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TRPM3/TRPV4 regulates Ca²⁺-mediated RANKL/NFATc1 expression in osteoblasts

Aran Son¹, Namju Kang^{1,2}, Jung Yun Kang^{1,2}, Ki Woo Kim¹, Yu-Mi Yang¹ and Dong Min Shin^{1,2}¹Department of Oral Biology, Yonsei University College of Dentistry, Seoul, Korea²BK21 PLUS Project, Yonsei University College of Dentistry, Seoul, KoreaCorrespondence should be addressed to Y-M Yang or D M Shin: ymyang@yuhs.ac or dmsin@yuhs.ac

Abstract

Mechanical stress plays an important role in the regulation of bone turnover. However, the mechanism underlying hypo-osmotic stress-induced cellular response in osteoblasts remains poorly understood. In this study, we investigated the effect of hypotonic stress on the expression of bone remodeling factors, including the receptor activator of nuclear factor- κ B ligand (RANKL) and the nuclear factor of activated T cells type c1 (NFATc1) in primary mouse osteoblasts and MC3T3-E1 cells. Hypo-osmotic stress induced significant increases in RANKL mRNA expression and intracellular Ca²⁺ concentration ([Ca²⁺]_i) from the extracellular space. Hypo-osmotic stress-induced effects on [Ca²⁺]_i and RANKL and NFATc1 protein expression were decreased by antagonists of transient receptor potential melastatin 3 (TRPM3) and vanilloid 4 (TRPV4). Agonists of TRPM3 and TRPV4 activated [Ca²⁺]_i and RANKL and NFATc1 protein expression. Furthermore, genetic suppression of *Trpm3* and *Trpv4* reduced hypo-osmotic stress-induced effects in mouse osteoblasts. These results suggest that hypo-osmotic stress induces increases in [Ca²⁺]_i through TRPM3 and TRPV4 to regulate RANKL and NFATc1 expression in mouse osteoblastic cells and that mechanical stress-activated TRP channels may play a critical role in bone remodeling.

Key Words

- ▶ hypo-osmotic stress
- ▶ bone remodeling
- ▶ RANKL
- ▶ osteoblast
- ▶ mechanosensitive TRP channels

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Introduction

Bone is a highly dynamic tissue that is constantly being renewed throughout the life through a process called remodeling. Interactions between signals regulating bone-forming osteoblasts and bone-resorbing osteoclasts are basically adaptive responses of the skeleton to its environment (Zaidi 2007, Crockett *et al.* 2011). Although remodeling is initiated by the resorptive action of osteoclasts, it is dependent on signals from osteoblasts. Osteoblasts modulate the expression of the receptor activator of nuclear factor- κ B ligand (RANKL) and osteoprotegerin (OPG) in response to factors that stimulate resorption. RANKL regulates osteoclast

precursors involved in differentiation into multinuclear osteoclasts, whereas OPG inhibits RANKL action by blocking the interaction between RANK and RANKL (Zaidi 2007, Boyce & Xing 2008, Crockett *et al.* 2011, Boyce 2013). The relative ratio of RANKL to OPG expression in osteoblasts is believed to be a key determinant of RANKL-mediated bone resorption. Nuclear factor-activated T cells (NFAT) are members of a transcription factor family that is dependent on calcineurin regulation. NFAT proteins are highly phosphorylated in the cytoplasm of resting cells, and NFAT proteins are then dephosphorylated by calcineurin activation. Dephosphorylated NFAT

proteins are translocated into the nucleus to regulate the transcription of NFAT-dependent genes (Hogan *et al.* 2003). Calcineurin and NFAT are the key factors of osteoclastogenesis and bone resorption (Takayanagi *et al.* 2002, Hirotani *et al.* 2004, Ikeda *et al.* 2004). The Ca^{2+} -calcineurin-NFAT pathway reportedly acts as a positive regulator of osteoblastogenesis and bone formation by maintaining a balanced regulation of osteoclastic and osteoblastic activities (Koga *et al.* 2005, Sun *et al.* 2005, Winslow *et al.* 2006).

Mechanical stress is one of the important regulatory factors for maintaining bone mass and integrity. Mechanical stresses include osmotic stress, shear stress, fluid flow stress and cell membrane stretch (Jin *et al.* 2011, Yang *et al.* 2015, Wittkowske *et al.* 2016). Among the various types of mechanical stress, alteration of osmo-mechanical stresses is often associated with early changes in Ca^{2+} signaling and downstream Ca^{2+} -dependent processes in osteoblastic cells (Weskamp *et al.* 2000, Sun *et al.* 2005, Mehrotra *et al.* 2006, Mizoguchi *et al.* 2008, Nakai *et al.* 2009). There has been much debate regarding the source of the mechanosensitive Ca^{2+} signals and the downstream effector pathways. Recently, there has been increasing evidence for a key role of Ca^{2+} -permeable transient receptor potential (TRP) channels in this process (Pedersen & Nilius 2007, Mizoguchi *et al.* 2008, Jin *et al.* 2011). Mechanosensitive TRP channels (TRPC1, TRPC5, TRPC6; TRPV1, TRPV2, TRPV4; TRPM3, TRPM7; TRPA1 and TRPP2) have been mainly investigated in the cardiovascular system (Christensen & Corey 2007, Pedersen & Nilius 2007). Abed and colleagues have shown that mRNAs of diverse mechanosensitive TRP channels (TRPV2, TRPV4, TRPM3, TRPM4 and TRPM7) are strongly expressed in osteoblastic cells (Abed *et al.* 2009). TRPV2 acts as a mechanosensor for membrane stretch and osmotic cell swelling in vascular smooth muscle cells (Muraki *et al.* 2003, Beech *et al.* 2004). TRPV4 is activated by osmotic cell swelling, excluding potential pathways, such as membrane stretch (Liedtke *et al.* 2000, Strotmann *et al.* 2000, Becker *et al.* 2005, Son *et al.* 2015). TRPM3 is stimulated by hypo-osmotic cell swelling (Grimm *et al.* 2003, Son *et al.* 2015), and TRPM4 has a role as a mechano-/stretch-sensitive channel involved in the control of pressure-induced smooth muscle cell depolarization (Pedersen & Nilius 2007, Guinamard *et al.* 2010). TRPM7 is activated by cell stretch and hypo-osmotic cell swelling; moreover, shear stress induces the translocation of TRPM7 to the plasma membrane and activation of TRPM7 currents (Oancea *et al.* 2006, Numata *et al.* 2007). However, the mechanisms through

which mechanosensitive TRP channels and intracellular Ca^{2+} are regulated have not been clearly elucidated.

In this study, we investigated the effect of hypo-osmotic stress on basal RANKL expression and the function of mechanosensitive TRP channels in primary mouse osteoblasts and MC3T3-E1 cells. We also demonstrated that hypo-osmotic stress induces large increases in the differentiation markers RANKL and NFATc1 and in intracellular Ca^{2+} concentration ($[\text{Ca}^{2+}]_i$) through the osmo-mechanosensitive TRP channels TRPM3 and TRPV4.

Materials and methods

Reagents

All cell culture media and supplements were purchased from Invitrogen. The acetoxymethyl-ester form of fura2 (fura2/AM) was purchased from Teflabs (Austin, TX, USA). Pluronic F127 was obtained from Molecular Probes. Anti-RANKL and anti-NFATc1 antibodies were obtained from Santa Cruz Biotechnology. Anti-actin antibodies were purchased from Sigma-Aldrich. Ruthenium red (RR), 4a-phorbol didecanoate (4a-PDD), ononetin and HC067047 were obtained from Tocris (Bristol, UK). All other chemicals were purchased from Sigma-Aldrich. Stock solutions of all drugs were made with distilled water, except for 2-aminoethoxydiphenyl borate (2APB) with ethanol and 4a-PDD and fura2/AM with DMSO.

Cell culture

All animal care and experimental procedures complied with institutional guidelines and were approved by the Institutional Animal Care and Use Committee (IACUC) of Yonsei University (IACUC approval no. 2014-0067). Primary osteoblast cell cultures were prepared as described previously (Hogan *et al.* 2003). In brief, the calvariae of 1-day-old ICR mice were digested with an enzyme solution containing 0.1% collagenase and 0.1% dispase. The isolated osteoblastic cells were cultured for 4 days in alpha-minimum essential medium (a-MEM) supplemented with 10% fetal bovine serum (FBS) and 1% antibiotic-antimycotic solution (100 U/mL penicillin and 100 µg/mL streptomycin) in a humidified incubator containing 5% CO_2 . MC3T3-E1 cells (Korean Cell Line Bank, Seoul, Korea) were maintained in a-MEM containing 10% FBS and 1% antibiotic-antimycotic solution. Osteoblastic differentiation of MC3T3-E1 cells was induced by 50 µg/mL ascorbic acid and 10 mM beta-glycerolphosphate.

Reverse transcription polymerase chain reaction (RT-PCR)

The expressions of RANKL, OPG, TRPM3, TRPV4 and beta-actin were evaluated by RT-PCR using total RNA isolated from murine osteoblastic cells. Cells were lysed in TRIzol reagent according to the instructions of the manufacturer (Invitrogen). AccuPower RT PreMix (BIONEER, Daejeon, Korea) with total RNA (1 µg) was used for cDNA synthesis. cDNA was amplified by PCR with HiPi Thermostable DNA polymerase (Elpis, Pusan, Korea) using the following primer sets: RANKL (forward): 5'-ATCAGAAGACAGCACTCACT-3' (reverse): 5'-ATCTAGGACATCCATGCTAATGTTTC-3'; OPG (forward): 5'-TGAGTGTGAGGAAGGGCGTT-3' (reverse): 5'-TTCCTCGTTCTCTCAATCTC-3'; TRPM3 (forward): 5'-CACCTGATGACCAAGGAATG-3' (reverse): 5'-CTTGTGTTTATCTTCTGGAGTG-3'; TRPV4 (forward): 5'-GCTGAA GGCAAAGTCTTG-3' (reverse): 5'-CTAGGGAACCCCAA CTGTGA-3'; beta-actin (forward): 5'-TGTGATGGTGGG AATGGGTCAG-3' (reverse): 5'-TTTGATGTCACGCACG ATTTCC-3'. PCR was performed under the following conditions: 94°C for 5 min, 94°C for 30 s, 40 s for annealing step (temperatures varied with primers), 72°C for 30 s, followed by 72°C for 10 min after 38 cycles were finished. The annealing temperature was 60°C for TRPM3, 56.5°C for TRPV4 and RANKL and 57.6°C for OPG and beta-actin. The PCR products were separated on 1.2% agarose gels and visualized with RedSafe nucleic acid staining solution (iNtRon Biotechnology, Gyeonggi-do, Korea).

[Ca²⁺]_i measurement

Osteoblasts were seeded on cover glass in 35 mm dishes (5 × 10⁴ cells/coverslip). The cells were loaded with 5 µM fura2/AM and 0.05% Pluronic F-127 for 30 min in physiological salt solution (PSS) containing (in mM): 140 NaCl, 5 KCl, 1 MgCl₂, 10 HEPES, 1 CaCl₂ and 10 glucose, adjusted to pH 7.4 and 310 mosmol. For hypo-osmotic stress, PSS was replaced with a hypo-osmotic solution (80 mM NaCl, 215 mosmol). Fura2 fluorescence intensity was measured using excitation wavelengths of 340 and 380 nm, and emitted fluorescence 510 nm (ratio = F₃₄₀/F₃₈₀) was collected and monitored at 2-s intervals using a CCD camera (Universal Imaging Co., Downingtown, PA, USA) as described previously (Son et al. 2009). Images were digitized and analyzed by MetaFlour software (Universal Imaging).

Electrophysiology

Whole-cell voltage-clamp recordings were performed at room temperature by the perforated patch-clamp

method. Currents were recorded with a MultiClamp 700B amplifier (Axon Instruments, Foster City, CA, USA), subsequently digitized at a sampling rate of 10 kHz and analyzed with pCLAMP10 software (Axon Instruments). Pipette resistance varied between 3 and 5 MΩ. Whole-cell currents were elicited by voltage ramps from -100 mV to +100 mV (400-ms duration) applied every 10 s from a holding potential of 0 mV. The bath solution contained (in mM) 140 NaCl, 5 KCl, 2 CaCl₂, 1 MgCl₂, 10 HEPES and 10 glucose, adjusted to pH 7.4 with NaOH. Pipettes for recording TRPM3 currents were filled with an internal solution containing (in mM) 140 CsCl, 5 MgCl₂, 10 BAPTA and 10 HEPES, adjusted to pH 7.2 with CsOH and the bath solution was changed to a K⁺-free external solution (Grimm et al. 2003). The internal solution for recording TRPV4 currents contained (in mM) 140 KCl, 5 EGTA and 10 HEPES, adjusted to pH 7.4 with KOH (Guler et al. 2002).

Western blot

Cell lysates of MC3T3-E1 cells were prepared using RIPA lysis buffer (20 mM Tris, pH 7.4, 250 mM NaCl, 2 mM EDTA, pH 8.0, 0.1% Triton X-100, 0.01 mg/mL aprotinin, 5 µg/mL leupeptin, 0.4 mM PMSF and 4 mM NaVO₄) and spun at 12,000 rpm for 10 min to remove insoluble material. Proteins (50–100 µg/well) were subjected to 8–12% SDS-PAGE and separated by size. The proteins were electro-transferred to a nitrocellulose membrane, blocked with 5% skim milk and probed with antibodies against RANKL (1:500), NFATc1 (1:1000) and actin (1:2000). Thereafter, the blots were washed, exposed to horseradish peroxidase-conjugated secondary antibodies for 1 h, and finally detected by chemiluminescence (Amersham Pharmacia Biotech, Arlington Heights, IL, USA).

Transfection of small interfering (si) RNAs

After a cell density of 70% confluence was reached, siRNA duplexes specific for mouse *Trpm3*, *Trpv4* or negative control (BIONEER) were transfected using Lipofectamine 2000 (Invitrogen) according to the manufacturer's instructions.

Statistical analysis

All data were expressed as means ± S.E.M. Statistical significance was determined by a paired Student's *t*-test. Statistical significance was set at the *P* < 0.05 level.

Results

Hypo-osmotic stress induces increased RANKL expression in osteoblasts

Various mechanical stresses have been identified as regulators of bone formation in the differentiation of bone cells and as inducers of RANKL expression in osteoblasts (Boyce & Xing 2008). In this study, we examined whether hypo-osmotic stress affects RANKL expression after treatment of hypo-osmotic solution (215 mosmol) for 3 h in primary mouse osteoblasts. The expression of RANKL mRNA was significantly increased by hypo-osmotic stress (~1.5-fold) compared with control. This increase was not different from the increase in expression of RANKL mRNA by 1,25(OH)₂D₃ (~1.6-fold), when the hormonally active form of vitamin D was used as a positive control (Takeda *et al.* 1999) (Fig. 1A and B). Furthermore, the expression of RANKL mRNA (Supplementary Fig. 1A, see section on [supplementary data](#) given at the end of this article) and the ratio of RANKL to OPG expression were significantly increased 3 h after stimulation by hypo-osmotic stress (by ~1.2-fold and 2.3-fold, respectively) (Fig. 1C and D), but the level of OPG mRNA was not altered by osmotic stimulation (Supplementary Fig. 1B). This suggests that increase in RANKL mRNA by hypo-osmotic stress initiates RANKL-mediated bone resorption.

Hypo-osmotic stress-induced increases of [Ca²⁺]_i and RANKL expression are dependent on extracellular Ca²⁺ entry

Mechanical stress generates the intracellular Ca²⁺ response in bone cells (Weskamp *et al.* 2000, Winslow *et al.* 2006, Huo *et al.* 2008). Therefore, we examined Ca²⁺ signaling by osmotic stress in osteoblasts. As shown in Fig. 2A, hypo-osmotic stress increased [Ca²⁺]_i and even the repeated application of hypo-osmotic stress did not significantly change the increase in [Ca²⁺]_i (87.3±0.4% of first application). To identify the Ca²⁺ influx-related pathways involved, we examined whether the increase in [Ca²⁺]_i promoted by hypo-osmotic stress was dependent on extracellular Ca²⁺ in osteoblasts. When hypo-osmotic stress was applied in Ca²⁺-free external solution, the hypo-osmotic stress-induced increase in [Ca²⁺]_i was completely inhibited (Fig. 2B). In addition, 100 μM lanthanum (La³⁺), as a non-specific blocker of Ca²⁺-permeable channels, or 10 μM gadolinium (Gd³⁺), as a non-specific blocker of stretch-activated and some voltage-gated ion channels, also significantly prevented the hypo-osmotic stress-induced increase in [Ca²⁺]_i (Fig. 2C). These results suggest that the major source of hypo-osmotic stress-induced [Ca²⁺]_i increase is Ca²⁺ influx from the extracellular space. To investigate whether the hypo-osmotic stress-induced increase in [Ca²⁺]_i affected RANKL expression, we examined the changes in RANKL mRNA induced by

Figure 1

Effect of hypo-osmotic stress on RANKL mRNA expression in primary mouse osteoblasts. (A) Primary mouse osteoblasts were stimulated with hypo-osmotic solution (215 mosmol) (hypotonic) as mechanical stimulation and active vitamin D, 1,25(OH)₂D₃ (10 nM), as a positive control for 3 h. RANKL mRNA expression was measured by RT-PCR. (B) The level of RANKL mRNA was quantified after the normalized value to beta-actin ($n=6$). (C) Cells were stimulated with the hypo-osmotic solution for the indicated times. Expression of mRNAs was measured. (D) The expression ratio of RANKL/OPG mRNA was quantified after the normalized value to beta-actin ($n=3$). Data are expressed as means±s.e.m. * $P<0.05$ compared with control.

Figure 2

Dependence on extracellular Ca²⁺ of hypo-osmotic stress-induced [Ca²⁺]_i increase and RANKL expression. (A) Treatment with hypo-osmotic solution (hypotonic) induced increases in [Ca²⁺]_i and also showed that [Ca²⁺]_i was increased similarly by repeated application of hypo-osmotic solution in primary mouse osteoblasts using fura2 fluorescence dye. The changes in [Ca²⁺]_i were quantified after the normalized value to the first hypo-osmotic stress (*n*=9). (B and C) The effects of increases in [Ca²⁺]_i by hypo-osmotic stress were inhibited by Ca²⁺-free external solution and 100 μM La³⁺, which block a wide range of Ca²⁺-permeable channels. The inhibitory effects on blockers were quantified after the normalized value to hypo-osmotic stress (*n*=6) and ionomycin (*n*=6). (D) Cells were incubated with Gd³⁺ or La³⁺ in the hypo-osmotic solution for 3 h. The increases in RANKL mRNA by hypo-osmotic stress were reduced by 10 μM Gd³⁺ and 100 μM La³⁺ and the level of RANKL mRNA was quantified after the normalized value to beta-actin (*n*=3). (E) Cells were treated with 1 μM thapsigargin in PSS (a presence of 1 mM CaCl₂) for the indicated times. The expression of RANKL mRNA was quantified after the normalized value to beta-actin (*n*=6). Data are expressed as means ± s.e.m. **P*<0.05, ***P*<0.01, and ****P*<0.001 compared with control.

hypo-osmotic stress after treatment with 100 μ M La^{3+} or 10 μ M Gd^{3+} for 3 h. The increase of RANKL mRNA by hypo-osmotic stress was significantly decreased by Gd^{3+} and La^{3+} (Fig. 2D). This result suggests that the increase in $[\text{Ca}^{2+}]_i$ from the extracellular space by hypo-osmotic stress leads to the increase in RANKL expression. Thapsigargin, an inhibitor of sarco/endoplasmic reticulum Ca^{2+} ATPase, can increase $[\text{Ca}^{2+}]_i$ by depletion of intracellular Ca^{2+} stores and subsequent influx of Ca^{2+} into the cytosol via activated plasma membrane Ca^{2+} channels. Thus, to investigate whether the increase in $[\text{Ca}^{2+}]_i$ by thapsigargin affected RANKL expression, we examined the level of RANKL mRNA after treatment with 1 μ M thapsigargin in PSS (a presence of 1 mM CaCl_2). The expression of RANKL mRNA was significantly increased 3 and 6 h (by ~ 1.8 -fold and 1.7-fold, respectively) after treatment with thapsigargin; the increase was not different from the increase in expression of RANKL mRNA by 1,25(OH) $_2\text{D}_3$ (~ 1.7 -fold) (Fig. 2E). This result suggests that the RANKL expression is related with the increase of $[\text{Ca}^{2+}]_i$ through extracellular Ca^{2+} entry as well as thapsigargin-mediated store-operated Ca^{2+} entry (SOCE).

Activation of TRPM3 and TRPV4 induces increases of $[\text{Ca}^{2+}]_i$ and the expression of RANKL-mediated NFATc1 in osteoblasts

Diverse mechanosensitive TRP channels have been identified in osteoblastic cells (Abed *et al.* 2009). Osmo-mechanosensitive TRP channels (TRPM2, TRPM3, TRPM7 and TRPV4) have been implicated in controlling the osmosensitive response in osmoregulation organs, such as kidney, sensory neurons, airway epithelia, bone and so on, to changes in osmolarity (Liedtke *et al.* 2000, Strotmann *et al.* 2000, Grimm *et al.* 2003, Koga *et al.* 2005, Sun *et al.* 2005, Bessac & Fleig 2007, Christensen & Corey 2007, Numata *et al.* 2007, 2012, Nakai *et al.* 2009, Jin *et al.* 2011). In particular, TRPM3 and TRPV4 are directly activated by hypo-osmotic stress (Strotmann *et al.* 2000, Alessandri-Haber *et al.* 2003, Grimm *et al.* 2003). To determine which osmo-mechanosensitive TRP channels play a role in the hypo-osmotic stress-induced increases in $[\text{Ca}^{2+}]_i$, we confirmed the expression and functional activities of osmo-mechanosensitive TRP channels in osteoblasts. As shown in Fig. 3A, both TRPM3 and TRPV4 are present in primary mouse osteoblasts. However, TRPM2 was scarcely expressed, and TRPM7 was not affected by hypo-osmotic stress for 3 h (Supplementary Fig. 2). To directly assess the functional activities of these channels, we examined the expression of RANKL and NFATc1 proteins after treatment

with agonists and antagonists of TRPM3 and TRPV4 for 24 h. The expression of RANKL and NFATc1 protein was markedly increased by pregnenolone sulfate (PS) (50 μ M) and 4a-PDD (10 μ M), which are agonists of TRPM3 and TRPV4. Both ononetin (5 μ M) and HC067047 (500 nM), which are antagonists of TRPM3 and TRPV4, inhibited the agonist-induced increases in protein expression (Fig. 3B). In addition, both PS and 4a-PDD enhanced the increase of $[\text{Ca}^{2+}]_i$ in osteoblasts, and these effects on increased $[\text{Ca}^{2+}]_i$ by activation of TRPM3 and TRPV4 were diminished by ononetin and HC067047 (Fig. 3C and D). Taken together, these results show that activation of RANKL and NFATc1 can be induced by increases in $[\text{Ca}^{2+}]_i$ via TRPM3 and TRPV4. Next, we examined the sensitivity of TRPM3 and TRPV4 to hypotonicity using a whole-cell patch-clamp technique. Hypo-osmotic stress activated TRPM3-mediated outwardly rectifying currents (-0.10 ± 0.01 nA at -80 mV and 0.22 ± 0.05 nA at $+80$ mV) (Fig. 3E) and TRPV4-mediated outwardly rectifying currents (-0.28 ± 0.07 nA at -80 mV and 0.40 ± 0.05 nA at $+80$ mV) (Fig. 3F), which reversed near 0 mV. These results suggest that the hypo-osmotic stress-evoked response is mediated by both TRPM3 and TRPV4. Therefore, we confirmed the effects of TRPV4 and TRPM3 on hypo-osmotic stress in subsequent experiments.

Hypo-osmotic stress activates the increases of $[\text{Ca}^{2+}]_i$ and expression of RANKL-mediated NFATc1 through osmo-mechanosensitive TRP channels (TRPM3 and TRPV4) in osteoblasts

To determine the specific activation of osmo-mechanosensitive TRP channels (TRPM3 and TRPV4) by osmotic stress in osteoblasts, we used blockers of TRPM3 and TRPV4. The hypo-osmotic stress-induced increase in $[\text{Ca}^{2+}]_i$ was partially inhibited by 100 μ M 2APB, a non-specific blocker of the TRPM3 channel, and 5 μ M ononetin, a selective inhibitor of the TRPM3 channel (Fig. 4A). Antagonists of TRPV4 (10 μ M RR as a non-specific blocker and 500 nM HC067047 as a potent antagonist) also partially inhibited the hypo-osmotic stress-induced increase in $[\text{Ca}^{2+}]_i$ (Fig. 4B). These results indicate that hypo-osmotic stress is capable of inducing Ca^{2+} influx from the extracellular space via TRPM3 and TRPV4 on plasma membranes. To investigate whether the expressions of RANKL and NFATc1 was affected by the increase of $[\text{Ca}^{2+}]_i$ via hypo-osmotic stress, we examined the changes in RANKL and NFATc1 protein levels by hypo-osmotic stress after treatments with antagonists of TRPM3 and TRPV4 for 24 h. The increases in RANKL and NFATc1 protein levels

Figure 3

Functional roles of TRPM3 and TRPV4 in $[Ca^{2+}]_i$ increase and RANKL expression. (A) TRPM3 and TRPV4 were expressed in mouse primary osteoblasts. Expression of TRPM3 and TRPV4 mRNA was examined by RT-PCR. (B) MC3T3-E1 cells were incubated with or without antagonists of TRPM3 or TRPV4 in pretreatment with agonists of TRPM3 or TRPV4 for 24 h. Increases in RANKL and NFATc1 proteins by 50 μ M PS as an agonist of TRPM3 or 10 μ M 4 α -PDD as an agonist of TRPV4 were significantly reduced by 5 μ M ononetin or 500 nM HC067047 in MC3T3-E1 cells. Expression of RANKL and NFATc1 proteins was quantified after the normalized value to control ($n=6$). (C) The effects of increases in $[Ca^{2+}]_i$ by 50 μ M PS were inhibited by 5 μ M ononetin ($n=6$). (D) The effects of increases in $[Ca^{2+}]_i$ by 10 μ M 4 α -PDD were also completely inhibited by 500 nM HC067047 ($n=6$). The inhibitory effects on blockers were quantified after the normalized value to ionomycin. (E) Hypo-osmotic stress activated TRPM3-mediated outwardly rectifying current by a voltage ramp (–100 to +100 mV at 400-ms intervals, $V_h=0$ mV), which was used to determine current–voltage relations in primary mouse osteoblasts currents (-0.112 ± 0.024 nA at –80 mV and 0.183 ± 0.038 nA at +80 mV, $n=10$). These effects disappeared with treatment by 5 μ M ononetin (-0.028 ± 0.007 nA at –80 mV and 0.037 ± 0.009 nA at +80 mV, $n=11$). (F) Hypo-osmotic stress also induced TRPV4-mediated outwardly rectifying currents (-0.292 ± 0.031 nA at –80 mV and 0.477 ± 0.048 nA at +80 mV, $n=18$). TRPV4-mediated currents were reduced significantly by 50 nM HC067047 (-0.084 ± 0.015 nA at –80 mV and 0.142 ± 0.020 nA at +80 mV, $n=16$). Data are expressed as means $\pm$ s.e.m. * $P<0.05$, ** $P<0.01$, and *** $P<0.001$ compared with control.

by hypo-osmotic stress were decreased by antagonists of TRPM3 and TRPV4 (Fig. 4C and D). This result suggests that the increase in $[Ca^{2+}]_i$ through TRPM3 and TRPV4 by hypo-osmotic stress causes activation of RANKL and NFATc1.

To investigate the role of osmo-mechanosensitive TRP channels in osteoblasts, we examined the functional

effects of TRPM3 and TRPV4 on Ca^{2+} influx and protein expression by hypo-osmotic stress using *Trpm3* and *Trpv4* siRNAs in MC3T3-E1 and primary mouse osteoblast cells. Upon knockdown with specific siRNA for *Trpm3* or *Trpv4*, their mRNA levels were completely suppressed (Fig. 5A). The hypo-osmotic stress-induced increases in RANKL and

Figure 4

Relationship between osmo-mechanosensitive TRP channels and hypo-osmotic stress. (A) The effects of increases in $[Ca^{2+}]_i$ by hypo-osmotic stress (hypotonic stress) were partially inhibited by 100 μ M 2APB as a non-specific blocker of TRPM3 and 5 μ M ononetin as an antagonist of TRPM3 in osteoblasts ($n=5$). (B) The effects of increases in $[Ca^{2+}]_i$ by hypo-osmotic stress were also partially inhibited by 10 μ M ruthenium red (RR) as a non-specific blocker of TRPV4 and 500 nM HC067047 as an antagonist of TRPV4 in osteoblasts ($n=4$). The inhibitory effect on blockers was quantified after the normalized value to hypo-osmotic stress. (C) Cells lysates were collected from MC3T3-E1 cells stimulated with hypo-osmotic solution and blockers of osmo-mechanosensitive TRP channels for 24 h. (D) Expression of RANKL and NFATc1 proteins was quantified after the normalized value to control ($n=4$). Data were expressed as means $\pm$ S.E.M. * $P<0.05$, ** $P<0.01$, and *** $P<0.001$ compared with control.

NFATc1 expression were diminished in *Trpm3*- and *Trpv4*-siRNA-transfected cells (Fig. 5B). Moreover, the amplitude of hypo-osmotic stress-induced increases in $[Ca^{2+}]_i$ was reduced by approximately 37 and 35%, respectively, in *Trpm3*- and *Trpv4*-siRNA-transfected cells, compared with negative control cells (Fig. 5C and D). These results indicate that activation of TRPM3 and TRPV4 is essential for hypo-osmotic stress-mediated extracellular Ca^{2+} influx and induction of RANKL and NFATc1 expression in osteoblasts.

Discussion

In this study, we demonstrated a novel regulatory mechanism between osmo-mechanosensitive TRP channels, especially TRPM3 and TRPV4, and differentiation markers of bone cells through entry of extracellular Ca^{2+} in mouse osteoblast cells (Fig. 6). These effects could be the result of the accelerating activity in $[Ca^{2+}]_i$ via TRPM3 and TRPV4 by mechanical stresses during bone remodeling, which is the previously reported physiological function of mechanical stresses such as endogenous RANKL synthesis, maintenance of bone mass and regulation of osteoclast differentiation in osteoblast cells (Tsuzuki et al. 2000, Takayanagi et al. 2002, Becker et al. 2005, Mehrotra et al. 2006, Mizoguchi et al. 2008, Abed et al. 2009, Nakai et al. 2009). However, the major signaling pathway and actions of mechanosensors in osteoblasts remain a mystery. Our results show that hypo-osmotic stress-induced increases

in RANKL mRNA expression depend on the increase of $[Ca^{2+}]_i$ in osteoblasts. Although other groups reported that extracellular Ca^{2+} entry as well as SOCE-elevated $[Ca^{2+}]_i$ effected expression levels of RANKL and OPG mRNA in primary osteoblasts (Takami et al. 2000, Hwang et al. 2012), these different results suggest that various stimuli can induce increase of $[Ca^{2+}]_i$ through activation of a different mechanism of bone remodeling. Hence, the increase of $[Ca^{2+}]_i$ in osteoblasts is necessary for the activation of differentiation markers of osteoblasts and the regulation of osteoclast differentiation (Takami et al. 2000, Weskamp et al. 2000, Romanello et al. 2005, Hwang et al. 2012, Robinson et al. 2012).

In the present study, hypo-osmotic stress-induced increases of RANKL and NFATc1 expressions were also dependent on Ca^{2+} influx through the extracellular space and osmo-mechanosensitive TRP channels (TRPM3 and TRPV4). In previous reports, various mechanical perturbations, such as mechanical load and fluid shear stress or pressure, played an important role in bone remodeling by activation of Ca^{2+} signaling and RANKL due to the influx of extracellular Ca^{2+} through mechanosensitive TRP channels, purinergic receptors, big conductance Ca^{2+} -activated K^+ channels in bone (Pavalko et al. 1998, Weskamp et al. 2000, Romanello et al. 2005, Mehrotra et al. 2006, Oancea et al. 2006, Masuyama et al. 2008, Suzuki et al. 2013). TRPV4 was the first mechanosensitive TRP channel shown to be a volume-activated and Ca^{2+} -permeable cation channel by

Figure 5

Effects of deleted osmo-mechanosensitive TRP channels on $[Ca^{2+}]_i$ increases and activation of differentiation factors mediated by hypo-osmotic stress. (A) Osteoblasts were transfected with siTRPM3 and siTRPV4 using Lipofectamine 2000. After 72 h of transfection, whole mRNA was collected using the Trizol method. beta-actin was used as loading control. (B) Transfected MC3T3-E1 cells were treated with the hypotonic solution (Hypotonic) for 24 h. Hypo-osmotic stress-induced increases in RANKL and NFATc1 expressions were decreased in *Trpm3*- and *Trpv4*-siRNA-transfected MC3T3-E1 cells. Expression of RANKL and NFATc1 proteins was quantified after the normalized value to negative control (NC) ($n=6$). (C and D) Application of hypo-osmotic solution induced increases in $[Ca^{2+}]_i$ in negative control osteoblasts (a). However, these effects were reduced in siTRPM3-treated cells (b) ($n=4$) and siTRPV4-treated cells ($n=4$). We identified the siRNA-transfected cells by GFP fluorescence (upper side) before a test, and increases in $[Ca^{2+}]_i$ in single cells were measured using fura2 fluorescence dye (lower side). Data are expressed as means $\pm$ s.e.m. * $P < 0.05$ and *** $P < 0.001$ compared with control.

hypo-osmotic stimuli (Liedtke *et al.* 2000, Strotmann *et al.* 2000). TRPV4 also regulates bone resorption, which is necessary for sustained Ca^{2+} signaling and NFATc1 activation by the Ca^{2+} /calmodulin signal pathway in osteoclasts, and bone formation, which is essential for flow-induced Ca^{2+} signaling during differentiation in osteoblasts (Boyce & Xing 2008, Masuyama *et al.* 2008, Mizoguchi *et al.* 2008, Suzuki *et al.* 2013). TRPM3 of osmo-mechanosensitive TRP channels also participates in hypo-osmotic cell swelling (Grimm *et al.* 2003, Harteneck & Schultz 2007). In our previous study, hypo-osmotic stress induced RANKL expression via TRPM3 and TRPV4 in human periodontal ligament cells (Son *et al.* 2015). However, the functional mechanism of TRPM3 and TRPV4 in bone remodeling is not fully understood. Our study is the first to show that TRPM3 and TRPV4 are expressed

in osteoblasts and that the activities of differentiation markers are regulated by hypo-osmotic stress. TRPM7 as a mechanosensor is also expressed in osteoblasts and plays an important role in various cell types. TRPM7 has been proposed to be directly activated by volume regulation on mechanical stimuli and magnesium regulation, and the increase in $[Ca^{2+}]_i$ via TRPM7 potentiates by hypo-osmotic cell swelling (Oancea *et al.* 2006, Numata *et al.* 2007, Abed *et al.* 2009, Boyce 2013, Kim *et al.* 2017, Won *et al.* 2018). According to our previous report, TRPM7 controls RANKL-induced osteoclastogenesis by regulation of Ca^{2+} oscillation and NFATc1 expression in osteoclast cells (Yang *et al.* 2013). In the present study, we had ruled out the possibility that TRPM7 was involved in osteoblast differentiation because of the inhibition of TRPM7 activity by regulation of Mg^{2+} in all buffers

Figure 6

Proposed model of the regulatory mechanisms of osteoblastic signal pathways for osmo-mechanosensitive TRPM3 and TRPV4. The model shows that mechanical stresses, such as hypo-osmotic stimuli, activate the expression of RANKL and NFATc1, which is related to the differentiation and proliferation of bone cells by increases in $[Ca^{2+}]_i$ through TRPM3 and TRPV4 in osteoblasts.

used. We also confirmed that hypo-osmotic stress had no effect on the TRPM7 expression in osteoblast cells (Supplementary Fig. 2). Therefore, activation of TRPM3 and TRPV4 by mechanical stress is essential for control of bone remodeling and osteoblastic differentiation.

Notably, osteoblasts can control osteoclast differentiation and activity and consequently bone remodeling through the modulation of RANKL/OPG and calcineurin/NFAT in osteoblastogenesis (Koga et al. 2005, Hadjidakis & Androulakis 2006, Winslow et al. 2006, Zaidi 2007, Hwang et al. 2012). NFATc1 must first be activated through dephosphorylation by the Ca^{2+} /calmodulin-regulated phosphatase calcineurin. Activated NFATc1 then binds to Osterix, a regulator of osteoblast transcription downstream of Runx2, to stimulate osteoblastogenesis and bone formation. Therefore, deletion of NFATc1 and A alpha isoform of calcineurin in the osteoblast impair bone formation (Koga et al. 2005, Sun et al. 2005, Winslow et al. 2006). Here, we have also shown that hypo-osmotic stress-induced NFATc1 expression is regulated by activation of TRPM3 and TRPV4 in osteoblasts. Hence, extracellular Ca^{2+} influx and increases in $[Ca^{2+}]_i$ through TRPM3 and TRPV4 as osmo-mechanosensitive TRP channels are essential for the activity of NFATc1 and the physiological activity of osteoblasts.

In conclusion, this is the first report showing that hypo-osmotic stress affects regulation of $[Ca^{2+}]_i$ and the

differentiation markers RANKL and NFATc1 via modulation of TRPM3 and TRPV4 in osteoblastic cells. These results potentially have a broad impact and clinical relevance for pathological conditions in the skeletal system. Based on our results, adjusting the activities of TRPM3 and TRPV4 by genetic modification and Ca^{2+} ingestion has to be considered as a novel way to protect against bone diseases caused by impaired bone remodeling, such as osteopenia and osteoporosis.

Supplementary data

This is linked to the online version of the paper at <https://doi.org/10.1530/JME-18-0051>.

Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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Author contribution statement

A S and D M S designed the study. N K and J Y K performed experiments. A S and Y M Y contributed to data acquisition and interpretation. Y M Y and D M S drafted the manuscript. K W K, Y M Y, and D M S revised the manuscript. All authors approved the final version of the manuscript.

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