The Plasma Membrane Ca^{2+} ATPase and the Plasma Membrane Sodium Calcium Exchanger Cooperate in the Regulation of Cell Calcium

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Calcium is an ambivalent signal: it is essential for the correct functioning of cell life, but may also become dangerous to it. The plasma membrane Ca^{2+} ATPase (PMCA) and the plasma membrane $\text{Na}^+/\text{Ca}^{2+}$ exchanger (NCX) are the two mechanisms responsible for Ca^{2+} extrusion. The NCX has low Ca^{2+} affinity but high capacity for Ca^{2+} transport, whereas the PMCA has a high Ca^{2+} affinity but low transport capacity for it. Thus, traditionally, the PMCA pump has been attributed a housekeeping role in maintaining cytosolic Ca^{2+} , and the NCX the dynamic role of counteracting large cytosolic Ca^{2+} variations (especially in excitable cells). This view of the roles of the two Ca^{2+} extrusion systems has been recently revised, as the specific functional properties of the numerous PMCA isoforms and splicing variants suggests that they may have evolved to cover both the basal Ca^{2+} regulation (in the 100 nM range) and the Ca^{2+} transients generated by cell stimulation (in the μM range).

Ca^{2+} controls critical cellular responses in all eukaryotic organisms. It controls both short-term biological processes that occur in milliseconds, such as muscle contraction, as well as long-term processes that require longer times, such as cell proliferation and organ development. The specificity of cellular Ca^{2+} signals is controlled by a sophisticated “toolkit” comprising numerous ion channels, pumps, and exchangers that drive the fluxes of Ca^{2+} ions across the plasma membrane and across the membranes of intracellular organelles (Berridge et al. 2003).

The plasma membrane contains several types of channels that mediate Ca^{2+} entry from the

extracellular ambient, and two systems for Ca^{2+} extrusion: a low affinity, high capacity $\text{Na}^+/\text{Ca}^{2+}$ exchanger (NCX), and a high-affinity, low-capacity Ca^{2+} -ATPase (the plasma membrane Ca^{2+} pump (PMCA)) (Fig. 1). The type of channels and the relative proportions of NCX and PMCA vary with the cell type, the NCX being particularly abundant in excitable tissues, e.g., heart and brain. The regulated opening of the Ca^{2+} channels by either voltage gating, interaction with ligands or the emptying of intracellular stores, allows a limited amount of Ca^{2+} to enter the cell to transmit signals to its designated targets. Thereafter, the Ca^{2+} transients must be dissipated: its extrusion from the

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Figure 1. A schematic representation of the structures involved in cellular Ca^{2+} homeostasis. The model shows a cell with its Ca^{2+} -transporting systems: Ca^{2+} -ATPases (plasma membrane and sarco/endoplasmic reticulum, PMCA and SERCA), plasma membrane (PM) Ca^{2+} channels, $\text{Na}^+/\text{Ca}^{2+}$ exchangers (NCX and NCLX), 1,4,5-triphosphate receptor (IP_3R) and ryanodine receptor (RyR), the electrophoretic mitochondrial uptake uniporter (U). Mitochondria are drawn as yellow ellipses, nucleus as orange circle and endoplasmic reticulum is colored in red. The different Ca^{2+} -transporting systems cooperate to maintain the Ca^{2+} concentration gradient between the extracellular and the intracellular ambient.

cell is mediated by the NCX and the PMCA pump, but Ca^{2+} is also restored to basal levels by sequestration in the endo/sarcoplasmic reticulum via the SERCA pump and in the mitochondria by the electrophoretic uniporter. The NCX has also been found at the inner membrane of the nuclear envelope (NE) and has been proposed to mediate Ca^{2+} flux between the nucleoplasm and the NE (Xie et al. 2002), and then to the ER (Wu et al. 2009) in neuronal and certain other cell types. Ca^{2+} binding proteins also contributed to Ca^{2+} buffering: In this review, we will not cover them, as we will only discuss the systems that extrude Ca^{2+} out of the cell.

The PMCA pump is a minor component of the total protein of the plasma membrane (less than 0.1% of it). Quantitatively, it is overshadowed by the more powerful NCX in excitable tissue like heart; however, even cells in which the NCX predominates, the PMCA pump is likely to be the fine tuner of cytosolic Ca^{2+} , as it can operate in a concentration range in which the low affinity NCX is relatively very inefficient.

The PMCA was discovered in erythrocytes (Schatzmann 1966), and was then described and characterized in numerous other cell types. It was purified in 1979 using a calmodulin affinity column (Niggli et al. 1979), and cloned about 10 years later (Shull and Greb 1988; Verma et al. 1988). It shows the same essential membrane topology properties of the SERCA pump. Molecular modeling work using the structure of the SERCA pump as a template (Toyoshima et al. 2000) predicts the same general features of the latter, with 10 transmembrane domains and the large cytosolic head-piece divided into the three main cytosolic A, N, and P domains. The $\text{Na}^+/\text{Ca}^{2+}$ cotransport process was discovered at about the same time as PMCA by two independent groups working on heart (Reuter and Seitz 1968) and on the squid giant axon (Baker et al. 1969). The exchanger was cloned in 1990 (Nicoll et al. 1990). The sequence was initially predicted to correspond to a protein with 11 transmembrane domains and one large cytosolic loop linking transmembrane domain five and six but a revised model

predicting only nine transmembrane domains is now generally accepted.

PLASMA MEMBRANE CALCIUM PUMP

Structural and Regulatory Characteristics

The PMCA pump belongs to the family of P-type ATPases, which are characterized by the temporary conservation of ATP energy in the form of a phosphorylated enzyme intermediate (hence P-type) formed between the γ – phosphate of hydrolyzed ATP and an invariant D-residue in a highly conserved sequence of the pump molecules: SDKTGT (L/I/V/M) (T/I/S). The pump operates with a 1:1 Ca^{2+} /ATP stoichiometry as a Ca^{2+} : H^{+} exchanger: the matter of Ca^{2+} / H^{+} stoichiometry is still controversial (Niggli et al. 1982; Hao et al. 1994). The pump has high Ca^{2+} affinity and a multitude of agents that modulate it. The K_d of the pump for Ca^{2+} , which is 10 to 20 μM in the resting state, decreases to less than 1 μM following calmodulin interaction (Brini and Carafoli 2009). Acidic phospholipids also increase the Ca^{2+} affinity of the pump. Even if the molecular mechanism of the activation by phospholipids is obscure, it could be physiologically meaningful: It has been calculated that in the membrane environment the pump is probably permanently activated by acidic phospholipids to about 50% of its maximal activity (Niggli et al. 1981).

Structurally, the pump is predicted to consist of 10 transmembrane domains, two large intracellular loops, and of amino- and carboxy-terminal cytoplasmic tails (Fig. 2). The 90-residue amino-terminal tail contains a consensus-binding site for the 14-3-3 protein, which plays an inhibitory role on pump activity (Rimessi et al. 2005). The first cytosolic loop between transmembrane domains two and three is considered the “transducer domain.” It contains one of the two sites that mediate the activation by acididic phospholipids (Zvaritch et al. 1990) and one of the two sites for the autoinhibitory interaction with the calmodulin binding domain. This loop also contains site A, which is one of the two main sites of the alternative

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Figure 2. (A) Topology model of PMCA. The pump is organized in the membrane with ten transmembrane domains connected on the external side by short loops. The cytosolic portion of the pump contains the catalytic center and other functionally important domains. The ATP binding site is indicated with a yellow circle. Acidic phospholipid binding domain (PL) and Calmodulin binding domain (CaMBD) are represented with pale blue boxes. (B) Deduced 3D structure of the PMCA pump. The three-dimensional structure of the PMCA pump has been obtained modeling it on that of the SERCA pump (Toyoshima et al. 2000). The image is a kind gift of Dr. M. Hilge (Nijmegen, Holland).

splicing that generates pump variants. Interestingly, the A-splice-site-insert also plays a key role in the targeting of PMCA to specialized portion of the plasma membrane of polarized cells (Chicka and Strehler 2003).

The cytosolic loop that connects transmembrane domains four and five contains the catalytic center that includes the ATP binding site and the aspartate residue that forms the acyl phosphate intermediate. It also contains the second binding site for the carboxy-terminal

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calmodulin-binding domain. The carboxy-terminal tail contains the main regulatory sites for the activity of the pump: the calmodulin binding domain (which also binds acidic phospholipids), consensus sites for protein kinases A (PKA), and C (PKC) and high affinity allosteric Ca^{2+} binding sites. The calmodulin-binding domain of the PMCA pump acts as an auto-inhibitory domain, binding to “receptor” sites in the first and second cytoplasmic loops of the pump. Calmodulin would remove the domain from the binding sites, freeing the pump from autoinhibition.

The carboxy-terminal region of the pump is also the target of isoform-specific phosphorylations by PKA and PKC. Phosphorylation could influence the regulation of the pumps either directly (e.g., by de-inhibiting it) or indirectly (e.g., via interference with calmodulin regulation). Recent studies have shown that PKC affects the various PMCA isoforms and splicing variants in different ways according to variations in the carboxy-terminal sequence: the phosphorylation may activate the pump, inhibit it, or have no effects (Strehler and Zacharias 2001). As for PKA, its consensus site has only been described in isoform one of the pump. The PMCA pump is also a substrate of intracellular proteases, as its carboxy-terminal tail contains target sequences for the Ca^{2+} dependent protease calpain (Guerini et al. 2003) and for caspase 1 and 3. Calpain cleaves the pump in the calmodulin binding sequence, leading to permanent activation of the pump. Both activation and inactivation have been described for caspases 1 and 3 action on PMCA2 and 4 (Schwab et al. 2002; Paszty et al. 2005).

The carboxy-terminal domain of the pump also contains an important site of alternative splicing, site C. C-site splicing occurs in all pump isoforms (albeit with varying degree of complexity) and causes the inclusion of one (or two) additional exons, or of portion of exons. The insertion has structural and functional consequences: it leads to the truncation of the pump because of a frame shift in the coding region, and alters, for instance, the pH sensitivity of the interaction of calmodulin with the calmodulin binding domain. In general, the

truncated isoforms have much decreased affinity for calmodulin (Enyedi et al. 1994). Surprisingly, however, the full length and the truncated variants expressed in living cells appear to have the same ability to restore the cytosolic Ca^{2+} transient generated by cell stimulation: the finding suggests that in vivo, calmodulin may not be the only factor regulating PMCA activity (Brini et al. 2003).

The carboxy-terminal domain also mediates the interaction with PDZ domains of different proteins, and mediates PMCA dimerization (which also activates the pump (Kosk-Kosicka and Bzdega 1988; Vorherr et al. 1991)). The interaction with PDZ domains is apparently limited to the full-length splice isoforms, as the carboxy-terminally truncated variants lack the consensus site for its binding. PMCA has been reported to interact with PDZ domain-containing proteins involved in the recruitment and maintenance of membrane proteins in specific membrane domains, such as members of membrane associate guanylate kinase (MAGUK) family, i.e., SAP, CASK, the Na/H exchanger regulatory factor 2, PISP and the Ania3/Homer protein (reviewed in (Brini and Carafoli 2009)). These proteins could contribute to targeting and retention of PMCA in specialized domains. The increased local pump concentration could regulate Ca^{2+} signaling. The PDZ binding domain also directs the interaction of one of the PMCA-isoforms (4b) with nitric oxide synthase 1 (NOS-1) in a ternary complex with α -syntrophin. The latter would bind to a domain further upstream in the pump sequence (Williams et al. 2006). The double interaction would thus down-regulate the production of NO by decreasing Ca^{2+} in the immediate vicinity of the synthase: i.e., the pump could function as a modulator of signal transduction, in addition to regulating cell Ca^{2+} (Cartwright et al. 2009).

Isoforms and Tissue Distribution

The four basic PMCA isoforms (the PMCA is the product of four separate genes) have tissue-specific expression (Brini and Carafoli 2009). Although PMCA1 and PMCA4 are expressed in most tissues, PMCA2 and 3 are found in a

restricted number of tissues, among them in brain, striated muscle and the mammary gland. In particular, PMCA2 is abundant in specialized cells such as cerebellar Purkinje neurons and cochlear hair cells, and it is also present in uterus and in lactating mammary glands. It is also expressed in significant amounts in liver and kidney. PMCA3 distribution is more restricted, the choroid plexus expressing significant amounts of it.

The transcript of each of the four genes encoding PMCA pumps is subject to alternative splicing: The sites in which it occurs are named A, B, and C. Of the large number of splice variants theoretically possible about 30 have been detected at the RNA or protein levels (Brini and Carafoli 2009). Alternative splicing at site A and C occurs for all four isoforms (however, PMCA1 is never spliced at site A). Alternative splicing at site B has been described only for human isoform one and four and it could be artifactual. Splice site A is located upstream of one of the regulatory binding sites for acidic phospholipids and involves up to three exons that can be optionally inserted or excluded. In PMCA3 and 4 a single exon may be included or excluded in the mature mRNA, thus generating the x and z splice variants, respectively. The situation in PMCA2 is more complex: in the human pump, three exons of 33, 60, and 42 nt, alternatively used, could in principle originate eight different variants. However, only four of them have so far been detected as mRNAs in a variety of tissues. Variant w includes all three exons, variant z excludes all three exons, variant x includes the 42 nt exon, and variant y includes the 33 and 60 nt exons. This alternative splicing was shown to be essential in the apical targeting of PMCA, and thus could also be involved in the regulation of the pump distribution in specialized portions of the plasma membrane (directly or through the interaction with other molecular determinants). Considering that splice site A is contiguous to one of the two phospholipid-binding domains, it could in principle effect the phospholipid regulation of PMCA isoforms by altering the accessibility of the binding domain to acidic phospholipids. As mentioned, the splicing

at site B may be an artifact, as its occurrence in vivo has been a point of controversy. It has been proposed to remove a carboxy-terminal 108 nt segment, leading to the loss of the 10th transmembrane domain and causing a reorganization of the pump topology (Preiano et al. 1996). Splice site C is located in the carboxy-terminal of PMCA protein, within the calmodulin binding domain: the inclusion of a large exon (154–175 nt) alters the carboxy-terminal half of the domain in all isoforms, and changes the reading frame of the remaining carboxy-terminal tail. This splicing is thus responsible for the generation of variant *a*, truncated after the first 12 residues of the calmodulin binding domain. The variant has reduced responsiveness to calmodulin with respect to the full-length variant (termed variant *b*). Lower Ca-calmodulin affinity and a significantly elevated basal pumping activity are general characteristics of the *a* splice variants in comparison with the *b* variants (Preiano et al. 1996; Elwess et al. 1997; Caride et al. 2007).

Interestingly, the expression of the PMCA pumps is transcriptionally regulated by Ca^{2+} itself (Zacharias and Strehler 1996; Carafoli et al. 1999). The phenomenon has been studied on cultured neurons and is correlated to their maturation. In cerebellar granule cells, as Ca^{2+} concentration in the cytoplasm increases, the pattern of expression of the PMCA pumps undergoes a dramatic rearrangement: isoform two and three become strongly up-regulated, whereas isoform one undergoes a splicing switch that generates high levels of the truncated *a* variant. In contrast, PMCA4 disappears in a process that is calcineurin dependent (Guerini et al. 2000). The reprogramming of the transcription of PMCA is essential to the survival of the developing neurons: the regulation of PMCA expression may thus be critical to the survival of cells exposed to pathological increases in intracellular Ca^{2+} . Parallel amplification of Ca^{2+} influx and efflux pathways may enable differentiated neurons to precisely localize Ca^{2+} signals in time and space (Usachev et al. 2001). Similarly, in rat hippocampal neurons, transcripts for all isoforms of the PMCA pump have been found to be strongly up-regulated during the second

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week in culture, the overall increase in Ca^{2+} extrusion systems being accompanied by changes in the expression and cellular localization of different isoforms. The accumulation of PMCAs in dendrites and dendritic spines coincided with the functional maturation of these neurons, underlining the importance of the proper spatial organization of the Ca^{2+} extrusion systems in synaptic function and development (Kip et al. 2006).

Role in Physiology and Pathology

PMCA pumps are expressed ubiquitously, thus, they are unlikely candidates as molecules that could mediate cell specific processes. However, the existence of so many PMCA variants (basic isoforms and alternative spliced forms), the finding on knockout mice for the specific PMCA isoforms, and studies on the mechanisms modulating their cell/tissue specific distribution indicate that PMCA pumps may play important roles as signaling molecules, in addition to having a constitutive role as Ca^{2+} housekeeping enzymes.

The ablation of isoforms two and four in mice has indeed revealed that they are important in specialized biological processes. PMCA2 plays an essential role in the hearing process, because it dissipates with peculiar kinetics the Ca^{2+} transients generated by the opening of the mechanoelectrical transduction (MET) channels in the stereocilia of cochlear hair cells. Mutations in the PMCA2 pumps in hereditary deafness impair the longer-term export of Ca^{2+} from hair cells, but not their ability to respond to a sudden increase of Ca^{2+} . This suggests that the mutated pumps reduce the extracellular Ca^{2+} gradient in the endolymph surrounding the hair cells, compromising the adaptation process responsible for the control of the opening of MET channels (Ficarella et al. 2007; Spiden et al. 2008). Interestingly, the PMCA2 isoform has another peculiar function in mammary gland: its activity is responsible for the high content of Ca^{2+} in the milk. PMCA2 is strongly up-regulated in lactating mammary gland, and PMCA2 null mice have 40% less Ca^{2+} in the milk (Reinhardt et al. 2004).

PMCA4 is crucial to male fertility: in mice, its ablation reduces sperm motility, probably resulting from Ca^{2+} overload and mitochondrial damage (Schuh et al. 2004). Interestingly, PMCA4 is also essential for the modulation of the activity of the calcium/calmodulin dependent neuronal nitric oxide synthase (nNOS) (Schuh et al. 2001) and thus, in turn, controls nitric oxide production, which is important in the regulation of excitation-contraction coupling of the heart. It has been shown that PMCA4b regulates cardiac contractility in vivo through its interaction with nNOS (Oceandy et al. 2007).

PMCA defects have been described in a large number of disease conditions, among them those linked to the oxidative stress in several tissues, particularly brain, in brain ischemia, diabetes, atherosclerosis, aging, neurodegenerative diseases, and various disturbances of Ca^{2+} metabolism. PMCA defects have also been reported in various cancer types (reviewed in Monteith et al. 2007), in line with accumulating evidence on the involvement of Ca^{2+} homeostasis dysfunction in the process of malignant transformation. The genetic diseases involving the PMCA pump are best characterized: a number of dysfunctions linked to the ablation, or to partial disruptions, including point mutations, of its genes, have been described in mice. So far, the only identified spontaneous human disease related to a PMCA pump (isoform 2) defect is the form of hereditary deafness described above (Schultz et al. 2005; Ficarella et al. 2007).

Genetic Manipulations

Knockout mice have been developed and the phenotypes studied for PMCA1, PMCA2, and PMCA4, but not yet for PMCA3. As this isoform is widely expressed in developing embryos, it may be essential for normal gestation.

The ablation of the PMCA1 gene has resulted in early embryonic lethality in homozygotes, underlying the essential role of PMCA1 in the early stages of development and in organogenesis, in line with its suggested housekeeping function (Okunade et al. 2004). Heterozygotes, by contrast, appeared healthy, however,

loss of one copy of the PMCA1 gene in mice with a PMCA4 null background led to increased propensity to apoptosis in the smooth muscles of blood vessels, possibly because of Ca^{2+} overload (Okunade et al. 2004). The ablation of the PMCA2 gene revealed its involvement in the hearing process: the phenotype of the PMCA2 knockout mice (Kozel et al. 1998) was characterized by balance impairment and hearing loss. The study of the vestibular inner ear revealed the absence of the otoconia and the progressive degeneration of the hair cells beginning 10 days after birth. The PMCA2 pump is probably also important to general neuronal Ca^{2+} homeostasis because PMCA2 null mice showed a significant reduction in the number of spinal cord motor neurons (Kurnellas MP 2005) and abnormalities in Purkinje neurons (Kurnellas MP 2007). However, the balance defect was apparently not due, as could in principle have been expected, or was only partially due, to a cerebellar dysfunction. It was apparently related to the absence of the Ca-carbonate crystals of the otoconia embedded in the otolithic membrane overlying the sensory epithelium that sense gravity and linear acceleration in the inner ear. In keeping with the high level of expression of PMCA2 in the mammary gland, the ablation of its gene, in addition to generating the hearing loss phenotype, also strongly reduced the concentration of Ca^{2+} in the milk (Reinhardt et al. 2004).

The ablation of the PMCA4 gene failed to cause a very evident general pathological phenotype (Schuh et al. 2004), but local defects were present. This is interesting, as PMCA4 is also widely expressed, and has also been proposed to play a housekeeping role. However, it has now become evident that PMCA4 plays more specialized roles, and is not only essential for the general function of controlling Ca^{2+} homeostasis in all cells. One prominent defect caused by PMCA4 dysfunction was male infertility, reflecting the dominance of the PMCA4 pump in the testis, where it represents more than 90% of the total PMCA protein (Prasad et al. 2004). The ablation of the PMCA4 gene produced other localized dysfunctions, for instance in the modulation of the Ca^{2+} signals in

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B-lymphocytes (Chen et al. 2004), and in the contractility of vascular or bladder smooth muscles (Okunade et al. 2004).

Studies on mice overexpressing PMCA4 isoform had revealed other important functions for this isoform. Early studies, in which the PMCA4 pump had been expressed specifically in the myocardium of the rat, and in vascular smooth muscle cells in mice, had indicated a role in the modulation of myocardial growth and hypertrophy (Hammes et al. 1998). They have also indicated a role in the regulation of the peripheral vascular tone, as the mice displayed increased peripheral blood pressure (Gros et al. 2003; Schuh et al. 2003).

SODIUM/CALCIUM EXCHANGER

Structural and Regulatory Characteristics

NCX accomplishes Ca^{2+} extrusion by using the electrochemical gradient of Na^{+} : during each cycle three Na^{+} ions enter the cell and one Ca^{2+} ion is extruded against its gradient (for comprehensive reviews see Blaustein and Lederer [1999]; Lytton [2007]). In addition to being transported, cytoplasmic Na^{+} and Ca^{2+} ions also regulate exchanger activity. Binding of Ca^{2+} ions to sites located in the cytosolic loop generally activate the exchanger, whereas binding of Na^{+} ions has been shown to deactivate it: the physiological importance and molecular mechanisms underlying the regulation remain unclear. The operation of NCX is fully reversible, and the direction of the movement of the transported ions depends entirely on the electrochemical gradients of Na^{+} and Ca^{2+} and on the number of ions that bind to the molecule and are transported. Under resting cellular ionic condition and membrane potential, the NCX acts to extrude Ca^{2+} from the cytoplasm. However, in the heart, when the plasma membrane becomes depolarized during systole, and the Na^{+} levels rise because of the opening of the plasma membrane voltage-operated Na^{+} channels, the exchanger reverses its operation and mediates Ca^{2+} entry. The influx of Ca^{2+} may play an important regulatory role in the excitation-contraction process by influencing

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the gating of voltage-operated Ca^{2+} channels and by altering the SR Ca^{2+} load.

A membrane topology model based on isoform one of the NCX now predicts nine transmembrane α -helices that divide the molecule in a amino-terminal portion composed of the first five transmembrane domains, and a carboxy-terminal portion composed of the last four transmembrane domains (Fig. 3A). These two portions are separated by a cytosolic loop

of about 500 residues that contains the site for NCX1 regulation, and is thus the target for the development of inhibitory compounds. NCX1 also contains two regions of internal repeats: the α repeats ($\alpha 1$ and $\alpha 2$ in Fig. 3A) are involved in ion binding and transport; the β repeats in the large intracellular loop are involved in binding of regulatory Ca^{2+} (CBD1 and CBD2 in Fig. 3). The β repeats have been structural defined by crystallization (Nicoll et al. 2006) and NMR studies (Hilge et al. 2006).

The first portion of the large cytosolic loop (close to transmembrane domain five) is an amphipathic sequence, called XIP (exchanger inhibitor peptide) because the addition of a peptide corresponding to this sequence inhibits the exchanger. This first portion of the region is responsible for the regulation by Na^+ and the acidic phospholipids. The second portion of the loop contains the two Ca^{2+} binding domains (CBD1 and CBD2) that are arranged in an antiparallel fashion, and connected through a third domain designated as CLD (α -catenin like domain) to the membrane portion of the NCX (Fig. 3B). The CBD1 is the primary Ca^{2+} sensor that detects small cytosolic Ca^{2+} increases and undergoes large structural changes that activate the exchanger. The CBD2 undergoes instead modest structural alterations and binds Ca^{2+} only at elevated concentrations (Hilge et al. 2006; Nicoll et al. 2006). The carboxy-terminal end of the large cytosolic loop contains a hydrophobic and proline-rich sequence that had been originally modeled as a transmembrane domain. Recent evidence has positioned it on the cytosolic side of the membrane.

In addition to the two transported species, other regulatory agents of the exchanger include the intracellular pH, metabolic components (e.g., ATP, phosphatidyl inositol 4,5 bisphosphate), protein kinases PKA and PKC, redox agents, hydroxyl radicals, H_2O_2 , dithiothreitol, O_2^- , Fe^{3+} , Fe^{2+} , Cu^{2+} , and OH^- (Doering and Lederer 1993; Matsuoka et al. 1995; Matsuoka et al. 1997; Iwamoto et al. 1998; He et al. 2000).

As is the case for the PMCA, a number of cytoskeletal proteins have been shown to interact

Figure 3. (A) Topology model of NCX. The nine transmembrane domains comprise the two putative transport repeat regions: α -1 and α -2. XIP region, CBD1 and CBD2 are indicated by yellow boxes. The red spheres indicate the Ca^{2+} binding sites position (B) Hypothetical structural model of the intact NCX. The nine transmembrane domains are shown as yellow α -helices, the CLD as blue α -helices and the CBD1 and CBD2 β repeats as red and green β strands, respectively. The red spheres indicate the Ca^{2+} binding sites position. The image is a kind gift of Dr. M. Hilge (Nijmegen, Holland).

with the NCX. The large cytosolic loop interacts with several proteins: the 14-3-3 protein, phosphorylated PLM (phospholemman, a member of a family of transport regulators, best known as modulators of Na, K-ATPase activity) and calcineurin: they all have inhibitory function on NCX activity (Katanosaka et al. 2005; Pulina et al. 2006; Wang et al. 2006; Zhang et al. 2006). Other proteins have also been shown to interact with the NCX: ankyrin (Li et al. 1993), the filamentous actin network (Condrescu and Reeves 2006), caveolin 3 (in the heart) (Bossuyt et al. 2002; Camors et al. 2006), and caveolin 1 and 2 (in neuronal cells) (Cha et al. 2004). These interactions have been proposed to regulate NCX activity by modulating the membrane NCX localization and by recruiting it to specialized portions of the plasma membrane as a part of macromolecular complexes (Schulze et al. 2003; Lencesova et al. 2004), similarly to PMCA, which can be recruited through the interaction with nNOS to form a specialized Ca^{2+} concentration microdomain.

The phosphorylation of NCX1 remains a topic of much controversy: several studies have claimed that PKA and/or PKC, operating through adrenergic stimulation, could modulate NCX function. The regulation could also be indirect and mediated by interactions with other proteins in macromolecular complexes.

Isoforms and Tissue Distribution

The NCX family belongs to a larger superfamily of related Ca^{2+} /cation antiporter genes that has five major branches. Three of them relate to the $\text{Na}^+/\text{Ca}^{2+}$ transport in mammals: NCX (SLC8 family), NCKX ($\text{Na}^+/\text{Ca}^{2+}$ - K^+ exchangers, SLC24 family) and the CCX (Ca^{2+} /anion exchangers), which contains the $\text{Na}^+/\text{Ca}^{2+}$ - Li^+ exchanger (NCLX). NCKX proteins play a crucial role in regulating Ca^{2+} fluxes during photoreceptors adaptation, synaptic plasticity, and skin pigmentation (for a comprehensive review see Lytton [2007]). NCLX is the single mammalian member of the phylogenetically ancestral CCX family. It has been initially characterized as a novel plasma membrane $\text{Na}^+/\text{Ca}^{2+}$ exchanger that catalyzes Na^+ or Li^+ -dependent

Ca^{2+} transport, but very recent evidence indicates that it is the long elusive mitochondrial $\text{Na}^+/\text{Ca}^{2+}$ exchanger (Palty 2009).

The SLC8 family includes three basic proteins, NCX1, NCX2, and NCX3, encoded by three separate genes. A fourth member, described in teleost fish, is not present in the mammalian genome (Marshall et al. 2005). NCX1 was originally characterized and cloned from heart, but its expression is almost ubiquitous, with high levels in brain and kidney. NCX2 and NCX3 are selectively expressed in the brain and skeletal muscle, and in some neuronal populations, respectively. NCX1 is the best-characterized member of the family. Its expression is controlled by alternative promoters in a tissue-specific and transcription factor-specific manner: one of them is specific to heart, one to kidney, and one to the other tissues. The transcripts of NCX1 gene are alternatively spliced at two sites: the first is located in the 5'-untranslated region and does not change the structure of the mature protein. The second is located within the coding region and leads to the introduction of two mutually exclusive exons (exon A in excitable tissues such as brain and muscle, and exon B in nonexcitable tissues), and of four other exons, thus potentially encoding a large number of NCX1 isoforms with differences in the cytosolic loop. Interestingly, giant excised patch clamp experiments have shown that exon A containing isoforms, in contrast to exon B containing variants, may not have intracellular Na^+ -dependent inactivation (Dyck et al. 1999). NCX3, but probably not NCX2, is alternatively spliced to generate variability in the cytoplasmic region corresponding to that of NCX1. No distinctive properties have been described for the different splicing isoforms, suggesting that their specific function is probably related to different spatial targeting.

Importantly, as is the case for the PMCA pump, the transcription of NCX genes is also differentially regulated by Ca^{2+} during development, and also in adult neurons (Carafoli et al. 1999). NCX2 and NCX3 transcripts are down-regulated and up-regulated in cerebellar granule neurons, respectively, under the conditions of intracellular Ca^{2+} -increase that reprogram the

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pattern of transcription of the PMCA pumps. The down-regulation of NCX2 transcription in cerebellar granules is calcineurin dependent (Li et al. 2000), whereas that of NCX3 in model neuroblastoma neurons is regulated by Ca^{2+} through the transcriptional repressor DREAM (Gomez-Villafuertes et al. 2005).

Role in Physiology and Pathology

As the NCX is not an enzyme, the qualitative measurement of its activity has always been difficult and the role of NCX in physiological conditions has been mostly inferred from theoretical considerations and from observations obtained on isolated systems in artificial conditions, or with invasive approaches such as giant patch techniques. More recently, the availability of selective inhibitors (Iwamoto 2007) and genetic manipulation approaches have permitted the dissection of many physiological aspects of NCX activity. The role of NCX has so far been best characterized in the heart: contraction is initiated by a small influx of Ca^{2+} from the extracellular ambient that induces a larger release of Ca^{2+} from the SR; relaxation requires Ca^{2+} removal, which is achieved mainly by the extrusion activity of NCX1 and the pumping back of Ca^{2+} in the SR lumen by the SERCA pump. Ca^{2+} extrusion through NCX1 also generates an inward depolarizing current (because of Na^+ entry and to the uneven charge translocation) which may contribute to the shaping of the action potential, and/or counteracts the Ca^{2+} entry pathway by reducing its driving force. The Ca^{2+} entry during the reverse mode operation of NCX1 (induced by signals that generated Na^+ entry, i.e., through ionotropic glutamate receptors) has been described to contribute to Ca^{2+} influx. These actions probably do not modify the single action potential, but they have been suggested to play a possible role in the refilling of the stores with Ca^{2+} , thanks to the formation of protein complexes and functional coupling between plasma membrane microdomains and the ER lumen mediated by local Ca^{2+} gradients (Blaustein and Golovina 2001). In addition to the heart, where its function has received great attention,

numerous studies have underlined the role of the NCX in other tissues as well. The NCX1 has been shown to play a special role in the kidney, where the Ca^{2+} filtered at the glomerulus is reabsorbed passively along the proximal nephron. However, at the distal nephron level, vitamin D_3 and parathyroid hormone regulate Ca^{2+} transport. NCX1 is highly expressed in the basolateral membrane of the distal nephron, and helps regulating the active transcellular Ca^{2+} reabsorption, thus contributing to the regulation of systemic Ca^{2+} levels rather than to the intracellular levels. The role of the exchanger in other tissues has not been analyzed in the same detail as it has in the heart (or kidney); however, in the last few years its role in the brain has received increasing attention, particularly as a means to protect neurons from ischemic damage. The specific contribution of each of the three basic NCX isoforms cannot be established with certainty, because they are coexpressed in brain neurons. However, the widespread and abundant expression of NCX2 has indicated a key role for it in neuronal Ca^{2+} homeostasis. NCX3 also appears to be important to neurons: its specific cleavage during brain ischemia and in neurons undergoing excitotoxicity has shown that it plays a critical role in protecting them from ischemic insults (Bano et al. 2005; Annunziato et al. 2007). However, the issue still has controversial aspects: evidence has been provided that during brain ischemia changes in membrane potential and unregulated Ca^{2+} entry could activate the NCX Ca^{2+} entry-mode, and thus contribute to Ca^{2+} overload and eventual neurodegeneration (Kintner et al. 2007). Other work has suggested instead that during excitotoxic insults, the activation of calpain leads to the cleavage of NCX and thus to the impairment of the Ca^{2+} extrusion process, culminating in excitotoxic death. Thus, the inhibition (or the down-regulation) of the exchanger could be neuroprotective (Luo et al. 2007), but could also transform a Ca^{2+} transient elicited by nonexcitotoxic stimulation into a lethal Ca^{2+} overload (Bano et al. 2005; Annunziato et al. 2007).

Considering the essential role of NCX in regulating Ca^{2+} homeostasis in a variety of

tissues, the importance of understanding how changes in its activity could contribute to pathological situations is obvious. No genetic disease has so far been associated to NCX mutations, but a number of pathologies have been related to its malfunctioning. Again, the best documented studies are related to heart dysfunctions: NCX deficiencies have been claimed to play a role both in cardiac arrhythmias and in the ischemia/reperfusion injury. They can contribute to the former in two ways: through changes in the balance between the forward-mode of Ca^{2+} efflux and the reverse-mode of Ca^{2+} influx consequent on the deregulation of intracellular Na^+ , or through conditions that could lead to Ca^{2+} overload in the SR, and thus to spontaneous Ca^{2+} release from it that could generate extrasystolic events: the extra Ca^{2+} release could enhance NCX activity, thus generating a depolarizing current that can contribute to the elongation of the action potential. Up-regulation of NCX expression at the transcriptional level has been described in cardiac hypertrophy, ischemia, and failure (Kent et al. 1993). However, it is difficult to establish whether the up-regulation is the cause of the dysfunction or rather an adaptive mechanism that, in the end, leads to cardiac hypertrophy and heart failure.

In ischemia/reperfusion injury the action of NCX is still linked to its reverse mode of operation: although other factors could contribute, the removal of H^+ ions that accumulate during the ischemic period by the Na^+/H^+ exchanger could enhance the intracellular Na^+ , thus activating Ca^{2+} entry, and eventually causing SR Ca^{2+} overload.

Genetic Manipulations

In the last few years, various NCX knockout mice have been generated. Homozygous NCX1 KO mice were not vital and died during embryonic development, possibly because of heart failure (Wakimoto et al. 2000). However, the cardiac-specific knockout of NCX1 was not lethal (Henderson et al. 2004) suggesting that the cause of lethality in NCX1 KO may have been extra-cardiac, and that NCX2 and 3 could

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not compensate for the absence of NCX1. The cardiac-specific NCX1 KO mice presented only a mild deficit in cardiac function that culminated in hypertrophy and heart failure only as the mice aged. Ca^{2+} measurements on myocytes revealed that the reduced Ca^{2+} clearance in the absence of NCX1 led to enhanced Ca^{2+} dependent inactivation of L-type Ca^{2+} channels. These myocytes also displayed a shorter action potential because of hyperpolarization caused probably by the increase in the expression of K^+ channel subunits. The outcome of these modifications was a reduction of about 80% of Ca^{2+} fluxes through the plasma membrane: however, this reduction did not impair the Ca^{2+} release from the SR and contractility because of compensation mechanisms involving the gain of excitation-contraction coupling.

NCX2 KO mice displayed impairment in several hippocampal-dependent learning and memory tasks (Jeon et al. 2003). Larger presynaptic Ca^{2+} transients evoked an increase in the neurotransmitter release, and the increased postsynaptic Ca^{2+} transient enhanced long-term potentiation: the findings are consistent with the predominant role of NCX2 in pre- and postsynaptic Ca^{2+} clearance.

Mice lacking the NCX3 gene, which is highly expressed in cerebellum and in the peripheral nervous system, showed reduced motor activity and weakness of forelimb muscles (Sokolow et al. 2004). In addition, skeletal muscle defects were also observed, in line with the finding of abundant NCX3 expression in skeletal muscle.

CONCLUDING REMARKS

To respond dynamically to the changing needs of Ca^{2+} signaling, cells must be able to precisely control the type, amount, localization, and activation of Ca^{2+} transporters. This control is performed in two ways: through the variable expression of Ca^{2+} transporters with different biochemical characteristics, and through the fine modulation of their function by expressing differently active isoforms in response to the local or temporal cell demands. The cell also modulates the activity of its transporters by choosing specific molecular partners. This

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increases the complexity of the controlling machinery, but protects the cells from “the Ca^{2+} damage” that would inevitably follow the general failure of the Ca^{2+} controlling operation. The absence, or malfunction, of specific Ca^{2+} transport proteins would induce a confined defect: although generating functional discomfort, it would frequently still be compatible with cell life.

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The Plasma Membrane Ca^{2+} ATPase and the Plasma Membrane Sodium Calcium Exchanger Cooperate in the Regulation of Cell Calcium

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