Magnesium prevents β-glycerophosphate-induced calcification in rat aortic vascular smooth muscle cells

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Abstract. Vascular calcification (VC), in which high serum phosphate plays a critical role, is one major problem in patients with chronic kidney disease. Clinical studies report that magnesium has a protective effect on VC. However, the studies regarding the impact of high serum magnesium on VC at a cellular level are few and require further investigation. Therefore, the present study explored the effect of magnesium on calcification induced by β-glycerophosphate (BGP) in rat aortic vascular smooth muscle cells (RAVSMCs). In the present study, the addition of magnesium decreased calcium deposition, which was increased by BGP. Higher magnesium levels inhibited BGP-induced alkaline phosphatase (ALP) activity and decreased the expression of core-binding factor α-1 (Cbfα1). In conclusion, higher magnesium levels prevented BGP-induced calcification in RAVSMCs and inhibited the expression of Cbfα1 and ALP. Thus, magnesium is influencing the expression of Cbfα1 and ALP associated with VC and may have the potential to serve as a role for VC in clinical situations.

Introduction

Chronic kidney disease (CKD) is becoming a major public health concern and is also highly prevalent in developing countries (1,2). Recently, based on a report published in the Lancet in 2012, China had a higher incidence and prevalence rate of CKD, estimating the overall prevalence of CKD to be 10.8% (3). The number of patients with CKD in China is estimated to be ~119.5 million (4). The incidence of complications in the CKD patients is increasing with the changes of lifestyle. Vascular calcification (VC) is a frequent complication in patients with CKD and directly or indirectly contributes to cardiovascular disease and increased mortality rates (5,6). It is reported that VC is an actively regulated process. Apart from the passive precipitation of calcium phosphate, the transdifferentiation of vascular smooth muscle cells (VSMCs) into osteoblast-like cells, as well as changes in the expression of bone-associated and mineralization regulating proteins, and apoptosis play critical roles in VC (7). However, the underlying pathophysiological mechanisms are not well understood. It is proposed that along with traditional cardiovascular risk factors, non-traditional risk factors, such as those associated with uremic status and a disturbed bone and mineral metabolism, are likely to control whether or not VC occurs in patients with CKD (8).

Magnesium is involved in numerous important enzymatic processes and also plays a role in skeletal and mineral metabolism and vascular tone (9,10). Data from observational clinical studies showed an inverse association between serum magnesium concentrations and the presence of VC or atherosclerosis (11,12). However, to date, only a limited number of experimental studies performed in animals have confirmed these findings, whereas the majority investigated the role of lowering dietary magnesium on VC (13,14). Thus far, only 2 studies have reported that magnesium has a favorable effect on bone matrix mineralization in an in vitro model (15,16). Therefore, the present study aimed to investigate whether increasing magnesium concentrations influence calcification of rat aortic vascular smooth muscle cells (RAVSMCs).

Materials and methods

Cell culture of RAVSMC. RAVSMCs were prepared as previously described with minor modifications (17-19). Briefly, RAVSMCs were isolated from the thoracic aorta of adult male Sprague-Dawley rats (80-100 g). After the rats had been anesthetized with chloral hydrate (400 mg/kg), the thoracic aorta was obtained under aseptic conditions and cut into small pieces following the removal of residual blood. Small pieces of tissue (1-2 mm²) were placed in a culture dish in Dulbecco's modified Eagle's medium (DMEM) supplemented with 15% fetal bovine serum (FBS; HyClone Corp., Logan, UT, USA), 4.5 g glucose, 100 U/ml penicillin and 100 µg/ml streptomycin in a 5% CO₂ incubator at 37ºC. Cells that migrated from explants were collected when confluent. The cells were maintained in DMEM supplemented with 15% FBS, and the medium was

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replaced twice a week. RAVSMCs were identified by the SMC typical hill and valley appearance, and immunocytochemistry using a monoclonal antibody against the α-smooth muscle actin protein 1A4 (Acta 2) (Santa Cruz Biotechnology, Inc., Dallas, TX, USA) further confirmed the purity of the primary cell culture (20). The cells between passages 6 and 12 were used in all the experiments.

Calcifications assays. RAVSMC calcification was induced as previously described (21). VSMCs were randomly divided into a negative control group, high phosphorus and magnesium intervention groups, a negative control group using low glucose DMEM medium containing 10% FBS and a high phosphorus group in which 10 mM β-glycerophosphate (BGP) was added to normal medium to generate high phosphorus medium. Magnesium chloride was added to the medium for the magnesium intervention groups on the basis of high phosphorus; the final concentrations of magnesium ions (Mg²⁺) were 1, 2 and 3 mM (magnesium intervention groups 1, 2 and 3, respectively), and the stimulating time was 7 days. Control medium was prepared equally but without BGP. The medium was replaced twice a week and Mg²⁺ was added from the beginning and at each medium renewal of the experiment. Supernatants were collected for further investigations. For precise biochemical Ca²⁺ measurements, calcium content in the supernatant was extracted with 6 N hydrochloric acid overnight. The α-cresolphthalein complexone method was conducted using a Calcium Assay kit (BioSino Biotechnology, Beijing, China) and normalized to the protein content of the same culture.

For alizarin red staining (AR), cells were washed twice with phosphate-buffered saline (PBS) and fixed with ethanol 95%. Subsequently, the samples were exposed to 40 mM AR (pH 4.2). Following two washing steps, images were captured of the wells to show the presence of induced mineralization.

Assay of alkaline phosphatase (ALP) activity. Cells were plated at a density of 2 × 10⁵ cells/well into 24-well plates. The cells were cultured for 7 days after intervention and rinsed 3 times with PBS. Subsequently, 500 μl of 0.1% Triton X-100 was added to each well and incubated for 12-24 h at 4°C. The protein assay was performed with the bicinchoninic acid protein assay reagent (Sigma, St. Louis, MO, USA). ALP activity was assayed by a method modified from that of Lowry et al (22). In brief, the assay mixtures contained 0.1 M 2-amino-2-methyl-1-propanol, 1 mM MgCl₂, 8 mM p-nitrophenyl phosphate disodium and the cell homogenates. After incubation for 4 min at 37°C, the reaction was stopped with 0.1 N NaOH and the absorbance was read at 405 nm. Each value was expressed as p-nitrophenol produced in nanomoles/min/microgram of protein.

RNA extraction and reverse transcription-polymerase chain reaction (RT-PCR). RT-PCR was performed to determine the expression levels of the target gene, Cbfα1. Total RNA was extracted from cultured cells using TRIzol reagent (Invitrogen Life Technologies, Grand Island, NY, USA) according to the manufacturer’s instructions. cDNA was synthesized using the advantage RT-for-PCR kit (Clontech, Palo Alto, CA, USA) with oligo (dT) priming according to the manufacturer’s instructions. The GAPDH gene was used as the endogenous control. The primers for Cbfα1 were: 5'-CCGCAGCAGCACAC CGACCAT-3' (sense) and 5'-CGCTCCGCCCCAAT CTC-3' (antisense). The primers for GAPDH were 5'-CAAGGT CATCCATGACAAAATTG-3' (sense), 5'-GTCCACCAC CCTGTTGCTGTA-3 (antisense). The generated PCR products were 289-bp for Cbfα1 and 496-bp for GAPDH. The PCR condition consisted of one incubation for 2 min at 95°C, followed by 35 cycles of a 30-sec denaturation at 95°C, a 30-sec annealing at 55°C and a 45-sec extension at 72°C, and a final extension at 72°C for 5 min. The PCR products were tested in 1% agarose gel in electrophoresis, visualized with ethidium bromide staining and quantified using an image analysis system (Gel work-2ID; Cell Signaling Technology, Shanghai, China). The optical density value of the mRNA expression was calculated.

Statistical analysis. Each experiment was repeated independently 3 times and all the data are expressed as mean ± standard deviation. Means were compared by the Student’s t-test. Means between multiple groups were performed using the analysis of variance (ANOVA). Intergroup comparison was made using ANOVA and the Student-Newman-Keuls test. All the statistical analyses were performed using the SPSS 17.0 software (SPSS, Inc., Chicago, IL, USA). For all the statistical tests, P<0.05 was considered to indicate a statistically significant difference.

Results

Effects of BGP on calcium content of cultured RAVSMCs. Biochemical Ca²⁺ measurements were performed on day 7 after the intervention with various media conditions. The calcium content of the BGP group was significantly higher than those of the control group (Fig. 1A).

Mg²⁺ inhibits BGP-induced calcification in RAVSMCs. To investigate the effects of different Mg²⁺ concentrations on VC, the calcium content was measured of the cells incubated with the indicated conditions. AR staining was also performed on day 7 to further confirm the occurrence of calcium/phosphate (Ca/P) deposition. In the presence of 10 mM BGP disodium, Mg²⁺ at a total concentration of 1, 2 and 3 mM in the medium was effective to significantly inhibit the calcium content on day 7. After 7 days of intervention, Mg²⁺ decreased the calcium content significantly in a dose-dependent manner (Fig. 1A). The effect of Mg²⁺ was evident at 1 mM, and became clearer as the concentration increased and reached the maximum at 3 mM. In addition to the biochemical analysis, Ca/P deposits were also identified by AR staining (Fig. 1B). The size and amount of the granular calcified formations appear to decrease with the addition of Mg²⁺ in a dose-dependent manner.

Effects of Mg²⁺ on Cbfα1 expression and activity of ALP in RAVSMCs. To further validate the effects of Mg²⁺ on the differentiation of RAVSMCs, the gene expression associated with osteoblast differentiation was examined on day 7 after intervention of Mg²⁺ by RT-PCR. RT-PCR showed that the mRNA expression of Cbfα1 was downregulated following intervention with 1, 2 and 3 mM Mg²⁺ (P<0.05) (Fig. 2). After 7 days of
The ALP activity of the BGP group was significantly higher than those of the control group (*P<0.05). Mg^{2+} decreased the activity of ALP significantly in a dose-dependent manner (Fig. 3). The effect of Mg^{2+} was evident at 1 mM, became clearer as the concentration increased and reached the maximum at 3 mM. These data demonstrated that Mg^{2+} plays an important role in preventing BGP-induced calcification in RAVSMCs.

Discussion

Results obtained from previous clinical studies have indicated that lower magnesium levels are associated with increased VC in uremic and non-uremic conditions, and higher serum magnesium results in a decreased risk of peripheral VC in non-diabetic haemodialysis patients (13,23-26). Furthermore, magnesium supplementation improved the media thickness in patients on hemodialysis (12). Notably, the dietary magnesium level influenced the ectopic mineralization in an animal model, which may have broader implications (13,27). However, the detailed mechanisms by which magnesium acts on VC were not clear and the relevant studies were few and limited. The aim of the present study was to provide in vitro evidence for the preventive effect of magnesium on calcification in a cell culture model of RAVSMCs.

Mg^{2+} is the second most abundant intracellular divalent cation and functions as an allosteric modulator of several enzymes, or bridges structurally distinct molecules (28). Previous clinical studies showed that a low magnesium status has an important role in the pathogenesis of cardiovascular disease (29), hypertension (30) and thrombosis (6) in subjects without kidney disease. In the present experimental model, BGP-induced calcification was strongly inhibited by increasing magnesium concentrations in the media of RAVSMCs in a dose-dependent manner and was stable over a period of ≤7 days. Increasing magnesium concentrations decreased calcium deposition, providing evidence that magnesium is able to impair hydroxyapatite crystal growth in in vitro experiments, which is in agreement with the previous studies. It has been suggested that the effect of magnesium on VC may be mediated via this mechanism (31-33). The present study also found that magnesium was not only able to prevent RAVSMC
calcification but also inhibit progression of already established calcification. Even partly reverted calcification was observed with longer incubation time, which renewed the attraction in the therapeutic applications of magnesium on VC. Although phosphate binders are administered to patients of CKD in order to lower serum phosphorus concentrations, as one aspect of VC development, VC progressed overtime. Therefore, magnesium administration or magnesium-containing phosphate binder supplementation in CKD patients may be beneficial as it may partially revert or at least inhibit the progression of VC.

VC is an active, cell-mediated multistep process that includes apoptosis, and the osteochondrogenic differentiation of VSMCs is similar to bone formation, with the role of deposition of calcium-phosphate in terms of matrix mineralization (7,8,34). The bone-related factors, particularly the specific osteogenic transcription factor Cbfa1, were shown to be upregulated in VSMCs undergoing BGP-induced transdifferentiation into osteoblast-like cells. Factors such as ALP, which may contribute to the mineralization process, were also upregulated by elevated phosphate concentrations in in vitro models (35). In the present study, high phosphorus induced the enhanced expression of Ca\(^{2+}\) content, Cbfa1 and ALP, while these decreased when administered with higher magnesium levels, clearly showing that the protective effect of magnesium on RAVSMCs calcification by not only decreasing the deposition of calcium in the cells but also affecting the osteogenic transdifferentiation. However, it appears unlikely, that the inhibitory effect of magnesium on calcification will also account in the bone, as it has been shown that the physiological magnesium concentrations present in bone are not able to inhibit crystal growth (32). The present study suggests that magnesium inhibits the pathological transdifferentiation process from VSMCs into osteoblast-like cells, which does not occur in the physiological process of calcification in the bone. However, detailed clinical investigations regarding the effect of magnesium on bone metabolism, specifically in renal failure patients, are rare and (24) contradictory (36,37). Well designed in vitro and clinical studies to further evaluate the influence of magnesium on bone are required.

In conclusion, higher magnesium levels prevented calcification and further progression of already established calcification in RAVSMCs. Increased magnesium concentrations not only reduced the deposition of calcium, but also inhibited osteogenic transdifferentiation, which are consistent with experimental animal studies and with clinical studies linking elevated magnesium levels to beneficial effects on VC (11,26,38). However, the detailed mechanism by which magnesium may influence this process is unclear. Thus, it appears to be that magnesium influences the molecular processes associated with VC, providing an important insight into the clinical application of magnesium. More clinical research is required to observe the serum magnesium levels in patients with renal failure and to confirm a possible protective effect of magnesium against VC. There is a large gap concerning controlled interventional studies providing unchallengable results to improve the understanding of its role.

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References


