

(+)-Cholesten-3-one induces osteogenic differentiation of bone marrow mesenchymal stem cells by activating vitamin D receptor

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Abstract. In our previous reports, it was revealed that steroids in traditional Chinese medicine (TCM) have the therapeutic potential to treat bone disease. In the present study, an *in vitro* model of a vitamin D receptor response element (VDRE) reporter gene assay in mesenchymal stem cells (MSCs) was used to identify steroids that enhanced osteogenic differentiation of MSCs. (+)-cholesten-3-one (CN), which possesses a ketone group that is modified in cholesterol and cholesterol myristate, effectively promoted the activity of the VDRE promoter. Phenotypic cellular analysis indicated that CN induced differentiation of MSCs into osteogenic cells and increased expression of specific osteogenesis markers, including alkaline phosphatase, collagen II and Runt-related transcription factor 2. Furthermore, CN significantly increased the expression of osteopontin, the target of the vitamin D receptor (VDR), which indicated that CN may activate vitamin D receptor signaling. Over-expression of VDR or knockdown studies with VDR-small interfering RNA revealed that the pro-differentiation effects induced by CN required VDR. Furthermore, the present study determined that the C-terminal region of the VDR is responsible for the action of CN. Taken together, the present findings demonstrated that CN induced osteogenic differentiation of MSCs by activating VDR. The present study explored the regulation of stem cells by using a series of similar steroids and provided evidence to support a

potential strategy for the screening of novel drugs to treat bone disease in the future.

Introduction

Mesenchymal stem cells (MSCs) are mesodermal cells derived from bone marrow, which may differentiate into osteoblasts, chondrocytes, myoblasts, osteoblasts, chondrocytes, myoblasts, stromal cells, adipocytes and other cells (1). Dysregulation of the differentiation of MSCs into osteoblasts is involved in several bone-related diseases, including osteoporosis (2). MSCs have the potential to be used as therapeutic agents since they are easily isolated and extracted from patients, without ethical or technical problems. The inherent potential of MSCs makes them ideal seed cells for cell transplantation therapy, which is expected to become an effective treatment of intractable orthopedic disease (3,4). Although MSCs are promising targets for the treatment of a number of bone diseases, the outcome of MSCs must be controlled before their therapeutic potential can be realized. Therefore, an improved understanding of the signaling pathways that control the differentiation of MSCs is required.

Nuclear receptors are ligand-activated transcription factors that perform a central function in the differentiation of stem cells (5). Among them, vitamin D receptor (VDR) has a critical role in the control of osteogenic differentiation of MSCs (6). VDR is composed of three distinct regions: An N-terminal region, containing a ligand-independent activation function-1; a highly conserved central region, containing the DNA binding domain; and the C-terminal region of the receptor, containing a multifunctional domain that harbors the ligand binding domain, the retinoid X receptor heterodimerization motif and ligand-dependent activation function-2 (7). VDR response elements (VDREs) have been identified in the promoters of vitamin D3-responsive genes, such as osteopontin (8). Previous studies have characterized MSCs as a target of VDR functions (9). Furthermore, the epigenetic regulation of the VDR may be key to rejuvenating

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osteoblastogenesis in MSCs from elders (10). The active form of vitamin D, $1\alpha, 25$ -dihydroxyvitamin D₃ [$1,25(\text{OH})_2\text{D}_3$; VD₃], is the ligand for VDR and has been revealed to enhance osteoblast-mediated mineralization (11). Moreover, vitamin D induced transcriptional networks in the osteoblast differentiation in a previous study (12). Thus, VDR is proposed as an important pharmacological target for signaling pathways involved in osteogenic differentiation of MSCs. Therefore, VDR can effectively screen for novel drugs from medicinal plants, particularly traditional Chinese herbs. We previously reported that steroids found in traditional Chinese herb were able to control the fate of MSCs (13,14).

In the present study, an *in vitro* model was established using a VDRE reporter gene assay, which constructed a VDRE promoter reporter and was transfected into MSCs. Then MSCs were treated with steroids to identify VDR signaling activators that may be involved in the osteogenic differentiation. Furthermore, based on the previous finding that steroids are able to enhance osteogenic differentiation of MSCs (13), various structural analogs of steroids were investigated. The analogs included VD₃, (+)-cholesten-3-one (CN), steraric acid ethyl ester (SE), cholesterol (CL), cholesterol myristate (CM), cyasterone (CE), β -sitosterol (SL) and oleanolic acid (OA). Among them, CN was the specific compound that most markedly promoted osteogenic differentiation of MSCs.

Materials and methods

Animal grouping and drug administration. A total of 10 male specific pathogen free Sprague-Dawley rats, aged 4 weeks (180–200 g), were acquired from the Animal Centre of Guangzhou University of Chinese Medicine (Guangzhou, China). All animals received humane care in accordance with the guidelines set out by the Care of Experimental Animals Committee of Guangzhou University of Chinese Medicine. Dulbecco's modified Eagle medium (DMEM) and fetal bovine serum (FBS) were purchased from Gibco (Thermo Fisher Scientific, Inc., Waltham, MA, USA); VDRE, VD₃, dimethyl sulphoxide (DMSO) and other chemical reagents were purchased from Sigma-Aldrich (Merck Millipore, Darmstadt, Germany); alizarin red stain solution, alkaline phosphatase (ALP; BA0632), osteopontin (OPN; PB0589), Runt-related transcription factor 2 (RUNX2; BA3613-2), VDR (PB0479) and collagen II (BA0533) antibody were provided by Wuhan Boster Biological Technology, Ltd., (Wuhan, China); and SE, CL, CM, CN, CE, SL and OA were provided by Tokyo Chemical Industry Co., Ltd., (Tokyo, Japan).

Culture of MSCs. The femur and tibia were harvested from rats, anaesthetized by chloral hydrate (330 mg/kg; Wuhan Boster Biological Technology, Ltd.) and sacrificed by cervical dislocation, in order to collect fresh marrow. The marrow was mixed with complete medium (low glucose DMEM supplemented with 10% FBS) and gradient centrifuged at $900 \times g$ for 30 min at room temperature with Percoll at a density of $1.073 \text{ g}\cdot\text{ml}^{-1}$. Cells of the appropriate density were collected, washed with PBS three times, manually counted using a light microscope and cultured at a density of $1 \times 10^6 \text{ cm}^{-2}$ on dishes supplemented with complete medium, in a humidified atmosphere at 37°C and 5% CO_2 , using an incubator. The medium

was refreshed and the suspension cells were removed every three days. Isolated MSCs were confluent after nine days, and were digested using 0.25% trypsin in order to promote separation. Cells were subsequently passaged at a density of $1 \times 10^4 \text{ cm}^{-2}$ onto dishes. MSCs that had been passaged three times were used for subsequent experiments. The surface antigen identification and differentiation ability of MSCs were verified by our previous study (13,14).

Plasmids, small interfering (si) RNA cell transfection and assay for luciferase activity. The purchased pGL3-basic vector and VDRE were digested using purchased *NheI* and *HindIII* (Takara Biotechnology Co., Ltd., Dalian, China), and both were combined to generate the VDRE promoter-Luc vector. Plasmid for cytomegalovirus (CMV) was cotransfected to normalize the variations in transfection efficiency. The pGL3-basic vector, VDRE vector, and pCMV-Myc-basic, pCMV-Myc-VDR, pCMV-Myc-VDR-N and pCMV-Myc-VDR-C plasmids were provided by the National Laboratory of Medical Molecular Biology (Beijing, China).

The sequence of the siRNA oligonucleotide that was used to silence the VDR was 955-GUGCCAUUGAGGUCAUCAUTT-975. The scramble siRNA with random nucleotides was 5'-UUCUCCGAACGUGUCACGUTT-3' (negative control siRNA). Both oligonucleotides were synthesized by Genepharma Corp., (Shanghai, China). siRNA were transfected into cells using Lipofectamine[®] 2000 (Invitrogen; Thermo Fisher Scientific, Inc.) according to the manufacturer's protocols.

Dual luciferase assay was performed to study the effects of VD₃, SE, CL, CM, CN, CE, SL and OA on VDRE promoter activity. VD₃ ($30 \text{ ng}\cdot\text{ml}^{-1}$), SE, CL, CM, CN, CE, SL and OA were dissolved in DMSO, respectively, and subsequently added to MSCs with an increasing concentration of 0–100 $\mu\text{g}\cdot\text{ml}^{-1}$. MSCs seeded in complete medium treated with an equal volume of DMSO without steroids served as the control group. Luciferase assays were performed using a Dual Luciferase Assay kit (Promega Corp., Madison, WI, USA) and were used to detect the luciferase activity. A total of 10 μl of cell lysate was added to MSCs to detect firefly luciferase activity. Following the standardization of *Renilla* luciferase activity, the activity of *Renilla* luciferase was detected.

Immunocytochemistry. Following treatment with CN, MSCs were fixed with 4% polyoxymethylene for 20 min at room temperature and permeabilized with 0.25% Triton X-100 (Sigma-Aldrich; Merck Millipore) and blocked with 0.1% bovine serum albumin (BSA; Roche Diagnostics, Basel, Switzerland). Subsequently, MSCs were incubated overnight at 4°C with the following primary antibodies at dilutions of 1:200: ALP, OPN, RUNX2 and collagen II. After washing three times with PBS, the cells were reacted with the horseradish peroxidase-conjugated secondary antibody (ab6721; 1:2,000; Abcam, Cambridge, UK) for 30 min at 37°C . The cell nuclei were stained with hematoxylin (Wuhan Boster Biological Technology, Ltd.) and visualized via light microscopy (Leica Camera AG, Wetzlar, Germany). Percentage of positive cells vs. total cells in 4 fields was calculated. Rat ALP, OPN, RUNX2 and collagen II primary antibodies (all 1:200), mixed with non-immune serum, were used to immunostain MSCs at 4°C overnight, respectively. Secondary

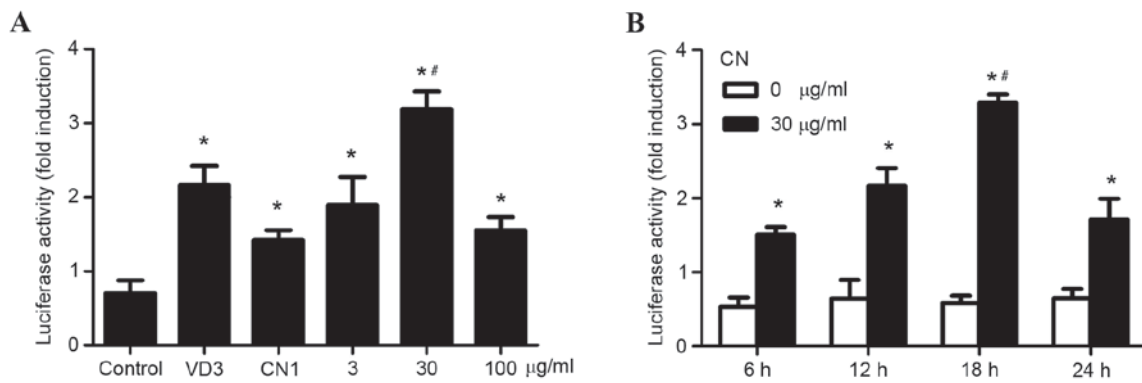


Figure 1. CN increases VDRE promoter activity. (A) Dose-response of CN on the VDRE promoter activity. * $P < 0.05$ vs. control; # $P < 0.05$ vs. all other groups. (B) Time-response of CN on the VDRE promoter activity. * $P < 0.05$ vs. 0 $\mu\text{g/ml}$; # $P < 0.05$ vs. all other groups. All data were expressed as mean \pm standard deviation. CN, (+)-cholesten-3-one; VDRE, vitamin D response element.

antibody was used to incubate for 30 min. The expression of ALP, OPN, RUNX2 and collagen II was detected after diaminobenzidine staining, followed by hematoxylin as the counterstain. Cytoplasm of positive cells were detected as deep brown. The same method was applied to the control group, except that the primary antibody with non-immune serum.

Calcium mineral deposition. MSCs were cultured for 14 days, as described. The level of calcium mineral deposition was revealed using alizarin-red staining (AR-S). Following three weeks in culture, cells were fixed with 70% ethanol, washed five times with deionized water and treated with 40 mM alizarin-red solution for 10 min at pH 4.2. Subsequently, cells were washed with PBS for 15 min before being treated with 10% cetylpyridinium chloride in 10 mM sodium phosphate for 15 min at room temperature. An AR-S standard curve, at an optical density of 540 nm, was used to calculate the AR-S concentration.

Western blot analysis. Total protein was extracted from MSCs with cell lysis buffer (Invitrogen, Thermo Fisher Scientific, Inc.) and the protein concentration was determined using a Bradford protein assay (bicinchoninic acid protein assay kit; Bio-Rad Laboratories, Inc., Hercules, CA, USA). SDS-PAGE (6-15%; Gude Biological Technology Co.) was created and separated proteins (20 μg) were transferred to a membrane. The membrane was blocked in 5% BSA at 24°C for 1 h, incubated with primary polyclonal antibodies against ALP, OPN, RUNX2, VDR, collagen II (all 1:200) and β -catenin (ab32572; 1:5,000; Abcam, Cambridge, UK) at 4°C overnight, followed by three washes in TBST for 5 min each. The membrane was then incubated with Alexa Fluor-conjugated goat anti-rabbit IgG secondary antibody (ab150077; 1:2,000; Abcam) for 30 min at 24°C, washed with TBST three times for 5 min each and protein bands were detected with DAB. The blot was scanned and optical density values of the targeted bands were analyzed with Image J software (1.48U; National Institutes of Health, Bethesda, MD, USA).

Statistical analysis. All data were expressed as mean \pm standard deviation. SPSS 20.0 software (IBM SPSS, Armonk, NY, USA) was used to analyze all data. $P < 0.05$ was considered to indicate a statistically significant difference.

Results

CN increases VDRE promoter activity. Induced expression of VDR provoked by CN stimulation of MSCs had been confirmed in our previous study (13,14). The present study examined whether CN increased VDRE promoter activity. VDRE promoter reporter construct was transfected into MSCs and cells were treated with 1, 3, 30 or 100 $\mu\text{g}\cdot\text{ml}^{-1}$ of CN, respectively. MSCs that did not receive treatment were considered the control group, while VD3 was used for positive control, due to its effective promotion in VDRE promoter activity, as verified by a previous study. The highest VDRE promoter activity was observed at a CN concentration of 30 $\mu\text{g}\cdot\text{ml}^{-1}$, which was statistically significant when compared to the other groups ($P < 0.05$; Fig. 1A). MSCs transfected with VDRE promoter reporter construct were treated with 30 $\mu\text{g}\cdot\text{ml}^{-1}$ of CN and compared to MSCs that did not receive CN treatment. VDRE promoter activity was measured after 6, 12, 18 and 24 h of exposure. The results indicated that the optimal time point of VDRE promoter activity of MSCs treated with 30 $\mu\text{g/ml}$ of CN was 18 h, as revealed by the significantly increased fold induction of luciferase activity ($P < 0.05$; Fig. 1B).

CN showed the highest activity on the VDRE, which may be associated with its ketone group. In our previous studies (13,14), it was revealed that steroids had a stimulating effect on VDRE promoter activity; however, the specific compound responsible for achieving this effect is still unclear. The present study aimed to investigate MSCs that were transfected with VDRE promoter reporter construct, followed by either no treatment with the analogs of steroid (negative control), treatment with either VD3 (30 $\text{ng}\cdot\text{ml}^{-1}$, positive control), or with SE, CL, CM, CN, CE, SL, and OA (30 $\mu\text{g/ml}$) for 18 h (Fig. 2A). A significantly increased activity level was observed in CN-treated MSCs when compared with MSCs treated with the other steroids ($P < 0.05$; Fig. 2B). This indicated that CN was the optimal compound in promoting the highest VDRE promoter activity, even higher than VD3, which was the positive control ($P < 0.05$). Compared with the other compounds, especially CL and SL with high similarity with CN, the highest activity was observed in CN-treated MSCs; however, the only structural difference between CN and CL is the presence of a ketone group and OH group, respectively, which suggested that the

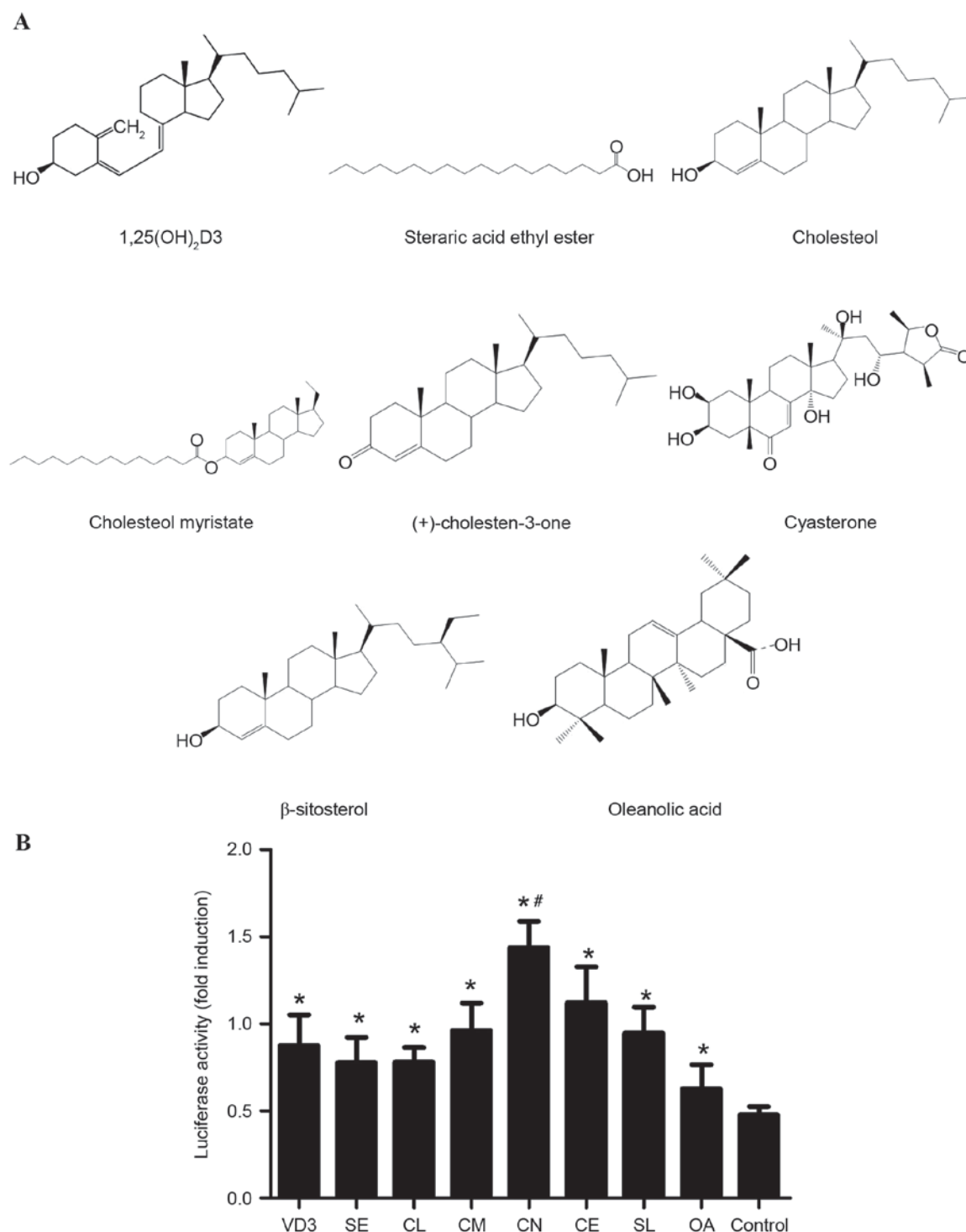


Figure 2. Effective activity of VDRE promoter is associated with the ketone group of CN that promotes the effective activity of the VDRE promoter. (A) Structure of VD3, SE, CL, CM, CN, CE, SL and OA. (B) Comparative effects of VD3, SE, CL, CM, CN, CE, SL and OA. All data were expressed as mean \pm standard deviation. * $P < 0.05$ vs. control; # $P < 0.05$ vs. all other groups. CN, VD3, 1 α , 25-dihydroxyvitamin D3; SE, steraric acid ethyl ester; CL, cholesterol; CM, cholesterol myristate; CE, cyasterone; SL, β -sitosterol; OA, oleanolic acid.

ketone group of CN may be involved in promoting VDRE promoter activity.

CN promotes osteogenic differentiation of MSCs. MSCs were stimulated by CN (30 μ g/ml) for seven days, immunocytochemically stained using osteogenesis markers and AR-S and further analyzed using western blotting. Immunocytochemical staining for osteogenesis markers containing ALP, OPN, RUNX2 and

collagen II indicated that CN promoted osteogenic differentiation of MSCs as CN-treated MSCs exhibited a significantly increased number of osteogenesis-positive cells compared with MSCs that did not receive CN treatment ($P < 0.05$; Fig. 3A). AR-S demonstrated that CN-treated MSCs exhibited an increased number of calcium mineral deposition derived from osteoblasts when compared with MSCs that were not treated with CN (Fig. 3B). Furthermore, western blotting revealed significantly higher

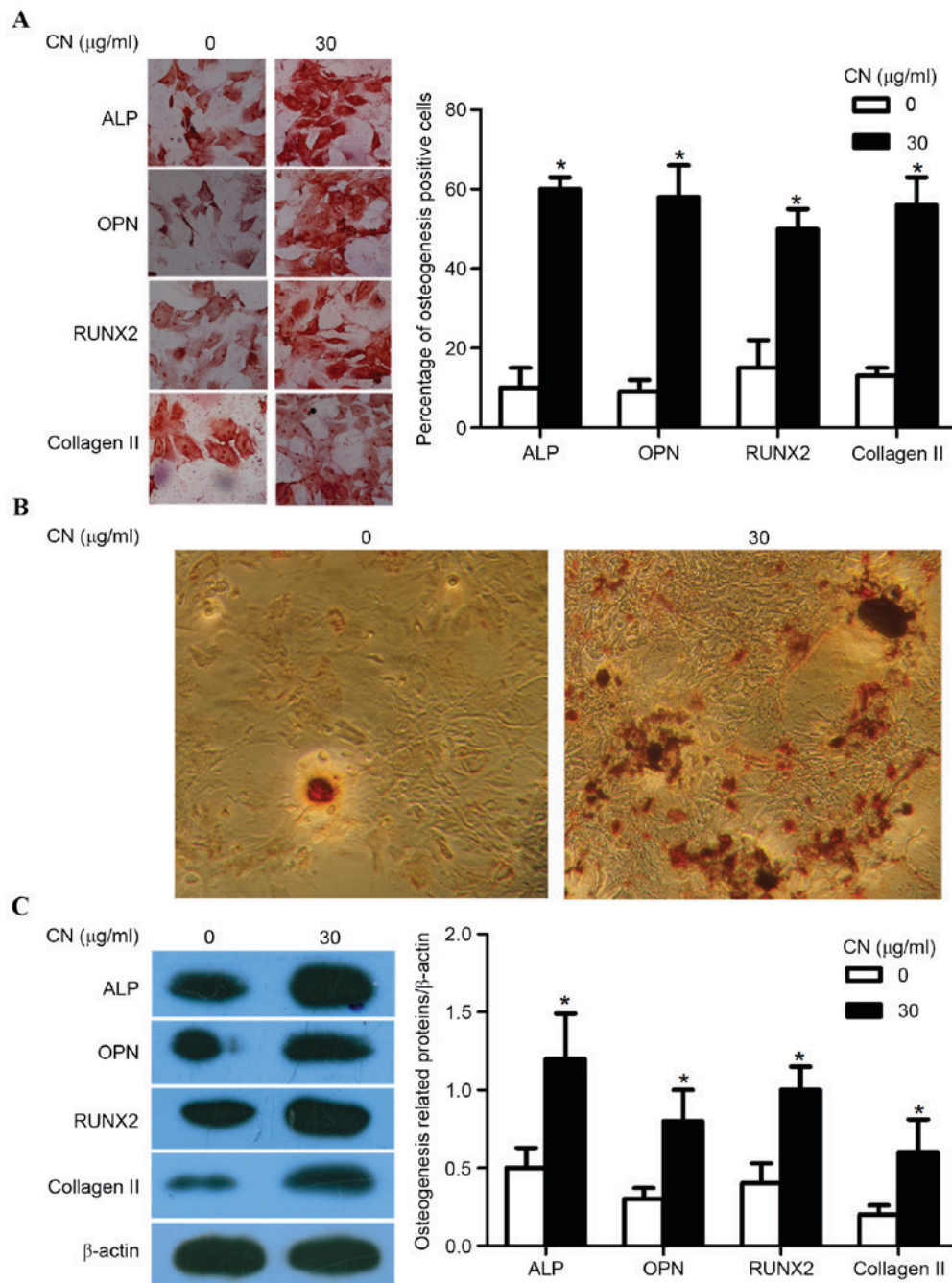
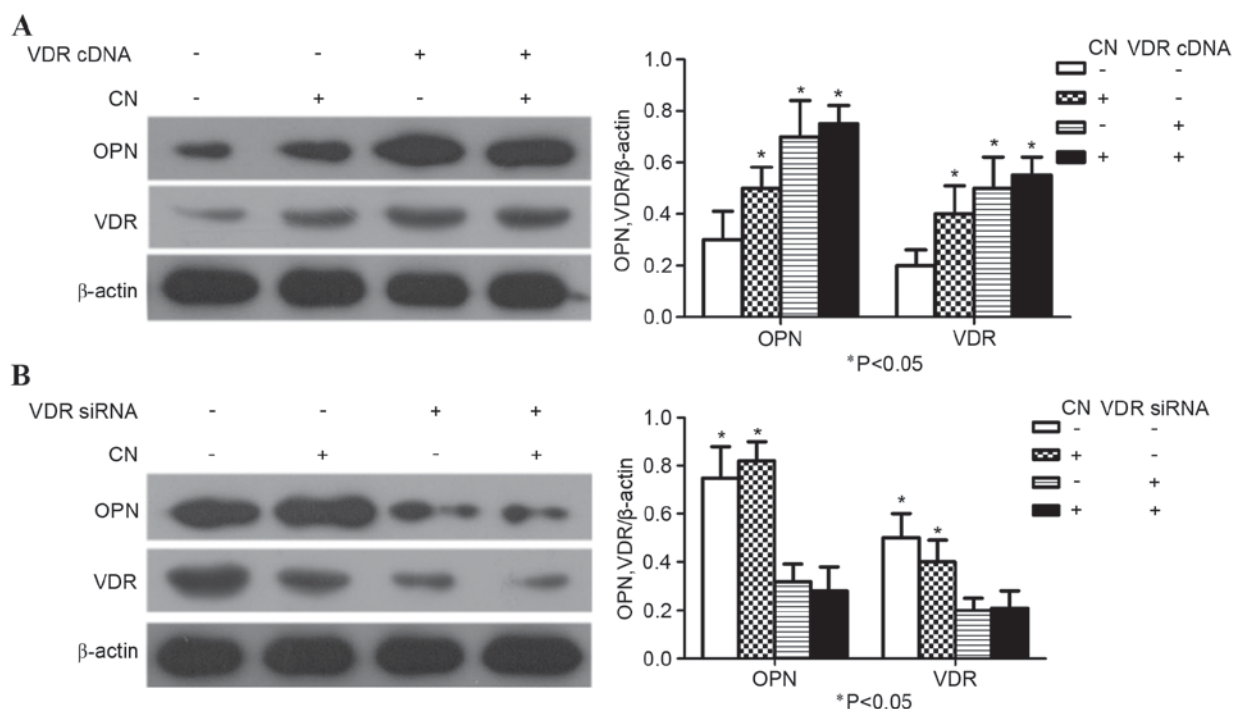


Figure 3. CN promotes the osteogenic differentiation of MSCs. (A) Photomicrographs were captured at a magnification of x200 and demonstrated the levels of ALP, OPN, RUNX2 and collagen II expressed in MSCs, where the cytoplasm was stained dark brown (left panel). Comparison of the percentage of ALP, OPN, RUNX2 and collagen II-positive cells after CN treatment for seven days (right panel). (B) Alizarin-red staining revealed the calcium mineral deposition of CN-treated MSCs compared with MSCs that were not treated with CN dyed by alizarin-red staining. (C) Western blot analysis indicated the expression of ALP, OPN, RUNX2 and collagen II (left panel) by comparing the relative density of ALP/β-actin, OPN/β-actin, RUNX2/β-actin and collagen II/β-actin band (right panel). All data were expressed as mean ± standard deviation. *P<0.05 vs. 0 μg/ml. MSCs, mesenchymal stem cells; ALP, alkaline phosphatase; OPN, osteopontin; RUNX2, runt-related transcription factor 2; CN, (+)-cholesten-3-one.

expression of ALP, OPN, RUNX2 and collagen II in CN-treated MSCs when compared to MSCs that did not receive CN treatment, indicating that CN-treated MSCs exhibited significantly increased osteogenic cell differentiation (P<0.05; Fig. 3C).

CN promotes osteogenic differentiation of MSCs and requires VDR. MSCs transfected with VDR cDNA construct were treated with optimal concentrations of CN. The expression of OPN and VDR were significantly increased in the group

that was comprised of the combination of VDR cDNA and CN compared with the group where VDR cDNA and CN were both absent. (P<0.05; Fig. 4A). MSCs were also transfected with a VDR siRNA construct and treated with optimal concentrations of CN. Furthermore, VDR siRNA inhibited the expression of OPN and VDR in MSCs compared with the groups where VDR siRNA was present and CN was absent (P<0.05; Fig. 4B). These experiments indicated that CN-induced differentiation of MSCs required VDR.



C-terminal region of VDR is responsible for the action of CN. VDR fusion proteins included the following: VDR 1-91, an N-terminal region containing activation function-1 (AF-1) domain and a DNA binding domain (DBD) and VDR 115-427, a C-terminal region with a ligand binding domain, and a ligand-dependent AF-2 domain (Fig. 5A). To confirm whether CN interaction with the binding domains of VDR affected the differentiation of MSCs into osteogenic cells, MSCs were stimulated by CN and transfected with either pCMV-Myc-basic, pCMV-Myc-VDR, pCMV-Myc-VDR-N or pCMV-Myc-VDR-C plasmids. After seven days, immunocytochemical staining demonstrated that groups transfected with pCMV-Myc-VDR-C plasmids exhibited significantly increased osteogenic-positive cells ($P<0.05$; Fig. 5B and C). Western blot analysis indicated the expression of ALP, OPN and RUNX2 increased when MSCs were treated with CN and transfected with pCMV-Myc-VDR-C plasmids (Fig. 6A and B). Moreover, AR-S identified that CN-treated MSCs transfected with pCMV-Myc-VDR-C plasmids exhibited increased numbers of osteogenic-positive cells when compared with the positive and negative control groups (Fig. 6C).

Discussion

MSCs are capable of self-renewal and differentiation into osteogenic cells; however, the efficiency of these functions is typically low (15) and the process is not fully understood. Thus, the present study aimed to elucidate the process of MSC differentiation into osteogenic cells by exploring its physiological mechanism and therapeutic potential. In the present study, a functional stem cell assay was used based on the

VDR pathway as this facilitated improved understanding of the identified compounds involved in this pathway. The major findings of the present study were: i) CN exhibited significantly higher VDRE promoter activity when compared with SE, CL, CM, CE, SL and OA; ii) CN-promoted osteogenic differentiation of MSCs that required VDR; and iii) the C-terminal region of the VDR is involved in the action of CN. The present study demonstrated a model that revealed structure-function relationships between steroids and VDR. The findings of the present study provide the incentive to develop a pioneering strategy to screen novel drugs for orthopedic disorders and to study a series of similar compounds involved in the regulation of stem cells.

In the present study, CN was identified as an inducer of MSC differentiation into osteoblasts, through cell phenotypic analysis. MSCs differentiate into all mesodermal cell types, including osteoblasts, chondrocytes, myoblasts, stromal cells and skeletal muscle cells, under appropriate conditions. Therefore, MSCs were chosen as a cell model in the present study. There is growing interest in the use of MSCs for the repair of bone damage; however, this requires efficient protocols for directing the differentiation of MSCs into the osteoblast lineage. Several studies have explored the use of bone morphogenetic proteins to induce MSCs toward the osteoblast lineage (16,17). Furthermore, these reports have described the positive effect of large molecules on the osteogenesis of MSCs; however, small molecules that may promote the osteogenic differentiation of MSCs have not been identified (16,17). Previous findings have also indicated that dexamethasone and VD3 were used to promote MSC differentiation *in vitro* (18). In the present study, immunohistochemical analysis and

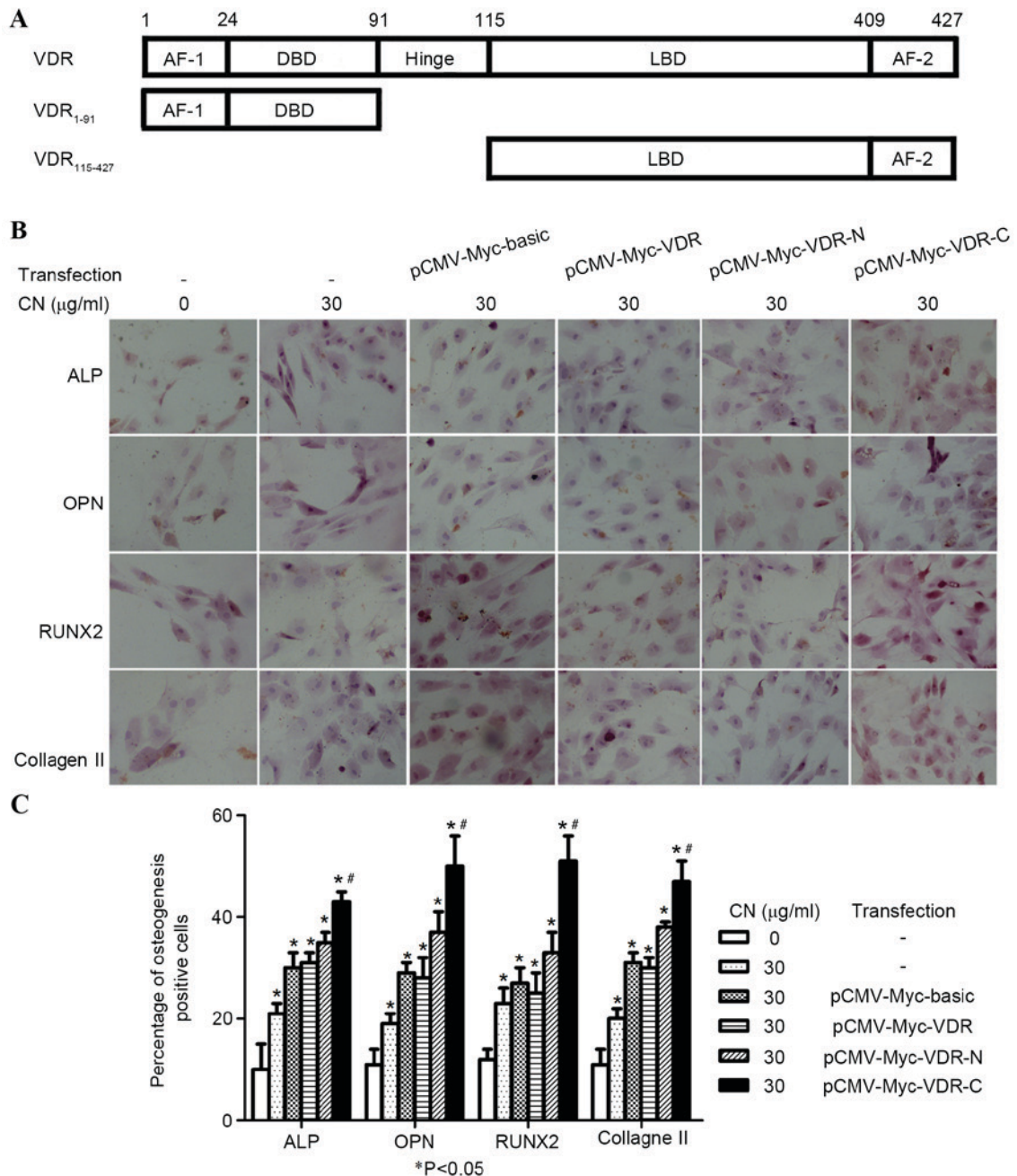


Figure 5. C-terminal region of the VDR is responsible for the action of CN. (A) Structure of VDR. (B) Photomicrographs showing ALP, OPN, RUNX2 and collagen II-positive cells in cytoplasm, which were dark brown (magnification, x200). (C) Comparison of the percentage of ALP, OPN, RUNX2 and collagen II-positive cells after CN treatment and transfection for seven days. All data were expressed as mean \pm standard deviation. *P<0.05 vs. CN and transfection were absent; #P<0.05 vs. all other groups. VDR, vitamin D receptor; OPN, osteopontin; CN, (+)-cholesten-3-one; ALP, alkaline phosphatase; RUNX2, runt-related transcription factor 2; CMV, cytomegalovirus.

western blot analysis established that CN induced MSCs to differentiate into osteoblasts. This result is consistent with our previous findings, which provided evidence that CN promotes the differentiation of neural stem cells (19). Therefore, CN may be a potential therapeutic molecule for treating differentiation-related bone diseases.

Another notable finding of the present study was that CN functions as an activator of VDR. VDR, which consists of an N-terminal DNA-binding domain, a C-terminal ligand-binding domain and an intervening hinge region, is the nuclear hormone receptor of the vitamin D endocrine system that is predominantly involved in the maintenance of calcium

and phosphate homeostasis and bone development (20-22). Furthermore, VDR is a ligand-induced nuclear transcription factor that regulates the expression of genes in critical mineral-regulating target tissues, such as bone, in order to maintain appropriate mineral homeostasis. Humans and mouse models have indicated that a lack of functional VDR is associated with severe bone diseases (20-22). Furthermore, it has also been suggested that VDR has a direct role in promoting osteoblast differentiation (23). Additionally, prior reports have identified that VDR-null mice exhibited a reduction in osteoblasts, accompanied by a decrease in trabecular bone volume (24), and MSC differentiation was impaired in

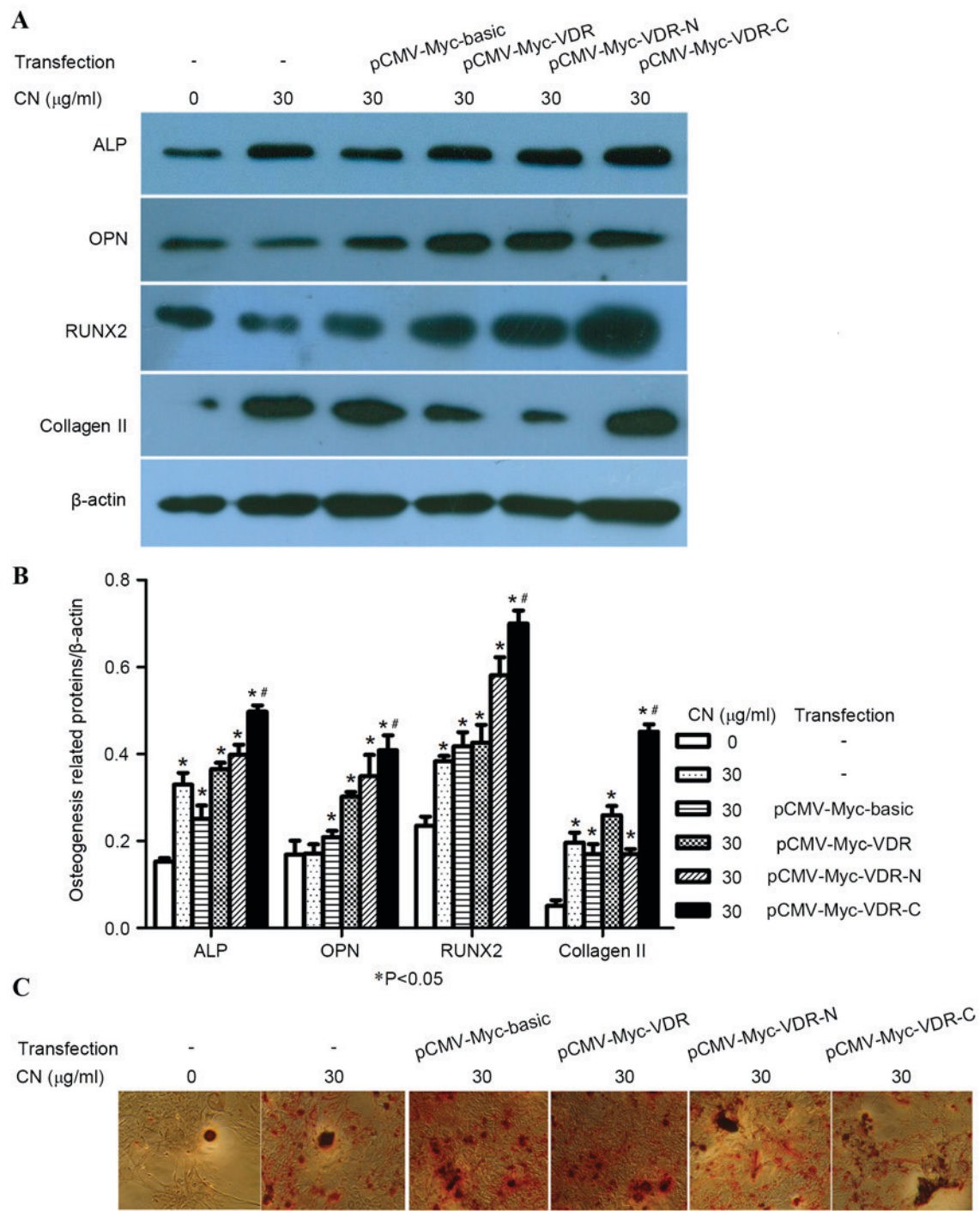


Figure 6. Western blot and alizarin-red staining established that the C-terminal region of the VDR is associated with activity. (A) Western blotting revealed the levels of ALP, OPN, RUNX2 and collagen II expressed in MSCs after transfection. (B) Comparison of the density between the experimental groups. (C) AR-S indicated the level of calcium mineral deposition. All data were expressed as mean ± standard deviation. *P<0.05 vs. CN and transfection were absent. #P<0.05 vs. all other groups. MSCs, mesenchymal stem cells; VDR, vitamin D receptor; OPN, osteopontin; ALP, alkaline phosphatase; RUNTX2, runt-related transcription factor 2.

VDR mice that presented with haploinsufficiency (25). Thus, strategies targeting VDR pathways for the stimulation of MSC differentiation are highly desirable.

We hypothesized that VDR may be one of the direct mediators of CN actions. Several experiments were performed to test this hypothesis, including the use of transcription assays, which employed a VDRE-firefly luciferase reporter plasmid and immunoblotting of CN-mediated induction of VDR target genes. Furthermore, over-expression of VDR and knockdown studies

with VDR-siRNA were performed, in addition to a series of deletion mutants of the VDR and a MSCs differentiation assay. Notably, compared with the other investigated compounds, CN showed the highest promotion for the activity of VDRE promoter; but CL and SL with the higher similarity with CN, exhibited reduced VDRE promoter activity. This indicated that the ketone group of CN may be a functional group that is associated with the activity of VDRE. The present study also revealed that CN induced the expression of the VDR target genes, such

as osteopontin, which suggested that the VDR pathway was activated by CN. This result is consistent with observations on the activity of the VDRE promoter from the present study. siRNA-mediated knockdown of VDR in the MSCs differentiation model system revealed that the pro-differentiation effects of CN required VDR. Furthermore, a series of deletion mutants of the VDR were generated and used to identify that the C-terminal region of the VDR is responsible for the action of CN. These results are consistent with previous reports that nuclear receptors are activated by small lipophilic ligands (5,7,9).

The present findings have promising clinical applications. With the prevalent-use of CN, it may be possible to improve the quality of engineered bone via MSC-mediated osteogenesis. This is an example of pharmacological control of MSC-mediated osteogenesis and highlights the need for an extensive screen of other families of small molecules both for bone formation and to regulate other differentiation pathways. Manipulation of MSC-mediated osteogenesis is an interesting alternative approach as the agents used *in vitro* were cleared from the graft before implantation.

VDR is an important target for osteoporosis and targeting VDR pharmacologically for the purpose of modulating stem cell function is a promising area in regenerative medicine. By identifying the role of VDR in MSCs and developing corresponding specific ligands as modulators, the therapeutic delivery of stem cells may become more controlled and efficient. Hence, the activation of vitamin D receptor by CN may provide a novel concept for the treatment of bone diseases, such as fracture repair and osteoporosis. Finally, systemic administration of CN may be used to treat patients by targeting either their endogenous or donated MSCs.

In conclusion, the present study demonstrated that a stem cell analysis system based on VDR signaling may be used to identify molecules that induce the osteogenic differentiation of MSCs and specifically revealed that CN significantly increased the efficiency of osteogenic differentiation of MSCs by activating the vitamin D receptor. These findings may have various applications in the field of bone and joint repair and raise interesting questions about the role of CN in osteogenesis.

Acknowledgements

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