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Activated \(NF-kB\) in Bone Marrow Mesenchymal Stem Cells from Systemic Lupus Erythematosus Patients Inhibits Osteogenic Differentiation Through Downregulating Smad Signaling

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Osteoporosis in patients with systemic lupus erythematosus (SLE) is thought to be the result of accelerated osteoclastogenesis induced by pro-inflammatory cytokines such as tumor necrosis factor (TNF). However, the molecular mechanisms involved in the osteoblastogenesis in SLE patients are not fully understood. We investigated the bone morphogenetic protein-2 (BMP-2)-induced osteoblastic capacity of bone marrow-derived mesenchymal stem cells (BMMSCs) from SLE patients and the TNF signaling system in determining BMP-2-induced regulatory pathways. It showed that the capacity of osteogenic differentiation of BMMSCs from SLE patients was reduced compared with that from healthy controls. The nuclear factor \(kappa\ B\) (\(NF-kappa\ B\)) signaling was activated while the BMP/Smad pathway was repressed in BMMSCs from SLE patients. TNF activated NF-\(kappa\ B\) pathway and inhibited the phosphorylation of Smad 1/5/8 and BMP-2-induced osteoblastic differentiation in BMMSCs from normal controls, while addition of pyrollidine dithiocarbamate (PDTC), an NF-\(kappa\ B\) inhibitor, to SLE-BMMSCs could partially reverse these effects. Thus, our findings have shown that the activated NF-\(kappa\ B\) pathway in SLE-BMMSCs inhibits the BMP-2-induced osteoblastic differentiation through BMP/Smad signaling pathway, suggesting that the impaired osteoblastic differentiation may participate in the pathology of osteoporosis in SLE patients.

Introduction

Systemic lupus erythematosus (SLE) is a chronic, multisystemic autoimmune disease that involves multiple organs including renal, cardiovascular, neural, musculoskeletal, and cutaneous systems. Osteoporosis (OP), a metabolic bone disease characterized by decreased bone mineralization, increased bone fragility, and increased risk of fracture, has been found to be prevalent in patients with SLE. Cross-sectional studies have shown an increased prevalence of OP in SLE patients, which is estimated to be between 4% and 22% [1–3]. The mechanisms of OP in SLE patients remain incompletely understood and may include the inflammatory disease itself, disease-related comorbidity, and the treatment of the disease including the corticosteroids (CS). Although exposure to CS is generally considered to be a major factor contributing to the development of OP, the systemic inflammation itself is recognized to be the most important risk factor of OP in SLE patients. Tumor necrosis factor (TNF) is one of the most potent pro-inflammatory cytokines and is known to be a catabolic factor in the inflammatory reaction of diseases such as rheumatoid arthritis (RA) [4]. SLE patients are reported to have increased TNF in its serum [5,6]. TNF induces osteoclastogenesis either by promoting the proliferation of osteoclast precursor cells or by causing the activation of the differentiated osteoclast through RANK/RANKL signaling pathway [7–9]. TNF stimulates osteoclasts to express RANK to interact with its ligand RANKL to accelerate the activity of osteoclasts.

Indeed, in physiologic remodeling, activation of bone resorption requires contact between cells of the osteoblast and osteoclast lineages. Osteoblasts play an essential role in the pathogenesis of OP. Osteoblasts produce RANKL, which activates the differentiation of osteoclasts and maintains their bone resorptive function. Osteoblasts also produce and secrete osteoprotegerin, a decoy receptor that can block RANKL/RANK interactions [10,11]. Since bone mass loss in SLE is believed to be associated with TNF system, it has also been hypothesized that TNF directly controls osteoblast survival and/or its function [12]. However, the role of TNF...
NF-κB inhibits osteogenic differentiation by Smad pathway

Signaling in modulating osteoblast’s function and differentiation remains controversial. Some groups reported that treatment of human mesenchymal stem cells (MSCs) with TNF during differentiation resulted in enhanced expression of osteogenic markers [13,14]. On the contrary, other studies have demonstrated that TNF blocked osteoblast differentiation in multiple model systems such as fetal calvaria, bone marrow (BM) stromal cells, and MC3T3-E1 cells [15–17]. Moreover, both spontaneous and bone morphogenetic protein (BMP)-induced osteoblast differentiation could be inhibited by TNF in vitro [18,19].

Osteoblasts are derived from MSCs, which can differentiate into osteoblasts both in vitro and in vivo [20,21]. We have previously reported that bone marrow-derived mesenchymal stem cells (BMMSCs) from SLE patients showed significantly decreased bone-forming capacity and impaired reconstruction of BM osteoblastic niche in vivo [22]; however, the cellular mechanism underlying this deficiency has not yet been elucidated.

BMPs, members of the transforming growth factor (TGF)-β superfamily, have been shown to play critical roles in governing various aspects of embryological development, including brain, heart, kidney, and eye [23]. BMPs are also known to promote the differentiation of MSCs into chondrocytes and osteoblasts and the differentiation of osteoprogenitor cells into osteoblasts through the Smad signaling transduction pathway via 2 transmembrane serine-threonine kinase receptors, BMP receptor (BMPR) type I and BMPR type II [24]. The activated receptor kinases, in turn, phosphorylate the transcription factors Smad 1, 5, and 8. The phosphorylated Smads then form a heterodimeric complex with Smad 4 in the nucleus and activate the expression of target genes. In this study, we investigated the nuclear factor κB (NF-κB) and BMP/Smad signaling pathways in the BMMSCs from SLE patients. We demonstrated the role of TNF activated NF-κB pathway in regulating osteoblastic differentiation and found that the activated NF-κB pathway of BMMSCs from SLE patients inhibited BMP-2-induced osteogenic differentiation through downregulating Smad signaling pathway. Together, our results have uncovered a novel mechanism that TNF participates in the pathology of OP in SLE patients.

Materials and Methods

Patients and controls

BM cells for cDNA microarray were obtained from 4 patients with SLE, according to the American College of Rheumatology [25]. (Supplementary Table S1; Supplementary Data are available online at www.liebertpub.com/sed). All the patients were female with the mean age of 37 ± 11 years (range 20–44). The normal controls were 1 male and 3 females, with the mean age of 39 ± 7 years (range 29–45). MSCs from 10 SLE patients (mean 37 ± 13 years, range 15–57 years) and 10 age- and sex-matched normal controls were examined for various osteoblastic markers by quantitative reverse transcriptase-polymerase chain reaction (qRT-PCR) analysis. Three to five of these 10 samples mentioned above were used for western blot analysis. All the SLE patients above had active disease with a Systemic Lupus Erythematosus Disease Activity Index (SLEDAI) [26] score of more than 8 at the time of BM aspiration. Serum from 16 patients (15 females and 1 male, mean age of 33 ± 14 years, range 13–57, SLEDAI: 2–21) and 15 female normal controls (mean age of 32 ± 13 years, range 16–57) were tested for BMP-2 levels (Supplementary Table S2). All participants gave written consent to the study, which was approved by the Ethics Committee of the Affiliated Drum Tower Hospital of Nanjing University Medical School [27].

Cell culture and flow cytometry

BM samples were taken from the iliac crest of SLE patients and healthy subjects. The BM mononuclear cells were plated at 10³/mL density in low glucose Dulbecco modified Eagle medium (L-DMEM; Gibco) supplemented with 10% heat inactivated fetal bovine serum (Invitrogen) and 1% antibiotic-antimycotic and incubated at 37°C in a 5% (vol/vol) humidified CO2 chamber as described previously [27]. Markers CD29, CD44, C105, CD34, CD45, and HLA-DR on the cells at passage 2 were consequently detected by flow cytometry [27].

Cells at passage 3–5 were plated on a 6-cm dish (2.5 × 10⁴/mL) and were grown to confluence for 3 days at 37°C in 5% CO2 and then stimulated with osteogenic medium including 10 mM β-glycerophosphate (GP; Sigma), 50 μg/mL ascorbic acid (AA; Sigma), and 300 ng/mL BMP-2 (Miltenyi Biotec) for 2, 14, or 21 days.

RNA extraction and quantitative real-time RT-PCR analysis

BMMSCs from SLE patients were cultured in a 6-well plate (2.0 × 10⁵ viable cells), and treated with indicated concentrations of TNF (R&D), BMP-2, pyrrolidine dithiocarbamate (PDTC), or both of them for indicated times; BMMSCs in the presence or absence of stimulation were placed in TRIZol (Invitrogen) and total cellular RNA was extracted. cDNA was synthesized using PrimerScript RT reagent Kit (TaKaRa). qRT-PCR was performed on an ABI 7500 FAST real-time PCR detection system (Applied Biosystems) using SYBR Green detection mix [25]. The following primers were used in this study: alkaline phosphatase (ALP) Sense 5′-GCACCGTCAAGGTGAGAAC-3′, Antisense 5′-TTGGTGAAGACCGCCA GTGGA-3′; Runx2 Sense 5′, CACTGCCGC TGCAACAAGA-3′, Antisense 5′-CATTCCGAGCTCACCG AG AATAA-3′; osteocalcin (OCN) Sense 5′-CGGTCAGA GT CCAGCAAAAG-3′, Antisense 5′-TACAGGTACCGCCCGT GTCCT-3′; COLIA2 Sense 5′-GAGGCAACAGC AGCTTCA CTAA-3′, Antisense 5′-TCA GCCACCCGATGCA CTTA-3′, and (GAPDH) sense: 5′-TGACTTCAACAGCGACACCCA-3′, antisense: 5′-CACCCTGTIG CTTAG CCAAA-3′.

The expression of the target genes in SLE samples as compared with that in controls was examined using 2⁻ΔΔCt method. Briefly, for each sample, a value for the cycle threshold (Ct) was determined, defined as the mean cycle at which the fluorescence curve reached an arbitrary threshold. The ΔCt for each sample was then calculated according to the formula Ct target gene − Ct GAPDH; ΔΔCt values were then obtained by subtracting the ΔCt of a reference sample (average ΔCt of the control group) from the ΔCt of the studied samples. Finally, the levels of expression of the target genes in the studied samples as compared with the reference sample were calculated as 2⁻ΔΔCt.
cells per ml, were used as a common reference in the 2 channel microarrays. The array was hybridized at 42°C overnight and washed with 2 consecutive washing solutions (0.2% SDS, 2 × SSC at 42°C for 5 min and 0.2% SSC) for 5 min at room temperature. Finally, arrays were scanned with a confocal LuxScan 10 KA scanner (CapitalBio).

The data of obtained images were extracted with LuxScan 3.0 software (CapitalBio). Genes with the signal intensity more than 800 (Cy3 or Cy5) were regarded as the expressed genes. In every 2 channel slides, the intensity ratio of the Cy3 to Cy5 of each spot was calculated after normalization with LOWESS regression. Statistical data and differential analysis files were generated by using SAM software 3.0 (Stanford University). The significantly changed genes were selected based on P-value < 0.05 and > 2-fold as criteria. All the differentially expressed genes were analyzed using a free web-based Molecular Annotation System 2.0 (MAS 2.0, http://bioinfo.capitalbio.com/mas).

All data are MIAME compliant and that the raw data have been deposited in a MIAME compliant database (GEO). The raw data can be seen at www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE21649 and the accession number is GSE 21649.

ALP staining

Fourteen days after stimulation, BMSCs were washed twice with phosphate-buffered saline, fixed with 0.5 mL/well formalin/methanol/H2O (1:1:1.5) for 15 min at room temperature, and washed thrice with distilled water. For staining, 1 FAST5-bromo-4-chloro-3-indolylphosphate/nitro blue tetrazolium tablet (Sigma) was dissolved in 10 mL of water, and 0.5 mL of substrate solution was added to the fixed cultures for 15 min at room temperature. After staining, cultures were washed thrice with distilled water and air-dried.

Alizarin Red S staining for mineralization

Twenty-one days after stimulation, BMSCs were washed twice in phosphate-buffered saline, fixed with 0.5 mL/well formalin/methanol/H2O (1:1:1.5) for 15 min at room temperature, and washed thrice. Saturated Alizarin Red S solution (pH 4.1) was filtered, and 1.5 mL/well was added and incubated for 5 min at room temperature. Cells were then washed 4–5 times and air-dried.

Western blotting analysis

Cells (2 × 10^5 viable cells) were precultured in 6-well plates in DMEM containing 10% fetal calf serum for 48 h. After preculture, the medium was replaced with serum-free fresh medium, and then indicated concentrations of BMP-2, TNF, or PDTC were added to the culture medium for different time intervals. After stimulation, cells were lysed with SDS-sample buffer containing 20 mM Tris-HCl (pH 7.6), 250 mM NaCl, 0.5% NP-40, 3 mM ethylenediaminetetraacetic acid, and 1.5 mM ethyleneglycoltetraacetic acid with 10 mg/mL Aprotinin, 10 mg/mL leupeptin, 1 mM DTT, 1 mM para-nitrophenylphosphate, and 0.1 mM Na2VO4 as protease and phosphatase inhibitor. Cell lysates were separated by sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and transferred to a polyvinylidene difluoride (PVDF) membrane (Millipore). Blots were probed by anti-phospho-Smad1/5/8 antibodies (Cell Signaling Technology, Inc.), anti-total-Smad1/5/8 antibodies (Santa Cruz Biotechnology, Inc.), anti-Smad4 antibodies (Cell Signaling Technology, Inc.), anti-inhibitor kB (IkB) antibody (Cell Signaling Technology, Inc.), and anti-GAPDH antibody (Millipore) before visualizing with horseradish peroxidase-conjugated secondary antibodies followed by development with FluorChem FC2 System (Alpha Innotech Corporation).

Immunoprecipitation analysis

BMSCs from patients and healthy controls were starved in serum-free fresh medium for 24 h and stimulated with BMP-2 for 30 min, then lysed with (SDS)-sample buffer, and diluted to 1 μg/μL. Anti-Smad1/5/8 antibodies were added to 1 mL cell lysates at 4°C overnight. The immunocomplex was captured by adding 100 μL Protein A Agarose, Fast Flow, bead slurry (Millipore). After incubation at 4°C for 2 h, the carrier beads were washed 4 times in buffer. The samples were boiled in Laemmli loading buffer for 10 min, applied to 10% SDS-PAGE gels, and electroblotted onto PVDF western blot membranes (Roche Diagnostics). The Smad4 protein was identified by western blot analysis.

Enzyme-linked immunosorbent assay analysis

Serum samples from 16 SLE patients and 15 healthy controls were collected, and the concentrations of BMP-2 of each individual were measured using commercial ELISA kit (R&D) according to the manufactory introduction.

Statistical analysis

All results are shown as mean ± standard error of the mean of data from at least 3 separate experiments, each performed with triplicate samples. Differences between groups were analyzed for statistical significance using analysis of variance. Differences between 2 independent groups were analyzed by student’s t-test (SPSS 16.0 software). P values < 0.05 were accepted as statistically significant.

Results

Characterization of osteoblastic differentiation in BMSCs from SLE patients

Consistent with our previous findings, BMSCs from both SLE patients and normal controls were positive for CD29,
CD44, and CD105, and negative for CD14, CD34, CD45, and HLA-DR. There was no significant difference between the phenotypic features of MSCs from SLE patients and normal controls [27]. To exclude the effect of senescence on the osteogenic capacity of BMMSCs, we compared the proliferation of BMMSCs during the culturing of osteogenic medium with BMP-2 (OMB) (AA, GP, and BMP-2). The growth curves showed that both the 2 groups rapidly grew from about the 9th day and reached the highest level at the 14th day. Cell amount decreased after 14 days during the osteogenic medium culture. There was no difference between the cell numbers of normal controls (n = 3) and SLE patients (n = 3) at every point (all \( P > 0.05 \); Fig. 1A). And then, we evaluated the osteogenic capacity of BMMSCs from SLE patients induced by

**FIG. 1.** Characterization of osteoblastic differentiation in BMMSCs from SLE patients. (A) The growth curves of BMMSCs during osteogenic differentiation from SLE patients and normal controls. Two groups of cells were cultured in the osteogenic medium with BMP-2 (OMB) for 0 ~ 18 days. (B) Stimulation with different doses of BMP-2 on the BMMSCs. Cells were starved for 24 h, and stimulated for 30 min with different concentrations of BMP-2. The phosphorylated Smad1/5/8 was detected by western blotting analysis. (C) Alkaline phosphatase (ALP) staining of cells for 14 days and Alizarin Red S staining for 21 days after treatment with OMB. *\( P < 0.05 \) and **\( P < 0.01 \) versus control groups. ALP-positive cells are stained as blue, and mineralization is visible as red spots on the photos shown. (D) qRT-PCR analysis of osteoblastic markers after osteogenic stimulus with BMP-2 for 2 days. The relative mRNA levels were analyzed by comparing with normal groups. (E) qRT-PCR analysis of ALP activity on day 14. **\( P < 0.01 \) versus indicated groups. (F) The area of mineralized nodules of BMMSCs from SLE patients and healthy controls. **\( P < 0.01 \) versus normal control groups (n = 3). Nor, normal controls; SLE, systemic lupus erythematosus; BMMSCS, bone marrow-derived mesenchymal stem cells; BMP-2, bone morphogenetic protein-2; qRT-PCR, quantitative reverse transcriptase-polymerase chain reaction.
Osteoblasts (BMMSCs) from SLE patients were treated with different concentrations of BMP-2 to find that BMP-2 activated the phosphorylation of Smad1/5/8 in a dose-dependent manner, so we selected 100 ng/mL BMP-2 in the following experiments. Before stimulation, BMMSCs from both SLE and control subjects showed a low expression of ALP, Runx2, OCN, and collagen II A2 (ColIIA2) at the transcriptional level. In response to BMP-2 treatment for 2 days, BMMSCs from both groups showed a significant increase in mRNA levels of ALP, ColIIA2, Runx2, and OCN. Notably, BMMSCs from SLE patients showed lower expression of ALP, ColIIA2, and Runx2 post-stimulation compared to normal controls, while no statistical difference could be found in OCN mRNA level between these 2 groups. In addition, after a longer exposure to osteogenic stimulus, the ALP activity and area of the mineralized nodules were lower in SLE-BMMSCs as detected by ALP and Alizarin Red S staining on day 14 and 21, respectively. These results suggested that BMMSCs from SLE patients display impaired osteoblastic differentiation.

Repressed BMP/Smad and activated NF-κB signaling pathways in BMMSCs from SLE patients

As many signaling pathways like Wnt, Notch, fibroblast growth factors, and BMP/Smad pathways [29–32] are involved in the osteoblastic differentiation of MSCs, we compared the gene expression profiles of BMMSCs between SLE patients and normal controls, and found that all the differentially expressed genes in BMP/TGF-β signaling pathway except for BMP5 were downregulated in the patients’ cells. Western blot analysis revealed that the Smad1/5/8 phosphorylation was higher in normal BMMSCs 30 min after BMP-2 stimulation. Moreover, the complex of BMP-2 and Alizarin Red S staining on day 14 and 21, respectively. These results suggested that BMMSCs from SLE patients display impaired osteoblastic differentiation.

Activation of NF-κB inhibited BMP-2-induced expression of osteogenic markers in BMMSCs from normal controls

Next, we investigated the effect of TNF on the phosphorylation of Smad1/5/8 induced by BMP-2 in a dose-dependent manner. TNF (20 ng/mL) alone effectively activated the NF-κB pathway as detected by the decreased IkB, but it had no effect on the activation of Smad1/5/8; nevertheless, it could inhibit the Smad1/5/8 phosphorylation induced by BMP-2. To further observe whether TNF inhibited the Smad1/5/8 phosphorylation through NF-κB pathway, PDTC, an NF-κB inhibitor was added to the BMP-2-TNF system. As shown in Fig. 3C, PDTC, at the concentration of 50 and 100 nM, reversed the inactivation of Smad1/5/8 caused by TNF. These data collectively indicate that the activation of NF-κB inhibits the BMP/Smad signaling pathway.

For further investigation of the mechanism by which NF-κB affects the BMP-2-induced osteoblastic differentiation, we examined the mRNA levels of ALP and Runx2 and the extracellular mineralization in normal BMMSCs treated with BMP-2 and/or TNF. Consistent with the western blot results, TNF inhibited ALP activity and Runx2 mRNA expression induced by BMP-2 for 48 h. Moreover, TNF inhibited BMP-2-induced extracellular mineralization on day 21.

PDT restored BMP-2-induced osteoblastic differentiation of BMMSCs from SLE patients

Next, we investigated whether inhibiting the activation of NF-κB could restore the osteoblastic differentiation of BMMSCs in SLE patients. Although PDTC alone had no effect on the phosphorylation of Smad1/5/8, it increased the phosphorylation of Smad1/5/8 induced by BMP-2 in the absence of exogenous TNF. Similarly, PDTC enhanced the levels of Runx2 and ALP mRNA treated with BMP-2 for 48 h. After exposure to osteogenic stimulus for 21 days, PDTC increased the mineralization of BMMSCs from SLE patients, although it did not restore it to the level of normal cells.

The circulating levels of BMP-2 in patients with SLE

BMP-2 is secreted by osteoblasts to bone matrix and also is one of the markers of osteoblastogenesis bone formation. Although we hardly detected BMP-2 in the BM of SLE patients, the level of BMP-2 in the serum of the patients (n = 16) was lower than normal controls (n = 15), suggesting a decreased osteoblastogenesis in SLE patients.

Discussion

BMMSCs from SLE patients demonstrated early signs of senescence [34,35]. SLE-BMMSCs were also observed to grow slower than the normal controls in ordinary cultural medium [27,34]. However, in this study, we did not find the difference in the proliferation rates between the 2 groups in the OMB. Different from the ordinary cultural medium, the numbers of the BMMSCs in OMB decreased after about 14 days, suggesting an enhanced trend to differentiation. The growth curve excluded the effect of senescence on the osteoblastic differentiation.

Both patients and normal BMMSCs cultured in ordinary medium showed extremely low levels of all the osteogenic markers, confirming that BMMSCs, even for a long period of
FIG. 2. Repressed BMP/Smad and activated NF-κB signaling pathways in BMMSCs from SLE patients. (A) Clustering analysis of genes in BMP/TGF-β signaling pathway in BMMSCs from SLE patients and healthy controls. (B) Western blotting analysis of phosphorylated Smad1/5/8 of BMMSCs from SLE patients and healthy controls. Cells were starved (the left) and incubated with BMP-2 (100 ng/mL) for 30 min (the right). (C) Immunoprecipitation (IP) analysis of BMMSCs from SLE patients and healthy controls. (D) Western blotting analysis of P IKKβ, t IKKβ, and IκB in BMMSCs from SLE patients and healthy controls. (E–H) Quantity analysis of western blotting analysis. *P < 0.05 and **P < 0.01 versus normal control groups. Nor, normal controls; SLE, systemic lupus erythematosus; Nor+BMP-2, normal BMMSCs stimulated with BMP-2; SLE+BMP-2, SLE-BMMSCs stimulated with BMP-2; TGF-β, transforming growth factor-β; NF-κB, nuclear factor κB; IκB, inhibitor κB; IKKβ, inhibitor κB kinase β.
FIG. 3. Effect of activating NF-κB signaling pathway on BMP-2-induced osteogenic markers in normal BMMSCs. (A) Effect of different concentrations of TNF on the Smad1/5/8 phosphorylation in BMMSCs revealed by western blotting analysis. BMMSCs were starved for 24 h and subsequently stimulated with TNF for 2 h and BMP-2 for 30 min. (B) Effect of TNF (20 ng/mL) on the levels of IκB and phosphorylation of Smad1/5/8 induced by BMP-2. (C) Effect of PDTC on the Smad1/5/8 phosphorylation in the presence of BMP-2 (100 ng/mL) and TNF (20 ng/mL). Starved cells were treated with TNF and PDTC for 2 h, followed by BMP-2 for 30 min. (D) qRT-PCR analysis of ALP and Runx2 mRNA levels. Cells were treated with TNF or BMP-2 for 48 h. *P < 0.05 and **P < 0.01 versus controls or between the indicated groups (n = 4). (E) Alizarin Red S staining of cells cultured in the OMB for 21 days in the presence or absence of TNF. (F) The quantity analysis of mineralized nodules of normal BMMSCs in the OMB for 21 days in the presence or absence of TNF. *P < 0.05 versus normal control groups (n = 3). TNF, tumor necrosis factor; PDTC, pyrroolidine dithiocarbamate.

FIG. 4. Effect of PDTC on BMP-2 induced osteogenic differentiation in SLE-BMMSCs. (A) Western blotting analysis of phosphorylated Smad1/5/8 stimulated with PDTC (100 nM) and/or BMP-2 (100 ng/mL). Cells were starved for 24 h and treated with PDTC for 2 h, followed by BMP-2 for 30 min. (B) qRT-PCR analysis of ALP and Runx2 mRNA levels. Cells were treated with PDTC and/or BMP-2 for 48 h. *P < 0.05 and **P < 0.01 versus controls or between the indicated groups (n = 3). (C) Alizarin Red S staining of SLE-BMMSCs cultured in the OMB for 21 days in the presence or absence of PDTC. (D) Quantity analysis of mineralization area. *P < 0.05 and **P < 0.01 versus normals or between the indicated groups (n = 3). Nor, normal controls; SLE, systemic lupus erythematosus.
time in culture, cannot differentiate into osteoblasts without osteogenic stimulation [36]. After stimulating with BMP-2 for 2 days, the mRNA levels of ALP, CollIA2, OCN, and Runx2 in BMMSCs were increased. Besides, the mRNA levels of aforementioned genes in BMMSCs from SLE patients were lower than that from normal controls except for OCN. The ALP activity (at day 14) and the mineralized level (at day 21) were also lower in SLE patients compared with normal subjects. All the results mentioned above indicated there was a defective capacity of SLE-BMMSCs to differentiate into osteoblasts. OCN, one of the later markers of osteoblast differentiation compared with ALP and Runx2, was reported to be induced at the third day on MC3T3 cells [37] or at the sixth day on RD-C6 cells [38] induced by BMP-2. Therefore, the difference in OCN expression between BMMSCs from SLE patients and normal controls may not be detected at day 2 of culture.

Our results differ from findings by Nie et al., which suggested a similar ability of MSCs differentiating into osteoblasts between SLE patients and healthy subjects [34]. The underlying reasons responsible for this discrepancy remains unclear, but several possibilities might exist: since MSCs in vitro have been recognized to represent heterogeneous group of progenitors with different self-renewal properties [39], MSCs of different passages may have distinct capacity of differentiation [40]; moreover [41], the osteoblastic differentiation may change with the alternations of cytokines in the osteoblastic medium [42]; some medications, such as glucocorticoid (GC) [43,44], methotrexate, and warfarin may affect the osteoblastic capacity of MSCs [45], at last, in the study of Nie et al. [34] the quantitative analysis of Alizarin Red S staining had not been measured.

In this study, cDNA microarray analysis revealed 19 differentially expressed genes in BMP/TGF-β signaling pathway. Among them, 18 genes were downregulated, including Smad1, Smad5, BMPRIA, and Id1 (inhibitor of differentiation or inhibitor of DNA binding-1). Moreover, the lower level of Smad1/5/8 phosphorylation and the decreased complex of Smad1/5/8 with Smad4 confirmed the decreased BMP/smad signaling pathway in SLE-BMMSCs.

Several possible mechanisms may be responsible for the reduction of Smad1/5/8 phosphorylation: First, the inactive status of BMPR (BMPR I and BMPR II) because of the weak stimulation of BMPs in BM or the intrinsic defect of the receptors; second, the regulation of cytokines or growth factors that directly inhibits the Smad1/5/8 phosphorylation. We found in this study that the protein level of BMPRI-IA did not change as analyzed by western blot (Supplementary Fig. S1), although we found the level of BMP-2 in the serum of SLE patients was lower than normal controls. It has been shown that many BMPs, including BMP2, BMP6, BMP7, and BMP9 may induce osteoblast lineage-specific differentiation of MSCs [46–48], and the total amount of BMP activity is more important than the activity of a specific BMP [49]; moreover, other BMPs may be able to compensate for the loss of one BMP [50]. Therefore, with the normal BMPRI-IA, the total amount of BMPs in SLE-BMMSCs might remain unchanged. Thus, our data suggest that the change in BMP/Smad signaling pathway likely results from the function of other cytokines or growth factors in the BM.

TNF is a potent inflammatory cytokine that contributes to local and systemic bone loss in inflammatory bone diseases such as RA, periodontitis, and multiple myeloma, and in estrogen deficiency [51–54]. It is now well known that TNF can induce osteoclastogenesis both in vivo and in vitro. However, the effect of TNF on the differentiation or maturation of osteoblast and their mechanisms remain unresolved: TNF may enhance the osteoblastic differentiation through NF-κB pathway [13,14], or it activates SAPK/JNK, ERK1/2, and Ras/Rho-MAPK signaling pathway to inhibit the spontaneous or BMP-2-induced osteoblast differentiation [18,19]. We found here that TNF inhibited BMP-2 induced osteoblastic differentiation of BMMSCs from normal controls. We further found that TNF inhibited the osteoblastic differentiation of BMMSCs by inactivating Smad1/5/8, and NF-κB inhibition restored the activation of Smad1/5/8. These data collectively indicate that the inhibitory effect of TNF is attributed to, at least in great part, the activation of NF-κB.

In the current study, we found that the NF-κB activity was increased while the BMP signaling induced by BMP-2 was inactive in BMMSCs from SLE patients. Importantly, NF-κB inhibitor, PDTC, restored the activation of BMP signaling and subsequent osteoblastic markers and mineralization of BMMSCs from SLE patients. Similar results were obtained in the addition of another NF-κB inhibitor, SN50 (Supplementary Fig. S2). It is interesting that PDTC failed to repair the osteogenic capacity of SLE-BMMSCs stimulated with dexamethasone (Dex), GP, and AA (Supplementary Fig. S3). These results suggest that an enhanced NF-κB activity by high level of TNF suppressed the BMP/Smad, rather than other osteogenesis-related pathways in SLE patients.

GC is used extensively for the treatment of SLE. In this study, we cannot eliminate the effect of GC received by these patients. It is reported that GC alters bone metabolism at the cellular and molecular levels not only by increasing osteoclastic action [55] but also by inhibiting osteoblastic growth and differentiation [56,57]. Dex suppressed the BMP-2 induced Smad1/5/8 activation in mouse myoblastic C2C12 cells in vitro [58]. We show here in the Supplementary Data that Dex, at the concentration of 0.1 and 1μM, reduced the phosphorylation of Smad1/5/8 induced by BMP-2, but it had no any effect on the NF-κB activation (Supplementary Fig. S4). Therefore, theoretically, the NF-κB inhibitory
treatment cannot restore the inactivation of pSmad1/5/8 induced by GC. From above, we indicate that in SLE patients, with the treatment of GC, the suppressed BMP signaling pathway is at least partially attributed to the activation of NF-κB.

The inhibitory mechanism of NF-κB to Smad1/5/8 is not fully elucidated. Inconsistent with our results, Yamazaki et al. did not observe an inhibitory effect of TNF activated NF-κB on the phosphorylation of Smad1, Smad5, and Smad8 or on the nuclear translocation of the Smad1-Smad4 complex. NF-κB inhibited Smad pathway by interfering with the DNA binding of Smad proteins to its target gene in MC3T3-E1 [33]. Although we could not explain this discrepancy, the discrepancy might depend on the stage of the osteoblast differentiation, the type of the cell, and the methods of the experiments and needed further investigation.

In conclusion, we have demonstrated an activated NF-κB activity and NF-κB inhibited BMP/Smad signaling pathways in BMMSCs derived from SLE patients. The activated NF-κB pathway inhibits BMP-2-induced osteogenic differentiation through downregulating Smad signaling. Thus, our results may suggest a new approach to increasing osteoblastic differentiation for the treatment of OP in SLE patients.

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Author Disclosure Statement

No competing financial interests exist.

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