I-BET151 inhibits osteoclastogenesis via the RANKL signaling pathway in RAW264.7 macrophages

JING CHENG, JIFU ZHENG, NINGHONG GUO and FUMING ZI

Department of Hematology, The Second Affiliated Hospital of Nanchang University, Nanchang, Jiangxi 330006, P.R. China

Received November 10, 2016; Accepted July 12, 2017

DOI: 10.3892/mmr.2017.7631

Abstract. Excessive bone resorption mediated by osteoclasts may lead to the risk of various lytic bone diseases. In the present study, the effects of I-BET151, a bromodomain and extra terminal domain protein inhibitor, on osteoclastogenesis in RAW264.7 cells and the underlying mechanism of this process was investigated. Cells were divided into 6 groups, including the control group, receptor activator of nuclear factor-κB ligand (RANKL) group and 4 other groups containing RANKL and I-BET151 at different concentrations. Tartrate-resistant acid phosphatase (TRACP) staining was used to observe the effect of I-BET151 on osteoclastogenesis and the number of TRACP positive multinucleated cells was calculated. Western blotting was used to evaluate the expression of tumor necrosis factor receptor associated factor (TRAF6), nuclear factor of activated T-cells cytoplasmic 1 (NFATc1), transcription factor p65 (p65), nuclear factor of κB inhibitor-α (IkB-α), extracellular signal-regulated kinase, Jun N-terminal kinase (JNK) and p38. mRNA expression levels of osteoclast-specific genes TRACP, matrix metalloproteinase-9 (MMP9), cathepsin K (CtsK) and proto-oncogene tyrosine-protein kinase Src (c-Src) were measured using the reverse transcription-quantitative polymerase chain reaction (RT-qPCR). TRACP staining results demonstrated that I-BET151 inhibited osteoclastogenesis induced by RANKL and the inhibition was dose dependent. TRACP multinucleated positive cells were significantly decreased when treated with I-BET151 compared with the RANKL group. The inhibitory effect on TRAF6 was significant when concentrations of 100 and 200 nM I-BET151 were used, and NFATc1 was significantly inhibited when a concentration of 200 nM was used compared with the RANKL group, in a dose-dependent manner. Nuclear translocation of p65 was significantly inhibited by I-BET151 at all concentrations. The degradation of IkB-α, and phosphorylation of JNK and p38 were also significantly inhibited by I-BET151, with the exception of the expression of IkB-α following treatment with 50 nM I-BET151. The RT-qPCR results revealed that osteoclast-specific genes TRACP, MMP9, CtsK and c-Src were all dose-dependently inhibited by I-BET151, except for CtsK. In conclusion, I-BET151 may significantly suppress the osteoclastogenesis of RAW264.7 cells via the RANKL signaling pathway.

Introduction

Bone formation and resorption are mediated by osteoblasts and osteoclasts, respectively, and regulate the balance of normal bone metabolism. Interruption of this balance may result in increased resorption compared with formation and lead to excess bone loss, causing a variety of diseases, including myeloma bone disease, osteoporosis and rheumatoid arthritis (1,2). As isolating and culturing bone cells is difficult, research and molecular analysis of osteoclastogenesis has stagnated for a long period. However, developments in techniques over the past decade have allowed for further investigation (3,4). There are multiple biomolecules involved in the signaling pathways of osteoclastogenesis. Among these, the most important factors are receptor activator for nuclear factor-κB ligand (RANKL) and macrophage colony-stimulating factor (M-CSF).

Research has demonstrated that RANKL and its receptor, RANK, are necessary for osteoclastogenesis (5). These proteins activate the reconstitution of dynamic differentiation processes, including cell fusion (5,6). In addition, the proteins are considered to be a possible factor involved in controlling the differentiation process. RANKL may also activate the expression of transcriptional factors such as c-Fos, microphthalmia-associated transcription factor and nuclear factor of activated T-cells cytoplasmatic 1 (NFATc1), which are important for osteoclastogenesis (7). Therefore, the RANKL-RANK signaling pathway is a key signaling component involved in osteoclastogenesis.

RANK mediates signaling by recruiting adaptor molecules, including proteins of the tumor necrosis factor (TNF) receptor-associated factor (TRAF) family (8,9). Studies have revealed that TRAF1 binds to RANK (10,11). TRAF6 also binds to RANK, which further induces trimerization of TRAF6 and subsequently activates nuclear factor (NF)-κB and...
mitogen-activated kinases (MAPKs) (12,13). NF-κB is associated with the biomolecular progress of osteoclastogenesis induced by RANK and is a basic component of osteoclastogenesis activated by TRAF6 (14).

M-CSF is another essential cytokine involved in osteoclastogenesis besides RANKL (15,16). Research has demonstrated that M-CSF serves an important role in the proliferation and survival of osteoclast precursor cells (17), upregulates the expression levels of RANK, and may participate in the progression of differentiation by activating c-Fos, protein kinase B and extracellular signal-regulated kinase (ERK) pathways, which may interact with RANKL signals (18,19).

Studies have demonstrated that bromodomain and extra terminal domain proteins (BET) serve an important role in different types of cancer, including cancer of the bones (20,21). I-BET151 is a quinoline class of BET protein inhibitors, which has been demonstrated to exhibit activity against several types of cancer (22). Studies revealed that I-BET151 induced B-cell lymphoma like 11-dependent apoptosis and cell cycle arrest of human melanoma cells (23), and suppressed expression of inflammatory genes and matrix degrading enzymes in rheumatoid arthritis synovial fibroblasts (24). Research has also demonstrated that I-BET151 suppresses pathologic bone loss in TNF-induced inflammatory osteolysis (21). Despite numerous reports of I-BET151, few have focused on the role of I-BET151 in osteoclastogenesis. In the present study, the effects of I-BET151 on osteoclastogenesis and the underlying molecular signaling pathways involved were investigated.

Materials and methods

Cell culture. A mouse macrophage cell line, RAW264.7 (TIB-71™), was purchased from American Type Culture Collection (Manassas, VA, USA). RAW264.7 cells were cultured in Dulbecco’s modified Eagle’s medium (Thermo Fisher Scientific, Inc., Waltham, MA, USA) supplemented with 10% fetal bovine serum (Gibco; Thermo Fisher Scientific, Inc.) and 100 μg/ml penicillin-streptomycin (Sigma-Aldrich; Merck KGaA, Darmstadt, Germany). RANKL (PeproTech, Inc., Rocky Hill, NJ, USA) was diluted to 100 ng/ml in aquae sterilisata. I-BET151 (MedChemExpress, Monmouth Junction, NJ, USA) was dissolved in dimethyl sulfoxide at a concentration of <0.1% (Sigma-Aldrich; Merck KGaA). RAW264.7 cells were seeded in 6-well plates (2x10^5 cells/well) in a humidified incubator at 5% CO_2 and 37°C, supplemented with 100 ng/ml RANKL, and treated with different concentrations of I-BET151 (25). Cells were cultured for 7 days to induce osteoclast differentiation. The study was divided into 6 groups: Control; RANKL (100 ng/ml); RANKL (100 ng/ml) and I-BET151 (50 nM); RANKL (100 ng/ml) and I-BET151 (100 nM); RANKL (100 ng/ml) and I-BET151 (200 nM); and RANKL (100 ng/ml) and I-BET151 (400 nM).

TRACP staining. TRACP staining was used to determine the effect of I-BET151 on osteoclastogenesis and the percentage of TRACP positive multinucleated cells was calculated. Cells were fixed using freshly made, refrigerated, 3% paraformaldehyde (PFA; Sigma-Aldrich; Merck KGaA) and 2% sucrose in phosphate-buffered saline (PBS; Sigma-Aldrich, Merck KGaA) for 10 min and stained for TRACP using a tartrate-resistant acid phosphatase stain kit according to manufacturer’s protocol (Nanjing Jiangcheng Bioengineering Institute, Nanjing, China), after 7 days culture in the presence of RANKL and different concentrations of I-BET151. TRACP positive multinucleated cells (>3 nuclei) were counted under 8 fields of view for each sample and regarded as osteoclasts (26). Cells were observed using a Zeiss Axio Observer D1 microscope with x100 magnification and images were captured and analyzed with Zeiss ZEN software version 2012 (Zeiss GmbH, Jena, Germany).

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR). RAW264.7 cells (2x10^5) were incubated with RANKL (100 ng/ml) and different concentrations of I-BET151 for 4 days. RT-qPCR was employed to determine the expression levels of osteoclastic-specific marker genes and GAPDH was used as a control. RNA extraction and reverse-transcription were performed as described previously (27). Total RNA was extracted using TRIzol reagent (Invitrogen; Thermo Fisher Scientific, Inc.) according to the manufacturer’s protocol. A High Capacity cDNA Reverse Transcription kit (Applied Biosystems; Thermo Fisher Scientific, Inc.) was used to convert RNA into cDNA. qPCR was conducted using a ABI 7500 real-time PCR instrument (Applied Biosystems; Thermo Fisher Scientific, Inc.) with SYBR Premix Ex Taq (Takara Biotechnology Co., Ltd., Dalian, China). Primers used in PCR were as follows: Primers for TRACP were: Forward, 5’-ACA CAGTGTACGTGTGTCGGCAACCT-3’ and reverse, 5’-CCA GAGCCTTCACATATAGTGG-3’; Primers for MMP9 were: Forward, 5’-AGTTTGTTGTCGCGAGGAC-3’ and reverse, 5’-TACATGAGCCCTCGCGAC-3’; Primers for CtsK were: Forward, 5’-GGGCAACTCAAGAGAAAAC-3’ and reverse, 5’-TCTTTGATGCCATCTGTTT-3’; Primers for TRAP were: Forward, 5’-CCAGCTTGGAGTGTGACT-3’ and reverse, 5’-CAGCTTGGCTATGTGAGT-3’; Primers for GAPDH were: Forward, 5’-AACCTTGAGCTTGGAAGGG-3’ and reverse, 5’-ACACATTGGAGGTAGAACA-3’. DNA was denatured at 94°C for 10 min, followed by initial denaturation with 30 cycles at 94°C for 1 min, 60°C for 1 min and 72°C for 2 min, and finally ended up with an extension step at 72°C for 5 min. Relative quantification of RT-qPCR product was performed using the comparative 2^{-ΔΔCt} method (28).

Western blotting. Western blotting was used to determine the expression levels of TRAF6, NFATo1, and the influence of I-BET151 on the NF-κB signaling pathway (p65 and IκB-α) and MAPK signaling pathway [ERK, Jun N-terminal kinase (JNK) and p38]. A Nuclear Extraction kit was purchased from Cayman Chemical Company (Ann Arbor, MI, USA) to use for measurement of p65. β-actin was used as a loading control. Samples extracted from the cells as previously described (29) were loaded on 10% SDS-PAGE, prior to transfer onto polyvinylidene difluoride membranes. Membranes were blocked with 5% non-fat milk in TBS containing 0.1% Tween-20 for 2 h at room temperature. Subsequently, membranes were probed with primary antibodies at 4°C overnight and a horseradish peroxidase (HRP) conjugated secondary antibody for 2 h at room temperature (Cell Signaling Technology, Inc., Danvers, MA, USA). The membranes were incubated with an Enhanced Chemiluminescence kit.
kit (Amersham Pharmacia Biotech AB, Uppsala, Sweden) and were exposed to X-ray film. The films were scanned and proteins were quantified using Quantity One software version 4.2.1 (Bio-Rad Laboratories, Inc., Hercules, CA, USA). Antibodies used in western blotting were as follows: Anti-TRAF6 (ab33915; 1:1,000; Abcam, Cambridge, UK); NFATC1 antibody (MA3-024; 1:2,000; Thermo Fisher Scientific, Inc.); anti-NF-κB p65 (ab16502; 1:1,000); anti-IκB-α (E130; ab32518; 1:1,000); anti-ERK1+ERK2 (ab17942; 1:1,000); anti-ERK 1/2 (phospho-Thr202/Tyr204; ab214362; 1:1,000); anti-JNK1+JNK2+JNK3 (ab179461; 1:1,000); anti-JNK1+JNK2+JNK3 (phospho T183+T183+T221; ab124956; 1:1,000); anti-p38 (ab31828; 1:1,000); anti-p38 (phospho Y182; ab47363; 1:1,000); anti-β-actin (ab8226; 1:1,000); and anti-mouse IgG VeriBlot for IP secondary (HRP-conjugated; ab131368; 1:2,000) (all from Abcam).

Statistical analysis. Data are expressed as the mean ± standard deviation. Independent continuous variables were compared using a Student’s t-test for comparison of two groups or a one-way analysis of variance followed by Tukey post hoc test for multiple comparison. P<0.05 was considered to indicate a statistically significant difference. All calculations were made using SPSS software version 18.0 (SPSS, Inc., Chicago, IL, USA).

Results

TRACP staining. TRACP staining was used to determine the effect of I-BET151 on osteoclastogenesis and the percentage of TRACP multinucleated positive cells was calculated. Results revealed that in the RANKL group, multiple cell fusion increased the cell volumes and cell synapse, and resulted in multinucleated cells, which was inhibited by I-BET151 (Fig. 1). In the control group, osteoclastogenesis proceeded normally; however, this was enhanced in the RANKL group. When treated with I-BET151, it was apparent that osteoclastogenesis induced by RANKL was dose-dependently inhibited (Fig. 1). Fig. 1B demonstrates the percentage TRACP positive cells. Compared with the control group, the percentage of TRACP positive cells in groups 2-5 was increased (Fig. 1). However, the percentage of TRACP positive cells was almost completely abolished following treatment with 400 nM I-BET151. The percentage of TRACP positive cells in all groups treated with I-BET151 was significantly decreased compared with the RANKL group (P<0.001; Fig. 1) and the inhibitory effect was dose dependent.

Effect of I-BET151 on the RANKL signaling pathway. The expression levels of TRAF6 and NFATC1, which are upstream and downstream effectors of the RANKL signaling pathway, respectively, were measured by western blotting (Fig. 2). TRAF6 and NFATC1 were dose-dependently inhibited by I-BET151. Densitometric analysis of the TRAF6/β-actin ratio in the I-BET151 100 and 200 nM groups were significantly decreased compared with the RANKL group, whereas the NFATC1/β-actin ratio was only significantly decreased compared with the RANKL group following treatment with 200 nM I-BET151. TRAF6/β-actin and NFATC1/β-actin in the 50 nM I-BET151 group demonstrated no significant difference compared with the RANKL group. The NFATC1/β-actin ratio and TRAF6/β-actin ratio in the RANKL group was significant increased compared with the control group. The inhibition effect of both TRAF6/β-actin and NFATC1/β-actin by I-BET151 were dose-dependent.

Effect of I-BET151 on the NF-κB and MAPK signaling pathways. The expression levels of p65 and IκB-α (involved in the NF-κB signaling pathway) and ERK, JNK and p38 (involved in the MAPK signaling pathway) were evaluated by western blotting. Results revealed that nuclear expression of p65 was significantly inhibited by I-BET151 (Fig. 3A and B) at concentrations of 50, 100 and 200 nM compared with the RANKL group, and the effect increased as the dose increased. In addition, under concentrations of 100 and 200 nM I-BET151, IκB-α was significantly inhibited compared with cells treated with RANKL alone (Fig. 3A and C). However, expression of IkB-α after treatment with 50 nM I-BET151 did not demonstrate a significant difference compared with the RANKL group (Fig. 3C). Phosphorylation of JNK and p38 was significantly inhibited by 50, 100 and 200 nM I-BET151 compared with the RANKL group (Fig. 3D, E and F). By contrast, only 50 nM I-BET151 markedly inhibited the expression levels of TRAF6 and NFATC1, which are upstream and downstream effectors of the RANKL signaling pathway, respectively.
Figure 2. (A) Western blotting of TRAF6 and NFATc1. (B) Quantified expression levels of TRAF6 and NFATc1 protein in RAW264.7 cells treated with or without RANKL, and with or without I-BET151 at 200, 100 or 50 nM. *P<0.05 vs. control group; **P<0.01 vs. RANKL group. TRAF6, tumor necrosis factor receptor-associated factor 6; RANKL, receptor activator of nuclear factor-κB ligand; NFATc1, nuclear factor of activated T-cells cytoplasmic 1.

Figure 3. (A) Western blotting of p65 and IκB-α. RAW264.7 cells treated with or without RANKL, and with or without I-BET151 at 200, 100 or 50 nM. Densitometric analysis of (B) nuclear/cytoplasmic p65 and (C) IκB-α. (D) Western blotting of total or phosphorylated ERK, JNK and p38. Densitometric analysis of (E) p-JNK/JNK, (F) p-p38/p38 and (G) p-ERK/ERK. *P<0.05, **P<0.01 vs. control group; *P<0.05, **P<0.01 vs. RANKL group. p65, nuclear factor of κ light polypeptide gene enhancer in B-cells inhibitor-α; IκB-α, nuclear factor of κB alpha; ERK, extracellular signal-regulated kinase; JNK, Jun N-terminal kinase; p-ERK, phosphorylated extracellular signal-regulated kinase; RANKL, receptor activator of nuclear factor-κB ligand; p-ERK, phosphorylated extracellular signal-regulated kinase.
of p-ERK compared with the RANKL group; however, no significant difference was observed following treatment with I-BET151 (Fig. 3D and G).

Effect of I-BET151 on the expression of osteoclast-specific genes. The mRNA expression levels of the genes encoding osteoclast-specific proteins TRACP, matrix metalloproteinase-9 (MMP9), cathepsin K (CtsK) and c-Src were measured using RT-qPCR. The results revealed that all genes were dose-dependently inhibited by I-BET151 compared with the RANKL group. With the exception of CtsK, the inhibitory effects on TRACP, MMP9, and c-Src at all concentrations of I-BET151 were statistically significant compared with the RANKL group (Fig. 4). For Ctsk, only 200 nM I-BET151 led to significant inhibition in expression compared with the RANKL group.

Discussion

A delicate balance exists in the process of bone remodeling between bone formation and resorption (30). Excessive bone resorption may lead to over activation of osteoclasts and be a risk of various lytic bone diseases, including rheumatoid arthritis, psoriatic arthritis and osteoporosis (31). RANKL serves a role in the process of osteoclastogenesis. In the RANKL signaling pathway, a number of biomolecules are involved and have their own role, including TRAF6, NF-kB and MAPKs. NF-kB is essential for the initial induction or autoamplification of NFATc1, which may activate other osteoclast-specific genes including TRACP, CtsK, calcitonin receptor and MMP9. Although there are various studies focusing on I-BET151, few report the effects of BET on the inhibition of osteoclastogenesis. Park-Min et al (21) reported that I-BET151 suppresses pathologic bone loss in TNF-induced inflammatory osteolysis and suppresses osteoclastogenesis; however, the detailed effects of I-BET151 on the RANKL signaling pathway, and its associated proteins and genes involved in this process, remain to be identified. In the present study, the effects of I-BET151 on osteoclastogenesis were investigated by determining the expression levels of TRAF6 and NFATc1 that are involved in RANKL pathway, and the effects of I-BET151 on osteoclast-specific gene expression.

According to the TRACP staining results, I-BET151 inhibited RANKL-induced osteoclastogenesis and the inhibition was demonstrated to be dose dependent. Also, the percentage of TRACP multinucleated-positive cells was reduced when treated with I-BET151, and the effect was significant compared with the RANKL group. Expression of TRAF6 and NFATc1, which are upstream and downstream effectors of the RANKL pathway were dose-dependently inhibited by I-BET151, which was consistent with the research by Park-Min et al (21). In the present study, at concentrations of 100 and 200 nM I-BET151, the inhibitory effect of TRAF6 was significant, as was the inhibitory effect for NFATc1 at...
The effects of both cysteine proteinase and collagenase inhibitors on dentine resorption by isolated osteoclasts-Bone. 44: 45-46, 2014.


