZL, JCQ, XJQ, BG, WJG, CJL, TQC, YXZ, AJL, and PQS conducted the experiments. XDW, TZL, and JCQ acquired the data. XDW, TZL, JCQ, DSH, and YP analyzed the data. XDW, DSH, and YP wrote the manuscript. All authors read and approved the final manuscript. Xudong Wang, Tongzhou Liang, and Jincheng Qiu contributed equally to this work.

Acknowledgments

This work was supported by the National Natural Science Foundation of China (No. 81572134) and the Natural Science Foundation of Guangdong Province, China (Nos. 2017A030311008, 2016A030313284, and 2018A030313096).
Supplementary Materials

Supplementary Figure 1: identification of isolated BMMSCs. Flow cytometric analysis of CD34 (a), CD45 (b), CD73 (c), CD90 (d), and CD105 (e). (Supplementary Materials)

References


