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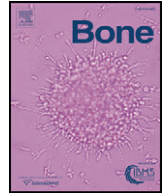
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Implication of the calcium sensing receptor and the Phosphoinositide 3-kinase/Akt pathway in the extracellular calcium-mediated migration of RAW 264.7 osteoclast precursor cells

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ABSTRACT

While the processes involved in the formation, maturation and apoptosis of osteoclasts have been investigated extensively in previous studies, little is known about the mechanisms responsible for the localization and homing of osteoclast precursor cells to the bone environment in order to initiate the bone remodeling process. Recent studies have suggested that the extracellular Ca^{2+} (Ca^{2+}_o) concentration gradient present near the bone environment may be one of the participating factors, producing a chemoattractant effect on osteoclast precursors. Using the murine osteoclast precursor cells of the monocyte–macrophage lineage, the RAW 264.7 cell line, we have shown that Ca^{2+}_o increases the migration of these cells in a directional manner. The participation of the calcium sensing receptor (CaR) in this effect was tested by knocking down its expression through RNA interference, which resulted in an abolition of the migratory response. By the use of specific pathway inhibitors and western blot analysis, the phosphoinositide 3-kinase (PI3K)/Akt and phospholipase $\text{C}\beta$ pathways were shown to be implicated in the migratory effect. The implication of the Akt pathway in the Ca^{2+}_o -induced chemoattraction of RAW 264.7 cells was also confirmed by transducing the cells with the fusion protein TAT-dominant negative-Akt, which decreased the migratory effect. In contrast, the MAPK pathways (ERK1/2, p38 and JNK) were not involved in the production of the migratory effect. We conclude that through the activation of the CaR and subsequent signaling via the PI3K/Akt pathway, Ca^{2+}_o produces a chemoattractant effect on the osteoclast precursor RAW 264.7 cells. These results suggest that the Ca^{2+}_o gradient present near the bone may be one of the initiating factors for the homing of osteoclast precursors to bone, thus possibly playing a role in the initiation of bone remodeling.

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Introduction

Throughout life a continuous process of bone turnover takes place, involving the degradation of old bone and formation of new bone. The bone remodeling process is the result of a collaborative effort of osteoclasts, cells responsible for bone resorption, and osteoblasts, cells in charge of bone formation [1]. These two cell types form temporary anatomical structures, the so-called Basic Multicellular Units (BMUs), which dissolve old bone and replace it with new bone [2]. About 1 million operational BMUs function at any instant in healthy human adults [3,4]. However, the factors underlying BMU initiation are currently not well understood. The first stage of the remodeling process is believed to involve the recruitment of mononucleated osteoclast precursors from the circulation [5]. These cells then

infiltrate the bone lining cell layer, followed by the fusion of osteoclast precursors, thus forming multinucleated osteoclasts, capable of resorbing bone [6]. The resorption phase ends with osteoclast apoptosis, which is then followed by bone formation phase, initiated by the recruitment of osteoprogenitor cells to the surface of resorption lacunae [7]. Osteoprogenitor cells are attracted to this environment due to the release of appropriate chemoattractant signals from bone, which then drive the differentiation of these cells to functional osteoblasts involved in bone formation [8,9].

In addition to its important roles in body movement and the protection of internal organs, the skeleton also serves as a reservoir of calcium and phosphorus and other factors. These bone derived products play an important role in directing various cell types to the bone environment. For example, in the case of osteoclast precursor cells, it has previously been described that peripheral blood monocyte cells (PBMC) are chemotactically attracted to resorbed products of the bone matrix [10,11]. Furthermore, a previous study by Olszak et al. (2000) has described that Ca^{2+}_o stimulates the migration of PBMC

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[12,13]. Since the concentrations of Ca^{2+}_o can reach very high levels near resorbing bone (up to 40 mM) [14,15], it is tempting to speculate that Ca^{2+}_o is one of the signals that initiates the invasion of bone by circulating osteoclast precursors in order to initiate the formation of a BMU and begin the bone remodeling process. While a number of studies have investigated osteoclast formation, differentiation and apoptosis, very little is known about the mechanisms that lead circulating osteoclast precursor cells to the bone environment to begin bone resorption.

For this reason, in this study we examined the effect of Ca^{2+}_o on the migration of a well established osteoclast precursor model, the RAW 264.7 cell line. Our results show that increased Ca^{2+}_o stimulates the migration of osteoclast precursor cells in a concentration-dependent manner. One mechanism by which cells can respond to variations in Ca^{2+}_o concentrations is through the calcium sensing receptor (CaR), a G protein-coupled receptor sensitive to divalent cations, aromatic L-amino acids and aminoglycoside antibiotics [16]. Using RNA interference, we have demonstrated the involvement of the CaR in the migratory effect, and we show that the phosphoinositide 3-kinase (PI3K)/Akt signaling pathway is implicated in the phenomenon. In this work we propose the CaR as a possible candidate for the initiation of the formation of a BMU and the bone remodeling process.

Materials and methods

Materials

RAW 264.7 cells were purchased from American Type Culture Collection (ATCC). Dulbecco's modified Eagle's medium (DMEM), phosphate buffered saline (PBS), trypsin-EDTA and penicillin/streptomycin were provided by Sigma, France. GlutaMAX™ was provided by Invitrogen, France. Wortmannin, U73122, U0126, SB 202190 and SP600125 were purchased from Sigma, France. Bovine serum albumin (BSA), Trypan blue, Giemsa stain, Tween 0.1% and bromophenol blue were provided by Sigma, France. Nitrocellulose membranes, SDS-polyacrylamide gel (SDS-PAGE), Whatman papers and fetal bovine serum (FBS) were purchased from Dominique Dutscher Laboratories, France. Nuclepore Track-Etched membranes (8 µm pore) were purchased from Whatman. ECL substrate solution was provided by Amersham Pharmacia Biotech, Orsay, France. The Lowry protein assay was purchased from BIO-RAD. Boyden Chambers and polycarbonate membranes were provided by Neuroprobe, Inc[®], USA. siPORT NéoFX came from Ambion, France and OPTI-MEM was bought from Gibco, Invitrogen, France. Small interfering RNA (siRNA) directed against CaR and scramble sequences came from Eurogentec, Belgium. Primary mouse anti-CaR antibody was purchased from Bioreagent; rabbit pERK antibody was obtained from Cell Signaling Technology, Boston, MA. Secondary anti-rabbit and anti-mouse peroxidase IgG antibodies were purchased from Sigma, France. Antibodies against P-Akt^{Ser473} and Akt came from Cell signaling Technology, Boston, MA. Antibodies against HA came from Roche, Basel, Switzerland. Phycoerythrin (PE) conjugated IgG mouse antibodies were obtained from BD Biosciences, Le pont de Claix, France. BL21 pLysS *Escherichia coli* came from Stratagene, Amsterdam, Netherlands. Ni-NTA agarose came from Qiagen. All others reagents were purchased from Sigma, France.

Cell culture and stimulation

Murine RAW 264.7 monocyte-macrophage cells were maintained in DMEM supplemented with 10% FBS, 2 mM GlutaMAX™, 100 U/ml penicillin/streptomycin. Cell viability was assayed using Trypan blue dye exclusion, and was typically greater than 95%. Cultured cells were maintained up to passage 20, and then discarded. Before each experiment, cells were serum starved for at least 4 h in DMEM containing 0.2% BSA, preincubated with or without pharmacological

inhibitors for 30 min. For western blot analyses, cells were stimulated for the indicated times and concentrations with Ca^{2+} . Stimulation was stopped by aspirating the medium and washing the cells with ice-cold PBS.

Migration assay

Boyden Chamber cell migration assays were performed using polycarbonate membranes with a pore size of 8 µm. Serum-free DMEM with different concentrations of Ca^{2+} (1.8–20 mM) was added to the lower chamber. 5×10^4 RAW 264.7 cells in 500 µl serum-free DMEM (with or without TAT-Akt proteins) were added to the upper chamber. Cells were allowed to migrate for 5 h at 37 °C in an incubator with 5% CO_2 . After the incubation period, the upper media were removed, chambers were disassembled and membranes were fixed in 96% methanol for 2 min. Filters were stained with Giemsa, then rinsed well with water. Cells on the upper side of the membrane were cleaned off with a cotton swab. The cells were then viewed at 20× magnification under an inverted microscope and counted. To allow for comparison between multiple assays, the data were normalized, and they are expressed as percentage of the number of cells present in the membrane at 1.8 mM Ca^{2+}_o .

CaR-siRNA transfection

Cells were grown in a 6-well plate to a confluence of 25%. They were then transiently transfected either with CaR-siRNA or scramble-siRNA (negative control) using 50 nM siRNA in OPTI-MEM medium and the siPORT NéoFX transfection reagent, following the manufacturer's instructions. The mixture was delicately applied to cells. After a 3 h incubation at 37 °C in an incubator containing 5% CO_2 , the cells received DMEM supplemented with 30% FBS and they received fresh DMEM with 10% FBS the next day. Transfected cells were maintained in culture for 24 h before use.

Preparation of cell lysates and Western blot analysis

After stimulation, cells were washed once with PBS and lysed with 100 µl lysis buffer (25 mM Tris-HCl, pH 7.6, 150 mM NaCl, 10% glycerol, 1% Nonidet P-40, 2 mM Na_3VO_4 and cocktail mini-protease inhibitors), placed on ice for 30 min and then centrifuged at $14,000 \times g$ for 20 min at 4 °C. Protein concentrations were measured using the Lowry protein assay. Following SDS-polyacrylamide gel electrophoresis, equal amounts of protein were electrophoretically transferred to protean nitrocellulose membrane. All SDS-polyacrylamide gel electrophoresis were performed under reducing conditions (β -mercaptoethanol). Blots were blocked, then probed with antibodies raised against the following proteins: CaR, P-Akt^{Ser473}, Akt, HA. Immunoblots were developed using appropriate secondary horseradish peroxidase-coupled antibodies before adding ECL substrate solution and exposing to Kodak X-Omat film. To ensure equal loading of proteins from whole extracts, the membranes were stripped by incubating the membrane for 30 min at 55 °C in 62.5 mM Tris-HCl, pH 6.7, 100 mM β -mercaptoethanol and 2% SDS. The blot was then washed twice with 50 mM Tris-HCl, pH 7.5, 150 mM NaCl and 0.1% Tween 20 and re probed as described above.

Flow cytometry analysis

5×10^5 cells were incubated with CaR monoclonal antibodies or with isotype control antibodies for 30 min on ice in the dark. After washes in PBS-3% FBS and 0.1% sodium azide, cells were incubated with PE-conjugated IgG mouse antibodies for 30 min. After several washes, expression of cell surface CaR was analyzed by FACS (FACS Aria cytometer).

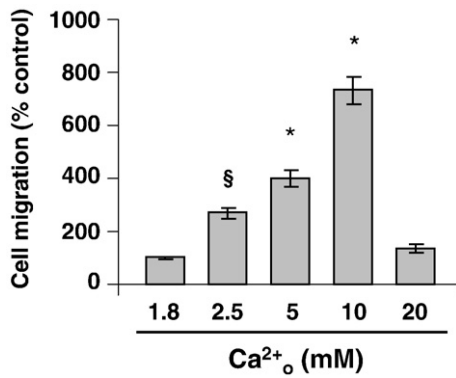


Fig. 1. The effect of Ca²⁺_o on RAW 264.7 cell migration. The effect of different concentrations of Ca²⁺_o (1.8–20 mM) on the migration of RAW 264.7 cells was tested in the Boyden Chamber migration assay. A Ca²⁺_o concentration-dependent increase in migration of RAW 264.7 cells was observed with a maximal effect at 10 mM Ca²⁺_o. Results from four independent experiments are shown. Data were analyzed by *t*-test analysis, § *p*<0.01 and * *p*<0.001 control (1.8 mM Ca²⁺_o) vs. migration at increased Ca²⁺_o.

Purification of TAT–Akt fusion proteins

The cDNAs encoding wild-type (wt) and dominant negative (dn)-Akt were cloned into pTAT-HA plasmid by C.B. and F.G. as previously described [17,18]. Akt fusion proteins were expressed by inducing BL21 pLysS *Escherichia coli* transformed with plasmids encoding the TAT–Akt fusion proteins with 100 μM IPTG for 4 h. Cells were harvested by centrifugation at 4000×*g* for 5 min. The bacterial pellets were resuspended in extraction buffer A (4 M urea, 20 mM HEPES pH 8.0 and 500 mM NaCl) followed by sonication. Lysates were clarified by centrifugation and adjusted to 15 mM Imidazole and loaded on a Ni-NTA agarose column. Afterwards, the resin was washed in buffer A + 15 mM imidazole and eluted with 250 mM imidazole. The eluate was dialyzed against PBS overnight at 4 °C. Purified fusion protein concentrations were determined by Coomassie blue staining of SDS-PAGE compared to BSA standards.

Statistical analyses

The results are expressed as the mean ± the standard error of measurement (S.E.M.) of the indicated number of experiments.

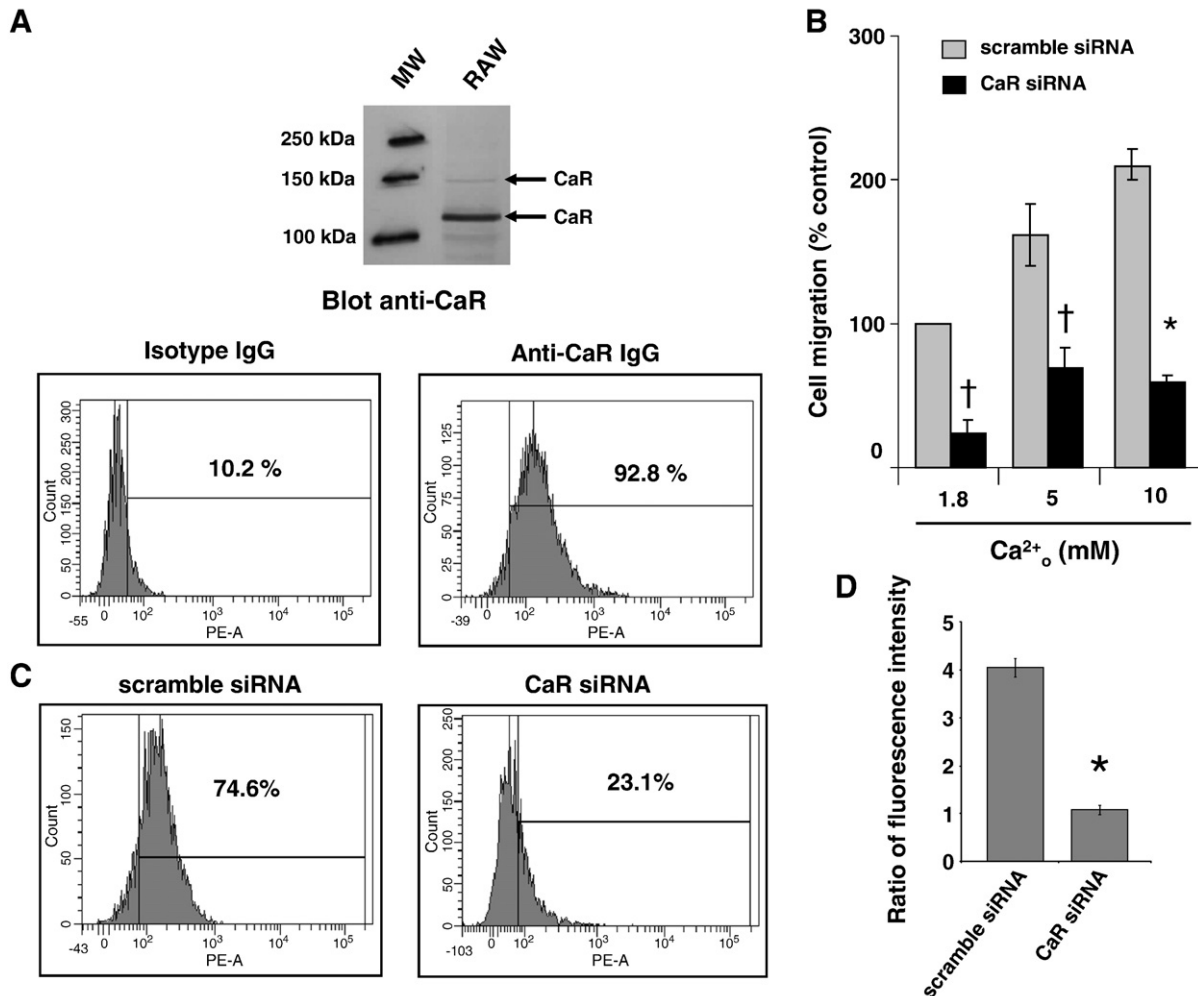


Fig. 2. The Ca²⁺_o-induced migratory effect on RAW 264.7 cells is mediated through the CaR. CaR expression in RAW 264.7 cells was confirmed by Western blot analysis using anti-CaR antibodies and flow cytometry analysis (A). MW = Molecular Weights. The involvement of the CaR in the Ca²⁺_o-induced migratory response of RAW 264.7 cells was assessed using cells transfected either with siRNA directed against the mRNA of this receptor, or scramble-siRNA (control). These cells were then used in Boyden Chamber migration assays to test their migratory ability in response to Ca²⁺_o (B). Results from three independent experiments are shown. Data were analyzed by *t*-test analysis, *, *p*<0.001: scramble-siRNA-transfected vs. CaR-siRNA-transfected cell migration at the same Ca²⁺_o concentration; †, *p*<0.05: scramble-siRNA-transfected vs. CaR-siRNA-transfected cell migration at the same Ca²⁺_o concentration. The efficacy of the siRNA sequences was confirmed by cytometry analysis. Results are expressed in number (count) per PE-conjugated antibody (C) or in ratio of fluorescence intensity (D). *, *p*<0.001: scramble-siRNA-transfected vs. CaR-siRNA-transfected cell.

Statistical analyses were performed using Student's *t*-test. A value of $p < 0.05$ was considered to be significant.

Results

Migration of RAW 264.7 cells increases at elevated Ca^{2+}_o concentrations

Since Ca^{2+}_o has previously been described to regulate the migration of different cell types [12,13,19,20], we studied the effects of Ca^{2+}_o on the migration of RAW 264.7 osteoclast precursor cells. Using the Boyden Chamber assay, we showed that Ca^{2+}_o strongly stimulates the migration of RAW 264.7 cells in the direction of elevated Ca^{2+}_o concentrations (Fig. 1). This effect was Ca^{2+}_o concentration-dependent, with a maximal increase in migration observed at 10 mM Ca^{2+}_o . Higher Ca^{2+}_o concentrations (20 mM) did not affect the migration of RAW 264.7 cells.

The Ca^{2+}_o -induced migration of RAW 264.7 cells is mediated through the CaR

Previous studies have reported the role of the CaR in the migration of various cells [12,13,19,20]. Firstly, we have demonstrated that the CaR is expressed in RAW 264.7 cells by Western blot analysis, corresponding to a 120 kDa protein, and cytometry analyses (Fig. 2A), confirming previous reports [21]. Cytometry analysis has demonstrated that cell surface CaR is expressed by 96% of the cells (Fig. 2A).

Next, to determine whether the CaR is involved in the Ca^{2+}_o -induced migration of RAW 264.7 cells, the expression of the CaR was decreased using specific siRNA targeting the CaR or control scramble-siRNA. With CaR-siRNA-transfected cells, the Ca^{2+}_o -induced migration was greatly reduced compared to cells transfected with scramble-siRNA (Fig. 2B). These results demonstrate that the Ca^{2+}_o -induced migratory effect is mediated through the activation of the CaR. The functionality of the siRNA sequences utilized in this study was confirmed by cytometry analysis that has demonstrated that while 74.6% of scramble-siRNA-transfected cells expressed cell surface CaR, only 23.1% of CaR-siRNA-transfected cells expressed the receptor (Fig. 2C). Moreover, the ratio of receptor expression between CaR-siRNA vs. scramble-siRNA-transfected cells was even greater when total fluorescence intensity was calculated, with receptor expression of around 1:4 (Fig. 2D).

Ca^{2+}_o -induced migration of RAW 264.7 cells is mediated via the PI3K/Akt pathway

To investigate the intracellular pathways involved in the Ca^{2+}_o -induced migration of RAW 264.7 cells, cells were incubated with 10 μM of various specific pathway inhibitors, including inhibitors of PI3K (wortmannin), phospholipase C β (PLC β) (U73122) and inhibitors of different MAPKs, ERK1/2 (U0126), p38 (SB 202190) and JNK (SP600125). This was followed by testing cell migration in a Boyden Chamber assay at different concentrations of Ca^{2+}_o , as described above. Fig. 3 shows that the Ca^{2+}_o -induced migration of RAW 264.7 cells was completely inhibited by wortmannin, demonstrating that the PI3K pathway is involved in the Ca^{2+}_o -induced migratory effect. Cell viability after wortmannin treatment remained unchanged compared to control when assessed by Trypan blue exclusion (data not shown). The inhibitor of PLC β also dramatically reduced the Ca^{2+}_o -induced migratory response (Fig. 4A). This was in contrast to the MAPK pathways, inhibition of which produced no effect on the migratory response mediated by Ca^{2+}_o (Figs. 4 B–D).

One of the known targets of PI3K is the serine–threonine kinase Akt. We studied the activation of Akt in RAW 264.7 cells in response to Ca^{2+}_o by western blot analysis. Ca^{2+}_o induced the phosphorylation of Akt in a concentration-dependent manner (Fig. 5A). This effect was maximal after 5 min of stimulation with Ca^{2+}_o and decreased after

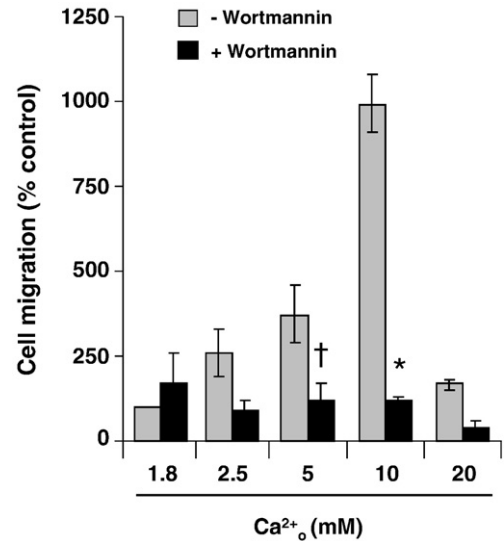


Fig. 3. The PI3K pathway is involved in the Ca^{2+}_o -induced migratory response of RAW 264.7 cells. RAW 264.7 cells were preincubated with or without the PI3K inhibitor wortmannin (10 μM), prior to their use in the Boyden Chamber migration assay, to assess their chemoattraction towards elevated Ca^{2+}_o . Results from three independent experiments are shown. Data were analyzed by *t*-test analysis. *, $p < 0.001$: untreated vs. wortmannin-treated cell migration at the same Ca^{2+}_o concentration; †, $p < 0.05$: untreated vs. wortmannin-treated cell migration at the same Ca^{2+}_o concentration.

30 min of stimulation (Fig. 5B). The phosphorylation of Akt did not change when cells were stimulated with 10 or 20 mM Ca^{2+}_o , suggesting that other molecular mechanisms may be responsible for the modulation of cell migration at high calcium concentrations. The involvement of the CaR in the Ca^{2+}_o -induced Akt activation was confirmed by siRNA directed against this receptor, which abolished Akt activation (Fig. 5C).

To assess the involvement of Akt in RAW 264.7 cell migration, these cells were transduced either with wt-Akt or mutant dn-Akt K179M. To prepare the cell permeable Akt fusion protein, an expression vector expressing Akt as a fusion protein with the protein transduction domain (PTD) of the HIV-1 TAT protein was constructed. Then, different recombinant TAT–Akt fusion proteins were generated (Fig. 6A). A wt- or a dn-form of Akt, containing a point mutation inactivating its kinase activity (K179M), was fused to the protein transduction domain of the TAT HIV protein. Proteins TAT–wt-Akt and TAT–dn-Akt were generated and purified by affinity chromatography on a nickel column. The purity of TAT–wt-Akt and TAT–dn-Akt fusion proteins was verified by Coomassie blue stained gel. The purified products were confirmed by western blotting with anti-HA and anti-Akt antibodies (Fig. 6B). Transduction efficiency of the different TAT–Akt proteins in RAW 264.7 cells was evaluated by western blot analysis with anti-Akt and anti-HA antibodies (Fig. 6C). We observed equal transduction of both proteins. The biological effects of the TAT–Akt proteins on the migration of RAW 264.7 cells in response to Ca^{2+}_o were then evaluated. As shown in Fig. 6D, transduction of TAT–dn-Akt completely inhibited cell migration induced by Ca^{2+}_o , whereas TAT wt-Akt had no effect. These results demonstrate that Akt plays an important role in RAW 264.7 cell migration in response to Ca^{2+}_o .

Discussion

In the parathyroid, the CaR plays a primary role in the regulation of systemic Ca^{2+}_o homeostasis due to its ability to sense minute changes in Ca^{2+}_o concentrations, leading to modulation of parathyroid hormone secretion [22]. Recent data have reported that the CaR is also expressed by other organs and tissues involved in Ca^{2+}_o homeostasis such as

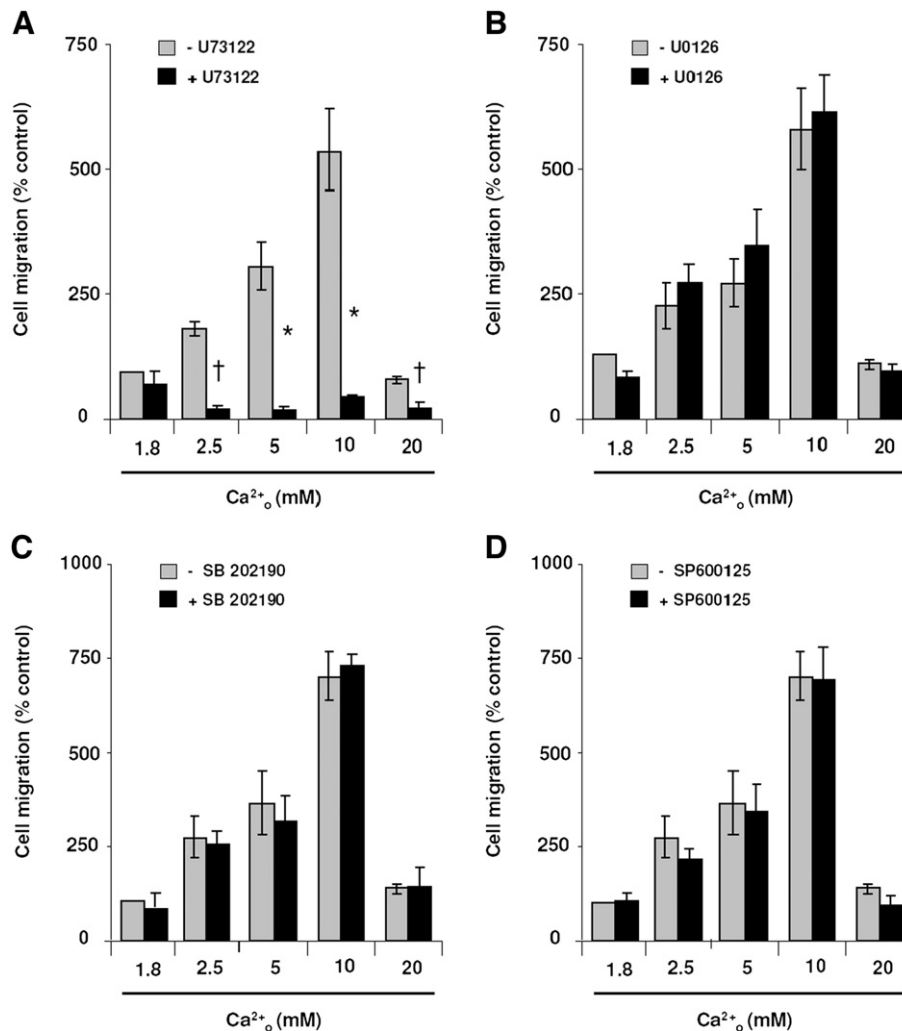


Fig. 4. PLC but not MAPKs activation is involved in the Ca²⁺_o-induced migratory response of RAW 264.7 cells. RAW 264.7 cells were preincubated with or without the inhibitors of PLC or MAPKs, *i.e.* ERK1/2 (U0126), p38 (SB 202190) and JNK (SP600125), prior to their use in the Boyden Chamber migration assay, to assess their chemoattraction towards elevated Ca²⁺_o. Results from three independent experiments are shown. Data were analyzed by *t*-test analysis, †, *p*<0.05 and *, *p*<0.001: untreated vs. inhibitor-treated cell migration at the same Ca²⁺_o concentration.

the bone [23,24] and kidneys [25,26], as well as in tissues not involved in Ca²⁺_o homeostasis, such as lens epithelial cells and antral gastrin cells of the stomach [27,28]. Expression of the CaR has been confirmed in the different cell types present in bone, including the osteoclasts [24], osteoblasts [23] and osteocytes [23] and their precursors [29]. In these cell types, the CaR is involved in numerous physiological functions, for example osteoclast differentiation and apoptosis [21].

In addition to the previously reported role of the CaR in differentiation, proliferation and apoptosis of bone cells, several recent studies have suggested that this receptor may also play a role in the migration of osteoclast precursor cells, as well as other cells [12,13,19,20]. For example, Adams et al. (2006) have demonstrated the involvement of Ca²⁺_o and the CaR in the homing of hematopoietic stem cells to the bone marrow [12]. For this reason, we hypothesized that Ca²⁺_o and the CaR may play a role in the initiation of the formation of a BMU by recruiting osteoclast precursor cells of the hematopoietic lineage from the circulation to their osseous niches in the bone environment, where high Ca²⁺_o concentrations are present as a result of nearby bone resorption. In order to address this issue, we investigated the effect of Ca²⁺_o on the migration of RAW 264.7 cells of the monocyte–macrophage lineage using the Boyden Chamber migration assay. We have demonstrated that the migration of the osteoclast precursor RAW 264.7 cells increases with elevated Ca²⁺_o

concentrations. This effect was shown to require the presence of the CaR, and knocking down CaR expression using siRNA abrogated the Ca²⁺_o-induced migration.

Furthermore, we wanted to decipher the intracellular signaling pathways responsible for the migratory effect. The CaR couples to G α _q and G α _i G proteins, leading to the activation of several intracellular responses, especially PLC β [30] and MAPKs [31,32], and inhibition of adenylate cyclase activity [33]. In order to investigate the involvement of these pathways in the migratory effect of RAW 264.7 cells in response to Ca²⁺_o, we have utilized different pathway inhibitors, demonstrating that the PI3K/Akt and PLC β pathways are involved in the migratory effect. Next, we addressed the involvement of the Akt, a downstream target of PI3K, in the migration of the RAW 264.7 cells. Ca²⁺_o-induced activation of Akt was demonstrated by western blot analysis and it was shown to be mediated through the CaR. Furthermore, the role of Akt in the migratory effect was shown by transducing the cells with dominant negative TAT–Akt, which decreased the migration of the cells in response to Ca²⁺_o. In contrast, inhibition of the p42/44, p38 and JNK MAPKs did not abolish the Ca²⁺_o-induced migration of RAW 264.7 cells, suggesting that these pathways are not required for the migratory response. It is not yet clear whether the CaR present in osteoclasts and osteoclast precursors couples to the MAPKs. While it has been demonstrated that the CaR present in

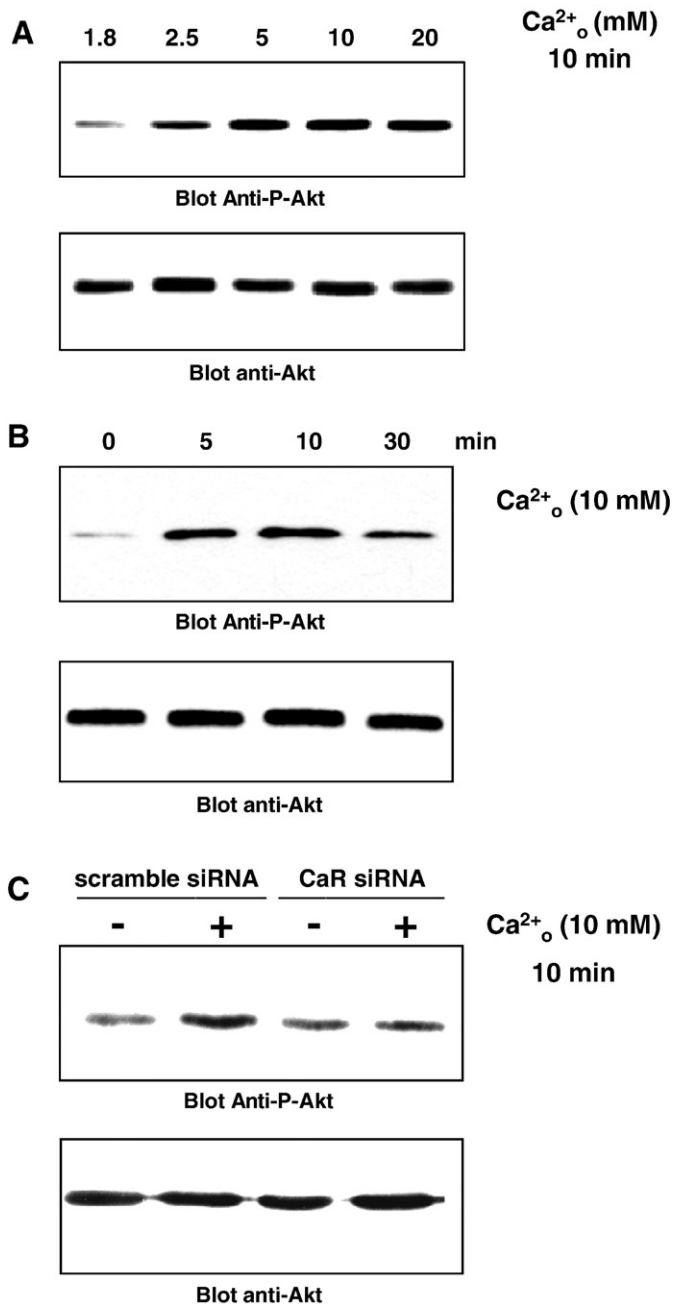


Fig. 5. Ca^{2+}_o -induced Akt phosphorylation is mediated through the CaR. RAW 264.7 cells were stimulated either with different Ca^{2+}_o concentrations (1.8–10 mM) for 10 min (A) or with 10 mM Ca^{2+}_o for 0–30 min (B). Cell lysates were analyzed by SDS-PAGE and western blotting using specific antibodies which recognized the phosphorylated form of the Akt (P-Akt). The blots was then stripped and reprobed with anti-Akt antibodies. The involvement of the CaR in the Ca^{2+}_o -induced Akt activation was demonstrated by the use of siRNA directed against the receptor (10 mM Ca^{2+}_o , 10 min) (C). Data shown are representative of three independent experiments.

osteoblasts does activate p38, p42/44 and JNK [31,32], the intracellular signaling pathways activated via the CaR in osteoclasts require further investigation. However, this study has nevertheless shown that the MAPKs are not involved in the Ca^{2+}_o -induced migratory effect, and importantly, we have clearly identified the PI3K/Akt pathway as being pivotal for the Ca^{2+}_o -induced migration of RAW 264.7 cells.

Activation of the PI3K pathway through the CaR has previously been described in different cell types [34,35]. It has been described that cell proliferation is mediated through the activation of this

pathway via the CaR [36,37]. Furthermore, Liu et al. (2003) have shown that the Ca^{2+}_o -induced secretion of serotonin from parafollicular cells is mediated through the CaR and subsequent activation of PI3K [38]. The involvement of the PI3K pathway in the migratory effect reported in our study is consistent with several recent findings. For example, it has been shown that osteoclasts derived from mice lacking the p85 α regulatory subunit of PI3K demonstrate a significantly reduced ability to migrate [39]. Additionally, in a study by Pilkington et al. (1998), the authors have proposed that the directional motility of mature osteoclasts in response to M-CSF is mediated through the PI3K pathway [40], and it has also been reported that the PI3K pathway is involved in the platelet derived growth factor-induced migratory effect [41,42]. However, contrasting findings have also been reported, for example Godwin and Soltoff (1997) have shown that Ca^{2+}_o -induced chemotaxis of MC3T3-E1 is mediated through a pertussis-toxin sensitive pathway and is independent of the PI3K pathway, shown by its resistance to wortmannin [41].

A possible source of the local increase in Ca^{2+}_o may originate from bone microdamages, as a result of daily activity. Through bone remodeling, bone cells are able to repair any microdamage, thus contributing to the maintenance of the structural integrity of bone and calcium homeostasis [43]. However, the mechanism by which microcracks are detected, and repair initiated, is not yet understood. From our data, it can be speculated that microdamage of the bone may result in the release of Ca^{2+}_o , stimulating the homing of cells specialised in bone-repair. It is also conceivable that this increase in Ca^{2+}_o is the result of an initial osteoclast activity which may lead to the recruitment of further osteoclast precursors, whose transformation into osteoclasts is promoted by the CaR [21]. While our study demonstrates that the migration of RAW 264.7 increases in response to elevated Ca^{2+}_o through the activation of the CaR/PI3K/Akt, thus possibly implicating this pathway in the homing of osteoclast precursor cells to bone and the initiation of the establishment of BMUs, the shortcomings of our study need to be stressed. Whether the CaR plays a pivotal role in the bone remodeling process in a physiological context still needs to be elucidated in future studies. The experimental model used in this study is limited and it is not suitable to address this issue. Therefore, to gain a better understanding of the precise mechanisms involved in the initiation of a functional BMU, a novel experimental approach needs to be developed, that maximally mimics the physiological conditions of the bone.

Interestingly, Garner et al. (2001) have reported that expected skeletal manifestations of PTH excess were not the main skeletal features in CaR knock-out mice, which rather showed growth plate calcification and disorderly deposition of mineral, excessive osteoid accumulation, and prolonged mineralization lag time in metaphyseal bone [44]. Given our results, certain outstanding questions still remain. Does the bone remodeling process take place in these animals? Do the osteoclast precursors in these animals target the correct sites in the bone in order to initiate bone remodeling? For all these reasons, in future investigations it would be very interesting to study the migration of osteoclast precursors derived from the bone marrow of $\text{CaR}^{-/-}$ mice. Our current study aimed to determine certain aspects of the intracellular signaling that takes place upon CaR activation in osteoclast precursor cells and the involvement of these pathways in the Ca^{2+}_o -induced chemoattraction. Given the results we obtained on the effects of high concentrations of calcium on the differentiation, bone resorption and apoptosis of osteoclasts, it can be hypothesized that the CaR is involved in the recruitment of osteoclast precursor cells from the circulation during the establishment of a functional BMU [21]. Furthermore, activation of the CaR may also lead to a negative feedback effect at higher concentrations of Ca^{2+}_o , released from bone during the bone resorption process. It is possible that the CaR is also involved in the infiltration of these cells through the bone lining cell layer and fusion of these cells into functional

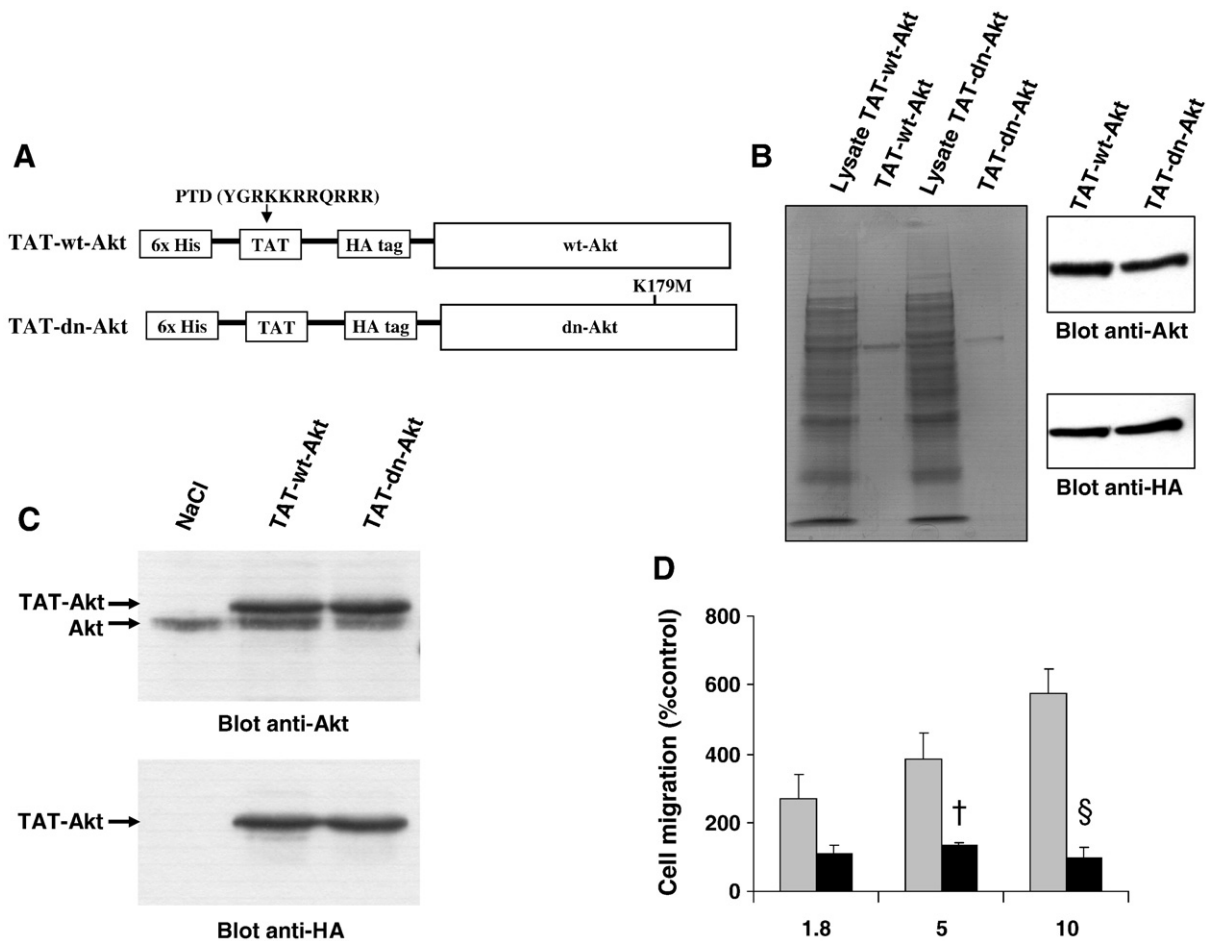


Fig. 6. The effect of recombinant TAT-Akt fusion proteins on Ca^{2+} -induced migration of RAW 264.7 cells. Schematic representation of the TAT-Akt constructs (A). Resulting recombinant Akt (wt and dn) proteins were fused at the N-terminal to 6 His-Tag sequence followed by the protein transduction domain (PTD) of the TAT protein and HA-Tag. Purity of recombinant TAT-Akt proteins was assessed by Coomassie blue staining of SDS-PAGE and western blot analysis of different TAT-Akt proteins with anti-HA and anti-Akt antibodies (B). RAW 264.7 cells were either transduced with different TAT-Akt proteins (100 nM) or treated with (PBS) for 24 h. After extensive washes, the presence of recombinant TAT-Akt proteins in RAW 264.7 cells was analyzed by western blotting with indicated antibodies (C). To determine the involvement of the Akt pathway in the migration of RAW 264.7 cells, the cells were transduced with 100 nM TAT-wt-Akt or TAT-dn-Akt, and cell migration was assessed in a Boyden Chamber assay (D). Results from three independent experiments are shown. Data were analyzed by *t*-test analysis, †, $p < 0.05$ and §, $p < 0.01$: TAT-wt-Akt- vs. TAT-dn-Akt-transduced cell migration at the same Ca^{2+} concentration.

resorbing osteoclasts. Clearly, these questions need to be addressed in future studies.

In conclusion, this study has demonstrated that the CaR and the PI3K/Akt pathway are involved in the chemoattraction induced by Ca^{2+} on osteoclast precursor RAW 264.7 cells. These findings suggest that the CaR may possibly have a role in the initiation of bone remodeling *in vivo*.

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