Clinical Study

The Impact of Vitamin D₃ Supplementation on Mechanisms of Cell Calcium Signaling in Chronic Kidney Disease

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Intracellular calcium concentration in peripheral blood mononuclear cells (PBMCs) of patients with chronic kidney disease (CKD) is significantly increased, and the regulatory mechanisms maintaining cellular calcium homeostasis are impaired. The purpose of this study was to examine the effect of vitamin D₃ on predominant regulatory mechanisms of cell calcium homeostasis. The study involved 16 CKD stages 2-3 patients with vitamin D deficiency treated with cholecalciferol 7000–14000 IU/week for 6 months. The regulatory mechanisms of calcium signaling were studied in PBMCs and red blood cells. After vitamin D₃ supplementation, serum concentration of 25(OH)D₃ increased (𝑃<0.001) and [Ca²⁺]ᵢ decreased (𝑃<0.001). The differences in [Ca²⁺]ᵢ were inversely related to differences in 25(OH)D₃ concentration (𝑃<0.01). Vitamin D₃ supplementation decreased the calcium entry through calcium release activated calcium (CRAC) channels and purinergic P2X₇ channels. The function of P2X₇ receptors was changed in comparison with their baseline status, and the expression of these receptors was reduced. There was no effect of vitamin D₃ on P2X₇ pores and activity of plasma membrane Ca²⁺-ATPases. Vitamin D₃ supplementation had a beneficial effect on [Ca²⁺]ᵢ decreasing calcium entry via CRAC and P2X₇ channels and reducing P2X₇ receptors expression.

1. Introduction

Vitamin D hormonal system has been classically implicated in the regulation of calcium homeostasis and bone metabolism. However, it has also noncalcitropic effects through the activation of tissue vitamin D receptors (VDR) [1]. Vitamin D insufficiency/deficiency is a significant risk factor for the development of various chronic diseases, and the deficiency of calcidiol (25(OH)D₃) as well as calcitriol (1,25(OH)₂D₃) is common in CKD patients [2]. Therefore, the supplementation of native vitamin D (cholecalciferol or ergocalciferol) or active vitamin D (calcitriol and VDR activators) in CKD is well established.

Free cytosolic calcium concentration ([Ca²⁺]ᵢ) is controlled by mechanisms that regulate Ca²⁺ entry from the extracellular space and Ca²⁺ release from intracellular stores and by the activity of ATP-dependent Ca²⁺ pumps and antiporters that move Ca²⁺ back into stores or out of cells [3]. Already in early stages of chronic kidney disease (CKD), [Ca²⁺]ᵢ, and calcium concentration of intracellular stores were significantly increased in comparison with healthy volunteers, and the regulatory mechanisms of calcium signaling were impaired by the disease [4–7].

Calcium enters into the cells by any of the general classes of calcium/cation channels. In nonexcitable cells like peripheral blood mononuclear cells (PBMCs), the predominant Ca²⁺ entry pathway is the store-operated one, in which the emptying of intracellular Ca²⁺ stores activates the Ca²⁺ influx. This type of the channel is known as the calcium release activated calcium (CRAC) channel in lymphocytes.

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The calcium entry through CRAC channels activates certain transcription factors which regulate the gene expression for cytokines responsible for immune responses [8, 9].

Another mechanism of calcium entry into the cell is represented by purinergic P2X receptors. At the present time, purinergic signaling is accepted as a crucial component of diseases and was found to mediate a vast array of biological processes. The P2X receptors are expressed primarily on cells of hemopoietic origin, where they participate in immune responses, cell proliferation, cell death, bone formation, and bone resorption [10]. The P2X receptor is a bifunctional purinoreceptor that opens a nonselective cation channel and consecutively forms a large, cytolytic pore. The key factor of P2X-dependent cytotoxicity is the massive intracellular Ca\(^{2+}\) increase triggered by its activation. This can lead to membrane blebbing and cell death by apoptosis or necrosis. There is an increasing body of evidence implicating P2X receptors in various pathological conditions [11–14].

The plasma membrane Ca\(^{2+}\)-ATPases (PMCA) responsible for removing excessive Ca\(^{2+}\) out of the cells to extracellular environment. The decreased PMCA activity increases [Ca\(^{2+}\)], and affects some intracellular processes.

To our knowledge, little information is available regarding the impact of vitamin D₃ supplementation on disturbed cell calcium homeostasis in CKD. Therefore, the aim of the present study was to examine the effect of vitamin D₃ supplementation on essential regulatory mechanisms of cell calcium homeostasis.

2. Materials and Methods

2.1. Patients. The study population consisted of 16 nondiabetic patients with CKD (9 patients CKD stage 2 and 7 patients CKD stage 3). All of them were screened and followed up in the outpatient department of nephrology at the Slovak Medical University. The diagnosis of CKD was based on clinical and laboratory examinations as defined by the K/DOQI criteria [15]. Causes of their renal disease were glomerulonephritis in 9 patients, tubulointerstitial nephritis in 3 cases, hypertensive nephrosclerosis in 2 patients, and other 2 causes. The glomerular filtration rate was estimated by the MDRD study formula [16]. Patients with acute impairment of renal function, nephrotic proteinuria, malignancies, and derangements in mineral metabolism of nonrenal origin were excluded from the study. Concurrent treatments interfering with mineral metabolism were not allowed. Previous therapy with vitamin D₂/D₃, calcitriol, or over-the-counter vitamin D preparations had to be cancelled at least 2 months before enrollment. Hypertension was the most common comorbidity present in all patients and treated with ACE inhibitors or angiotensin II receptor blockers in 14, diuretics in 8, beta blockers in 6, and calcium channel blockers in 8 cases. Dihydropyridine calcium channel blockers were allowed as they do not interfere with studied parameters and effects in PBMCs [17]. All patients had vitamin D deficiency (serum 25(OH)D₃ concentration <30 ng/mL) and were supplemented with cholecalciferol 7000–14000 IU/week for 6 months; the dose (approximately 1000–2000 IU/day) was chosen as a common supplementary dose for the treatment of vitamin D deficiency in general population.

The study was approved by the Ethics Committee of the Slovak Medical University and all participants gave their written informed consent.

2.2. PBMCs Isolation. Human PBMCs were isolated by the Ficoll gradient centrifugation, diluted 1:1 with RPMI-1640 medium, layered onto an equivalent volume of medium LSM-1077, and centrifuged at 700 g for 20 min at 22°C, as previously reported [5]. The PBMC layer was washed in 40 mL RPMI, resuspended in 10 mL RPMI and 10% fetal bovine serum (FBS), and centrifuged at 300 g 10 min at 22°C and the pellet was resuspended in 2 mL aliquots of physiological salt solution. Final concentration of PBMCs was adjusted to 2.5 × 10⁶ cells/mL. Our preparation contained lymphocytes (94–96%), monocytes (3–4%), and natural killer cells (the rest), as determined by flow cytometry (Coulter Epics XL, Ireland). The cell viability was quantified using a 0.8% solution of trypan blue and estimated to be 96–98%.

2.3. Red Blood Cell Membranes Isolation. Isolated red blood cell (RBC) membranes were used to assess the PMCA function. RBC membranes were obtained by hemolytic fragmentation in hypotonic media using standard method of Hanahan and Ekholm [18] modified in our laboratory to achieve a higher quality of the ghosts. RBCs from previous isolation were diluted 1:5 with physiological salt solution and centrifuged at 1270 g for 20 min at 4°C. The supernatant was removed and the procedure was repeated one more time. RBCs were then diluted 1:5 with tris(hydroxymethyl)amino methane (TRIS) medium (20 mmol/L, pH = 7.4) and centrifuged at 7700 g for 35 min at 4°C. This step was repeated twice for each TRIS medium with decreasing concentrations (20, 10, and 5 mmol/L).

2.4. Intracellular Ca\(^{2+}\) Measurements. The population of 2 × 10⁶ PMBCs/mL was loaded with fluorescence dye Fluo-3 AM at a final concentration of 2 μmol/L for 40 min at 22°C in a physiological salt solution. After incubation, the cells were centrifuged at 300 g, washed three times with a physiological salt solution, and kept at room temperature for 10 min before use. The Fluo-3 fluorescence was measured at 37°C in FluoroLog 3–11 spectrophotometer (HORIBA Jobin Yvon Inc., Edison, NJ, USA) with an excitation at 488 nm (bandpass 3 nm) and an emission at 526 nm (bandpass 5 nm). Each experiment was followed by [Ca\(^{2+}\)]i calibration to estimate the actual free cytoplasmic calcium concentration from the measured fluorescence signal (F) in each cell population. [Ca\(^{2+}\)]i was quantified in nmol/L according to the following equation:

\[
[Ca^{2+}]_i = K_d \cdot \frac{F - F_{min}}{F_{max} - F} , \tag{1}
\]

where \(K_d = 400 \text{ nmol/L} \) at 37°C [19]. The maximal fluorescence intensity (\(F_{max}\)) was assessed by the addition of Triton X-100 (0.1%) with Ca\(^{2+}\) (5 mmol/L), and the minimum
fluorescence level ($F_{\text{min}}$) was determined after the addition of 25 mmol/L EGTA (pH = 9). Digitonin (20 μmol/L) was used to answer for minimal compartmentalization [20]. To assess the role of CRAC channels in Ca$^{2+}$ entry, 2-aminoethyl-diphenyl borate (2APB), a widely used inhibitor of these channels was applied [21]. Although there are already more potent and selective inhibitors of these channels [22], 2APB was used for the possibility of comparing the results with our previous studies [4, 6]. The action of 2APB (50 μmol/L) was studied in cells where Ca$^{2+}$ entry through these channels was stimulated by thapsigargin (Tg) (1 μmol/L), a specific inhibitor of endoplasmic reticulum Ca$^{2+}$-ATPase. To examine the function of P2X<sub>7</sub> receptors, AZ11645373 (50 nmol/L), a highly selective antagonist of human P2X<sub>7</sub> receptors [23], and 2',3'-O-(4-benzoyl)benzoyl ATP (BzATP) (50 μmol/L), the most potent and selective agonist of these receptors, were used [24].

2.5. Cell Surface P2X<sub>7</sub> Receptors Expression. PBMCs were stained with P2X<sub>7</sub> polyclonal antibody labelled with fluorescein isothiocyanate (FITC) according to the protocol provided by the manufacturer. Briefly, 100 μL of PBMCs were incubated either with FITC-conjugated anti-P2X<sub>7</sub> (2 μg/mL) or FITC-conjugated IgG2a (2 μg/mL, control) for 20 min at 22 °C in the dark. PBMCs were simultaneously stained with phycoerythrin- (PE-) conjugated CD14 antibody in order to exclude monocytes from the examined population. After the incubation, cells were washed three times, dissolved in 500 μL of PBS, and subjected to the analysis on flow cytometer (Cytomics FC 500 cytometer, Beckman Coulter, USA). Values of control samples stained with FITC-conjugated IgG2a were subtracted from the evaluated results.

2.6. P2X<sub>7</sub> Receptors Visualization by Fluorescence Microscopy. To visualize the surface P2X<sub>7</sub> receptors, PBMCs were stained with P2X<sub>7</sub> (extracellular) antibody. Cell imaging was performed by using the Axiovert 200 inverted microscope (Carl Zeiss, Germany) with mercury lamp HBO-100 and FluoroArc driver. The 450–490 nm bandpass filter and FT510 dichroic mirror were used for excitation, and emission was detected with a 515 nm long-pass filter. Images were recorded by the PentaMax cooled CCD camera (Roper Scientific) with KODAK KAF-1400 chip. Images were taken with the C-Apochromat 40x/1.2 water immersion objective (Zeiss, Germany).

2.7. Ethidium Bromide Uptake by Flow Cytometry. The ethidium bromide uptake by PBMCs was measured on the Cytomics FC 500 cytometer (Beckman Coulter, USA) and results were processed with the CXP software. PBMCs at concentration of 10⁶ cells/100 μL were kept at room temperature when ethidium bromide at 30 μmol/L final concentration was added to the sample. After a 5 min incubation, the mean channel ethidium fluorescence was assessed in FL3 sensor after excitation with a 488 nm laser beam. Up to 10000 events were involved in the data analysis. For the investigation of P2X<sub>7</sub> pore function, ethidium fluorescence was measured after a 10 min incubation with either BzATP (50 μmol/L) or AZ11645373 (50 nmol/L) at 37 °C.

2.8. PMCA Activity Measurement. The RBCs membrane suspension was added to working medium (mmol/L): 100 TRIS, 80 KCl, 3 MgCl₂, 0.2 ethylenediaminetetraacetic acid (EDTA), and 1 ouabain (pH = 7.4) in the presence or absence of CaCl₂ (5 mmol/L). The reaction was started by the addition of 40 mmol/L of ATP, conducted for 60 min at 37 °C and stopped by the addition of a 15% trichloroacetic acid (TCA). The amount of liberated inorganic phosphate was determined using a phosphat colorimetric assay kit (BioVision) on the UV–VIS spectrophotometer Shimadzu UV-1700 (Shimadzu Corp., Japan). The estimated PMCA activity was calculated as the difference between the activities of the enzyme incubated in the presence and absence of CaCl₂ and was expressed as nmol of Pi/mg protein/h. The protein concentration was measured by the method of Lowry [25].

2.9. Reagents. The physiological salt contained (mmol/L) 140 NaCl, 5.4 KCl, 1 CaCl₂, 1 Na₂HPO₄, 0.5 MgCl₂, 5 glucose, and 5 HEPES (pH = 7.4). Thapsigargin (Tg) was procured from Calbiochem (San Diego, CA). 2-Aminoethyl-diphenyl borate (2APB), ethidium bromide, AZ11645373, and 2',3'-O-(4-benzoyl) benzoyl ATP (BzATP) were obtained from Sigma-Aldrich (St. Louis, MO, USA). Fluo-3 aceotomethylster (Fluo-3 AM) was from Molecular Probes (Eugene, OR), lymphocyte separation medium LSM-1077 was from PAA Laboratories GmbH (Pasching, Austria), and fetal bovine serum (PBS) and RPMI-1640 medium were from GIBCO (Grand Island, NY, USA). PE-conjugated anti-CD14, FITC-conjugated IgG2a, and anti-P2X<sub>7</sub> (extracellular) FITC were from Sigma-Aldrich (St. Louis, MO, USA). The phosphate colorimetric assay kit was from BioVision (Hayward, CA, USA). All other chemicals were purchased from Sigma-Aldrich.

2.10. Analytical Procedures. Serum calcium and creatinine were measured by the Vitros 250 Analyzer, Johnson & Johnson, Rochester, NY, USA. Intact parathormone and 25-hydroxyvitamin D<sub>3</sub> were determined by the electrochemical luminescence immunoassays (ECLIA) (Roche-Diagnostics, Mannheim, Germany).

2.11. Statistical Analyses. All values are expressed as means ± SD. Statistical analysis was carried out by the SPSS 15.0 (SPSS Inc., Chicago, IL, USA). The Shapiro-Wilk test was used to evaluate a sample normality distribution. The statistical significance of differences was tested by the independent 2-population Student’s t-test for normally distributed data and the Wilcoxon’s test for a nonparametric analysis. A P value < 0.05 was considered significant. The Spearman’s correlations between variables were used as a measure of association.

3. Results
3.1. Effect of Cholecalciferol Treatment on Main Laboratory Variables. Effect of cholecalciferol treatment was evaluated
Table 1: Main laboratory variables at baseline and after the cholecalciferol treatment.

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Baseline</th>
<th>6 months</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total serum calcium (mmol/L)</td>
<td>2.28 ± 0.08</td>
<td>2.34 ± 0.08</td>
</tr>
<tr>
<td>[Ca(^{2+})] (nmol/L)</td>
<td>120 ± 6</td>
<td>105 ± 3**</td>
</tr>
<tr>
<td>iPTH (pg/mL)</td>
<td>43 ± 17</td>
<td>43 ± 20</td>
</tr>
<tr>
<td>25(OH)D(_3) (ng/mL)</td>
<td>18 ± 2</td>
<td>36 ± 9***</td>
</tr>
<tr>
<td>Serum creatinine (μmol/L)</td>
<td>102 ± 21</td>
<td>101 ± 28</td>
</tr>
<tr>
<td>eGFR (mL/min)</td>
<td>64.8 ± 5.4</td>
<td>63 ± 4.2</td>
</tr>
</tbody>
</table>

Values are expressed as mean ± SD. **P < 0.001, n = 16 for comparison with baseline.

3.3. P2X\(_7\) Receptors

3.3.1. P2X\(_7\) Channels. The application of P2X\(_7\) receptors antagonist AZI1645373 (50 nmol/L) led to reduction in [Ca\(^{2+}\)] in CKD patients from 120 ± 6 to 112 ± 8 nmol/L (P < 0.001, n = 16). On the other hand, after the vitamin D\(_3\) supplementation, AZI1645373 had no effect on [Ca\(^{2+}\)], (106 ± 4 versus 104 ± 6 nmol/L; ns, n = 16) (Figure 3(a)). Differences in [Ca\(^{2+}\)], after the inhibition of P2X\(_7\) receptors were significantly decreased after the vitamin D\(_3\) supplementation (P < 0.001, n = 16) (Figure 3(b)). In BzATP stimulated cells, AZI1645373 (50 nmol/L) decreased the calcium influx before and also after the vitamin D\(_3\) supplementation (Figures 3(c) and 3(d)), but the effect of an inhibitor was attenuated (P < 0.01, n = 16) (Figure 3(e)). The 6-month vitamin D\(_3\) supplementation had an inhibitory effect on function of P2X\(_7\) channels and thereby decreased the Ca\(^{2+}\) entry. The agonist of purinergic P2X\(_7\) receptors, BzATP (50 μmol/L), caused a sustained increase in [Ca\(^{2+}\)], in CKD patients at baseline 120±6 to 155 ± 12 nmol/L (P < 0.001, n = 16) and also after the vitamin D\(_3\) supplementation 106 ± 4 to 136 ± 13 nmol/L (P < 0.001, n = 16). However, the P2X\(_7\) receptors activation after the vitamin D\(_3\) supplementation did not reach the values of [Ca\(^{2+}\)], before supplementation (Figure 4(a)). Furthermore, the effect of BzATP (50 μmol/L) on AZI1645373-inhibited calcium influx in PBMCs was evaluated. Under these conditions, a rising calcium influx through P2X\(_7\) channels was found at baseline and after supplementation (Figures 4(b) and 4(c)). All these results demonstrate the inhibitory effect of vitamin D\(_3\) supplementation on calcium entry through P2X\(_7\) channels.

3.3.2. P2X\(_7\) Pores. The uptake of ethidium bromide into PBMCs was measured by flow cytometry at basal conditions and with BzATP (50 μmol/L) stimulation or AZI1645373 (50 nmol/L) inhibition. The permeability of ethidium bromide through P2X\(_7\) pores in PBMCs of CKD patients was significantly increased in comparison with healthy volunteers [4, 5] and remained unchanged after the vitamin D\(_3\) supplementation. The treatment did not change the permeability of P2X\(_7\) pores after the application of either BzATP (50 μmol/L) or AZI1645373 (50 nmol/L) (Figures 5(a) and 5(b)).

3.3.3. Expression of Cell Surface P2X\(_7\) Receptors. The expression of cell surface P2X\(_7\) receptors was 1.5-fold greater on PBMCs from CKD patients compared to healthy donors [4]. We assessed a decreased expression of these receptors after vitamin D\(_3\) supplementation in the whole population of PBMCs (P < 0.001) (Figures 6(a), 6(b), and 6(c)).
Intracellular calcium concentration (nmol/L)

Time (s)

0 100 200 300 400 500

(a)

Intracellular calcium concentration (nmol/L)

0 20 40 60 80

CKD Vit. D CKD

(b)

**

3.4. Plasma Membrane Ca$^{2+}$-ATPases. The PMCA activity of RBCs membranes is decreased by 25% in patients with early stages of CKD when compared to healthy subjects [4]. Vitamin D$_3$ supplementation did not increase the PMCA activity (Figures 7(a) and 7(b)). The concentrations of total plasma membrane proteins were significantly enhanced (7.7±1.6 versus 11.8±2.7 mg/mL, P < 0.001) which may indicate increased expression of PMCA.

4. Discussion

Disturbances in intracellular calcium homeostasis in patients with CKD represent a complex process which aggravates with CKD progression. The mechanisms of cell calcium influx and efflux are impaired in renal disease. Already in early stages of CKD, cytosolic calcium concentration ([Ca$^{2+}$]) and calcium concentration of intracellular stores are increased [6, 7]. The elevated calcium entry through CRAC channels and P2X$_7$ receptors, increased expression of P2X$_7$ receptors, and decreased PMCA activity contribute to this state [4]. We have previously shown that vitamin D$_3$ supplementation in CKD patients led to a decline in [Ca$^{2+}$], to values comparable with healthy people [6]. The aim of this study was to examine the effect of vitamin D$_3$ supplementation on predominant regulation mechanisms of cell calcium homeostasis in nonexcitable cells from patients with early stages of CKD. The principal finding of the present study is that vitamin D$_3$ supplementation affected the mechanisms of intracellular calcium homeostasis as follows: the Ca$^{2+}$ entry through CRAC and P2X$_7$ channels was decreased, while no effect was found on the permeability and functionality of P2X$_7$ pores, and the expression of P2X$_7$ receptors was reduced. Finally, the activity of PMCA was not increased after treatment.

4.1. Vitamin D$_3$ Supplementation. All clinical studies of vitamin D supplementation in CKD patients reported a significant improvement in 25(OH)D$_3$ concentrations, although several studies did not reach a mean 25(OH)D$_3$ concentration in optimal range (≥30 ng/mL). In our study, vitamin D$_3$ supplementation with weekly cholecalciferol dosing significantly increased 25(OH)D$_3$ to the recommended levels. PTH values were in normal range and did not change after treatment, and no correlation was observed between 25(OH)D$_3$ and PTH concentrations. Total serum calcium was not changed but [Ca$^{2+}$]$_i$ significantly decreased after supplementation. Linear regression analysis demonstrated that changes in 25(OH)D$_3$ were significantly and inversely correlated with that of [Ca$^{2+}$]$_i$ (R = 0.617, P < 0.01). Besides our previous work [6], no studies have investigated the relationship between vitamin D status and [Ca$^{2+}$]$_i$ in CKD.

4.2. Ca$^{2+}$ Entry through CRAC Channels. Dysregulation of Ca$^{2+}$ homeostasis involving the endoplasmic reticulum (ER) and store-operated calcium channels has been manifested in patients with neurodegenerative disorders, immunodeficiency, acute pancreatitis, polycystic kidney disease, and cardiac hypertrophy [9, 26–29]. In nonexcitable cells, CRAC channels are the main pathway of Ca$^{2+}$ entry. These channels are composed of two proteins Orail and STIM1 (stromal interaction molecule). Orail protein is located in the plasma membrane and forms the channel pore. It is activated by STIM1 located in the membrane of ER. STIM1 has a dual function of sensing the Ca$^{2+}$ concentration in ER and activating CRAC channels. A decrease in the ER Ca$^{2+}$ concentration induces STIM1 translocation close to the plasma membrane where it binds to and activates the Orail channel. Alterations in STIM1/Orail system may contribute to several pathophysiological conditions including cardiovascular [30, 31] and pulmonary diseases [32], hypertension [33], immunodeficiency, and autoimmune and lymphoproliferative diseases.
Figure 3: The effect of vitamin D₃ supplementation on an inhibitory effect of AZI1645373. (a) The comparison of the AZI1645373 (50 nmol/L) effect on [Ca²⁺], in CKD patients at baseline and after vitamin D₃ supplementation (**P < 0.01, ***P < 0.001, n = 16). After vitamin D₃ supplementation, AZI1645373 (50 nmol/L) had no effect on [Ca²⁺]. (b) Differences in [Ca²⁺], after an inhibition with AZI1645373 (50 nmol/L) at baseline and after vitamin D₃ supplementation (**P < 0.001, n = 16). (c) The representative trace where AZI1645373 (50 nmol/L) inhibited the [Ca²⁺] rise induced by BzATP (50 μmol/L). (d) The comparison of the AZI1645373 (50 nmol/L) effect on BzATP- (50 μmol/L) activated calcium influx before and after supplementation (**P < 0.001, n = 16). (e) Differences in [Ca²⁺], in BzATP- (50 μmol/L) activated PBMCs after the AZI1645373 (50 nmol/L) inhibition at baseline and after vitamin D₃ supplementation (**P < 0.01, n = 16).
In our previous study, we have demonstrated that the function of CRAC channels is altered in PBMCs of patients in early stages of CKD [6]. It is not known what changes in the STIM1/Orai1 system develop over the course of CKD. In the current study, a 6-month vitamin D₃ supplementation significantly reduced the increased Ca²⁺ entry through CRAC channels which contributed to the decrease in [Ca²⁺]ᵢ. In our previous study, a 12-month vitamin D₃ supplementation decreased the Ca²⁺ entry through CRAC channels insignificantly, which could be due to less rigorous patient selection (patients with diabetes mellitus and polycystic kidney disease were included). To our best knowledge, the effects either 25(OH)D₃ or 1,25(OH)₂D₃ on CRAC channels were not studied in CKD.

### 4.3. Ca²⁺ Entry via P2X₇ Channels

The most recent advances provide compelling evidence for P2X receptors playing a key role in regulating physiological and pathophysiological
processes in the kidney [36]. In our study, P2X7 receptors were involved in the disrupted calcium homeostasis in PBMCs of CKD patients. We have shown that the Ca2+ entry via P2X7 channels and pores was increased and also the permeability of P2X7 pores was higher. In addition, the function of P2X7 channels and pores was altered [5]. It is known that 1,25(OH)2D3, an active metabolite of vitamin D, prevents Ca2+ increase through P2X7 channels and reduces the plasma membrane permeability through P2X7 pores in human PBMCs of healthy subjects [17]. 1,25(OH)2D3 can also up- or downregulate the expression of several genes in many cell types. To our knowledge, the effect of 1,25(OH)2D3 on P2X7 receptor expression has not been studied. The data of this study disclosed that vitamin D3 supplementation reduced Ca2+ influx through P2X7 channels and affected their functionality. Moreover, the supplementation had no effect on permeability of P2X7 pores, and the differences in responses to stimulation or inhibition were not found. Different effects of vitamin D3 on P2X7 channels and pores in CKD patients may have been made due to different sensitivity to these channels and pores. It is known that various agonists and antagonists of P2X7 receptors may have a distinct effect on the function of P2X7 channels and/or pores [37]. Not only the function but also the expression may be altered in some pathological conditions [38–41]. In our recent study, we found a 1.5-fold increase in the expression of surface P2X7 receptors on PBMCs from CKD patients compared to healthy subjects [4]. In the current study, the flow cytometric measurement revealed that vitamin D3 decreased the expression of P2X7 receptors by 45%.

4.4. Activity of PMCA. The PMCA is critical for the maintenance of resting [Ca2+]i in nonexcitable cells and may be the last gatekeeper for the control of low [Ca2+]i. We have observed a decreased PMCA activity in early CKD patients [4]. This finding is consistent with other studies and points out that the PMCA activity decreases with kidney disease progression [42]. The decline in PMCA activity may be caused by numerous factors such as calmodulin deficiency, an activity of other endogenous protein regulators, and inhibition of mitochondrial or glycolytic metabolism [43]. The Ca2+ influx through CRAC channels is also an important regulator of PMCA activity, and an altered communication between them may be another cause of PMCA malfunction [44]. 1,25(OH)2D3 is known to increase PMCA expression and activity in Ca2+ transporting tissues such as the intestine, as well as in osteoblasts and Madin-Darby bovine kidney epithelial cells [45–47]. Several experimental studies have observed direct and/or indirect effects of 1,25(OH)2D3 and/or 24,25(OH)2D3 on activity and expression of PMCA in different cells, but the effects of 25(OH)D3 itself have not been studied. The studies targeting at the effects of native vitamin D3 supplementation on PMCA activity in patients with early stages of CKD are also missing. In the present
Figure 6: The expression of surface P2X$_7$ receptors. (a) Representative flow cytometry histograms of PBMCs immunostained with primary antibody for the extracellular domain of the P2X$_7$ receptor (gray peak) and an isotype-matched control (IgG2a, white peak). (b) Immunofluorescence staining of P2X$_7$ receptors in plasma membrane using anti-P2X$_7$ (extracellular) antibody of lymphocytes visualised by fluorescence microscopy. Images represent surface P2X$_7$ receptors in four different samples. (c) Decreased expression of surface P2X$_7$ receptors after vitamin D$_3$ supplementation in a whole population of PBMCs ($^\ast\ast\ast P < 0.001$, $n=16$).

In conclusion, we have shown that vitamin D$_3$ supplementation reduces elevated $[\text{Ca}^{2+}]_i$ via CRAC and P2X$_7$ channels and decreases the expression of cell surface P2X$_7$ receptors in early CKD. We did not find the effect on P2X$_7$ pores and PMCA activity. Thus, vitamin D$_3$ supplementation had a beneficial effect on disturbed cell calcium homeostasis in early CKD.

Conflict of Interests

The authors declare that there is no conflict of interests regarding the publication of this paper.
Figure 7: The assessment of PMCA activity. (a) The representative absorbance spectrum of a CKD patient PMCA in the medium with (black line) and without (grey line) Ca²⁺. (b) The comparison of PMCA activity in CKD patients at baseline and after a 6-month vitamin D₃ treatment.

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