

ORIGINAL ARTICLE



Targeting a therapeutically relevant concentration of alendronate for *in vitro* studies on osteoblasts

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ABSTRACT

Objective: Bisphosphonates like alendronate mainly exert their effects on osteoclasts. However, osteoblasts are also affected, but exposed to a much lower concentration *in vivo* than the osteoclasts. Given that the effects are dose-dependent, the intention of the study was to identify a therapeutically relevant concentration of alendronate for *in vitro* studies on osteoblasts.

Materials and methods: Primary human osteoblasts were incubated with alendronate (5, 20 and 100 µM) for 1, 3, 7 and 14 days. Proliferation and viability were assessed, and the effects on cellular growth and function were evaluated by multianalyte profiling of selected proteins in cell culture media using the Luminex 200™.

Results: The viability was not affected by any of the dosages. Exposure to 5 µM alendronate had a neutral effect on osteoblast proliferation, and on secretion of osteogenic and inflammatory markers, while enhancing synthesis of a marker of angiogenesis. 20 µM alendronate induced a decline in proliferation and affected angiogenic and osteogenic biomarkers adversely. 100 µM alendronate reduced proliferation dramatically, and this dosage was excluded from further experiments.

Conclusion: A concentration of 5 µM alendronate exerted effects on human osteoblasts that may translate to those observed *in vivo* and could therefore be relevant for *in vitro* studies.

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Introduction

The bisphosphonate (BP) alendronate (ALN) is the most widely used drug in treatment of osteoporosis [1]. BPs have demonstrated to be efficacious by increasing bone mineral density (BMD) and reducing fracture rates [2]. These effects are mainly attributed to inhibition of bone resorption mediated by the osteoclasts [3]. However, the effect on BMD cannot completely explain the substantial reduction in fracture incidence in patients treated with BPs. There is a body of evidence that BPs also interact with the osteoblasts, and a stimulatory effect on osteoblast proliferation and maturation has been shown *in vitro* [4]. Moreover, both *in vitro* and *in vivo* studies have demonstrated that BPs are capable of preventing osteoblast and osteocyte apoptosis [5].

A multitude of *in vitro* studies have been performed to explore the mechanisms of action of BPs on osteoblasts and other cell types and how these potentially may explain positive or adverse effects observed *in vivo* [6,7]. These effects are dose-dependent, and it is challenging to identify a concentration that reflect the *in vivo* conditions. To approach

this, it is essential to understand bone metabolism and the behaviour of BPs in the body, here exemplified by ALN. The BP is rapidly eliminated from the circulation, binds with high affinity to hydroxyapatite and tends to concentrate in the vicinity of active osteoclasts, rather than at sites of bone formation [8]. In a rat study, more than 70% of the osteoclast surface was densely labelled 24 h after administration of a single dose of [³H]-ALN (0.4 mg/kg), in contrast to only 2% of the bone forming surface [9]. When osteoclasts secrete proteolytic enzymes and hydrochloric acid for the purpose of resorption, ALN is released and rapidly engulfed by the osteoclasts [10]. Hence, osteoclasts are obviously the cells exposed to the highest concentration of ALN. The level of ALN that can be achieved in the osteoclast resorption lacuna has been calculated to be as high as 10⁻⁴ M to 10⁻³ M (100–1000 µM) in newborn rats [9].

The *in vivo* and *in vitro* actions of BPs are well-described for osteoclasts [11,12], whereas the effects on osteoblast function remain to be fully elucidated. As elaborated on above, the osteoclasts are exposed to substantially higher concentrations than osteoblasts *in vivo* [13]. Notably,

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concentrations of BPs as low as 10^{-11} M have been shown to have an effect on osteoblasts, whereas concentrations above 10^{-5} M appear to be toxic [14]. In addition, Li et al. found that the minimum inhibitory concentration of the BP zoledronic acid (ZA) on mouse osteoclasts was 10^{-6} M [15].

In order to achieve a more comprehensive understanding of how BPs affect the cells involved in osteogenesis, the concentration of the drug applied *in vitro* should ideally be as close to relevant therapeutic concentrations as possible. Accordingly, the *in vitro* dosage that promotes similar effects in osteoblasts as described *in vivo* would be the preferable. The aim of the present study was to delineate the effects of different concentrations of ALN on human osteoblasts *in vitro*, and relate this to *in vivo* effects of ALN in order to identify a therapeutically relevant concentration or range of concentrations for *in vitro* studies.

Materials and methods

Study design

Primary human osteoblasts at passage 4 from tibia of a one-day old female donor (Cambrex BioScience, Walkersville, MD, USA) were grown in Lonza Osteoblast Growth Media (OGM) (Cambrex BioScience), containing ascorbic acid, foetal calf serum and gentamicin. Cells were subcultured at 37 °C in a humidified atmosphere of 5% CO₂ prior to confluence, according to manufacturers' instructions.

Cells were seeded in 12-well plates and incubated with ALN (Sigma-Aldrich Biotechnology, Saint Louis, MO, USA) dissolved in OGM at concentrations of 5, 20 and 100 µM. For protein quantification, cells and cell culture media were harvested after 1, 3, 7 or 14 days of incubation, with the last change of medium with or without ALN 24 hrs prior to harvest. Unexposed cells at each time point were used as control. Cells and cell culture supernatants were collected and stored at –80 °C until analysis.

Cell viability and proliferation

Cell viability was confirmed by monitoring the activity of lactate dehydrogenase (LDH) in cell culture medium. LDH was measured using the microplate-based Cytotoxicity Detection Kit (LDH; Boehringer, Mannheim, Germany). In accordance with the manufacturers' protocol, 50 µL aliquots of cell culture medium was used, and the absorbance was read using a microplate reader (ELx800, BioTek, Bad Friedrichshall, Germany) at 450 nm.

The proliferation rate of the cells was measured by [³H]-thymidine incorporation and the MTT colorimetric assay. In the [³H]-thymidine incorporation assay, the cells (1.7×10^4 cells/well in 12-well plates) were incubated with cell culture medium containing 5, 20 and 100 µM ALN for 1 and 3 days ($n = 3$). The cells were pulsed with 1 µCi [³H]-thymidine/well 12 h prior to harvest, and upon harvest, the medium was removed, and the cells were washed twice with PBS and twice with 5% trichloroacetic acid (TCA) to remove unincorporated [³H]-thymidine. The cells were solubilized in 500 µL

of 1 M sodium hydroxide (NaOH), and 200 µL of the solubilized cell solution was transferred to 4 mL scintillation fluid (Lumagel LSC BV; GE Groningen, Netherlands) and counted for 3 min in a liquid scintillation counter (Packard 1900 TR, Packard Instruments, Meriden, CT, USA).

The MTT colorimetric Cell Growth Assay (CT02 Chemicon, Merck KGaA, Darmstadt, Germany) was performed according to manufactures' instruction. Cells (4×10^3 cells/well in 96-well plates) were harvested after 3 days of incubation with the various concentrations of ALN ($n = 3$). Unexposed cells were used as control. The absorbance was measured using an ELISA plate reader (ELx800, BioTek, Vermont, USA) at a test wavelength of 570 nm with reference wavelength of 630 nm.

Protein quantification in cell culture medium

Multianalyte profiling was performed using the Luminex 200TM system (Luminex Corporation, Austin, TX, USA), and acquired fluorescence data were analysed by the xPONENT 3.1 software (Luminex).

Prior to analysis, aliquots of the cell culture medium were concentrated 5 times using Microsep Centrifugal tubes with 3 kDa cut-off (Pall Life Science, Ann Armour, MI, USA). A simultaneous quantification of 29 cytokine and chemokine biomarkers in 25 µL of cell culture media were ascertained using the 29-Milliplex Human Cytokine Immunoassay kit (Millipore, Billerica, MA, USA). The evaluated biomarkers included epidermal growth factor (EGF), eotaxin, granulocyte colony-stimulating factor (G-CSF), granulocyte-macrophage colony-stimulating factor (GM-CSF), interferon alpha-2 (IFN-α2), IFN-γ, interleukin-1a (IL-1a), IL-1b, IL-1ra, IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, IL-8, IL-10, IL-12p40, IL-13, IL-15, IL-17, interferon gamma-induced protein 10 (IP-10), monocyte chemoattractant protein-1 (MCP-1), macrophage inflammatory protein-1a (MIP-1a), MIP-1b, tumour necrosis factor-α (TNF-α), TNF-β and vascular endothelial growth factor (VEGF). In addition, the levels of osteoprotegerin (OPG), osteocalcin (OC), leptin, TNF-α, sclerostin, and fibroblast growth factor 23 (FGF-23) were simultaneously determined in 25 µL of cell culture media using Milliplex Human Bone Panel Immunoassay kit, HBNMAG-51K-7plex (Millipore).

Luminex Multiplex Bead Immunoassays are solid phase sandwich immunoassays, which are designed to be analysed with a Luminex instrument. All the reagents and tools needed were provided in the kit and analyses were performed according to the manufacturers' protocols. In brief, in each of the above-described multiplex kits, 25 µL cell culture medium was applied in 96 cell plates, diluted with assay buffer, and incubated over night at 2–8 °C with antibody coated fluorescent magnetic beads. After washing, analyte-specific biotinylated detector antibodies are added and incubated with the beads. Excess biotinylated detector antibodies were removed by washing and streptavidin conjugated to the fluorescent protein, R-Phycoerythrin (Streptavidin-RPE), was added and the mixture incubated. After the final washing step acquired fluorescence data were quantified based on the spectral properties of the beads and the amount of

Table 1. The impact of ALN on osteoblast viability (LDH) and proliferation (^3H -thymidine and MTT).

Method	Day	5 μM	20 μM	100 μM
LDH	1	100.5 $\pm$ 4.5	90.8 $\pm$ 44.5	110.4 $\pm$ 69.1
	3	103.4 $\pm$ 5.1	75.2 $\pm$ 17.8	84.6 $\pm$ 12.8
^3H -thymidine	1	84.1 $\pm$ 37.0	65.9 $\pm$ 19.8	37.3 $\pm$ 20.7
	3	128.9 $\pm$ 12.2	124.5 $\pm$ 21.1	7.2 $\pm$ 2.5 ^{††§}
MTT	3	95.5 $\pm$ 7.8	83.2 $\pm$ 6.0 [†]	34.9 $\pm$ 15.2 ^{††§}

[†] $p \leq .05$; ^{††} $p \leq .001$ compared to 5 μM .

[§] $p \leq .001$ compared to 20 μM .

LDH, lactate dehydrogenase; ^3H -thymidine incorporation in proliferating cells; MTT, abbreviation for the dye compound 3-(4,5-Dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide. Data are calculated relative to unexposed control cells at each time point and presented as percentages ($n = 3$).

associated R-Phycoerythrin (RPE) fluorescence in a Luminex 200TM. The concentration (pg/mL) of each biomarker in the samples was determined based on standard curves for each of the individual analytes in the kit.

Statistical analysis

Statistical evaluation was performed using the software SigmaPlot 13.0 and 14.0 (Systat Software, San Jose, CA, USA); statistical significance was assessed by Student's *t*-test and *p*-value set to .05, given a passed test of normality and equality.

Results

Viability and proliferation

There was no significant difference in the cell viability (LDH) between the cultures after exposure to 5, 20 or 100 μM of ALN at any of the time points tested (Table 1). Evaluation of thymidine incorporation indicated that there was a marked reduction in proliferation of the osteoblasts after three days of exposure to 100 μM ALN compared to 5 and 20 μM ALN ($p \leq .001$). This correlates with the findings in the MTT assay, where incubation with 100 μM ALN for three days resulted in a diminished proliferation compared to incubation with 5 and 20 μM ALN ($p \leq .001$). Moreover, the MTT assay disclosed a reduced proliferation of the cells exposed to 20 μM ALN for three days, compared to 5 μM ALN ($p \leq .05$) (Table 1).

The effect of ALN on secretion of biomarkers relevant to osteogenesis

As a consequence of the abolished proliferation after exposure to 100 μM ALN for 3 days, we chose to focus on 5 and 20 μM for the remainder of the study. Following an initial rise in secretion of sclerostin from cells exposed to 5 μM ALN ($p = .008$), there was no change in the concentration compared to control after 14 days (Figure 1). Sclerostin secretion increased gradually up to 14 days in cells incubated with 20 μM ALN ($p = .044$), however, not significantly compared to 5 μM ALN exposed cells. The release of granulocyte colony-stimulating factor (G-CSF) was transiently enhanced after incubation with 5 μM ALN ($p = .026$ day 3 and $p = .025$ day 7) (Figure 2a). After 14 days of exposure to 20 μM ALN, G-CSF

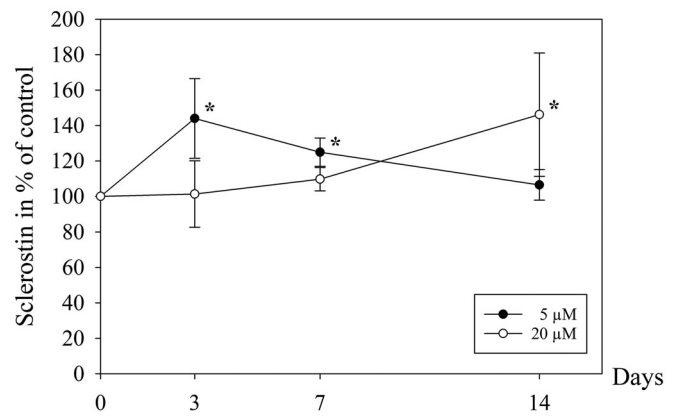


Figure 1. Sclerostin in cell culture media from human osteoblasts. The cells were exposed to 5 μM or 20 μM ALN. Data are presented in % relative to unexposed control at each time point. * $p \leq 0.05$ compared to control, $^{\dagger} p \leq .05$ compared to 5 μM .

was reduced by 50% compared to the cells incubated with 5 μM ALN ($p = .020$) and unexposed cells ($p = .017$).

The effect of ALN on secretion of biomarkers relevant to angiogenesis

There was a non-significant initial reduction in the release of VEGF from osteoblasts after exposure to both 5 and 20 μM ALN. This was followed by an increase of VEGF to control level after 14 days of incubation with 5 μM ALN, and a reduction of VEGF by 50% in cells exposed to 20 μM ALN compared to 5 μM ALN ($p = .003$) and unexposed cells ($p = .006$) (Figure 2b). The secretion of MCP-1 rose to 155% of control after 3 days of incubation with 5 μM ALN ($p = .019$) (Figure 2c), with a subsequent decline to control level after 14 days. In cells exposed to 20 μM ALN, the level of MCP-1 was reduced by 70% compared to cells incubated with 5 μM after 14 days ($p \leq .001$).

The effect of ALN on secretion of biomarkers relevant to inflammation

Compared to unexposed cells, there was a significant rise in the secretion of IL-8 after 1 day of exposure to 5 μM of ALN ($p = .002$) (Figure 2d). 20 μM ALN promoted a drop in IL-8 from day 3, resulting in 50% and 75% lower levels after 7 days ($p = .007$) and 14 days ($p \leq .001$), respectively, compared to 5 μM . A transient non-significant increase in IFN- $\alpha 2$ was observed in cells incubated with 20 μM ALN (Figure 2e), thereafter IFN- $\alpha 2$ decreased, and was 30% lower after 3 days and 35% lower after 14 days when compared to 5 μM ($p = .022$ and $.015$, respectively).

Most of factors tested were not found to change significantly at any time point (leptin, OC, OPG, IFN- γ , and IL-1 α), or the concentrations were lower than the set levels of detection for the analyses (FGF-23, EGF, eotaxin, IL-10, IL-12p40, IL-13, IL-15, IL-17, IL-1 α , IL-1 β , IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, IP-10, MIP-1 α , MIP-1 β , TNF- α and TNF- β).

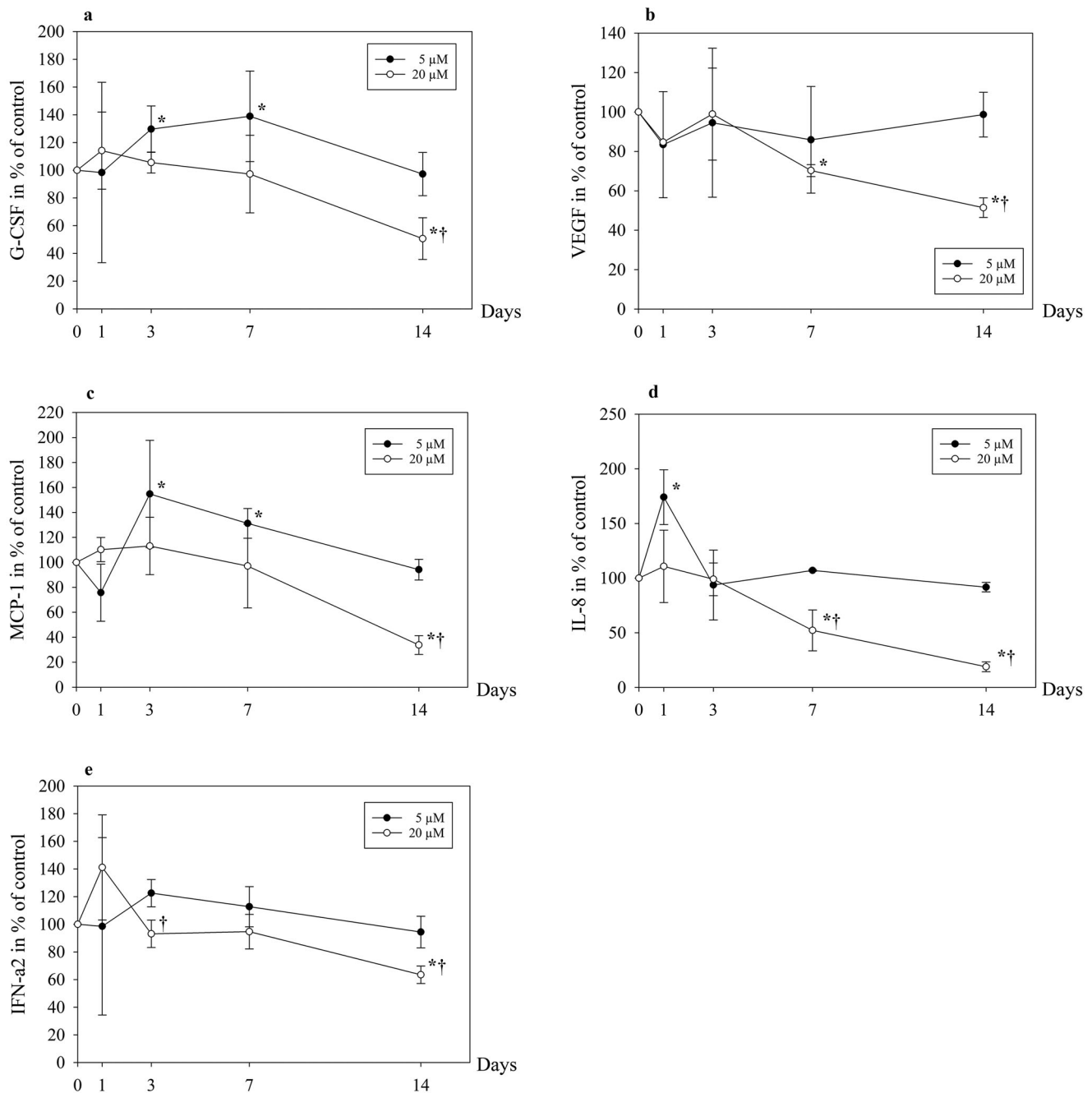


Figure 2. Granulocyte colony-stimulating factor (G-CSF) (a), vascular endothelial growth factor (VEGF) (b), monocyte chemoattractant protein (MCP-1) (c), interleukin-8 (IL-8) (d) and interferon- α 2 (IFN- α 2) (e) in cell culture media from human osteoblasts. The cells were exposed to 5 μ M or 20 μ M ALN. Data are presented in % relative to unexposed control at each time point. * $p \leq .05$ compared to control, † $p \leq .05$ compared to 5 μ M.

Discussion

In this study, exploring the *in vitro* effects of different concentrations of ALN on human osteoblasts, the viability was not affected by any of the dosages. Exposure to 5 μ M ALN had a neutral effect on osteoblast proliferation, and on secretion of osteogenic and inflammatory markers, while enhancing synthesis of MCP-1, a marker of angiogenesis. 20 μ M ALN induced a decline in proliferation and affected angiogenic and osteogenic biomarkers adversely after 14 days of exposure. ALN at a concentration of 100 μ M reduced proliferation dramatically, and this dosage was excluded from further experiments.

Higher concentrations of ALN reduce proliferation

As previously mentioned, the positive skeletal effects of BPs are mainly mediated by suppression of osteoclastic activity. This is achieved through inhibition of the mevalonate pathway, ultimately leading to apoptosis of the osteoclasts [16]. *In vivo* studies indicate that BPs also may affect osteoblasts and osteocytes [17,18]. As elaborated on in the introduction, osteoclasts are exposed to higher concentrations of BPs than cells of the osteoblastic lineage [13]. Thus, when exploring the effects of BPs on osteoblasts *in vitro*, it is of importance to identify a dosage that is close to the *in vivo* concentration.

ALN in concentrations of 10^{-5} M or higher have been reported to affect osteoblasts adversely, inducing apoptosis and inhibiting cell differentiation [19]. We observed no effect on viability after exposure to 5, 20 and 100 μ M ALN for three days, as assessed by LDH activity. The effect on proliferation was, however, dose-dependent. In accordance with the findings of Garcia-Moreno et al. [20], incubation with 100 μ M ALN resulted in a steep decline in osteoblast proliferation after 3 days. The proliferation assessed with an MTT-assay was also reduced after 3 days of exposure to 20 μ M ALN, compared to 5 μ M. We did, however, not observe enhancement of osteoblast proliferation by 5 μ M ALN. Probably an even lower dosage of ALN would be needed to exert this effect.

Notably, ALN has been shown to abolish the glucocorticoid-induced apoptosis in vertebral cancellous bone osteocytes and osteoblasts in mice [21]. These findings suggest that the effect of BPs in glucocorticoid-induced osteoporosis and osteoporosis in general may be due, in part, to their ability to prevent osteocyte and osteoblast apoptosis [3], this has also been confirmed in other studies [22]. Preservation of the bone-forming function of mature osteoblasts and maintenance of the osteocytic network is decidedly of significance for normal function of bone. Thus, to mimic *in vivo* conditions, a dosage of 5 μ M is clearly the better option.

Low dose of ALN has the least impact on biomarkers affecting osteogenesis

The Wnt signalling pathway is essential in bone formation, and bone homeostasis, as well as bone repair and regeneration following injury [23]. This signalling pathway plays an important role in controlling the differentiation of mesenchymal stem cells (MSCs), in favour of the osteoblasts, as well as promoting osteoblast maturation and survival [24]. Bone remodelling is constantly ongoing to replace old bone tissue by new bone tissue. Activation of the Wnt/ β -catenin signalling pathway leads to increased proliferation and differentiation of osteoblast precursor cells, reduces apoptosis of mature osteoblasts, and promotes the ability of differentiated osteoblasts to inhibit osteoclast differentiation [25]. Sclerostin, which is a protein predominantly secreted by osteocytes, is a potent antagonist to the Wnt pathway [26], and as such potentially inhibits bone formation and stimulates bone resorption [24]. We found that 5 μ M ALN induced a temporary rise in secretion of sclerostin after 3 days of exposure, whereas 20 μ M ALN induced a gradual rise in sclerostin throughout the duration of the experiment. In postmenopausal osteoporotic women treated with ZA, an early rise in sclerostin serum levels was observed [27]. On the other hand, no significant alteration in circulating sclerostin levels was seen in a retrospective observational study after long-term treatment of postmenopausal women with the oral BPs ALN and risedronate [28].

Furthermore, we found that ALN at a dosage of 5 μ M promoted a transient increase in release of G-CSF from osteoblasts, followed by a decline to control level after 14 days. 20 μ M ALN induced a pronounced decrease in G-CSF, with a

50% reduction after 14 days of exposure. A rise in G-CSF has also been reported in mice in response to a BP [29]. G-CSF has been mainly associated with the recruitment of stem cells from bone marrow [30,31]. Effects on osteoclast and osteoblast activity have also been reported, the data are, however, diverging [32–34]. The significance of our findings regarding G-CSF under *in vivo* conditions is uncertain.

Low dose of ALN promotes angiogenesis

The development of a microvasculature and microcirculation is critical for the homeostasis and regeneration of living bone, without which, the tissue would simply degenerate and die [35,36]. Thus, it is essential that drugs used in the treatment of osteoporosis do not inhibit angiogenesis. This complex process is orchestrated by multiple factors and mechanisms, which when in balance will contribute to insure a sufficient supply of nutrients and minerals. Osteoblasts have been shown to secrete VEGF, an essential angiogenic growth factor which is critical in the initial stages of wound healing and bone repair [37]. ALN in the concentration range 10^{-5} – 10^{-3} M has been reported to enhance VEGF secretion from osteoblasts [38], whereas 10^{-12} – 10^{-6} M of ZA and ALN reduced the VEGF secretion from osteoblastic cell lines in a dose-dependent manner [19]. We observed no significant changes in VEGF secretion from osteoblasts exposed to 5 μ M ALN, however a pronounced decline occurred after exposure to 20 μ M ALN, which reduced VEGF by 50% compared to 5 μ M ALN.

MCP-1 has been recognized as another important angiogenic chemokine, which is involved in induction of VEGF-A gene expression [39]. We noted a substantial rise in MCP-1 after 3 days of incubation with 5 μ M ALN, thereafter levelling off to control values after 14 days. In contrast, there was a marked drop in secretion of MCP-1 after 14 days of exposure to 20 μ M ALN. Accordingly, our data indicate that ALN in a dosage of 5 μ M could favour angiogenesis, whereas 20 μ M seemed to affect it negatively.

5 μ M ALN with marginal impact on inflammatory biomarkers after 14 days of exposure

The osteoblasts are capable of producing a wide range of cytokines and growth factors that are involved in bone damage and repair [40,41]. These substances include among others IL-1, IL-6, IL-8, TNF- α , IFN- α and IFN- γ which interact in balancing bone metabolism [42]. The expression of several proinflammatory cytokines from osteoblasts and osteoblast-like cells are affected by lower concentrations of BPs *in vitro* [43,44]. We have previously shown that 5 μ M ALN stimulated release of proinflammatory cytokines [45]. In the present study, an initial spike in the expression of IL-8 was seen after exposure to 5 μ M ALN, whereas 20 μ M induced a decline after 7 and 14 days. This is at odds with another study, showing that 100 μ M ALN promoted an increase in the secretion of IL-8 [46]. IL-8 has for decades been known as an important cytokine regarding development of inflammation, mainly through its activation of neutrophils [47].

Release of IFN- α 2 was also affected differently by the low and high dosages, 20 μ M evoking an initial rise, followed by a decline, whereas the cells exposed to 5 μ M ALN responded in a similar manner as control cells. The cytokine IFN- α 2 is mainly accredited as an important factor in the immune response to a microbial infection [48], but has also been applied in the treatment of several malignant diseases [49]. Moreover, it has been suggested as an inhibitor of osteoclasts differentiation [50].

In this study we have compared the effects of 5, 20 and 100 μ M of ALN on osteoblasts *in vitro*, addressing effects on viability, proliferation, and the release of factors affecting inflammation, angiogenesis and osteogenesis. We intended to identify a therapeutically relevant concentration of ALN for *in vitro* studies on osteoblasts. Taken together, we observed that osteoblasts exposed to 5 μ M ALN responded in a similar manner as control cells after 14 days, whereas the higher dosages negatively affected proliferation and factors regarding angiogenesis and osteogenesis. Therefore, it is reasonable to propose that ≤ 5 μ M ALN *in vitro* are concentrations that best mimic observations made *in vivo*.

Osteonecrosis of the jaw (BRONJ) and atypical femur fractures are rare adverse effects of BPs. Under these circumstances, the concentration of ALN is probably higher. These adverse effects tend to occur after long-term treatment and subsequent accumulation of ALN in bone. Schaudinn et al. applied energy-dispersive X-ray spectroscopy (EDS) to estimate the concentrations of ALN in jaw bone by measurement of percent nitrogen incorporation [51]. They observed that ALN concentrations correlated with both duration of therapy and BRONJ stage. Notably, in an inflammatory setting, peripheral blood mononuclear cells seem to increase their release of pro-inflammatory cytokines (TNF- α and IFN- γ) as a response to internalizing BPs [52]. Moreover, a local acidic environment, i.e. periodontitis, has been reported to amplify the dose-dependent cytotoxic effects of BPs, potentially triggering further osteolysis and an increased concentration of BPs [53]. It should be recalled that there may be differences between individuals and between skeletal sites that affect the *in vitro* response to ALN. Hence, when conducting *in vitro* studies on primary human osteoblasts, both skeletal site of origin and donor age are variables of significance [54].

Conclusion

With only marginal changes in viability, proliferation and secretion of factors of growth, angiogenesis and inflammation, incubation with 5 μ M ALN had the least negative impact on the osteoblasts of the concentrations tested in this study. Altogether, based on the current available literature and our own findings, we suggest that ≤ 5 μ M is a therapeutically relevant concentration of ALN for *in vitro* studies on primary human osteoblasts.

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Disclosure statement

No potential conflict of interest was reported by the author(s).

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