Reciprocal Interferences of TNF-α and Wnt1/β-Catenin Signaling Axes Shift Bone Marrow-Derived Stem Cells Towards Osteoblast Lineage after Ethanol Exposure

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Key Words
Ethanol • Multipotential mesenchymal stem cells (MSC) • Bone-marrow stem cells (BMSC) • Osteoblast • Osteogenesis • Adipogenesis • TNF-α • Wnt1 • β-catenin • siRNA

Abstract
**Background/Aims:** We have reported in a separate study that alcohol exposure triggers activation of the TNF-α signaling pathway leading to an adverse shift of multipotential mesenchymal stem cells in bone marrow (BMSCs) away from osteogenesis towards adipogenesis. However, inhibition of TNF-α signaling only yielded moderate inhibition of adipogenesis. Here we showed that in addition to promoting the TNF-α signaling, alcohol also suppressed the Wnt1/β-catenin signaling pathway. **Methods:** We treated primary BMSCs from human subjects with alcohol for 24 days. We measured changes of genes related to endoplasmic reticulum (ER) stress, adipogenic markers and osteogenic markers using quantitative real-time RT-PCR and Western blot analysis. We performed Alizarin red staining for osteogenesis. We also conducted assays for osteogenic biomarkers alkaline phosphatase, collagen-I and osteocalcin. **Results:** Wnt/β-catenin signaling was markedly activated in BMSCs treated with osteogenic inducers relative to the control cells, as indicated by the increased levels of nuclear β-catenin along with reduced levels of cytosolic β-catenin, as well as increased protein levels of Wnt1. Activation of Wnt/β-catenin signaling was significantly suppressed in BMSCs exposed to alcohol, which was reflected by downregulated expression of osteogenic marker genes Osf2/Cbfa1, osteopontin and osteocalcin, upregulated adipogenic marker PPARγ2 and aP2, and reduced number of calcification nodules. In contrast, activation of Wnt/β-catenin signaling by BIO favored osteogenesis even in the presence of alcohol. Simultaneous activation of Wnt1 by BIO and inhibition of TNF-α by 3,6′-dithiothalidomide
produced synergetic suppression of ethanol-induced adipogenic lineage compared to interference with either of them alone. **Conclusion:** This remarkable shift of BMSCs towards osteoblast lineage suggests the superiority of concordant and reciprocal interferences of the TNF-α and Wnt/β-catenin pathways for promoting osteogenesis.

**Introduction**

Habitual consumption of excessive quantities of alcohol is commonly recognized as a major factor for osteopaenia and increased incidence of bone fracture in alcoholics [1–3]. The current consensus of both clinical and experimental studies is that alcohol-induced bone loss is mainly due to the suppression of new bone formation with only relatively small changes (increase or decrease) occur in bone resorption [4]. In adults, the osteoblast, which is responsible for bone formation, is derived from multipotential mesenchymal stem cells in bone marrow (BMSC) [5–8]. The capacity and dynamics of these BMSCs to differentiate into osteoblasts plays a critical role in the cellular processes involved in the maintenance of the adult human skeleton.

The BMSCs have been proven to differentiate into multiple lineages and generate progenitors committed to one or more cell lines [9–11]. Among these multiple lineages, those of adipogenesis and osteogenesis are the most closely related: Several studies have provided substantial evidence that osteoblasts and adipocytes share a common progenitor: multipotential BMSCs [5–8, 12–14]. An inverse relationship has been demonstrated between osteogenesis and adipogenesis suggesting the possible reciprocal relationship between the two differentiation pathways [15–17]. There has been evidence for the differentiation switching between these two cell lineages, suggesting a large degree of plasticity between osteoblasts and adipocytes [18, 19]. Because the relationship between adipogenesis and osteogenesis is reciprocal and the adipocytic and osteogenic cells share a common lineage, it is possible that inhibition of adipogenesis may provide an approach to prevent or treat osteoporosis or other bone diseases. Thus, the signal transduction pathways implicated in these processes are evaluated as potential targets for therapeutic intervention of osteopenic disorders.

Wnt1 was originally described as a proto-oncogene in mouse mammary tumor induced by mouse mammary tumor virus [20]. Genetic evidence indicates that Wnt signaling is critically involved in bone homeostasis and is essential for normal osteogenesis [21–25]. Enhancement of Wnt signaling either by Wnt overexpression [26] or deficiency of Wnt antagonists [27] is associated with increased bone formation in mice and humans. A recent study by Chen et al. [28] has elucidated the critical role of Wnt/β-catenin signaling in mediating the ethanol-induced adipogenic lineage commitment of mesenchymal stem cells: ethanol inhibits bone formation through stimulation of oxidative stress to suppress Wnt signaling.

Several studies [29–33] have provided evidence that alcohol shifts lineage commitment and differentiation of BMSCs from osteogenesis to adipogenesis by upregulation of gene expression for peroxisome proliferator-activated receptor gamma (PPARγ); however, the molecular mechanism underlying the reciprocal relationship is not yet well understood. We have found in a separate study that alcohol exposure triggers ER stress in BMSCs where ATF4 and CHOP are excessively upregulated and activated. These changes result in activation of the TNF-α signaling pathway leading to an unwanted shift of BMSCs away from osteogenesis towards adipogenesis. Here we confirmed the results reported by Chen et al. [28] that alcohol-induced ER stress suppresses the Wnt1/β-catenin signaling pathway favoring adipogenic remodeling of BMSCs. We therefore proposed the following ethanol-induced signaling pathway leading to inhibition of osteogenesis: Ethanol → ER stress↑ → ATF4 and CHOP↑↑ / TNF-α↑↑ / Wnt1/β-catenin↓↓ → Osteoblasts↓↓↓↓. These data explain why ATF4 and CHOP which are normally required for osteogenic shift of lineage enter into adipogenic remodeling when overexpressed during ER stress. Obviously, targeting ATF4 and CHOP may not be an ideal strategy for promoting osteogenesis as while silencing ATF4
and CHOP may alleviate ER stress this maneuver may well damage the osteogenic potential of BMSCs. Moreover, targeting either TNF-α or Wnt signaling may not yield optimal effects in preserving osteogenic lineage of BMSCs. We therefore examined the potential benefit to osteogenesis via targeting the downstream components of the signaling pathway shown above by simultaneous silencing TNF-α and activating Wnt1.

Materials and Methods

**Human BMSCs preparation and culture**

Bone marrow samples for isolating BMSCs were collected from cancellous bony fragments of patients (age range 35–50 years; n=6) undergoing routine total hip replacement surgery in the Orthopaedic Hospital, the People’s Liberation Army General Hospital of Guangzhou Military Command. Written informed consent was obtained from all patients. The study protocol was approved by the Ethics Committee of the People’s Liberation Army General Hospital of Guangzhou Military Command. None of the patients were alcoholics, showed any evidence of concurrent illness, and received any medications that could affect bone metabolism.

The procedures for preparing primary BMSCs from human subjects were essentially the same as described by Huang et al [18], which was a modified method of Percoll density gradient centrifugation [34]. Briefly, when BMSC cultures became nearly confluent, cells were trypsinized and plated in 75 cm² flasks or 6-well plates for protein analysis or RNA analysis and cytochemical analysis, respectively. Only cells at passage two or three were used. Cell surface markers were evaluated by flow cytometry (Cytomics FC-500, Beckman Coulter, CA, USA) using the monoclonal antibodies conjugated with fluorescein isothiocyanate (FITC) or phycoerythrine (PE): CD34-FITC, CD105-PE, CD166-PE.

**Alcohol treatment and adipogenic induction**

BMSCs at passage two or three were divided into three groups: control group, osteogenic group, and alcohol group. The control group was cultured in Dulbecco’s modified Eagle’s medium (DMEM) with 10% FBS, the osteogenic group in DMEM containing osteogenic inducers (10 mM β-glycerophosphate, 50 μg/ml ascorbic acid, and 0.1 μM dexamethasone) supplemented with 10% FBS [18], and the alcohol group in DMEM containing 10% FBS, 100 mM alcohol, and osteogenic inducers. All cultures were maintained for 24 days prior to experimental measurements and treatments were implemented as to be specified. This concentration of alcohol is approximately twice as much as its IC₅₀ for inhibiting osteoblast proliferation in vitro and also represents a level within the physiological range observed in actively imbibing alcoholics [18, 35]. To supplement the potential loss of alcohol because of its volatile nature and maintain the constant alcohol concentration, we followed the procedures described by Huang et al. [18] by placing a reservoir containing the same concentration of alcohol in a chamber during culture, which was replenished daily.

**Drug treatment**

Approximately 4×10⁵ cells were treated in a plastic dish with a selective inhibitor of TNF-α biosynthesis 3,6'-dithiothalidomide (10 μM; C₁₀H₁₂F₂N₂O₂; Santa Cruz Biotechnology) [3, 4]. XAV939 (2-(4-(trifluoromethyl)phenyl)-7,8-dihydro-5H-thiopyran[4,3-d]pyrimidin-4-ol) is a cell permeable, small molecule inhibitor of the Wnt/β-catenin pathway [1, 2]. It inhibits tankyrase 1 (IC₅₀=11 nM) and tankyrase 2 (IC₅₀=4 nM), stabilizes axin and stimulates β-catenin degradation. XAV-939 was solubilized in DMSO at 55°C to make a 10 mM stock solution which may be diluted later to a working concentration of 100 μM.

To activate Wnt/β-catenin signaling pathway, BMSCs were treated with 20 μM BIO (Sigma) for 24 h [36, 37].

**Alizarin red staining**

To detect mineral deposition, alizarin red staining was performed on BMSCs after cultured in chamber slides for 28 days. Cells were fixed in 4% formaldehyde in PBS at 4°C for 45 min. Next, the cells were washed and exposed to alizarin red (2% aqueous, Sigma) for 10 min at RT, and washed again with distilled water. Finally, cells were examined under a phase contrast microscope.
Real-time PCR

Total RNA was prepared from cultures on day ten, using the Trizol reagent (Invitrogen) according to the manufacturer’s instructions. Quantitative real-time RT-PCR analysis was performed with the SYBR Green RT-PCR Kit (Qiagen, Valencia, CA) using the GeneAmp 5700 sequence detection system (Applied Biosystems) according to the manufacturer’s instructions. Levels of mRNAs of osteogenic markers Osf2/Cbfa1 [38], osteopontin [39] and osteocalcin [40], adipogenic markers (PPARγ2 and aP2) [41], and Wnt1 were determined. Samples were assayed in triplicates, and the values were normalized to the relative amounts of β-actin, and the results were interpreted using the ΔCT method.

Immunoblots

Cells were lysed in a RIPA lysis buffer containing inhibitors for proteases and phosphatases. Protein concentration was determined by the BCA Protein Assay Kit using bovine serum albumin as the standard (Pierce, USA). Equal amounts of protein samples (~25 µg) were fractionated by SDS-PAGE (12% polyacrylamide gels) and electrophoretically transferred to PVDF membrane (Millipore, Bedford, MA) using a Mini Trans-blot (Bio-Rad Laboratories, Shanghai). The membranes were blocked for 1 h in defatted milk (10% in Tris-buffered saline with Tween-20 buffer), and incubated with primary antibodies (1:1000) at 4°C overnight. The primary antibodies included rabbit polyclonal anti-PPARγ2, anti-Osf2/Cbfa1, and anti-Wnt1, all purchased from Santa Cruz Biotech (CA, USA). Next day, the membrane was incubated with horseradish peroxidase-conjugated secondary antibody (1:1,000; Molecular Probes) diluted in PBS for 2 h at RT. Finally, the membrane was rinsed with PBS before scanning using the Infrared Imaging System (LI-COR Biosciences). β-actin was used as an internal control for equal input of protein samples, using anti-β-actin antibody. Western blot bands were quantified using the LI-COR Image Studio Lite software by measuring the band intensity (Area × OD) for each group and normalizing to β-actin. The final results are expressed as fold changes by normalizing the data to the control values.

Assays for osteogenic biomarkers alkaline phosphatase, collagen-I and osteocalcin

Alkaline phosphatase activity was determined by measuring the formation of p-nitrophenol from p-nitrophenyl phosphate with the ALP Optimised clorimetric test kit according to the manufacturer’s instructions. C-terminal propeptide of type I collagen was measured by enzyme-linked immunosorbent assay with a Metra CICPEIAkit (Quidel, San Diego, CA). For assessing osteocalcin, 1,25-dihydroxycholecalciferol (0.1 µM) was added into the culture and incubated for 24 h. A Metra Osteocalcin EIA kit (Qudiel) was used for osteocalcin production measurement in the culture media on day ten. The values of alkaline phosphatase activity, type I procollagen and osteocalcin production were normalized to total protein determined with Bio-Rad total protein assay reagents (Bio-Rad, Hercules, CA).

Data analysis

Results are presented as mean±S.E.M. Multiple comparisons were performed using one-way ANOVA and Bonferroni after test comparison (Prism software; GraphPad, San Diego, CA). A P value <0.05 was considered significant.

Results

Characterization of BMSCs

Expression of CD34, CD105, and CD166 in the BMSCs were determined by flow cytometry using the monoclonal antibodies CD34-FITC, CD105-PE, and CD166-PE in the undifferentiated states. The BMSCs consistently expressed BMSC markers CD105 and CD166, but did not express hematopoietic cell marker CD34. The cells prepared by Percoll separation were composed of 98.6%±5.5% of CD34 negative cells, 94.9%±7.2% of CD105 positive cells and 93.7%±6.3% of CD166 positive cells. These results indicated the purity of BMSCs in our cell preparation.

Alcohol-induced dissipation of osteogenic lineage in BMSCs

We first compared the osteogenic and adipogenic status among the three groups by assessing the relative expression of osteogenic marker genes Osf2/Cbfa1 and osteocalcin,
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Quantitative analysis of the mRNA levels by real-time RT-PCR (qPCR) revealed that Osf2/Cbfa1 and osteocalcin mRNAs in the osteogenic group were significantly upregulated \((P<0.001)\), whereas PPARγ2 and aP2 mRNAs were downregulated \((P<0.05)\), compared to the control group (Fig. 1A). These changes were all inverted by alcohol: in the alcohol group, expression of Osf2/Cbfa1 and osteocalcin was markedly downregulated \((P<0.01)\), along with robust increases in the levels of PPARγ2 and aP2 mRNAs \((P<0.001)\), relative to the osteogenic group (Fig. 1B).

Similar results were obtained from Western blot analysis. Alcohol reversed the elevated protein levels of Osf2/Cbfa1 and the diminished levels of PPARγ2 and aP2 observed in the osteogenic group (Fig. 1C). Additionally, alkaline phosphatase activity was enhanced, and...
type I collagen (procollagen I) and osteocalcin productions were increased by osteogenic inducers, which were all abolished by alcohol exposure (Fig. 1D).

Moreover, Alizarin red staining showed that throughout the 28-day culture period, the number of calcification nodules in the osteogenic group was 79.3±6.5/cm², which was significantly decreased by nearly 50% by alcohol (P<0.001; Fig. 2).

Fig. 3. Evidence for the role of Wnt/β-catenin signaling in osteogenesis and alcohol-induced adipogenic lineage shift of BMSCs. A, Enhanced nuclear translocation of β-catenin, as indicated by the relative increase of β-catenin in the nucleus and decrease in the cytoplasm in cells treated by osteogenic inducers, and the reversal of these changes by alcohol exposure (100 mM for 24 days). B, Upregulation of Wnt1 protein in the Osteogenic group and downregulation in the Alcohol group. 1: the Control group; 2: the Osteogenic group; 3: the Alcohol group in the presence of osteogenic inducers; C: cytoplasm; N: nucleus. ***P<0.001 Osteogenesis vs Control; **P<0.01 Alcohol vs Osteogenesis; +++P<0.001 Alcohol vs Osteogenesis; n=4.

Fig. 4. Inhibition of Wnt/β-catenin signaling by XAV939 (100 μM) suppresses osteogenesis of BMSCs. A, XAV939 decreased the osteogenic marker Osf2/Cbfa1 protein level, alkaline phosphatase activity, and procollagen I and osteocalcin productions. B, XAV939 increased the adipogenic markers PPARγ2 and aP2 protein levels. C, XAV939 reduced the number of calcified nodules. O: the Osteogenic group as control; X: the Osteogenic group treated with XAV939. **P<0.01 XAV939 vs Control; ***P<0.001 XAV939 vs Osteogenesis; n=5.
Inhibition of Wnt1/β-catenin and alcohol-induced dissipation of osteogenic lineage

Wnt1 has been shown to play a crucial role in bone homeostasis and normal osteogenesis [21–28]. To investigate if Wnt1 is involved in alcohol-induced dissipation of osteogenic lineage of BMSCs, we evaluated the effects of Wnt/β-catenin inhibition and activation. As illustrated in Fig. 3, Wnt/β-catenin signaling was markedly activated in the osteogenic group relative to the control cells, as indicated by the increased levels of nuclear β-catenin along with reduced levels of cytosolic β-catenin (Fig. 3A), as well as increased protein levels of Wnt1 (Fig. 3B). Activation of Wnt/β-catenin signaling was significantly suppressed in BMSCs exposed to alcohol. It was noted that not only was the nuclear translocation of β-catenin enhanced but also was its expression level elevated, as indicated by the increased sum of cytosolic and nuclear β-catenin protein.

Moreover, in the osteogenic group treated with a Wnt/β-catenin signaling inhibitor XAV939, the osteogenic marker genes were downregulated (Fig. 4A), whereas the adipogenic markers were upregulated (Fig. 4B). The number of calcification nodules in the osteogenic group was also decreased by 30% by XAV939 (P<0.001; Fig. 4C).

On the contrary, activation of Wnt/β-catenin signaling by BIO in alcohol-treated BMSCs caused the opposite changes: upregulation of the osteogenic markers (Fig. 5A), downregulation of adipogenic markers (Fig. 5B), and increase in the number of calcification nodules (Fig. 5C). Osf2/Cbfa1 by ~1.8 fold, alkaline phosphatase activity by ~2.1 fold, procollagen I by ~1.9 fold, and osteocalcin production by ~1.6 fold (Fig. 5A). PPARγ2 and aP2 were decreased by ~28% and ~33%, respectively, by BIO (Fig. 5B). The number of calcification nodules was increased by ~1.5% (Fig. 5C).
Osteogenic lineage shift of BMSCs induced by simultaneous knockdown of TNF-α and upregulation of Wnt1

It was noted that the efficacy of promoting osteogenesis or suppressing adipogenesis by BIO (20 μM) was moderate and suboptimal based on the data shown in Figure 5. Increasing BIO concentration to achieve a greater extent of Wnt/β-catenin signaling activation might improve the efficacy; however, this maneuver could cause changes of other signaling pathways that could obscure and confound the interpretation of data. It is therefore more desirable to take alternative approaches. As already mentioned, we have shown in our separate study that alcohol exposure upregulated TNF-α signaling favoring adipogenic lineage of BMSCs. These findings prompted us to test if simultaneous inhibition of TNF-α signaling and activation of Wnt/β-catenin signaling could yield more powerful suppressing effects on alcohol-induced dissipation of osteogenic lineage. The results presented in Fig. 6 provide supportive evidence for the notion. Concordant treatment of BMSCs with 3,6'-dithiothalidomide (DT; 10 µM) and BIO (20 μM) in the alcohol group generated superior promotion of osteoblast lineage shift, or inhibition of adipocytes lineage, after ethanol exposure. Osf2/Cbfa1 was upregulated by ~3.2 fold, alkaline phosphatase activity enhanced by ~3.9 fold, and procollagen I and osteocalcin productions by ~2.6 fold and ~2.9 fold, respectively (Fig. 6A). PPARγ2 and aP2 were decreased by ~45% and ~63%, respectively, by BIO (Fig. 6B). The number of calcification nodules was increased by ~2.9% (Fig. 6C).

Discussion

Both TNF-α and Wnt1 have been shown to play a crucial role in bone formation and homeostasis with Wnt1 being essential for normal osteogenesis and TNF-α being anti-
osteogenic. The main goal of this study was to examine our hypothesis that concordant reciprocal interventions of TNF-α and Wnt1 signaling produce more thorough inhibition of ethanol-induced adipogenesis in human BMSCs. The main findings of the present study are (1) Wnt/β-catenin signaling in BMSCs was significantly suppressed by ethanol, indicating its involvement in the impaired osteogenesis; and (2) simultaneous activation of Wnt/β-catenin signaling and inhibition of TNF-α signaling produced synergetic suppression of ethanol-induced adipogenic lineage compared to intervention of either of them alone.

Wnt signaling has been known to participate in osteogenesis and bone homeostasis [21–28] and stimulation of Wnt signaling either by Wnt overexpression [27] or deficiency of Wnt antagonists [28] is implicated in increased bone formation in mice and humans. Chen et al. [28] provide the first evidence for the role of Wnt/β-catenin signaling in the regulation of osteogenic and adipogenic lineages of mesenchymal stromal cells with prolonged exposure to alcohol. Our results are consistent with this study demonstrating the importance of Wnt/β-catenin signaling against alcohol-induced adipogenic lineage commitment. Moreover, we showed that Wnt/β-catenin signaling was substantially activated in BMSCs treated with osteogenic inducers relative to untreated control cells, as indicated by the enhanced nuclear translocation of β-catenin, as well as increased protein levels of Wnt1. Artificial inhibition of Wnt/β-catenin signaling inhibited osteogenesis even in the presence of osteogenic inducers. On the other hand, Wnt/β-catenin signaling was significantly suppressed in BMSCs exposed to alcohol. Artificial activation of Wnt/β-catenin signaling in cells treated with alcohol alleviated the adipogenesis. This study did not get into the mechanisms for the activation of Wnt/β-catenin signaling during osteogenesis and inhibition during adipogenesis induced by alcohol. It is possible that it is ER stress induced by alcohol that inhibited Wnt/β-catenin signaling, as it has been shown that ER stress suppresses Wnt/β-catenin signaling [41, 42] and in our separate study, we have addressed the critical role of ER stress in adipogenic lineage shift of BMSCs after exposure to ethanol. Furthermore, it has been shown that oxidative stress downregulates Wnt/β-catenin signaling, and the antioxidant N-acetylcysteine is able to reverse this downregulation and block bone loss [28].

It is noted that either activation of Wnt/β-catenin signaling alone or inhibition of TNF-α signaling alone allowed for only moderate recovery of osteogenesis impairment induced by alcohol. There are several possible explanations for this observation. It may be that the activation of Wnt/β-catenin signaling or inhibition of TNF-α signaling was incomplete or suboptimal. Alternatively, it may be that multiple signaling pathways including Wnt/β-catenin signaling and TNF-α signaling (and likely others as well) are involved in the regulation of differentiation of BMSCs after ethanol insult and intervention with any one of these pathways could elicit only a fractional change of cell fate as the remaining pathways may well compensate for it. Or it is more likely that these above-mentioned two possibilities co-exist. To strengthen the efficacy of recovery of ethanol-induced osteogenesis impairment, one might simply elevate the concentrations of Wnt/β-catenin activator or TNF-α inhibitor; however, this could well be in the face of generating non-specific and unwanted effects. An alternative approach is to simultaneously interfere Wnt/β-catenin signaling and TNF-α signaling to achieve improved effects. The results in the present study indeed provided evidence in support of this latter strategy: concomitant activation of Wnt/β-catenin signaling and inhibition of TNF-α signaling give rise to superior effects in terms of inhibition of ethanol-induced adipogenesis over interference of any one these pathways alone. This is expected if Wnt/β-catenin signaling and TNF-α signaling are two separate signaling pathways. However, all these notions need to be examined by future studies.

Collectively, our study supports the finding that downregulation Wnt/β-catenin signaling as an important determinant of ethanol-induced adipogenic shift of BMSCs [28]. Activation of Wnt/β-catenin signaling partially reverted the impairment of osteogenesis. Moreover, simultaneous reciprocal interferences of Wnt/β-catenin signaling and TNF-α signaling yielded a significant improvement of effects on the recovery of osteogenesis. This strategy may therefore be considered as an alternative approach for the treatment of ethanol-induced impairment of osteogenesis.
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Conflicts of Interest

The authors declare no conflicts of interest for this study.

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