The Antagonist of Retinoic Acid Receptor α, ER-50891 Antagonizes the Inhibitive Effect of All-Trans Retinoic Acid and Rescues Bone Morphogenetic Protein 2-Induced Osteoblastogenic Differentiation

Siqian Wang, 1,* Wenjuan Bi, 2,* Yi Liu, 3,‡ Jiayi Cheng, 4 Wei Sun, 5 Gang Wu, 6 Xin Xu 1

1Department of Implantology, School and Hospital of Stomatology, Shandong University & Shandong Key Laboratory of Oral Tissue Regeneration & Shandong Engineering Laboratory for Dental Materials and Oral Tissue Regeneration, Jinan, Shandong Province, People’s Republic of China; 2College of Stomatology, North China University of Science and Technology, Tangshan, Hebei Province, People’s Republic of China; 3Key Laboratory of Oral Medicine, Guangzhou Institute of Oral Disease, Stomatology Hospital of Guangzhou Medical University, Guangzhou Medical University, Guangzhou, People’s Republic of China; 4Department of Oral Implantology and Prosthetic Dentistry, Academic Centre for Dentistry Amsterdam (ACTA), University of Amsterdam (UvA) and Vrije Universiteit Amsterdam (VU), Amsterdam, The Netherlands; 5The Affiliated Stomatology Hospital, Zhejiang University School of Medicine, Hangzhou, Zhejiang Province, People’s Republic of China

*These authors contributed equally to this work

Correspondence: Gang Wu
Department of Oral Implantology and Prosthetic Dentistry, Academic Centre for Dentistry Amsterdam (ACTA), University of Amsterdam (UvA), Gustav Mahlerlaan 3004, Amsterdam 1081LA, The Netherlands
Tel +31 20 5980866
Fax +31 20 5980333
Email g.wu@acta.nl

Xin Xu
Department of Implantology, School and Hospital of Stomatology, Shandong University & Shandong Key Laboratory of Oral Tissue Regeneration & Shandong Engineering Laboratory for Dental Materials and Oral Tissue Regeneration, No. 44-1, Wenhua Xi Road, Jinan, Shandong 250012, People’s Republic of China
Tel/Fax +86-531-88382923
Email xinxu@sdu.edu.cn

Abstract

Background: Hypervitaminosis A, alcoholism or medical treatment for acute promyelocytic leukaemia may cause unphysiologically high accumulation of all-trans retinoic acid (ATRA), which could inhibit osteoblastogenesis, thereby triggering osteoporosis. We have shown that bone morphogenetic protein-2 (BMP-2) can only partially antagonize the inhibitive effects of ATRA. In this study, we hypothesized that antagonists of retinoic acid receptors (RARs) could further antagonize the inhibitive effect of ATRA and rescue BMP2-induced osteoblastogenesis.

Materials and Methods: We first screened the dose-dependent effects of the specific antagonists of RAR α, β and γ and transforming growth factor-beta receptor (ER-50891, LE-135, MM11253, and SB-43142, respectively) on ATRA-inhibited phosphorylated Smad1/5. Neither ATRA, nor ER-50891 or MM11253, and SB-43142 significantly antagonized the inhibitive effects of ATRA. In this study, we hypothesized that antagonists of retinoic acid receptors (RARs) could further antagonize the inhibitive effect of ATRA and rescue BMP2-induced osteoblastogenesis.

Results: ER-50891 but not LE-135, MM11253, or SB-43142 significantly antagonized the inhibitive effect of ATRA and enhanced the total cell metabolic activity and proliferation of preosteoblasts. We selected ER-50891 and tested its effects on osteoblastogenesis with the presence or absence of 1 μM ATRA and/or 200 ng/mL BMP-2. We measured the following parameters: Alkaline phosphatase activity (ALP), osteocalcin (OCN) expression and extracellular matrix mineralization as well as the level of phosphorylated Smad1/5.

Conclusion: The antagonist of RARα, ER-50891 could significantly attenuate ATRA’s inhibitive effects on BMP 2-induced osteoblastogenesis.

Keywords: bone morphogenic protein 2, all-trans retinoic acid, retinoic acid receptor, osteoblastogenesis, transforming growth factor beta

Introduction

Bone tissues with sufficient quantity and quality are highly important for the proper functions of musculoskeletal systems and therein-implanted medical devices, such as dental implants.1 As a paramount biological process to maintain bone tissue and repair bone defects, mesenchymal stem cells are osteogenically committed to become a preosteoblast and thereafter undergo osteoblastogenesis.2 Osteoblastogenesis comprises a series of sequential cellular events, such as proliferation, alkaline phosphatase (ALP) expression (early differentiation marker), osteocalcin (OCN) expression (late

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Apart from breast cancer cells/well in 24-well plates, we also show a metabolic cells/well in 96-well plates for measuring which is at least partially, due to the significantly enhanced level phosphorylated Smad1/5 (p-Smad1/5). By forming a complex with Smad4, P-Smad1/5 translocates into nucleus and binds to DNA promoters so as to induce the expression of osteogenic genes, leading to osteoblastogenesis. Recently, we report that 1 μM ATRA inhibits osteoblastogenesis of both bone marrow stromal cells and mouse calvarial preosteoblast cell (MC3T3-E1 cell line). We also show that ATRA can antagonize the promoting effects of various BMPs on the expression of key osteogenic genes (such as Runx2, Osterix, Dlx5 and Smad1) irrespective of the BMPs’ potencies. However, hitherto, it is unclear which receptor plays a critical role in the inhibition of ATRA on BMP-2-induced osteoblastogenesis.

In this study, we adopted specific antagonists to different RARs and also TGF-β type I receptor kinase with an aim to assess the roles of different receptors in the inhibitive effects of ATRA on BMP-2-induced osteoblastogenesis.

### Materials and Methods

#### Cell Culture

A preosteoblast cell line (MC3T3-E1, subclone 4, CRL-2593) was purchased from the American Type Culture Collection (ATCC). Cells were cultured in growth medium consisting of α-MEM with 10% (v/v) FBS (fetal bovine serum) (Gibco Invitrogen, Grand Island, NY), 100 U/mL streptomycin and penicillin at 37°C and 5% CO₂ in a humidified atmosphere. Media change was performed every 3 days. When 80% confluence was reached, cells were plated at a final concentration of at 5×10⁴ cells/well in 24-well plates for measuring total cell metabolic activity and DNA content; for at 2.5×10⁴ cells/well in 48-well plates for measuring ALP activity, total protein, OCN and extracellular matrix mineralization (alizarin red staining); or 1×10⁴ cells/well in 96-well plates for measuring p-Smad1/5 using cell-based enzyme-linked immunosorbent assay (ELISA). Except for extracellular matrix mineralization, cells were seeded in plates in a low-serum medium (2% FBS); 24 hrs after seeding, cells were first starved in the low-serum media (0.5% FBS) for another 24 hrs and then followed by drug treatment also in media containing 0.5% FBS.
Treatment of Preosteoblasts

ATRA was purchased from Sigma-Aldrich. All the receptor antagonists and BMP-2 (CHO derived) were purchased from R&D Systems, Inc. Cultured preosteoblasts were treated with RARα antagonist ER-50891, RARβ antagonist LE-135, TGF-beta antagonist MM-11253 (0, 1, 2 and 3 (μM)), or TGF-beta antagonist SB-431542 (0, 3, 6 and 9 (μM)), in the absence and presence of 1 μM ATRA, during the entire period of differentiation of MC3T3-E1, respectively. For further studies, cells were treated with 200 ng/mL BMP-2.

Screening Receptor Antagonists

The parameters of total cell metabolic activity and proliferation (DNA content) were measured in order to identify which specific receptor ATRA ligand binds to. After 24 hrs starving culture, the cells were treated with ATRA or antagonist for 3 days. The results show that the ER-50891, RAR-alpha antagonist group had a significant difference in total cell metabolic activity and DNA content. Instead, the other receptor antagonists LE-135, MM-11253, SB-431542 made no significant difference when comparing with the ATRA group, respectively, with a similar effect level. As a consequence, the receptor in this study was determined as RAR-alpha.

ER-50891 Concentration Selection Test

Furthermore, to assess the concentration range for how RAR-alpha antagonist ER-50891 involves in the ATRA signaling cascade, a series of dose–response tests on cell metabolic activity, DNA content, ALP activity, total protein expression, and osteocalcin release were performed as described in the following sections 4 days post-treatment. ER-50891, RAR-alpha antagonist group on the concentration of 0.25 μM, 0.5 μM, 1 μM, 1.5 μM, 2 μM and 3 μM significantly decreased the ALP activity. These results showed that the ER-50891 antagonized the enhancing effect of ATRA on ALP activity and decreased ALP activity to the similar level as control when 2–3 μM ER-50891 was applied. Besides, ER-50891, RAR-alpha antagonist group has a significant increase in cell proliferation and osteocalcin release. Consequently, the concentrations in this study were determined at 1.5 μM and 3 μM.

Total Cell Metabolic Activity and Proliferation

To analyze the effect of ATRA on MC3T3-E1 proliferation in response to various concentrations of different RAR antagonists ER-50891, LE-135, MM-11253, and TGF-beta antagonist SB-431542, the total cell metabolic activity in the absence or presence of 1 μM ATRA was gauged using the alamar Blue™ cell viability assay (Invitrogen Corporation, Carlsbad, CA, USA) following the manufacturer’s instruction. In short, after a 3-day treatment, the alamar Blue™ reagent was added into the growth medium for an incubation of 4 hrs. After the incubation period, fluorescence intensity, was measured at EX 540 nm/EM 590 nm using SpectraMax M5 Molecular Devices (Sunnyvale, CA, USA). Proliferation of cells were measured using a DNA quantification assay. The QuantiTMT PicoGreen dsDNA Assay kit (Invitrogen, OR, USA) was used according to the instruction. The fluorescent intensity of each sample was measured using SpectraMax M5 Molecular Devices as well with a setting of EX480 nm/EM520 nm.

ALP Activity and Total Protein Expression

After a 3-day treatment, ALP activity and protein expression were measured using para-Nitrophenylphosphate as a substrate using LabAssay™ ALP colorimetric assay kit (Wako Pure Chemicals, Osaka, Japan) according to the manufacturer’s protocol. Briefly, 10 μL aliquots of cell lysates were added to 96-well plates. Thereafter, 50 μL of 2-amino-2-methyl-propanol buffer (pH 10.3) and 50 μL of 15.2 mM paranitrophenylphosphate (PNPP) in a 2 mM solution of magnesium chloride were added and incubated at 37°C for 1 hrs. The reaction was stopped by adding 200 μL stop solution (1 N solution of sodium hydroxide). The amount of produced paranitrophenol was gauged spectrophotometrically at 405 nm using the plate reader SYNERGYMs (BioTek®, Winoski, VT) with Gen5 1.09 software. The total protein content was measured using a BCA Protein Assay kit (Beyotime, China). Results are expressed as folds of ALP activity. ALP activity was calculated by dividing the amount of paranitrophenol by protein content. Results were expressed by calculating the fold changes in comparison with the control.

Osteocalcin Expression

Osteocalcin (OCN), referred to as a late marker of osteoblastic differentiation, was analyzed using an enzyme-linked immunosorbent assay (ELISA) for mouse OCN (EIA kit, Biomedical Technologies, Stoughton, MA, USA) according to the manufacturer’s protocol. After a 3-day treatment, supernatants of cell culture were collected and centrifuged (20,000 rpm, 4°C, 15 min) before analysis. The absorbance was measured at 450 nm. Results are expressed as the fold changes of OCN in comparison with control.
Extracellular Matrix Mineralization

We compared the area of mineralized nodules in samples stimulated by 200 ng/mL BMP-2 in the absence or presence of 1 µM ATRA and ER-50891 (1.5 µM and 3 µM). The mineralized nodules formation was analyzed using Alizarin red staining at the time points of 19 days after induction. Quadruplicate cell cultures were prepared and then treated with an osteogenic medium (10% FBS, 50 µg/mL L-ascorbic acid, and 10mM beta-glycerophosphate; Sigma-Aldrich, St. Louis, MO, USA) containing 200 ng/mL BMP-2 plus 1 µM ATRA, as well as 1.5 µM or 3 µM ER-50891. The medium was replaced every 3 days. On Day 19, extracellular matrix mineralization was determined by alizarin red staining at pH 4.2 (Sigma-Aldrich, St. Louis, MO, USA). Each well was photographed using NIS-Elements F2.20 (Nikon Eclipse 80i, Tokyo, Japan), and the mineralized areas were then quantified using Image-Pro Plus 6.0 software.

Cell-Based ELISA

After a 3-day treatment, P-Smad1/5 was measured using a cell-based ELISA kit (#KCB7660, R&D Systems, Inc., Minneapolis, MN, USA) kit and compared to glyceraldehyde 3-phosphate dehydrogenase (GAPDH) in the nuclear according to the instructions.

Statistical Analysis

All results are shown as means ± standard deviations (SD). Individual experiments were performed at least three times. The data were analyzed using the GraphPad Prism 6 version 6.0 (GraphPad Software Inc, La Jolla, CA). For comparison among groups, one-way analysis of variance (ANOVA) was conducted. For comparison between two groups, unpaired t-test was conducted. p<0.05, was considered a statistically significant difference.

Results

ATRA Inhibited the Total Cell Metabolic Activity and Proliferation of Preosteoblasts, Which Could Be Rescued by the Antagonist of RARα

We first evaluated the influence of the antagonists on the total cell metabolic activity and proliferation of preosteoblasts. 1 µM ATRA caused almost 50% reduction of total cell metabolic activity compared to the control group. The 1 µM or 2 µM RARα antagonist ER-50891 alone significantly promoted the total cell metabolic activity of pre-osteoblasts (Figure 1A). In contrast, RARβ antagonist LE-135 alone at all concentrations resulted in significantly lower total cell metabolic activity (Figure 1B). Neither RARγ nor TGF-β antagonists alone could significantly influence the total cell metabolic activity of preosteoblasts (Figure 1C and D). The inhibitive effect of ATRA could be relieved only by the antagonist of RARα (Figure 1A) but neither RARβ nor RARγ nor TGF-β (Figure 1B–D). A very similar pattern was also found in the effects of the antagonists on the DNA content (Figure 2–D). ATRA significantly inhibited the DNA content. Antagonist of RARα but neither RARβ nor RARγ nor TGF-β significantly antagonized the inhibitive effects of ATRA on the DNA content of preosteoblasts. Since the total cell metabolic activity and proliferation of osteogenic cells are highly important for osteogenesis and only RARα antagonist ER-50891 could antagonize the inhibitive effects of ATRA, we adopted ER-50891 in the following experiments.

ER-50891 Dose-Dependently Antagonized the Promoting Effect of ATRA on ALP Activity and Also Its Inhibitive Effects on Cell Viability, Proliferation, Total Protein and OCN Expression

We performed dose-dependent assay to identify the antagonistic effect of ER-50891 on the effects of ATRA. From 0.5 to 1.5 µM, ER-50891 significantly antagonized the inhibitive effect of ATRA and dose-dependently increased the total cell metabolic activity of preosteoblasts (Supplementary Figure 1A). In the concentration range of 1–3 µM, the total cell metabolic activity was elevated to a similar (1 µM and 3 µM) or significantly higher level (1.5 µM and 2 µM) compared to the control (Supplementary Figure 1A). ER-50891 of all the selected concentrations resulted in a significantly higher level of DNA content than the group of ATRA alone, which reached a similar level as the control (Supplementary Figure 1B). ER-50891 decreased total protein content to 48.0% of the control group. In contrast, ER-50891 dose-dependently increased the total protein content of preosteoblasts (Figure 3A) with protein content recovered to 80.6% of the control group. ATRA increased OCN expression to a similar level as the control (Supplementary Figure 1B). ER-50891 dose-dependently increased OCN expression to a similar level as the control (Supplementary Figure 1B). ER-50891 decreased OCN expression to 80.6% of the control group. In contrast, ER-50891 dose-dependently increased OCN expression to a similar level as the control (Supplementary Figure 1B).
relieved by ER-50891. ER-50891 at 2 and 3 µM almost completely rescued the OCN expression (Figure 3C).

ATRA Promoted ALP Activity and Inhibited OCN Expression Induced by BMP-2, Which Could Be Antagonized by ER-50891

ATRA or 200 ng/mL BMP-2 alone could promote ALP activity by 2.1 and 1.9 folds, respectively, while ER-50891 alone did not significantly influence ALP activity (Figure 4A). ATRA and BMP-2 synergistically promoted ALP activity by 18.9 folds in preosteoblasts, which could be significantly inhibited by ER 50891. BMP-2 significantly enhanced OCN by 5.9 folds in comparison with the control, which was not influenced by ER-50891. ATRA could significantly suppress OCN expression by 84.2%, which could be significantly relieved to 41.1% of the control group by ER-50891 (Figure 4B). ATRA also inhibited BMP-2-induced OCN expression by 90.0%. ER-50891 could antagonize such an effect of ATRA, but only rescue the OCN expression to a similar level as the control (Figure 4B).

ER-50891 Attenuated the Inhibitive Effects of ATRA on BMP-2-Induced Extracellular Matrix Mineralization

Nineteen days after treatment, treatment with BMP-2 significantly promoted extracellular matrix mineralization than the control (no ATRA, no ER-50891; Figure 5A). In
contrast, ATRA significantly decreased this parameter both in the absence (by 65.1%) and presence (by 93.9%) of BMP-2. ER-50891 alone did not significantly influence extracellular matrix mineralized nodules (Figure 5B). ER-50891 could rescue the formation of extracellular matrix mineralization to a level that was similar to the BMP-2 group (Figure 5B).

Phosphorylated-Smad1/5 Was Not a Target of Either ATRA or ER-50891
BMP-2 could significantly update the level of phosphor-Smad1/5, while ATRA alone did not influence phosphor-Smad1/5 (Figure 6). Furthermore, ATRA could not significantly influence BMP-2 induced elevation of phosphor-Smad1/5. Similarly, ER-50891 did not influence phosphor-Smad1/5 both in the absence or presence of BMP-2 (Figure 6).

Discussion
ATRA has long been found as an important morphogen in regulating various biological events.\textsuperscript{40} Alcoholism, hypervitaminosis A or APL treatment can result in unphysiologically overdosed ATRA that might lead to osteoporosis by, at least partially, decreasing osteoblastogenesis.\textsuperscript{37,38} BMPs are a group of paramount growth factors for inducing physiological osteoblastogenesis and also widely used to promote bone regeneration in clinic.\textsuperscript{35,36} In our previous study, we demonstrate that ATRA can significantly antagonize the promoting effect of heterodimeric BMP2/7, a more potent form of BMPs, on osteoblastogenesis of preosteoblast cell line\textsuperscript{18}.
Figure 3 Fold changes of total protein content (A), ALP activity (B), and osteocalcin (C) of murine calvarial pre-osteoblasts (MC3T3-E1 cells) treated with or without 1 μM ATRA in presence or absence of different concentrations of RAR-alpha antagonist ER-50891. All data are presented as mean values together with the standard deviation. *p<0.05; ***p<0.001.

Figure 4 Fold changes of ALP activity (A) and osteocalcin (B) of murine calvarial pre-osteoblasts (MC3T3-E1 cells) treated with or without 1 μM ATRA and 200 ng/mL BMP-2 in presence or absence of 1.5 μM RAR-alpha antagonist ER-50891. All data are presented as mean values together with the standard deviation. *p<0.05; ***p<0.001.
and primary BMSCs. We further show that the inhibitive efficacy of ATRA is independent on the potencies of BMPs. However, it remains unveiled which receptor mediates the inhibitive effect of ATRA osteoblastogenesis in the absence or presence of BMP-2. In this study, we, for the first time, reported that the antagonist of RARα but not RARβ or RARγ or TGF-β receptor significantly suppressed the inhibitive effects of ATRA and rescued osteoblastogenesis. This finding suggested that it was RARα that mediated the inhibitive effects of ATRA.

ATRA exerts a large variations of effects through different RARs. RARα, also known as NR1B1, is encoded by the RARA gene and involved in many biological events, such as multicellular development, proliferation, differentiation and apoptosis. Enhanced expression of RARA could induce epithelial-to-mesenchymal transition. After decade’s efforts, ATRA has been developed to aim at PML-RARα and newly discovered NLS-RARα oncoproteins as direct molecular targets based on epigenetic discoveries for APL involved HHEX and S100A10. RARβ is encoded by RARB gene and has been considered to play a distinct role in the tumor growth and development. The RARα and RARβ can bind to RAREs to modulate target gene expression. RARγ, encoded by RARG gene, is associated with skin, bone and teratogenic toxicity and is critical for hematopoiesis in modulating the balance between self-renewal and differentiation of hematopoietic stem cells (HSCs). RAR signaling directly regulates osteoblastogenesis and adipogenesis. RARγ activity in mesenchymal stem cells plays a role in regulating endochondral bone. Green et al report that RARγ is a negative regulator of osteoclastogenesis in vivo and in vitro. However, hitherto, there are few publications reporting the involvement of different receptors in the ATRA’s inhibitive effects of BMP-induced osteoblastogenesis.

Under the inhibitive effects of ATRA, mesenchymal cells or pre-osteoblasts are reported to exhibit a partially differentiated non-proliferating state. This character is
symbolized by enhanced ALP activity (early osteoblastogenic marker) as well as suppressed cell proliferation/viability, OCN expression and extracellular matrix mineralization.\(^6,37,38,52\) However, hitherto, it remains unclear how ATRA leads to these phenomena. In this study, we showed that ATRA suppressed both total cell metabolic activity but also the proliferation (DNA content). ATRA resulted in a 30–50% reduction of DNA content and a 45–55% reduction of total cell metabolic activity. Therefore, the ATRA-induced reduction of total cell metabolic activity should be largely attributed to the inhibition of cell proliferation. It has long been established that retinoids inhibit cell proliferation by causing G1 arrest.\(^32,53\) Such an effect of ATRA may be mediated via many mechanisms, among which endogenous TGF-βs have been shown to mediate the anti-proliferative effect of ATRA.\(^32–34\) In this study, we showed that SB-431542, a specific inhibitor of TGF-β type I receptor kinase, alone did not significantly influence cell viability or proliferation and could not antagonize the inhibitive effect of ATRA. This was complied with a previous publication that SB431542 showed no significant influence in the proliferation and osteogenic differentiation of pre-osteoblasts.\(^54\) These results suggested that endogenous TGF-β was not a mediator for ATRA’s effect. Similar effect pattern was also found to MM-11253, RARγ antagonist. LE-135, antagonist of RARβ alone even also suppressed total cell metabolic activity and proliferation, which suggested that endogenous RARβ-mediated signaling was a positive regulator for total cell metabolic activity and proliferation.

In contrast, RARα antagonist alone significantly promoted the total cell metabolic activity and proliferation of preosteoblasts, which suggested that endogenous RARα-mediated signaling was a negative regulator for cell proliferation and total cell metabolic activity. Furthermore, under the presence of ATRA, RARα antagonist also not only restored but also even promoted these two parameters to a level higher than the control, which suggested that RARα mediated the suppressive effects of ATRA on cell proliferation and total cell metabolic activity. In our subsequent dose-dependent assays, RARα antagonist significantly antagonized the effects of ATRA on both promoting ALP activity and also suppressing OCN expression and mineralization. These findings suggested that RARα was one of the critical mediators for ATRA-induced inhibition of osteoblastogenesis.

In our previous publication,\(^2\) we reported that the heterodimeric BMP2/7 induced osteoblastogenesis at significantly lower optimal concentrations than homodimeric BMP2. BMP2/7 reached the maximal effect and was advantageous over BMP2 at 50ng/mL. Therefore, in our further publication, we adopted BMP2/7 to antagonize the inhibitory effects of ATRA on osteoblastogenesis.\(^37,38\) However, in our most recent publication, we showed that although BMP2/7 was not advantageous over BMP2 in antagonizing the inhibitory effects of ATRA.\(^39\) Consequently, we adopted 200ng/mL BMP2 in this study. The other reason to adopt BMP2 is that BMP2 combined with collagen membrane is an FDA-approved medical device and can be used in the clinic. Therefore, BMP2 is of more significance than BMP2/7 in clinical treatment. Hitherto, it remains largely unknown how ATRA suppresses BMP-induced osteoblastogenesis. One potential mechanism is that ATRA significantly decreased the signal duration of p-Smad1/5 by promoting ubiquitination and proteasomal degradation.\(^55\) However, in this study, we showed that neither ATRA nor RARα nor their combination significantly affected the level of p-Smad1/5 induced by BMP-2. Recently, we have reported that ATRA could significantly inhibit the directly-upregulating genes (such as DLX5) of BMP signaling.\(^39\) Furthermore, we also found that the translocation of p-Smad1/5 to nuclei was not significantly affected by ATRA (data not shown). All these findings suggested that ATRA inhibited BMP-2 signaling and its-induced osteoblastogenesis in translational level, such as the binding of p-Smad1/5 to its targeting gene promoters. Further studies should be performed to investigate potential mechanisms.

In this study, we showed that ATRA enhanced ALP activity. This was consistent with previous studies that ATRA showed either no or promotive effect on ALP activity on various osteogenic cells, such as MC3T3-E1 preosteoblasts, rat UMR-201-10B preosteoblast cell line, C3H10T1/2 mesenchymal stem cells and primary mouse bone marrow stromal cells.\(^37,56–58\) Please note that the effect of ATRA on ALP gene expression is not through the classic osteogenic signaling pathways, such as promoting endogenous BMPs. Instead, this modulation of ATRA was shown to be highly dependent on a retinoid acid response element in the promoter of the ALP gene.\(^58\) Consequently, the ALP activity under the treatment of ATRA should be not considered as a conventional early osteogenic differential maker. In this study, our data showed that ATRA and BMP could synergistically enhance ALP activity to 18 folds of the control. The synergistic effect of ATRA and BMP is consistent with the previous reports from our and other researcher.\(^37,59\) However, in the later report, RARγ (but not
RARRα or RARRβ-dependent signaling can interact with BMP2 to regulate osteoblastic differentiation of C2C12 cells. In contrast, in our results, the antagonist of RARRα but not RARRγ suppressed the effects of ATRA. The inconsistency may be due to the different functions of RARRs in different cell types. Please note that the promoting effect of ATRA on ALP gene expression is realized not by activating the classic osteogenic signaling pathways, such as enhancing the expression of endogenous BMPs. Instead, such an effect of ATRA is shown to be highly related to a retinoid acid response element (RARE) in the promoter of the ALP gene. Consequently, ALP promoted by ATRA or the combination of ATRA and BMP-2 might not be generally regarded as a marker of osteoblastogenic differentiation. Also, the statement that ATRA induces a partially differentiated status of cells might not be very precise. Although ATRA-induced ALP activity, it failed to enhance OCN expression and extracellular matrix mineralization. Thus, ATRA inhibited osteogenesis.

Interestingly, RARRα antagonist could only partially restore BMP-induced OCN expression to the same level as the control group. In fact, the osteocalcin level associated with the treatment of ATRA, ER and BMP-2 was significantly higher than that associated with the treatment of ATRA and BMP-2. However, such an effect of ER on OCN seemed not to be as dramatic as on ALP activity, which might suggest the involvement of other signaling mechanisms in the inhibition of OCN by ATRA. In this study, RARRα antagonist was selected largely basing on its antagonistic effect on the inhibited cell proliferation by ATRA rather than on osteogenic differentiation. There might be also a possibility that other RAR receptors than RARRα were involved in the inhibited osteogenic differentiation by ATRA. For example, RARRγ was also shown to negatively influence BMP-2-induced bone formation. Consequently, further studies should be performed to reveal the inhibitory mechanisms of ATRA on osteocalcin expression. This might be one of the reasons that RARRα antagonist could not completely restore extracellular matrix mineralization. Consequently, in future studies, OCN may be used as a key parameter to check the effects of different receptor antagonists. Probably, a combination of two antagonists can completely nullify the inhibitory effect of ATRA on BMP-induced osteogenesis.

Previous cell studies show conflicting results regarding the role of retinoids on osteoblast differentiation. It is suggested that, to some extent, the effect of ATRA is dependent on the RA’s concentration. Nanomolar ATRA is reported to inhibit osteoblastogenic differentiation and functions, whereas micromolar ATRA stimulates osteoblastogenic differentiation. However, the dose-dependent mechanisms are not sufficient to explain all the phenomena observed in the previous studies.

One limitation in this study was that we only adopted a murine-derived preosteoblast cell line. Current results need to be further validated in primary cells. Besides, in-vivo study is also needed to show the interaction of ATRA and RAR-alpha antagonist in modulating in vivo BMP2-induced new bone formation.

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Disclosure

The authors declare no conflict of interests.

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