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Role of Calcineurin B Homologous Protein in pH Regulation by the Na^+/H^+ Exchanger 1: Tightly Bound Ca^{2+} Ions as Important Structural Elements[†]

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ABSTRACT: We studied the role of the interaction of calcineurin homologous protein 1 (CHP1) with the Na^+/H^+ exchanger 1 (NHE1), particularly its EF-hand Ca^{2+} binding motifs, in the intracellular pH (pH_i)-dependent regulation of NHE1. We found that $^{45}\text{Ca}^{2+}$ binds to two EF-hand motifs (EF3 and 4) of the recombinant CHP1 proteins with high affinity (apparent $K_d = \sim 90$ nM). Complex formation between CHP1 and the CHP1 binding domain of NHE1 resulted in a marked increase in the Ca^{2+} binding affinity ($K_d = \sim 2$ nM) by promoting a conformational change of the EF-hands toward the tightly Ca^{2+} -bound form. This suggests that CHP1 always contains two Ca^{2+} ions when associated with NHE1 in cells. Interestingly, overexpression of GFP-tagged CHP1 with mutations in EF3 or EF4 significantly reduced the exchange activity in the neutral pH_i range and partly impaired the activation of NHE1 in response to various stimuli, such as growth factors and osmotic stress. Furthermore, we found that, in addition to reducing the activity (V_{max}), a CHP1 binding-defective NHE1 mutant had a marked reduction in pH_i sensitivity (~ 0.7 pH unit acidic shift), which consequently abolished various regulatory responses of NHE1. These observations suggest that the association of NHE1 with CHP1 is crucial for maintenance of the pH_i sensitivity of NHE1 and that tightly bound Ca^{2+} ions may serve as important structural elements in the “ pH_i sensor” of NHE1.

The Na^+/H^+ exchanger (NHE1¹) proteins in the plasma membrane and various organellar compartments of mammalian cells catalyze the electroneutral countertransport of Na^+ for H^+ . Nine distinct isoforms of the Na^+/H^+ exchanger (NHE1 to NHE9) have been isolated to date, and these molecules have been shown to exhibit similar membrane

topologies with 12 predicted N-terminal membrane-spanning helices and a large C-terminal cytoplasmic region (1–10). They show considerable differences in their tissue expression patterns, membrane localization, and kinetic and pharmacological properties. The plasma membrane exchangers (NHE1–5) are primarily involved in regulation of intracellular pH and Na^+ concentration, but they also participate in a broad range of physiological processes, such as cell volume regulation, transepithelial transport of electrolytes, cell proliferation, apoptosis, and differentiation (1–3).

Of the nine isoforms identified to date, NHE1 has been characterized in the most detail. NHE1 is ubiquitously expressed in essentially all tissues and cell types and plays a major role in maintaining intracellular pH and cell volume homeostasis. The activity of NHE1 is controlled by various extrinsic factors, including growth factors, hormones, and mechanical stimuli (1–3). A variety of signaling molecules regulate the NHE1 protein, such as calcineurin B homologous protein (CHP) (11–13), Ca^{2+} /calmodulin (14, 15), the low molecular weight GTPases Ras and Rho (16), p42/44 mitogen-activated protein kinases (17), p90 ribosomal S6 kinase (18), 14-3-3 protein (19), Nck-interacting kinase (20), phosphatidylinositol 4,5-bisphosphate (21), and carbonic anhydrase II (22). Recently, we have focused on the role of CHP in regulation of the activities of the Na^+/H^+ exchangers (12, 13).

CHP was initially discovered as a protein (p22) involved in vesicular transport (23), as well as a molecule that interacted with NHE (11). Since then, CHP has been reported

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¹ Abbreviations: NHE, Na^+/H^+ exchanger; CHP, calcineurin B homologous protein; GFP, green fluorescent protein; CaN, calcineurin; CaM, calmodulin; pH_i , intracellular pH; EIPA, 5-(N-ethyl-N-isopropyl)-amiloride; DMEM, Dulbecco's modified Eagle's medium; HEPES, 2-[4-(2-hydroxyethyl)-1-piperazinyl]ethanesulfonic acid; Tris, Tris(hydroxymethyl)aminomethane; EGTA, O,O'-bis(2-aminoethyl)ethylene glycol-N,N,N',N'-tetraacetic acid; PBS, phosphate-buffered saline; PDGF-BB, platelet-derived growth factor-BB; PMA, phorbol 12-myristate 13-acetate; SDS-PAGE, sodium dodecyl sulfate–polyacrylamide gel electrophoresis.

to exhibit multiple functions, including inhibition of calcineurin phosphatase activity (24), as well as interaction with microtubules (25), DRAK2 (death-associated protein kinase related apoptosis inducing protein kinase 2) (26) and KIF1B β 2 (kinesin-family 1B β 2) (27). Previously, we reported that the ubiquitous CHP isoform (designated as CHP1) is an essential cofactor for the physiological activity of the Na⁺/H⁺ exchanger by interacting with the juxtamembrane region in the C-terminal cytoplasmic domain of plasma membrane exchanger isoforms (12). Furthermore, we reported that the second CHP isoform (CHP2) might be involved in maintenance of the abnormally high pH_i in malignant transformed cells (13). CHP2 is expressed at a relatively high level in the rat small intestine (28), suggesting that it plays a specific role in this tissue. These CHP proteins contain four EF-hand Ca²⁺ binding motifs and are myristoylated at the N-terminus (Gly²). In addition, CHP1 is phosphorylated in cells in a serum-dependent manner (11). However, the roles of these posttranslational modifications of CHP proteins in the pH_i-dependent regulation or acute activation of NHE in response to extracellular stimuli are largely unknown, although this protein family appears to be essential for the physiological exchange activity of plasma membrane NHEs.

In this study, we focused on the EF-hand Ca²⁺ binding motifs of CHP1. We found that the affinity of CHP1 for Ca²⁺ markedly (approximately 40-fold) increases upon complex formation with NHE1, probably by promoting a change in the conformation of the EF-hand motifs. The extremely low Ca²⁺ dissociation constant (~2 nM) of CHP1 suggests that Ca²⁺ ions remain tightly bound to CHP1 when it is complexed with NHE1 in the plasma membrane. On the basis of properties of various CHP1 and NHE1 mutant proteins in cells, we suggest that CHP1 is important for pH_i-dependent regulation of NHE1 and that tightly bound Ca²⁺ ions play an important role in maintaining a structure that is critical for this function of CHP1.

EXPERIMENTAL PROCEDURES

Materials. The amiloride derivative EIPA was a gift from the New Drug Research Laboratories of Kanebo, Ltd. (Osaka, Japan). ⁴⁵CaCl₂, ²²NaCl, and ¹⁴C-benzoic acid were purchased from Dupont-NEN (Boston, MA). The rabbit polyclonal antibodies against CHP1 and NHE1 were described previously (12, 14). All other chemicals were of the highest purity available.

Cells, Culture Conditions, and Stable Expression. The exchanger-deficient cell line PS120 (29) and corresponding transfectants were maintained in DMEM (Life Technologies Inc., Rockville, MD) containing 25 mM NaHCO₃ and supplemented with 7.5% (v/v) fetal calf serum, penicillin (50 units/mL), and streptomycin (50 μ g/mL). Cells were maintained at 37 °C in the presence of 5% CO₂. PS120 cells (5 \times 10⁵ cells/100-mm dish) were transfected with each plasmid construct (20 μ g) by the calcium phosphate coprecipitation technique. Cell populations stably expressing wild-type or mutant human NHE1 were selected by the H⁺-killing procedure as described previously (30). Cells stably overexpressing GFP-tagged CHP1 were first selected with G418, and then single colonies were selected by monitoring GFP fluorescence.

Construction of Expression Vectors. All the constructs were produced by means of a polymerase chain reaction (PCR)-based strategy. For construction of GFP-tagged CHP1 or its mutant forms with mutations in Ca²⁺ binding motifs or in the myristoylated glycine (G2A), a cDNA encoding CHP1 was cloned into the mammalian expression vector pEGFP-N1 (Clontech, Palo Alto, CA). The plasmids carrying cDNAs for the wild-type or mutant NHE1s were all cloned into the mammalian expression vector pECE. Constructs were confirmed by sequencing plasmids with an ABI-PRISM DNA sequencer model 3100 (Applied Biosystems, Foster City, CA).

Purification of Recombinant Proteins. Recombinant histidine-tagged CHP1 proteins were produced in *Escherichia coli* (BL21-Star; Invitrogen, San Diego, CA) transformed with pET11 carrying the cDNA encoding CHP1 containing the C-terminal six histidine residues as described previously (12). Myristoylated CHP1 was produced using the same bacteria except they also contained the vector pBB131, which carries the yeast *N*-myristoyltransferase cDNA (kindly provided by Dr. J. I. Gordon, Washington University). Myristoylation of CHP1 (or p22) produced by this method was previously confirmed (23). For production of the complex of CHP1 and the CHP1 binding region of NHE1, the cytoplasmic region (aa 503–545) of NHE1 was cloned into the vector pET24 and coexpressed with His-tagged CHP1 in *E. coli* in the presence of ampicillin and kanamycin. Myristoylated and nonmyristoylated CHP1 proteins and CHP1/NHE1 (aa 503–545) complex proteins were all recovered in the soluble fraction and partially purified by passage through a Ni²⁺ affinity resin column (ProBond, Invitrogen) according to the manufacturer's protocol. Partially purified CHP1 proteins were found to be ~70% pure. We did not carry out further purification of CHP1 because of aggregation during storage. The complexes consisting of CHP1 or its mutant variants complexed with the NHE1 fragment were further purified to more than 95% by diethylaminoethyl-Sepharose column chromatography. All the proteins were dialyzed overnight against 60 mM KCl and 10 mM HEPES/Tris (pH 7.2).

Measurement of Equilibrium ⁴⁵Ca²⁺ Binding. ⁴⁵Ca²⁺ binding to the proteins was measured by a filtration method as described previously (31). Purified proteins (0.1–0.2 mg/mL) were incubated for 1 h at 25 °C in a solution containing 60 mM KCl, 5 mM MgCl₂, 50 μ M CaCl₂, 0.02 μ Ci/mL ⁴⁵CaCl₂, 10 mM HEPES/Tris (pH 7.2), and different concentrations of EGTA (0–58 mM), giving a free Ca²⁺ concentration of 0.1 nM to 50 μ M. Aliquots (1 mL) of the reaction mixture were transferred onto 0.22- μ m Millipore filters (Millipore, Bedford, MA) and filtered under vacuum. As controls, the same reaction mixtures without proteins were filtered to measure the background binding of ⁴⁵Ca by the filters. More than 95% of the proteins were retained in the filters. After the filters were dried, ⁴⁵Ca radioactivity was measured by scintillation counting.

Measurement of ⁴⁵Ca²⁺ Release from Proteins. ⁴⁵Ca²⁺ release from proteins was measured using a rapid filtration apparatus as described previously (31). After preincubation of proteins with a solution containing 50 μ M ⁴⁵CaCl₂ for 1 h, aliquots (1 mL) of reaction mixtures were filtered through Millipore filters. Filters were washed at a constant rate (0.2–2 mL/s) for the indicated periods (0.2–30 s) with

0.4–6 mL of 60 mM KCl, 5 mM MgCl₂, 10 mM HEPES/Tris (pH 7.2), and 10 mM EGTA. After the filters were dried, ⁴⁵Ca radioactivity was measured by scintillation counting.

Immunoprecipitation and Immunoblotting. Immunoprecipitation and immunoblotting were performed essentially as described previously (14). Briefly, cells were solubilized with 1% Triton X-100 in a solution of 150 mM NaCl, 10 mM HEPES-Tris (pH 7.4), and protease inhibitors. Cell lysates were incubated with respective antibodies and protein A Sepharose. After centrifugation, precipitated materials were separated on 7.5% or 12% polyacrylamide gels and electrophoretically transferred to Immobilon membranes (Millipore). After blocking, incubation with antibodies and washing, protein signals were visualized by enhanced chemiluminescence (Amersham, Buckinghamshire, U.K.). The signal intensity was measured using a photonic microscope system (ARUGUS-100, Hamamatsu photonics).

Measurement of ²²Na⁺ Uptake. ²²Na⁺ uptake activity and its pH_i dependence were measured by the K⁺/nigericin pH_i clamp method essentially as described previously (32). Serum-depleted cells in 24-well dishes were incubated for 30 min at 37 °C in Na⁺-free choline chloride/KCl medium containing 20 mM HEPES/Tris (pH 7.4), 1.2–140 mM KCl, 2 mM CaCl₂, 1 mM MgCl₂, 5 mM glucose (or 5 mM 2-deoxyglucose plus 2 μg/mL oligomycin under conditions of ATP depletion), and 5 μM nigericin. ²²Na⁺ uptake was started by adding the same choline chloride/KCl solution containing ²²NaCl (37 kBq/mL, final concentration = 1 mM), 1 mM ouabain, and 100 μM bumetanide. In some wells, 0.1 mM EIPA was added to the solution. After 1 min, cells were washed four times with ice-cold PBS to terminate ²²Na⁺ uptake. The pH_i was calculated from the imposed K⁺ concentration gradient by assuming the equilibrium [K⁺]_i/[K⁺]_o = [H⁺]_o/[H⁺]_i and an intracellular K⁺ concentration of 120 mM. Data were normalized according to the protein concentration as measured by the bicinchoninic assay (Pierce Chemical Co., IL) using bovine serum albumin as a standard.

Measurement of pH_i. Changes in pH_i were measured by the [¹⁴C]-benzoic acid equilibration method (30). For this measurement, serum-depleted cells were incubated for 30 min in bicarbonate-free HEPES-buffered DMEM (pH 7.0) and then incubated in the same medium containing [¹⁴C]-benzoic acid (37 kBq/mL) for 20 min at 37 °C. After the cells were washed four times with ice-cold PBS, ¹⁴C-radioactivity taken up by cells was measured. Changes in pH_i were calculated as described previously (30).

Statistics. Data of the pH dependence of EIPA-sensitive ²²Na⁺ uptake were simulated by fitting the values to the sigmoidal dose-response equation, rate of EIPA-sensitive ²²Na⁺ uptake = V_{max}/(1 + 10^{log(pK - pH_i)ⁿ) (pK = pH_i giving half-maximal ²²Na⁺ uptake; n = Hill coefficient), using the simulation program included in Graphpad Prism (Microsoft Corp., Redmond, WA). Equilibrium ⁴⁵Ca²⁺ binding was fitted to the dose-response equation, ⁴⁵Ca²⁺ bound = maximal ⁴⁵Ca²⁺ bound/(1 + (K_d - [Ca²⁺])ⁿ) (K_d = apparent dissociation constant for Ca²⁺; n = Hill coefficient). Kinetic parameters were expressed as the best fit values with standard errors, whereas other data were expressed as the means ± SD for at least three determinations.}

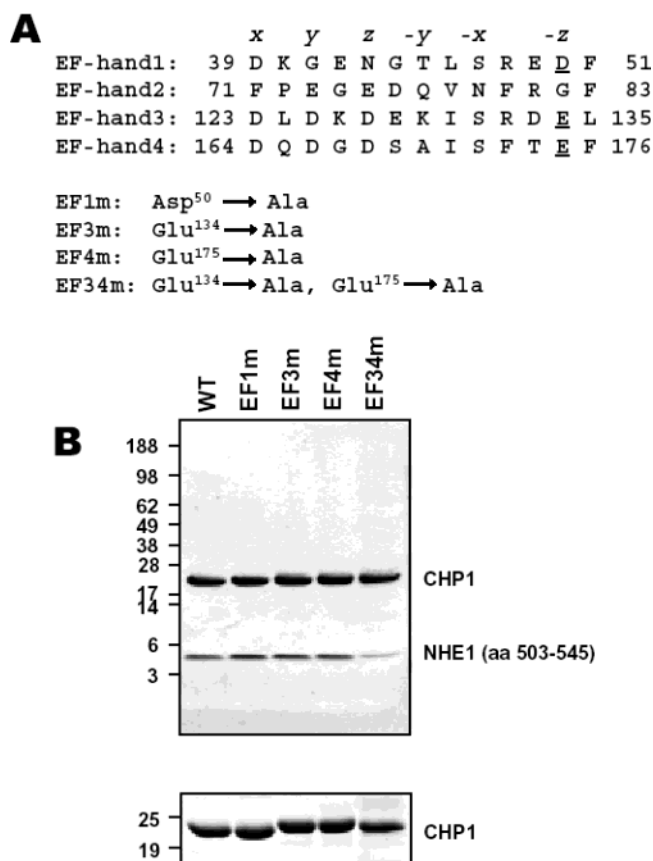


FIGURE 1: Amino acid sequences of EF-hand motifs and purified proteins of various CHP1 mutants. Panel A shows amino acid sequences of four EF-hand motifs present in CHP1. In four mutant CHP1s (EF1m, EF3m, EF4m, and EF34m), Asp⁵⁰, Glu¹³⁴, Glu¹⁷⁵, and both Glu¹³⁴/Glu¹⁷⁵ were replaced by alanine. In panel B, the purified complex of His-tagged CHP1 with the NHE1 segment (aa 503–545) (10 μg) was separated by electrophoresis on a 4–15% gradient (upper panel) or 12% SDS-PAGE gel (lower panel) and then visualized by Coomassie Brilliant Blue staining.

RESULTS

Characterization of Ca²⁺ Binding Motifs in CHP1. We first analyzed ⁴⁵Ca²⁺ binding to EF-hand motifs of CHP1 using recombinant CHP1 and its complex with the binding domain in NHE1. CHP1 interacts with NHE1 at the juxtamembrane region of the carboxyl-terminal cytoplasmic domain of NHE1. Hydrophobic residues of NHE1, such as Phe⁵²⁶, Leu⁵²⁷, Leu⁵³⁰, and Leu⁵³¹, were shown to be important for the interaction of CHP1 with NHE1 (12). CHP1 contains four potential EF-hand Ca²⁺ binding motifs, of which two ancestral sites (EF1 and -2) may not bind Ca²⁺ due to substitution of critical acidic residues (Figure 1A). The canonical EF-hand consists of 29 consecutive residues with two flanking helices and a 12-residue loop (Figure 1A). The chelating loop residues in positions 1 (+x), 3 (+y), 5 (+z), 7 (-y), 9 (-x), and 12 (-z) ligate Ca²⁺ through seven oxygen atoms arranged three-dimensionally on the axes of a pentagonal bipyramid (33, 34). The -z position, providing the only side chain oxygen atoms, is crucial for Ca²⁺ binding (33–35). To characterize these Ca²⁺ binding motifs, we introduced mutations into EF1, -3, and -4 in which acidic residues (aspartic acid or glutamic acid) at the -z position were replaced by alanine (Figure 1B). We coexpressed the wild-type or mutant CHP1s together with the juxtamembrane region of NHE1 (aa 503–545) in *E. coli*. We confirmed that

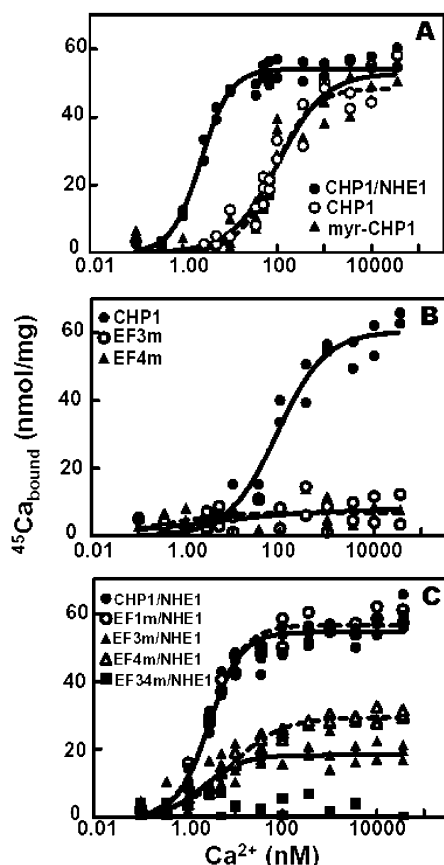


FIGURE 2: Equilibrium $^{45}\text{Ca}^{2+}$ binding to various CHP1 mutant proteins. In panels A–C, CHP1 or its various mutant proteins and the complex of CHP1 variants with the NHE1 segment (aa 503–545) (0.1–0.2 mg/mL) were incubated for 1 h in solutions containing $50\ \mu\text{M}$ $^{45}\text{CaCl}_2$ and various concentrations of EGTA, which produce 0.2 nM to $50\ \mu\text{M}$ free Ca^{2+} . Symbols corresponding to each protein variant were indicated in figures. The solutions were filtered through Millipore filters, and $^{45}\text{Ca}^{2+}$ bound to the CHP1 proteins was measured.

the purified complex proteins (>95% pure) of the wild-type CHP1, EF1m, EF3m, or EF4m with aa 503–545 of NHE1 were retained as a single peak on gel filtration chromatography and contained the CHP1 variant and the NHE1 peptide at a 1:1 molar ratio (data not shown). In addition, using 4–15% polyacrylamide gradient gels, we confirmed that the purified samples mostly contained comparable molar amounts of the CHP1 variant and the NHE1 fragment (Figure 1B, upper panel). However, in EF34m with double mutations at EF3 and -4, the amount of the NHE1 fragment was significantly reduced, suggesting that this double mutation impairs the interaction of CHP1 with NHE1. On 12% SDS–PAGE, EF3m, EF4m, and EF34m proteins were found to migrate more slowly than the wild-type or EF1m proteins (Figure 1B, lower panel), suggesting that a mutation-induced conformational change occurred in these three mutant proteins that had impaired Ca^{2+} binding (see below).

We measured $^{45}\text{Ca}^{2+}$ binding to various CHP1 mutant proteins by a membrane filtration procedure. We found that $^{45}\text{Ca}^{2+}$ bound to the partially purified CHP1 proteins with an apparent K_d of $\sim 90\ \text{nM}$ (Figure 2A and Table 1). The maximal amount of $^{45}\text{Ca}^{2+}$ bound to CHP1 corresponded to $\sim 2\ \text{mol}$ of Ca^{2+} bound/mol of CHP1, assuming that the CHP1 sample was 70% pure. Myristoylation did not significantly affect the apparent affinity for Ca^{2+} nor the

Table 1: Parameters for Equilibrium $^{45}\text{Ca}^{2+}$ Binding

proteins	apparent K_d $\pm\ \text{SE}$ (nM) ^a	Hill coefficient $\pm\ \text{SE}$
CHP1	89.9 ± 9.3	0.77 ± 0.12
myr-CHP1	86.4 ± 8.9	0.94 ± 0.17
CHP1/NHE1	2.32 ± 0.18	1.22 ± 0.15
EF1m/NHE1	2.17 ± 0.39	0.98 ± 0.14
EF3m/NHE1	2.89 ± 0.28	1.27 ± 0.55
EF4m/NHE1	2.24 ± 0.26	0.76 ± 0.09

^a The data shown in Figure 2 were fitted to the equation for equilibrium $^{45}\text{Ca}^{2+}$ binding as described in Experimental Procedures.

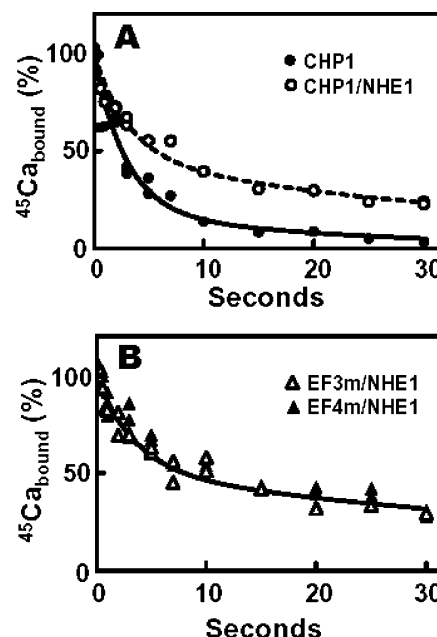


FIGURE 3: Time courses of $^{45}\text{Ca}^{2+}$ release from CHP1 proteins: (A) $^{45}\text{Ca}^{2+}$ release from CHP1 alone (●) or CHP1/NHE1 complex (○); (B) $^{45}\text{Ca}^{2+}$ release from EF3m/NHE1 (△) or EF4m/NHE1 complex (▲). CHP1 proteins or the complex of CHP1 variants with the NHE1 segment (aa 503–545) (0.1–0.2 mg/mL) were incubated for 1 h in a solution containing $50\ \mu\text{M}$ $^{45}\text{CaCl}_2$, applied to Millipore filters, and washed with a solution containing EGTA. $^{45}\text{Ca}^{2+}$ remaining on the filters was measured.

maximal level of $^{45}\text{Ca}^{2+}$ binding (Figure 2A and Table 1). Interestingly, when CHP1 formed a complex with the NHE1 fragment, the binding affinity for $^{45}\text{Ca}^{2+}$ increased markedly (~ 40 -fold, Figure 2A and Table 1). The extremely low apparent dissociation constant ($\sim 2\ \text{nM}$) deviates substantially from the physiological cytosolic Ca^{2+} concentration of cells (0.1 – $10\ \mu\text{M}$). The maximal level of Ca^{2+} binding on the complex again corresponded to $\sim 2\ \text{mol}$ of Ca^{2+} bound/mol of CHP1. Mutation of either of Ca^{2+} binding motifs EF3 or EF4, but not EF1, resulted in loss of approximately 1 mol of $^{45}\text{Ca}^{2+}$ bound to the complex (Figure 2C). On the other hand, $^{45}\text{Ca}^{2+}$ binding was completely blocked when the experiment was carried out using EF3m and EF4m proteins but without the NHE1 fragment (Figure 2B) or when two sites were simultaneously mutated (EF34m) (Figure 2C). Together, these results indicate that CHP1 binds two Ca^{2+} ions, one at EF3 and the other at EF4.

To determine how complex formation increases the Ca^{2+} binding affinity, we measured $^{45}\text{Ca}^{2+}$ release from CHP1 proteins by rapid filtration. As shown in Figure 3A, most of the $^{45}\text{Ca}^{2+}$ bound to CHP1 without the NHE1 fragment was released rapidly ($t_{1/2} = \sim 2\ \text{s}$). In contrast, $^{45}\text{Ca}^{2+}$ release from

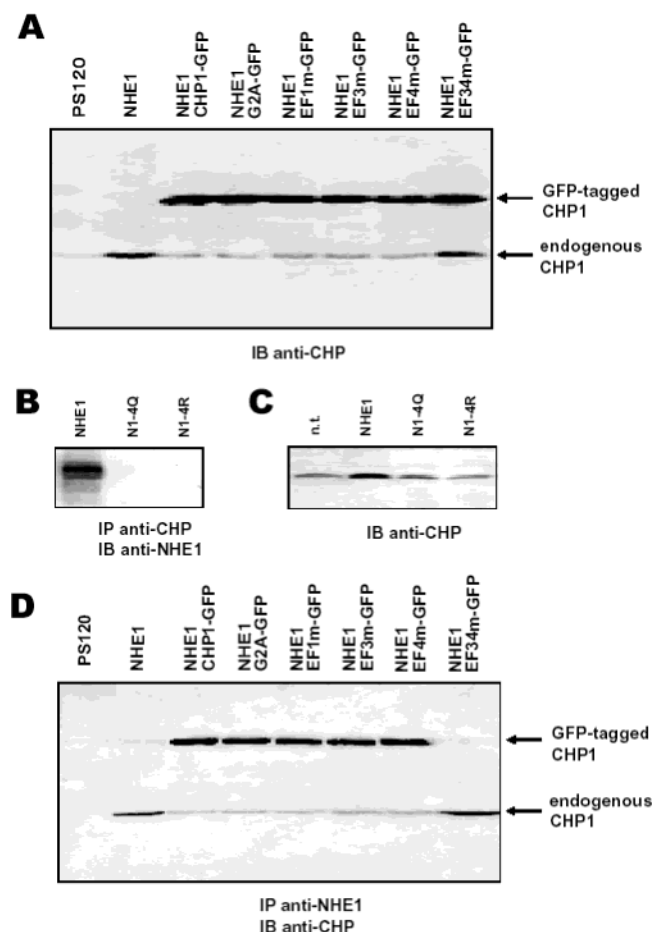


FIGURE 4: Expression of various GFP-tagged CHP1 proteins and their coimmunoprecipitation with NHE1. Panel A shows the expression level of GFP-tagged CHP1 and its variants (indicated at the top of the figure). Cell lysates (50 μ g) from stable transfectants were subjected to SDS-PAGE, and expression of endogenous and exogenous CHP1 proteins were detected by immunoblotting (IB) with an anti-CHP1 antibody. A result for untransfected PS120 cells is shown in the first lane. Panel B shows coimmunoprecipitation of the wild-type or mutant (4Q and 4R) exchangers with endogenous CHP1. Lysates from cells stably expressing these exchangers were subjected to immunoprecipitation with anti-CHP1 antibody followed by immunoblotting with anti-NHE1 antibody. Panel C shows the expression level of endogenous CHP1 in cells expressing the wild-type or mutant NHE1s: n.t., no transfection. Panel D shows coimmunoprecipitation of CHP1 proteins with NHE1. Lysates from cells stably expressing various proteins were subjected to immunoprecipitation with anti-NHE1 antibody followed by immunoblotting with anti-CHP1 antibody. Note that in lanes from cells not transfected with GFP-tagged CHP1 (left two lanes), IgG protein bands were visible at the same positions as GFP-tagged CHP1.

CHP1 complexed with the NHE1 fragment ($t_{1/2} = \sim 7$ s) was much slower. A slow release of $^{45}\text{Ca}^{2+}$ also occurred in two mutant CHP1 proteins, EF3m and EF4m, complexed with the NHE1 fragment (Figure 3B), suggesting that Ca^{2+} binds tightly to each EF hand.

Effects of CHP1 Mutations on NHE1 Regulation. To study the role of Ca^{2+} binding in NHE1 regulation by CHP1, we transfected GFP-tagged CHP1 into cells expressing NHE1 and obtained cells stably coexpressing these proteins. The results indicated that GFP-tagged CHP1 and its mutant derivatives were highly coexpressed in NHE1 transfectants (Figure 4A). Interestingly, expression of NHE1 markedly increased the level of expression of the endogenous CHP1

Table 2: Relative Amounts of Expressed GFP-Tagged CHP1 and Endogenous CHP1

transfected proteins	relative amount of GFP-tagged CHP1 ^a	relative amount of endogenous CHP1 ^b
untransfected		1.00 \pm 0.08
NHE1		3.63 \pm 0.81 ^c
NHE1 + CHP1-GFP	1.00 \pm 0.11	1.11 \pm 0.13
NHE1 + G2A-GFP	1.08 \pm 0.11	1.03 \pm 0.16
NHE1 + EF1m-GFP	0.93 \pm 0.06	1.07 \pm 0.11
NHE1 + EF3m-GFP	0.97 \pm 0.11	1.15 \pm 0.16
NHE1 + EF4m-GFP	1.06 \pm 0.07	1.09 \pm 0.10
NHE1 + EF34m-GFP	0.94 \pm 0.12	3.85 \pm 0.45 ^c
NHE1-4Q		1.02 \pm 0.09
NHE1-4R		1.03 \pm 0.06

^a The density of visualized protein bands on immunoblots (cf. Figure 4, panels A and C) is represented as values normalized according to the band density from cells expressing CHP1-GFP. Data are means \pm SD ($n = 3$). ^b The band density is represented as values normalized according to that from untransfected PS120 cells. Data are means \pm SD ($n = 3$). ^c $P < 0.05$ versus control.

(3.6-fold), while coexpression of various GFP-tagged CHP1 variants, with the exception of CHP1-EF34m-GFP, reduced it (Table 2).

We further examined the effect of expression of CHP1 binding-defective NHE1 mutants 4Q and 4R on the amount of endogenous CHP. These mutant exchangers do not bind CHP1 as shown by coimmunoprecipitation studies (Figure 4B). The level of expression of the endogenous CHP1 did not increase on coexpression of these mutant exchangers (Figure 4B,C, Table 2). Thus, the amount of endogenous CHP1 in cells is highly dependent on expression of NHE1 and GFP-tagged CHP1.

Figure 4D shows the results for coimmunoprecipitation experiments using NHE1- and CHP1-specific antibodies to determine interactions of the expressed CHP1-GFP with NHE1. Anti-NHE1 antibody immunoprecipitated endogenous CHP1 from cells expressing NHE1. In cells coexpressing GFP-CHP1 and NHE1, the same antibody coimmunoprecipitated large quantities of GFP-CHP1 or its derivatives, and at the same time, the amount of immunoprecipitated endogenous CHP1 was markedly reduced. In cells coexpressing EF34m-GFP and NHE1, anti-NHE1 antibody coimmunoprecipitated the endogenous CHP1 but not exogenous GFP-tagged mutant CHP1, consistent with the findings of *in vitro* binding studies indicating that double mutation at EF3 and EF4 impairs the interaction of CHP1 with NHE1.

We next examined the subcellular localization of GFP-tagged CHP1. As reported previously (12), the GFP-tagged CHP1 is localized in the plasma membrane in cells coexpressing NHE1 (Figure 5A). Consistent with the *in vitro* binding data (Figure 1B), the GFP fluorescence was observed in the plasma membrane in cells coexpressing GFP-tagged CHP1 mutants except EF34m with NHE1 (Figure 5A; data not shown for G2A and EF1m). These results, together with the data from coimmunoprecipitation experiments, indicate that the endogenous CHP1 bound to NHE1 was efficiently replaced by expressed GFP-tagged wild-type or CHP1 mutants. However, the double mutant EF34m was not localized at the plasma membrane (Figure 5A) because of the weak interaction of this mutant protein with the juxtamembrane region of NHE1. We observed that GFP fluorescence was still observed in the plasma membrane after addition of phorbol ester, serum, thrombin, lysophosphatidic

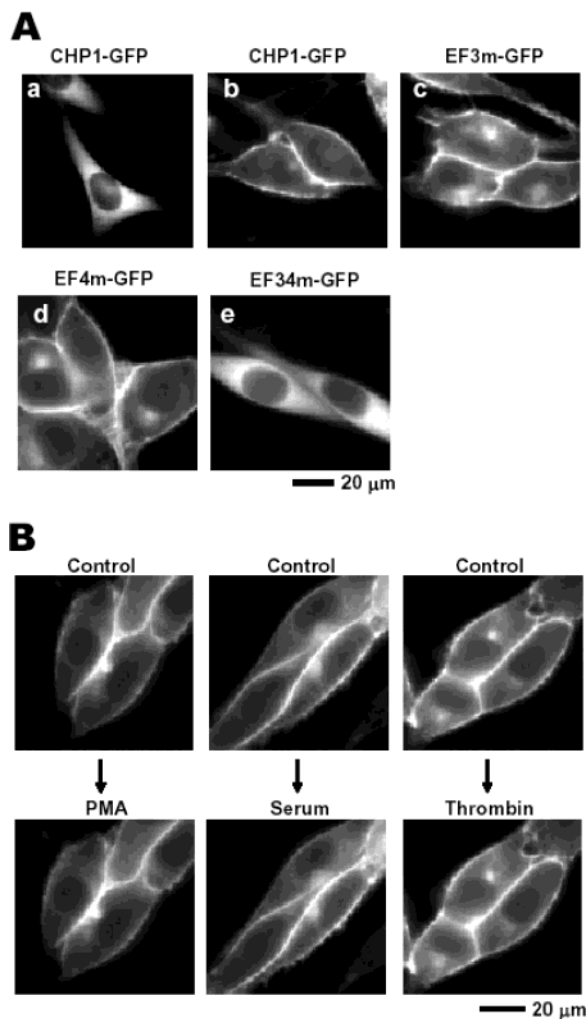


FIGURE 5: Subcellular localization of GFP-tagged CHP1: (A) subcellular localization of GFP-tagged wild-type CHP1 (a, b), EF3m (c), EF4m (d), and EF34m (e) expressed in PS120 cells (a) or in their stable transfectants of the wild-type NHE1 (b–e); (B) effect of various agents on the subcellular localization of the wild-type GFP-tagged CHP1 in NHE1 transfectants. Cells were placed in serum-free Dulbecco's modified Eagle's medium without phenol red for 5 h, and then 1 μ M PMA, 10% serum, or 2 units/ml thrombin were added. GFP fluorescence was observed under a fluorescent microscope equipped with a CoolSNAP imaging system (RS Photometrics) before (control) and 20 min after addition of the various agents.

acid, or PDGF-BB, which are all known to activate the exchange activity (Figure 5B, data not shown for some experiments). We also found that the plasma membrane localization of GFP fluorescence did not change upon addition of metabolic inhibitors (2-deoxyglucose plus oligomycin) that cause cell ATP depletion, thus inhibiting exchange activity (data not shown). Furthermore, we found no changes in the plasma membrane localization of GFP-tagged CHP1 mutants EF3m and EF4m after these various treatments (data not shown). These observations suggest that CHP1 is tightly associated with NHE1 in the plasma membrane and that this interaction is not affected by various stimuli.

All the cells expressing CHP1–GFP or its mutant derivatives exhibited high Na^+/H^+ exchange activity. The $^{22}\text{Na}^+$ uptake activity in cells clamped at acidic pH_i (5.6) by the $\text{K}^+/\text{nigericin}$ technique was in the range of 20–50 nmol/

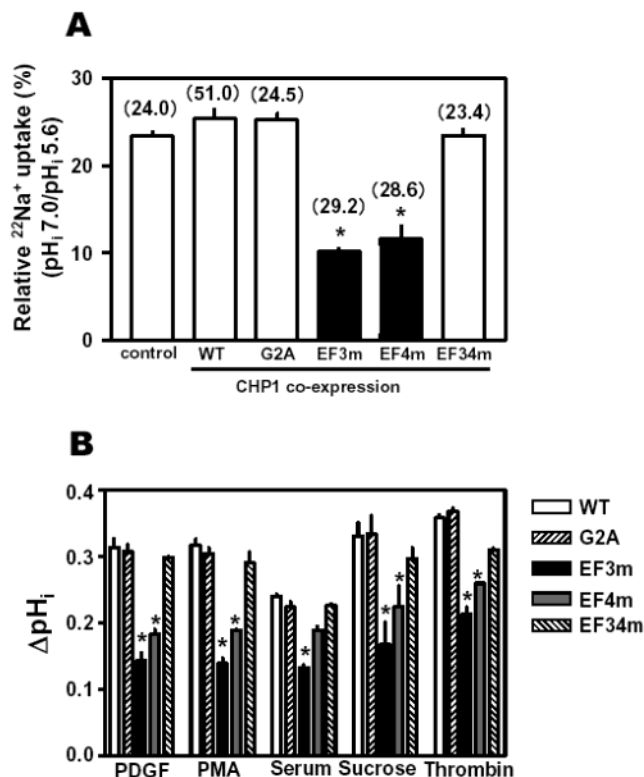


FIGURE 6: Exchange activity and regulation of NHE1 transfectants expressing various CHP1 mutants. Panel A shows ratios of EIPA-sensitive $^{22}\text{Na}^+$ uptake activities of cells coexpressing wild-type NHE1 and various CHP1 mutants at pH_i 7.0 and 5.6. Numbers (nmol/mg/min) in parentheses represent $^{22}\text{Na}^+$ uptake activity at pH_i 5.6. Control cells were not transfected with CHP1 but stably expressing NHE1. Data are means \pm SD ($n = 3$; *, $P < 0.05$ versus cells not expressing exogenous CHP1). Panel B shows changes in pH_i measured using the [^{14}C]benzoic acid equilibration method. The cells coexpressing NHE1 and various CHP1 variants were stimulated for 15 min at 37 $^\circ\text{C}$ with 10 ng/mL PDGF-BB, 1 μ M PMA, 10 $\mu\text{g}/\text{mL}$ lysophosphatidic acid, or 200 mM sucrose (hyperosmotic stress). Data are means \pm SD ($n = 6$; *, $P < 0.05$ versus cells expressing wild-type CHP1).

mg/min (data not shown). We compared the $^{22}\text{Na}^+$ uptake activities in cells expressing various CHP1 variants in the physiological pH_i range. As shown in Figure 6A, the ratio of $^{22}\text{Na}^+$ uptake at pH_i 7.2–5.6 was not significantly altered by expression of wild-type CHP1. Although a previous study (11) indicated that overexpression of CHP1 inhibits the NHE1 activity in the presence of serum, we observed no such CHP1-induced inhibition of the exchange activity. The reason for this discrepancy is unknown. Unlike the wild-type CHP1, the $^{22}\text{Na}^+$ uptake ratio was significantly reduced by EF3 or EF4 mutants. Consistent with this finding, we observed that mutations of EF3 or EF4 significantly reduced the cytoplasmic alkalinization in response to PDGF-BB, thrombin, phorbol ester, serum, or hyperosmotic stress (sucrose) (Figure 6B). These observations suggest that mutation of EF3 or EF4 partly impairs the regulation of NHE1 by reducing pH_i sensitivity. In contrast, double mutation (EF34m) of CHP1 at EF3 and EF4 did not reduce the $^{22}\text{Na}^+$ uptake ratio or cytoplasmic alkalinization (Figure 6A,B), consistent with the finding that this mutant CHP1 is not able to replace the endogenous CHP1 because of its weak interaction with NHE1. Finally, mutation of the myristoylation site (G2A) or EF1 did not affect pH_i -dependent regulation of NHE1.

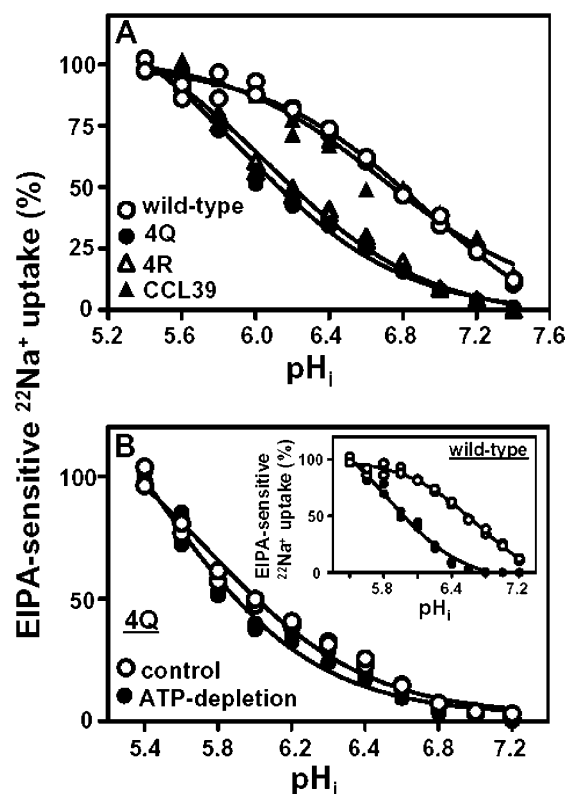


FIGURE 7: pH_i dependence of exchange activity in cells expressing some NHE variants. Panel A shows the pH_i dependence of ²²Na⁺ uptake in PS120 cells expressing wild-type NHE1 or CHP1 binding-defective mutants 4Q and 4R and CCL39 fibroblastic cells (the parental cell line of PS120). pH_i was clamped at various values with K⁺/nigericin. The maximal EIPA-sensitive ²²Na⁺ uptake activity measured at pH_i = 5.4 was high in cells expressing wild-type NHE1 (~50 nmol/mg/min), while it was lower but similar in cells expressing 4Q or 4R or in CCL39 cells (4.2, 4.2, or 4.1 nmol/mg/min, respectively). Data were normalized according to the maximal activity at pH_i = 5.4. Panel B shows the effects of ATP depletion on pH_i dependence of exchange activity in cells expressing 4Q or wild-type NHE1 (inset). Cells were depleted of ATP by treatment with the metabolic inhibitors 2-deoxyglucose (5 mM) and oligomycin (2 μg/mL). Data were normalized according to the maximal activity at pH_i = 5.4.

Properties of NHE1 Mutants Lacking CHP1 Binding. As described above, mutations of CHP1 partly impair pH_i-dependent regulation of NHE1. Therefore, it is of interest to determine how CHP1 binding affects the pH_i sensitivity of NHE1. Previously, we described two CHP1 binding-defective mutant exchangers, 4Q or 4R, in which Phe⁵²⁶, Leu⁵²⁷, Leu⁵³⁰, and Leu⁵³¹ of NHE1 were replaced by four glutamine or arginine residues, respectively (12) (see Figure 4B). In this study, by using extensive H⁺-killing selection, we obtained cells overexpressing 4Q or 4R and exhibiting relatively high activity (~4 nmol/mg/min at pH_i 5.4), thus allowing reliable measurement of the pH_i dependence of ²²Na⁺ uptake. As shown in Figure 7A, these mutations caused a marked acidic shift in the pH_i dependence (Figure 7A). As a control, we confirmed that CCL39 cells (the parental cell line of PS120) that exhibit exchange activity (*V*_{max}) comparable to 4Q or 4R show a pH_i dependence of exchange similar to that of PS120 cells overexpressing NHE1. In cells expressing these mutant exchangers, ATP depletion did not change the pH_i sensitivity of ²²Na⁺ uptake (Figure 7B). In addition, cytoplasmic alkalization in response to extracel-

lular stimuli, such as thrombin, PDGF-BB, hyperosmolarity, LPA, and PMA, was not observed in cells expressing 4Q or 4R (data not shown), consistent with the finding that these mutants exhibit an acidic shift of pH_i dependence.

DISCUSSION

In this study, we examined the role of CHP1, particularly its EF-hand Ca²⁺ binding motifs, in the pH_i-dependent regulation of NHE1. Our results indicated that a Ca²⁺ ion binds to each of EF3 and EF4 in CHP1 with an overall apparent *K*_d of ~90 nM and a Hill coefficient of ~1.0 (Table 1). This Ca²⁺ binding affinity was close to that of another family member, CaN-B (apparent *K*_d ≈ 70 nM) (36), although the apparent *K*_d values for Ca²⁺ in other EF-hand Ca²⁺ binding proteins vary widely (0.01–10 μM) (36–39). Although CHP1 potentially has four Ca²⁺ binding motifs, the two ancestral sites EF1 and EF2 do not bind Ca²⁺. This is in sharp contrast to CaN-B in which all four EF-hand motifs are able to bind Ca²⁺, although the two N-terminal sites, EF1 and EF2, have lower affinity for Ca²⁺ than the C-terminal sites, EF3 and EF4 (36). Intriguingly, the Ca²⁺ affinity of CHP1 increased markedly upon complex formation with the NHE1 fragment (aa 503–545). Consistent with this finding, ⁴⁵Ca²⁺ release from the complex was much slower than that from CHP1 alone. The extraordinarily high affinity of CHP1 for Ca²⁺ (~2 nM) suggests that the CHP1/NHE1 complex always contains two Ca²⁺ ions under physiological conditions.

The high affinity for Ca²⁺ was also observed in mutant CHP1 proteins EF3m and EF4m, which have a single Ca²⁺ binding site, complexed with the NHE1 fragment. Increases in the affinity for Ca²⁺ by interaction with target proteins have also been reported for other Ca²⁺ binding proteins. For example, the Ca²⁺ binding affinity for calmodulin was increased 16- to 38-fold upon interaction with myosin light chain kinase (40), 2.6-fold with myristoylated alanine-rich protein kinase C substrate peptide (41), and 75-fold with the calmodulin binding peptide in CaN-A (36). CHP1 was reported to interact with other proteins, such as microtubules (25), CaN-A (24), DRAK2 (26), and KIF1B/β2 (27), as well as members of the NHE1 family. Therefore, interaction with these proteins may also modify the Ca²⁺ binding affinity of CHP1.

Although mutation of CHP1 at either EF3 or EF4 impaired binding of 1 mol of Ca²⁺, it did not appear to affect the interaction of CHP1 with NHE1 as shown by *in vitro* binding of these mutant proteins, coimmunoprecipitation, and the plasma membrane localization of GFP-tagged CHP1. Therefore, these mutations do not appear to induce marked structural distortions. However, double mutation (EF34m) at both EF3 and EF4 impaired the interaction of CHP1 with NHE1. Consistent with this finding, Ca²⁺ removal by EGTA from the wild-type CHP1 reduced the interaction with NHE1 in a pull-down assay through the amylose resin column (data not shown). Thus, the tight association of NHE1 with CHP1 requires binding of at least one Ca²⁺ on either EF3 or EF4. Furthermore, it should be noted that the effect of double mutation (EF34m) on regulation of NHE1 cannot be properly analyzed in cells that express endogenous CHP1.

We found that expression of NHE1, but not the CHP1 binding-deficient mutant derivatives 4Q and 4R, significantly

increased the amount of endogenous CHP1. However, coexpression of GFP-tagged CHP1 proteins (wild-type, EF3m, and EF4m) preserving the strong interaction with NHE1 greatly reduced the amount of endogenous CHP1. Thus, the amount of endogenous CHP1 in cells is strongly dependent on the number of available CHP1 binding sites provided from NHE1. Although the precise reason for this is unknown, it is likely that interaction with target proteins is required for stable expression of CHP1. That is, dissociation from the target proteins may promote CHP1 degradation.

In this study, we found that CHP1 binding-defective mutants of NHE1 (4Q and 4R) caused a marked acidic shift in the pH_i dependence of Na^+/H^+ exchange activity and completely impaired ATP depletion-induced inhibition and cytoplasmic alkalization in response to various stimuli. As we reported previously (32), not only mutation of the CHP1 binding region, but also deletions of different regions in the amino-terminus (subdomain I, amino acids 515–595) of the NHE1 cytoplasmic domain also markedly reduced pH_i sensitivity. Thus, subdomain I with bound CHP1 appears to be a key structure that permits the putative “pH-sensor” to maintain a physiologically relevant conformation.

We found that mutation of EF3 or EF4 in CHP1 significantly reduced the Na^+/H^+ exchange activity in the physiological neutral pH_i range and reduced the cytoplasmic alkalization in response to various extracellular signals by decreasing the pH_i sensitivity of NHE1. Thus, mutation of each EF-hand in CHP1 somehow affects the pH_i -sensing of NHE1, although we could not evaluate the function of the double mutant CHP1 (EF34m) because of its weak interaction with NHE1. We found that EF3m, EF4m, and EF34m proteins migrated slowly on SDS–PAGE, suggesting that significant conformational changes of CHP1 occur upon mutation of each EF-hand. Such conformational changes appear to be due to removal of Ca^{2+} rather than the amino acid substitution itself, because incubation with EGTA resulted in similar slow migration of the wild-type CHP1 on SDS–PAGE (data not shown). Thus, the bound Ca^{2+} may play an important role in maintaining the CHP1 structure, thereby preserving the physiological pH_i sensitivity of NHE1.

According to the structural model of CHP1 deduced from the three-dimensional structure of CaN-B (43), EF-hand Ca^{2+} binding motifs would be located on the surface opposite the side where CHP1 binds to NHE1. It is likely that the surface of CHP1 with tightly bound Ca^{2+} controls the pH_i -sensing by interacting with other region(s) of NHE1. We observed that the pH_i sensitivity of NHE1 was markedly reduced by insertion of one amino acid residue (alanine) just to the N-terminal side (position aa 504 or 508) of the CHP1 binding site of NHE1, while the CHP1 binding ability was preserved (our unpublished observations). Therefore, the correct spatial orientation of CHP1 would be important for regulation of NHE. Recently, we reported that mutation of Arg⁴⁴⁰ in intracellular loop 5 (IL5), which connects transmembrane helices 10 and 11, markedly reduces the pH_i sensitivity of NHE1 (44). Thus, IL5 may interact with the CHP1 surface with tightly bound Ca^{2+} .

Many EF-hand Ca^{2+} binding proteins are known to regulate the functions of their target proteins in response to cytosolic Ca^{2+} mobilization. However, it is unlikely that CHP1 functions as such a Ca^{2+} sensor in the regulation of NHE1 because the affinity for Ca^{2+} ($K_d = \sim 2$ nM) for the

CHP1/NHE1 complex differs substantially from the range of intracellular Ca^{2+} concentrations (0.1–10 μ M). Instead, two EF-hand motifs of CHP1 together with tightly bound Ca^{2+} would serve as structurally important elements for preserving the normal function of NHE1, as discussed above. Such a structural role has also been suggested in the C-terminal EF-hand motifs in CaN-B (42) and in CaM (41). On the other hand, we reported previously that CaM interacts in a Ca^{2+} -dependent manner with the middle of the cytoplasmic domain of NHE1, which in the unstipulated state serves as an auto inhibitory domain decreasing the pH_i sensitivity of NHE1 (14, 15). The interaction of NHE1 with CaM is strictly Ca^{2+} -dependent, although it is much weaker than that with CHP1 (14). Previously, we proposed that NHE1 may be activated by Ca^{2+} -dependent interaction of CaM in response to Ca^{2+} -mobilizing signals (15). Our previous (14, 15) and several recent reports (45–48) reinforced the idea that CaM serves as an important regulatory protein in activation of NHE1 in response to hyperosmotic stress or Ca^{2+} -mobilizing agonists. NHE1 thus appears to be dually regulated by two Ca^{2+} binding proteins, CHP1 and CaM, similar to CaN-A. The former would preserve the physiological pH_i sensitivity of NHE1, whereas the latter would play a role in sensing cytosolic Ca^{2+} .

In summary, our current results suggest that the interaction of CHP1 with NHE1 is crucial for preserving the physiological pH_i sensitivity of NHE1 and that tightly bound Ca^{2+} serves as an important structural element that is required for this role. The significant effects of mutations in EF-hands on NHE1 regulation prompted us to generate a more efficient dominant negative mutant CHP1. In addition, the functional difference between CHP1 and CHP2, which we reported recently (13), provides important information for identification of the critical residues of CHP1. Further studies including analyses of the functions of mutated or chimerical CHP1 and determination of the crystal structure of CHP1/NHE1 complex are required to elucidate the molecular mechanism of CHP regulation of NHE1 and other NHE family members.

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