

Hivep3-dependent Alg2 Expression Inhibits Osteogenesis

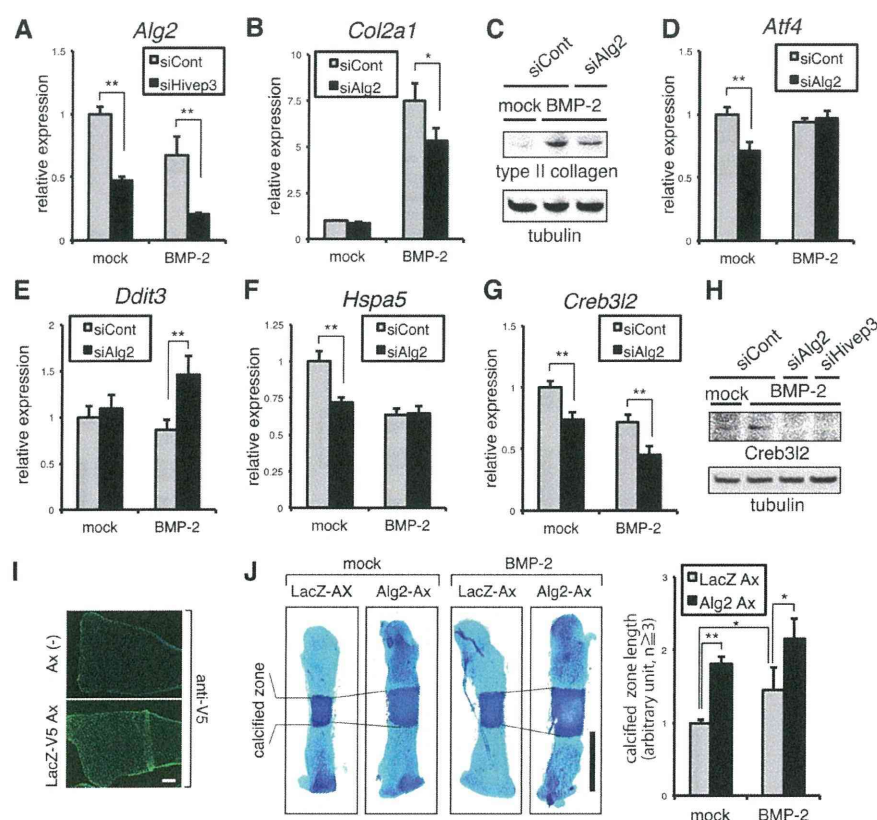


FIGURE 8. Hivep3 is essential for Alg2 expression in ATDC5 chondrocytes, although Alg2 promotes chondrogenesis. A, ATDC5 cells were transfected with siRNA for *Hivep3* and treated with BMP-2 (300 ng/ml) for 4 days. Expression of *Alg2* was evaluated by qRT-PCR. B, D, E, F and G, ATDC5 cells were transfected with siRNA for *Alg2* and treated with BMP-2 (300 ng/ml) for 4 days. The expression of the indicated genes was analyzed by qRT-PCR. C, ATDC5 chondrocytes were transfected with siRNA for *Alg2* with BMP-2 treatment (300 ng/ml) for 7 days. Cell lysates were analyzed by immunoblotting with an anti-type II collagen antibody. Tubulin served as a loading control. H, ATDC5 chondrocytes were transfected with siRNA for *Alg2* or *Hivep3* with BMP-2 treatment (300 ng/ml) for 3 days. Cell lysates were analyzed by immunoblotting with an anti-Creb3l2 antibody. Tubulin served as a loading control. I and J, metatarsal cartilages of E17.5 mouse embryos were infected with indicated adenovirus for 16 h. Immunostaining using FITC-linked anti-V5 antibody on bone coronal sections was performed at day 2 of culture (I). Nuclei were stained with Hoechst dye. Merged images are presented. Scale bar, 100 μ m. Alcian blue/alizarin red staining was performed at day 3 of BMP-2 treatment (J). The length of calcified zone (matured hypertrophic chondrocytes) was measured. Scale bar, 500 μ m. *, $p < 0.05$; **, $p < 0.01$.

Mutations in the *ALG2* gene cause the rarest form of congenital disorders of glycosylation (CDG) in humans, CDG type-Ii (CDG-Ii) (25). A single patient of CDG-Ii has been identified, who was only mildly affected with developmental delay, seizures, poor vision, coagulopathy, and delayed myelination (25), with no remarkable bone phenotype. However, because the reported patient was 3 years old, and *Hivep3* knock-out mice have been reported to show adult-onset osteosclerosis (10), skeletal disorders may develop during the adulthood of the CDG-Ii patient. The glycosyltransferase enzymes of the ALG pathway use nucleotide- or dolichol-activated monosaccharides as donor substrates (41), which are biosynthesized by the cytosolic enzyme PMM2. CDG-Ia is the largest group of CDG cases with more than 800 patients who have been identified as having mutations in the *PMM2* gene (41). It is noteworthy that

CDG-Ia patients show a variety of skeletal phenotypes, including osteopenia, rhizomelia, and ossification delay of bones (42). Fibrillar collagens, such as type I or type II collagens, are found predominantly in the ECM of bone or cartilage, respectively, and mature through ALG of the C-terminal pro-collagen, resulting in cleavage of the N- and C-terminal pro-peptide domains. Therefore, dysregulation of ALG may affect the maturation and function of collagens in the skeleton. Conversely, during the differentiation of osteoblasts and chondrocytes, nascent ECM protein is delivered in amounts that exceed the capacity of the ER, whose machinery processes the post-translational modifications and folding of proteins; ALG proteins play important roles for these steps. Such events of "overload" trigger mild ER stress (physiological ER stress) (43). In ST-2 cells, loss of *Hivep3* did not influence mild ER stress during

FIGURE 7. Hivep3 potentiates the physiological mild ER stress to promote chondrocyte differentiation. A, E, G, I, K, M, and O, ATDC5 cells were transfected with siRNA for *Hivep3* and treated with BMP-2 (300 ng/ml) for 4 days. The expression of the indicated genes was evaluated by qRT-PCR. B, ATDC5 chondrocytes were transfected with siRNA for *Hivep3* with BMP-2 treatment (300 ng/ml) for 7 days. Cell lysates were analyzed by immunoblotting with an anti-type II collagen antibody. Tubulin served as a loading control. C, F, H, J, L, and N, ATDC5 cells were transfected with a human *HIVEP3* expression vector and further stimulated with BMP-2 (300 ng/ml) for 4 days. The expression of the indicated genes was evaluated by qRT-PCR. D, ATDC5 cells were stably transfected with a pEF-Shn3 (*Hivep3*) expression vector. Micromass culture of the transfectants was treated with BMP-2 (300 ng/ml) for 21 days and stained with Alcian blue dye. *, $p < 0.05$; **, $p < 0.01$.

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osteoblast differentiation. In contrast, Hivep3 increased not only the level of *Creb3l2*, but also of other canonical ER stress-related genes in ATDC5 chondrocytes, although loss of *Alg2* decreased the expression of *Creb3l2*. These results suggest that the Hivep3-*Alg2* pathway is important for physiological ER stress in chondrocytes.

Although Hivep2 and Hivep3 showed cooperative roles in decreasing bone formation and bone volume *in vivo* (17), Saita *et al.* (16) reported results similar to ours where gain of Hivep2 expression enhanced osteoblast differentiation, and Hivep2 knock-out osteoblasts failed to support efficient osteoclastogenesis *in vitro*. In addition, they showed that bone formation, as well as bone resorption, decreased in the bones of Hivep2-null mice, resulting in osteopenia due to the low turnover of bone remodeling. Recently, Hivep3 was also found to promote osteoclastogenesis (14, 15). Taken together, the defects in osteoclastogenesis induced by loss of the Hivep genes seem to be mainly responsible for the additively increased bone volume in *Hivep2/Hivep3* double knock-out mice. In contrast, we showed that each of the three Hivep genes was essential for chondrogenesis *in vitro*, suggesting that the chondrodysplasia observed following combined ablation of Hivep2 and Hivep3 in mice (17) was a result of the loss of the cell autonomous action of the Hivep proteins. Although we found the physiological ER stress and the level of *Creb3l2* to be increased by the Hivep3-*Alg2* axis, the precise molecular mechanisms are unclear. Moreover, it is still unclear why the three Hivep genes exhibit diverse actions in osteogenesis. Because expression of *Alg2* was not altered by loss of Hivep1 or Hivep2, other molecular targets downstream of Hivep1 or Hivep2 should be evaluated. Although Hivep proteins are known to act as transcription factors as well as scaffolds, they may harbor some other unknown properties, as most of the area of the large proteins has not been classified into any known functional domains.

In conclusion, *Alg2* is a downstream mediator of Hivep3 in osteoblasts and chondrocytes. In addition to initiation of Runx2 protein degradation, Hivep3 interfered with the function of Runx2 via *Alg2*-mediated disturbance of localization and activity. In ATDC5 chondrocytes, Hivep3 and *Alg2* enhanced mild ER stress to promote differentiation. Thus, our results are the first to link the ALG gene to differentiation of skeletal cells. Future studies on mice with knock-out of ALG genes, as well as detailed clinical research with corresponding CDG patients, may provide more information regarding the roles of ALG proteins in osteogenesis, chondrogenesis, and bone remodeling.

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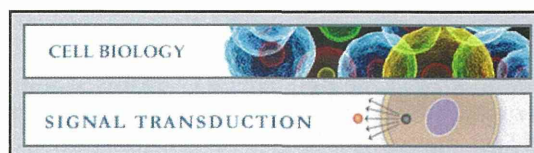
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Cell Biology:

**Bone Morphogenic Protein (BMP)
Signaling Up-regulates Neutral
Sphingomyelinase 2 to Suppress
Chondrocyte Maturation via the Akt
Protein Signaling Pathway as a Negative
Feedback Mechanism**

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Bone Morphogenic Protein (BMP) Signaling Up-regulates Neutral Sphingomyelinase 2 to Suppress Chondrocyte Maturation via the Akt Protein Signaling Pathway as a Negative Feedback Mechanism^{*[S]}

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Background: It is not clear how BMP-induced chondrocyte maturation is cell-autonomously terminated.

Results: BMP-2 induced the ceramide-generating enzyme neutral sphingomyelinase 2 (nSMase2) in chondrocytes, whereas silencing of nSMase2 enhanced maturation in an Akt signaling-dependent manner.

Conclusion: nSMase2 signaling regulates BMP-induced chondrocyte maturation as a negative feedback mechanism.

Significance: This study elucidated the novel link between BMP and lipid signaling in chondrogenesis.

Although bone morphogenic protein (BMP) signaling promotes chondrogenesis, it is not clear whether BMP-induced chondrocyte maturation is cell-autonomously terminated. Loss of function of *Smpd3* in mice results in an increase in mature hypertrophic chondrocytes. Here, we report that in chondrocytes the Runx2-dependent expression of *Smpd3* was increased by BMP-2 stimulation. Neutral sphingomyelinase 2 (nSMase2), encoded by the *Smpd3* gene, was detected both in prehypertrophic and hypertrophic chondrocytes of mouse embryo bone cartilage. An siRNA for *Smpd3*, as well as the nSMase inhibitor GW4869, significantly enhanced BMP-2-induced differentiation and maturation of chondrocytes. Conversely, overexpression of *Smpd3* or C_2 -ceramide, which mimics the function of nSMase2, inhibited chondrogenesis. Upon induction of *Smpd3* siRNA or GW4869, phosphorylation of both Akt and S6 proteins was increased. The accelerated chondrogenesis induced by *Smpd3* silencing was negated by application of the Akt inhibitor MK2206 or the mammalian target of rapamycin inhibitor rapamycin. Importantly, in mouse bone culture, GW4869 treatment significantly promoted BMP-2-induced hypertrophic maturation and calcification of chondrocytes, which subsequently was eliminated by C_2 -ceramide. *Smpd3* knockdown decreased the apoptosis of terminally matured ATDC5 chondrocytes, probably as a result of decreased ceramide production. In addition, we found that expression of hyaluronan synthase 2 (*Has2*) was elevated by a loss of *Smpd3*, which was restored by MK2206.

Indeed, expression of Has2 protein decreased in nSMase2-positive hypertrophic chondrocytes in the bones of mouse embryos. Our data suggest that the *Smpd3*/nSMase2-ceramide-Akt signaling axis negatively regulates BMP-induced chondrocyte maturation and Has2 expression to control the rate of endochondral ossification as a negative feedback mechanism.

Over 95% of bone formation during the embryonic and developmental stages is achieved through endochondral ossification. This process is primed by the condensation of mesenchymal progenitor cells expressing the chondrogenic master regulator Sox9 (1), after which cells further differentiate into proliferating chondrocytes that are able to express a specific marker, *Col2a1*, encoding type II collagen (2). These chondrocytes then mature to hypertrophic chondrocytes, which eventually mineralize the surrounding cartilage matrix to be replaced by bone-forming osteoblasts (3). The maturation of chondrocytes into hypertrophic chondrocytes, which are able to express the type X collagen-encoding gene *Col10a1*, is mainly governed by the Runx2 transcription factor (4, 5). The rate of proliferation and differentiation of chondrocytes *in vivo* is tightly regulated by a signaling network between Indian hedgehog, parathyroid hormone-related protein, fibroblast growth factor (FGF), and bone morphogenetic protein (BMP)³ signaling (6).

BMPs belong to the transforming growth factor- β (TGF- β) family, which transduces signals through type II and type I receptors to activate receptor-regulated Smads. Upon ligand binding, BMP type I receptors phosphorylate Smad1/5/8 in the cytoplasm. Phosphorylated Smads form a trimeric complex

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[S] This article contains supplemental Table S1.

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³ The abbreviations used are: BMP, bone morphogenic protein; nSMase2, neutral sphingomyelinase 2; MSC, mesenchymal stem/stroma cell; CHX, cycloheximide; RTK, receptor tyrosine kinase; OA, osteoarthritis; ITS, insulin/transferrin/selenium; PP2A, protein phosphatase 2A; rpS6, ribosomal protein S6.

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with Smad4 (co-Smad) to translocate into the nucleus to directly or indirectly regulate the transcription of target genes (7, 8). BMPs and their receptors are expressed throughout the growth plate and perichondrium of developing bone (9). BMPs maintain expression of *Sox9* and subsequent cartilage matrix production (10, 11). BMPs are also required *in vitro* for the induction of *Col2a1* and *Col10a1* to promote chondrocyte maturation at later stages (12–14). Chondrocyte-specific overexpression of the extracellular BMP antagonist, Noggin, in transgenic mice resulted in no cartilage formation (15). Similarly, mice with cartilage-specific combined deletions of two BMP type I receptor genes (*Bmpr1a* and *Bmpr1b*), or with a double knock-out of *Smad1* and *Smad5*, showed severely impaired chondrogenesis (16, 17). At a later maturation stage, BMP signaling directly accelerates the expression of *Col10a1* in concert with Runx2 (13, 18, 19). Forced expression of constitutively active *Bmpr1a* in cartilage promoted the maturation and hypertrophy of chondrocytes in mice (20). The evidence clearly demonstrates the accelerating roles of BMP signaling in chondrocyte commitment, proliferation, and hypertrophic maturation both *in vivo* and *in vitro*.

Chondrogenesis in the endochondral ossification process is a promising cellular event for application in cartilage and bone regeneration, and it is a process that can be artificially engineered from human mesenchymal stem/stroma cells (MSCs) (21). Regarding the treatment of articular cartilage defects, the engineered chondrocytes must arrest maturation processes, because abnormally matured hypertrophic chondrocytes, expressing *COL10A1*, mineralize cartilage matrix to cause pathological conditions such as osteoarthritis (OA) (22–24). This is a major problem of cartilage tissue engineering in an *in vitro* culture system, where MSCs rapidly express *COL10A1* in a monolayer or pellet culture before the cells express hypertrophic phenotypes, suggesting that the mechanism by which MSCs induce type X collagen expression *in vitro* is different from that in chondrogenesis *in vivo* (25, 26). In bone engineering events, MSCs have formed bone trabeculae *in vivo* only when they had developed hypertrophic chondrocyte structure *in vitro* prior to implantation (21). Therefore, controlling the maturation and hypertrophy of chondrocytes is crucial for regenerative medicine of both cartilage and bone.

Although the effects of BMP signaling in promoting chondrocyte maturation should be eliminated for cartilage regeneration, they may be artificially enhanced to promote the efficiency of bone engineering. Although the BMP-Smad pathway directly induces the inhibitory Smad, Smad6, to inhibit the phosphorylation of Smad1/5/8 as a negative feedback mechanism (27), and Smurf1 targets Smad6 and the BMP type I receptors to block BMP signaling (28, 29), these inhibitory molecules are not specifically expressed in maturing chondrocytes. Recently, we reported that expression of the transcriptional repressor SnoN gradually increases in BMP-induced differentiating chondrocytes to suppress BMP signaling and the subsequent chondrocyte hypertrophic maturation (30). However, because SnoN partially blocked chondrocyte maturation, the mechanism regulating the rate of BMP signaling-driven chondrocyte differentiation in a cell-autonomous manner remains unclear.

The membrane-bound enzyme neutral sphingomyelinase 2 (nSMase2), encoded by the sphingomyelin phosphodiesterase 3 (*Smpd3*) gene, cleaves sphingomyelin to generate the lipid second messenger ceramide, which affects a variety of cellular processes, including proliferation, apoptosis, and differentiation (31–34). A deletion mutation in the *Smpd3* gene was identified in fragilitas ossium (*fro*), a mouse model of osteogenesis imperfecta (35). *fro/fro* mice are characterized by retarded skeletal growth and a severely hypomineralized skeleton with normal osteoblast differentiation; bone mineralization can be rescued by osteoblast-specific overexpression of *Smpd3*, suggesting an important role of nSMase2 in the mineralization of the extracellular matrix (36). Interestingly, the number and area of hypertrophic chondrocytes were significantly increased, and conversion of cartilage into bone was retarded in *fro/fro* bone. *Smpd3* knock-out mice also showed fragile bones and dwarfism, with retarded transition of proliferating chondrocytes into hypertrophic chondrocytes (37, 38). Importantly, the knee joint cartilage of adult *Smpd3*-null mice showed severe deformity with exostosis, the phenotype of OA (38). These findings, from two lines of nSMase2 loss-of-function mice, suggest that *Smpd3*/nSMase2 plays an important role in suppressing hypertrophic maturation of chondrocytes and OA initiation. However, there is currently no information regarding the molecular mechanism for the function of nSMase2 in chondrocyte differentiation. In C2C12 myoblasts, BMP-2-induced Runx2 directly binds to the *Smpd3* promoter to up-regulate expression, suggesting that BMP signaling could elevate *Smpd3* level in a certain context (39).

In this study, we report that *Smpd3* is continuously up-regulated by BMP-2 stimulation of chondrocytes during the maturation stage to suppress the late differentiation step via the Akt pathway. We found that this induction of *Smpd3* by BMP-2 was Runx2-dependent and that the transcription factor is crucial for the hypertrophic maturation of chondrocytes. Loss- and gain-of-function experiments revealed that *Smpd3*/nSMase2 or ceramide signaling cell-autonomously suppressed expression of *Col2a1*, *Col10a1*, and hyaluronate synthase 2 (*Has2*), as well as accumulation of glycosaminoglycan. The Akt-S6 pathway was found to be responsible for the action of *Smpd3*/nSMase2. Importantly, application of an inhibitor compound for nSMase into the mouse bone organ culture system significantly increased the hypertrophic conversion and extracellular matrix calcification of cartilage.

EXPERIMENTAL PROCEDURES

Cell Culture and Differentiation—The mouse chondrogenic cell line ATDC5, established from a differentiated culture of the teratocarcinoma stem cell line AT805 on the basis of chondrogenic potential (40), was obtained from the RIKEN BioResource Center. Human chondrocyte cell line C28/I2 was a kind gift from Dr. Mary Goldring (41). The cells were cultured in Dulbecco's modified Eagle's medium (DMEM)/Ham's F-12 (1:1) (Invitrogen), containing 5% fetal bovine serum (FBS) with 100 units/ml penicillin G and 100 μ g/ml streptomycin. The mouse C3H10T1/2 cell line was obtained from the ATCC. The cells were cultured in Eagle's basal medium (Sigma) with 2 mM L-glutamine, 10% FBS, 100 units/ml penicillin G, and 100 μ g/ml

streptomycin. Primary chondrocytes were harvested from 2-day-old mice with a C57BL/6J background. Briefly, articular cartilage of the femoral heads, femoral condyles, and tibial plateau was isolated from mice and digested by 3 mg/ml collagenase D (Roche Applied Science) for 45 min, followed by 0.5 mg/ml collagenase D overnight. The chondrocytes were filtered through a sterile 40- μ m cell strainer and cultured in DMEM/Ham's F-12 (1:1) containing 10% FBS, 100 units/ml penicillin G, and 100 μ g/ml streptomycin. Differentiation of the cells, cultured in monolayer, was induced by the addition of recombinant human BMP-2 (PeproTech) at a concentration of 300 ng/ml, with or without insulin/transferrin/selenium (ITS) supplement (Sigma) on collagen type I-coated culture plates (Iwaki).

Embryonic Bone Organ Culture—Metatarsal bone rudiments were harvested from C57BL/6J mouse embryos at 16.5 days post-coitum (E16.5) and cultured in DMEM/Ham's F-12 (1:1) supplemented with 10% FBS, and 100 units/ml penicillin G, and 100 μ g/ml streptomycin, as described (42). Cultured bones were stained with Alcian blue and alizarin red dyes according to a standard protocol for skeletal preparation. Briefly, bones fixed in 96% ethanol were stained with 0.015% Alcian blue 8GX (Sigma) in a mixture solution of 96% ethanol/acetic acid (4:1) for 1 day, followed by a dehydration step in 100% ethanol. Dehydrated bones were immersed briefly in 1% potassium hydroxide (KOH), followed by staining in 0.001% alizarin red S (Sigma) in 1% KOH for 1 day. Images were captured with stereomicroscope M165FC (Leica). Four bones per group were analyzed. Animal experiments were approved by the Institutional Animal Care and Use Committee of Kagoshima University (number MD12043).

RNA Interference—Dharmacon siRNA ON-TARGETplus SMARTpool, a mixture of four independent siRNAs against mouse *Runx1*, *Runx2*, *Runx3*, *Smpd3*, or *Has2*, and the control reagent were purchased from Thermo Scientific. siRNAs were transfected into cells using Lipofectamine RNAiMax (Invitrogen). BMP-2 and compounds were added to the culture simultaneously after an overnight transfection of siRNA.

Plasmids and Adenovirus—Mouse *Smpd3* cDNA was cloned from ATDC5 by employing an RT-PCR-based technique, subcloned into the entry vector, pENTR, and further transferred into the C-terminally V5-tagged expression vector, pEF-DEST51, by attL-attR recombination (Invitrogen). To generate adenovirus-carrying *Smpd3* cDNA, the *Smpd3* gene in the pENTR-*Smpd3* vector was transferred into the C-terminally V5-tagged adenovirus expression vector pAd/CMV/V5-DEST by attL-attR recombination (Invitrogen) and further transfected into the adenovirus-producing cell line 293A according to the manufacturer's protocol. pAd/CMV/V5-GW/lacZ adenovirus expression vector was used as a control to express β -galactosidase. Adenovirus infection into ATDC5 cells was performed at a multiplicity of infection of 10. These experiments were approved by the Kagoshima University safety control committee for gene recombination techniques (number 22053).

Chemical Inhibitor Compounds and C_2 -ceramide—All of the following agents were resolved in dimethyl sulfoxide (DMSO), and DMSO was used as the mock control. Cycloheximide was

purchased from Sigma and applied at a concentration of 10 μ M for 2 h, prior to BMP-2 stimulation. The nSMase inhibitor, GW4869 (Sigma), or C_2 -ceramide (Enzo Life Sciences) was applied at the same time of BMP-2 induction, at 1 or 10 μ M, respectively. The PI3K inhibitor, LY294002 (Sigma), Akt inhibitor, MK2206 (Chemie Tek), and the mammalian target of rapamycin inhibitor (Sigma) were applied at the same time of BMP-2 induction, at the indicated concentrations.

Immunoblotting and RTK Signaling Antibody Array Analysis—Cells were lysed in M-PER lysis buffer (Thermo Scientific) supplemented with aprotinin, sodium orthovanadate, and phenylmethylsulfonyl fluoride (PMSF) and subjected to SDS-PAGE, membrane transfer, and chemiluminescence, using standard protocol. Blots were incubated with the following: anti-Runx2 (1:1,000, 8G5, MBL); anti-nSMase2 (1:500, H-195, Santa Cruz Biotechnology); anti-aggrecan (1:500, H-300, Santa Cruz Biotechnology); anti-collagen type II α 1 (1:1,000, LSBio); anti-phospho-Akt (Ser-473) (1:1,000, 587F11, Cell Signaling); anti-Akt (1:1,000, 5G3, Cell Signaling); anti-phospho-S6 ribosomal protein (Ser-235/236) (1:1,000, D57.2.2E, Cell Signaling); anti-S6 ribosomal protein (1:1,000, 54D2, Cell Signaling); anti-phospho-PI3K (1:1,000, number 4228, Cell Signaling); anti-PI3K (p85) (1:1,000, number 610045, BD Transduction Laboratories); anti-phospho-Smad1/5/8 (1:1,000, number 9511, Cell Signaling); anti-Smad1 (1:1,000, number 9743, Cell Signaling); and horseradish peroxidase (HRP)-conjugated anti-rabbit secondary antibody, anti-mouse secondary antibody (1:10,000) (Cell Signaling), or anti-tubulin antibody (1:1,000, DM1A, T9026, Sigma). Signals were detected using the LAS 4000 Mini Image Analyzer (Fujifilm). We employed PathScan[®] RTK signaling antibody array kit (Cell Signaling) to analyze the signaling pathway influenced by loss of *Smpd3*/nSMase2 function. This system is a slide-based antibody array, founded upon the sandwich immunoassay principle and allowing for the simultaneous detection of 28 receptor tyrosine kinases and 11 important signaling nodes when phosphorylated at tyrosine or other residues, was employed to analyze the signaling pathway influenced by loss of *Smpd3*/nSMase2 function. The experiment was performed according to the manufacturer's manual, and the chemiluminescent readout was performed with LAS 4000 Mini Image Analyzer.

Immunocytochemistry and Immunohistochemistry—For immunocytochemistry, cells were fixed with 4% paraformaldehyde in PBS for 30 min and treated with 0.2% Triton X-100. CAS block (Zymed Laboratories Inc.) was used for blocking. For immunohistochemistry, formalin-fixed mouse E17.5 embryo humeri were embedded in paraffin blocks, which were sliced at a 4- μ m thickness. The antigen was retrieved by the Liberate Antibody Binding solution (Polysciences). A CAS block was used for blocking. Cells or bone sections were incubated with anti-aggrecan (1:100, H-300, Santa Cruz Biotechnology), anti-collagen type II α 1 (1:100, LSBio), anti-nSMase2 rabbit polyclonal antibody (1:100, H-195, Santa Cruz Biotechnology), and anti-Has2 mouse monoclonal antibody (1:100, D-8, Santa Cruz Biotechnology). Anti-mouse Alexa Fluor 488 (1:200, A11001) or anti-rabbit Alexa Fluor 568 (1:200, A11011) (Invitrogen) was used to detect signals. Normal rabbit or mouse IgG was used as negative control. Hyaluronan (HA) detection

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involved the following protocol. Samples were blocked with a streptavidin/biotin blocking kit (Vector Laboratories), according to the manufacturer's protocol, and digested sequentially with 0.05% of trypsin and 1 unit/ml of chondroitinase ABC (Sigma). Samples were further blocked with CAS block and sequentially incubated with biotin-conjugated HA-binding protein (1:1,000, Hokudo) and 1 μ g/ml Alexa Fluor 488-conjugated streptavidin (Invitrogen). Images of immunocytochemistry and immunohistochemistry were captured with microscope AX80 and digital camera DP70 (Olympus). Animal experiments were approved by the Institutional Animal Care and Use Committee of Kagoshima University (number MD12043).

Real Time Quantitative PCR Assay—Cells were lysed with the TRIzol reagent (Invitrogen) to purify RNA, and 1 μ g of RNA was reverse-transcribed into cDNA using the Verso cDNA kit (Thermo Scientific). The relative amount of gene transcripts was determined by real time PCR using the SYBR premix Ex TaqII (Takara) and the Thermal Cycler Dice TP850 (Takara). PCRs were performed in duplicate per sample, and the measured expression level of each gene was normalized to that of *Hprt1*. ΔC_t values were calculated by subtracting C_t values of *Hprt1* from C_t values of target genes. Sequence information of primers used is listed in supplemental Table 1.

TUNEL Assay—For detecting ATDC5 cells that underwent apoptosis, we used the ApopTag[®] peroxidase *in situ* apoptosis detection kit (Merck), which detects apoptotic cells *in situ* by labeling and detecting DNA strand breaks by the TUNEL method. The apoptotic cells were visualized by immunoperoxidase staining. Four independent experiments were performed, and four fields per well were evaluated for the number of apoptotic cells.

Statistical Analysis—Data in this study are expressed as mean \pm S.D. of three independent experiments, unless otherwise noted. The statistical comparisons between the different treatments were performed using an unpaired Student's *t* test in which $p < 0.05$ was considered significant and $p < 0.01$ was highly significant.

RESULTS

***Smpd3* Is Induced by BMP-2 in Chondrocytes and Is Detected in Mature Chondrocytes in Vivo**—We hypothesized that BMP signaling cell-autonomously activated unknown mechanisms to terminate BMP-induced chondrocyte differentiation through a negative feedback pathway. We focused on the *Smpd3* gene, because it could be up-regulated by BMP-2 in C2C12 myoblasts of mesenchymal origin (39); loss-of-function models for *Smpd3* in mice showed an increased number of hypertrophic chondrocytes or retarded transition of proliferative chondrocytes into hypertrophic chondrocytes. Molecular mechanisms for both cartilage phenotypes are unclear (37, 38), suggesting a relationship between BMP signaling and *Smpd3*/nSMase2 in chondrogenesis. To investigate the possible cell-autonomous roles of *Smpd3* in chondrocyte maturation, we mainly employed the clonal chondrogenic mouse cell line ATDC5 because it is an excellent *in vitro* model for skeletal development, which can be stimulated by BMP-2 (43, 44). We also used the normal human chondrocyte cell line C28/I2, mouse chondrogenic cell line C3H10T1/2, and mouse primary

articular chondrocytes. We used BMP-2 to stimulate chondrogenic differentiation because, in addition to existing evidence, we and others have observed strong expression of the BMP-2 protein in proliferating and mature chondrocytes in the developing bones of rodents (45, 46). We first checked if these cell lines could differentiate into chondrocytes and mature into hypertrophic chondrocytes and not fibrochondrocytes. Upon BMP-2 induction, ATDC5 cells could produce chondrocyte-specific proteins, aggrecan and type II collagen, in the extracellular matrix (Fig. 1A). The early chondrocyte differentiation marker aggrecan (*Acan*) was elevated after day 1 and was maintained at elevated levels for 2 weeks, whereas *Col10a1* increased mildly from day 3 and was strongly up-regulated after day 7. *Col1a1* and *Col3a1* gene expression was not elevated, but rather decreased, suggesting that these cells did not differentiate into fibrochondrocytes (Fig. 1B). Similar results were obtained in C28/I2 cells (Fig. 1, C and D) and C3H10T1/2 cells (Fig. 1E). We also confirmed chondrogenic differentiation of primary chondrocytes by BMP-2 (Fig. 1F). ΔC_t values of quantitative PCR data indicated that primary chondrocytes express higher levels of chondrocyte marker genes than the three chondrogenic cell lines, suggesting that primary chondrocytes are already committed and differentiated into chondrocytes without BMP-2 induction, whereas the cell lines are relatively immature to differentiate into chondrocytes upon BMP stimulation. From these expression profiles, we considered these cell lines to be suitable models for chondrocyte differentiation. In ATDC5 cells, *Smpd3* showed a varied expression pattern from that of *Acan* and *Col10a1*, in that it was strongly elevated from day 1 and showed a second peak of increase after day 7 (Fig. 1G). We also observed the BMP-2-induced increment of the *Smpd3* gene in C28/I2, C3H10T1/2, and primary chondrocytes (Fig. 1H). Importantly, the primary chondrocytes also expressed higher levels of *Smpd3*, suggesting its functional importance in chondrocytes. If *Smpd3*/nSMase2 has an *in vivo* role in chondrogenesis or cartilage maintenance, it would be expressed in cartilage, although its expression in cartilage has not been characterized in detail. Quantitative PCR analysis on a panel of cDNAs from multiple tissues generated from 3-month-old mice revealed that *Smpd3* was expressed at the highest levels in the thymus, intestine, skin, fat, and bone, although it was almost absent in the heart, liver, and kidney (Fig. 1I). *Smpd3* was moderately expressed in cartilage tissue. This expression profile of *Smpd3* is essentially similar to the results reported for 2-week-old mice (36). In embryonic bone, we detected prominent expression of *Smpd3*-coding nSMase2 protein in the bone collar and trabecular bone by immunofluorescence (Fig. 1J). In the cartilage, little nSMase2 expression was noted in resting and columnar proliferating chondrocytes, whereas relatively high expression was observed in the prehypertrophic layer and hypertrophic chondrocytes (Fig. 1J), with an expression pattern resembling that of Runx2 (4).

BMP-2-induced Increase of *Smpd3* Expression Is Runx2-dependent—We wondered if this BMP-2-induced expression of *Smpd3* in ATDC5 was directed by the BMP-Smad signaling axis, although the expression pattern was not the typical rapid pattern of a gene that is directly targeted (47). To test this, we treated ATDC5 cells with cycloheximide (CHX), an inhibitor of

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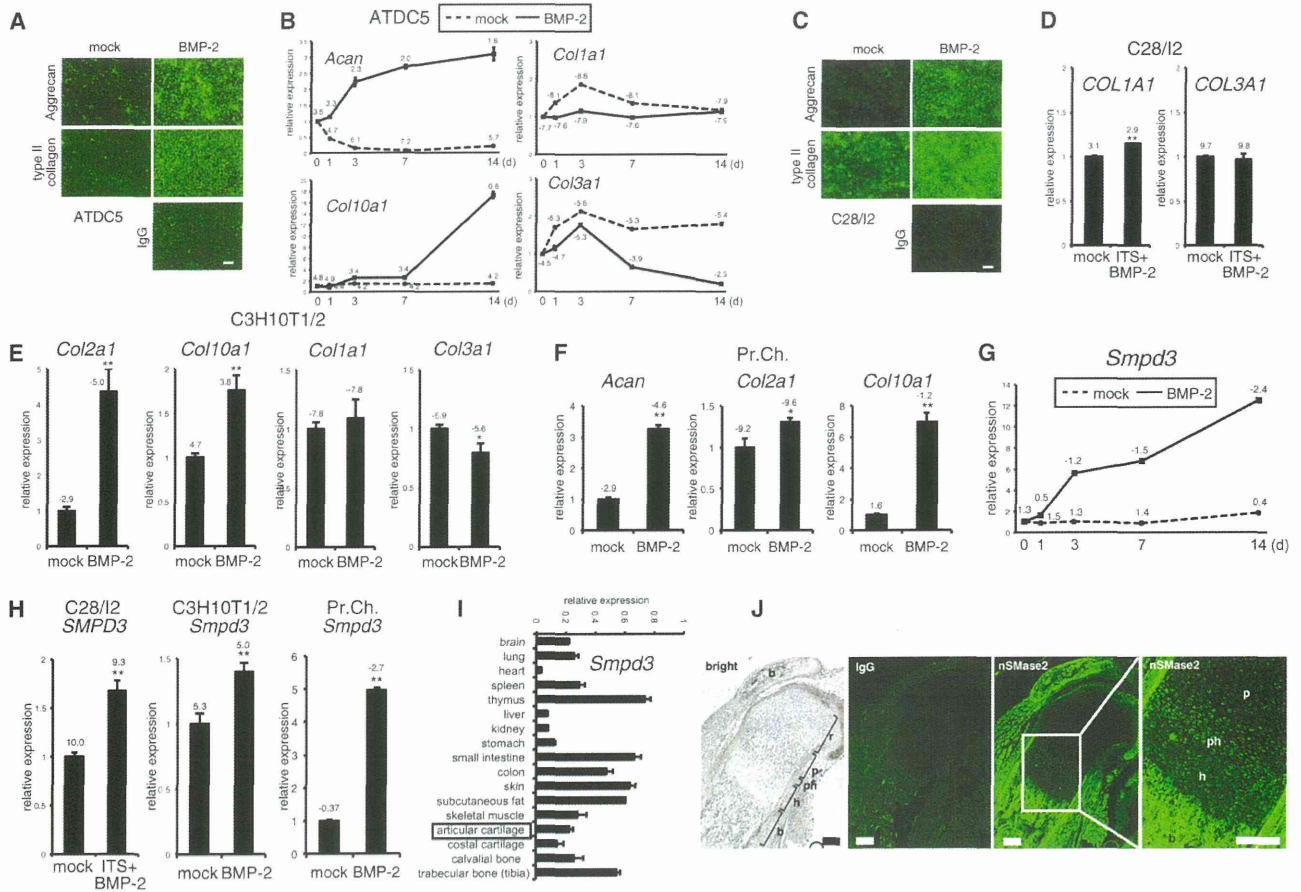


FIGURE 1. Expression of *Smpd3*/nSMase2 is promoted by BMP-2 treatment in maturing chondrocytes *in vitro* and is increased in prehypertrophic and hypertrophic chondrocytes in a growth plate *in vivo*. *A*, chondrogenic differentiation of mouse ATDC5 chondrocytes was induced by application of BMP-2 (300 ng/ml) for 7 days (*d*). Immunofluorescence for aggrecan or type II collagen was performed, with normal IgG as negative control. Scale bar, 100 μ m. *B*, ATDC5 cells were cultured in the presence of BMP-2 (300 ng/ml) for the indicated periods. Expression levels of *Acan*, *Col10a1*, *Col1a1*, and *Col3a1* were examined by quantitative RT-PCR. *C*, chondrogenic differentiation of human C28/12 chondrocytes was induced by the application of BMP-2 (300 ng/ml) for 7 days. Immunocytochemistry for aggrecan or type II collagen was performed, with normal IgG as negative control. Scale bar, 100 μ m. *D*, C28/12 chondrocytes were cultured in presence of BMP-2 (300 ng/ml) and ITS supplement for 14 days. Expression of *COL1A1* or *COL3A1* was evaluated by quantitative RT-PCR. *E*, mouse C3H10T1/2 cells were cultured with BMP-2 (300 ng/ml) for 48 h. Expression levels of *Col2a1*, *Col10a1*, *Col1a1*, and *Col3a1* were evaluated by quantitative RT-PCR. *F*, mouse primary chondrocytes (*Pr.Ch.*) cells were cultured in the presence of BMP-2 (300 ng/ml) for 6 days. Expression levels of *Acan*, *Col2a1*, and *Col10a1* were evaluated by quantitative RT-PCR. *G*, ATDC5 cells were cultured in presence of BMP-2 (300 ng/ml) for the indicated periods. Expression of *Smpd3* was examined by quantitative RT-PCR. *H*, quantitative RT-PCR for *Smpd3* was performed on samples in *D*, *E*, and *F*. *I*, real time PCR for *Smpd3* was performed on a tissue cDNA panel of a 3-month-old mouse. *J*, expression of nSMase2 in mouse E17.5 humerus cartilage was evaluated by immunofluorescence. Normal IgG was used as negative control. *r*, resting chondrocytes; *p*, proliferating chondrocytes; *ph*, prehypertrophic chondrocytes; *h*, hypertrophic chondrocytes; *b*, bone. Scale bar, 200 μ m. *, $p < 0.05$; **, $p < 0.01$. ΔC_t values (C_t target – C_t Hprt1) of quantitative RT-PCR are indicated in the graphs.

de novo protein synthesis, before adding BMP-2. As expected, 24 h after BMP-2 stimulation, the induction of *Id1*, a representative direct-target gene of the BMP-Smad pathway, was maintained, even in the presence of CHX (Fig. 2A). Induction of *Id1* was higher in CHX-treated cells, likely because of the suppression of inhibitory Smad6 synthesis (27). However, expression of *Col2a1* was eliminated by CHX treatment (Fig. 1A). This basal suppression by CHX probably results from an inhibition of the constitutive expression of Sox9 and downstream Sox5 and Sox6, transcription factors that cooperatively activate the promoter of *Col2a1* (2, 48). However, existing Sox proteins should be responsible for the partially increased expression of *Col2a1* by BMP-2. The basal expression level of *Smpd3* was also suppressed by CHX treatment. Unlike *Col2a1*, however, the CHX-eliminated basal expression of *Smpd3* was not up-regulated by BMP-2 (Fig. 2A) suggesting that a *de novo* protein other than

the Sox trio was necessary for BMP-induced expression of *Smpd3* in ATDC5 cells. Because the Runx2 protein is a master regulator of chondrocyte maturation (4, 5), and its direct interaction with the *Smpd3* promoter is important for its expression in myoblasts (39), we investigated the relationship between Runx2 and expression of *Smpd3* in chondrocytes. We confirmed the increment of the *Runx2* gene during BMP-induced maturation of ATDC5 chondrocytes (Fig. 2B). To address this question, we transfected ATDC5 cells with *Runx2* siRNA. At day 2 of BMP-2 application, *Runx2* was weakly induced, and its expression was knocked down by the siRNA to about 40% that of the control (Fig. 2C). Expression of *Id1* was not decreased by siRunx2, but rather it was increased, suggesting that Runx2 is inhibitory of the BMP pathway during this early differentiation stage. Interestingly, loss of Runx2 not only suppressed basal expression of *Smpd3* but also completely blocked induction by

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