Inhibition of the Na\(^+\)/Ca\(^{2+}\) Exchanger NCX\(_1\) Expressed in Xenopus Oocyte by Glycyrrhizic Acid and Cyclophylin A

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Abstract

The Na\(^+\)/Ca\(^{2+}\) exchanger plays an important role in regulation of airway smooth muscle contraction by regulating intracellular calcium, and is a potential target for treatment of asthma. To test modulation of exchanger activity, we used Xenopus oocytes as model system. Na\(^+\)/Ca\(^{2+}\) exchanger was expressed in the cells by microinjection of cRNA of the exchanger isoform NCX\(_1\). The activity of NCX\(_1\) was determined as Ni\(^2+\)-sensitive current under voltage clamp in low Cl\(^-\) medium and in the presence of the Cl\(^-\)-channel inhibitor niflumic acid. Only this composition of solution allowed determining NCX\(_1\)-mediated current with sufficient accuracy. Among a few tested Chinese herbal drugs, glycyrrhizic acid turned out to be a potent inhibitor of NCX\(_1\) with an apparent IC\(_{50}\) value of 40 \(\mu\)M. Previous work had revealed elevated cyclophylin A concentration in serum of asthmatic rats after receiving acupuncture treatment. Extracellular incubation of the oocytes in cyclophylin A for one day led to significant inhibition with an apparent IC\(_{50}\) value of about 1 \(\mu\)M. We suggest that effects of acupuncture and application of glycyrrhizic acid as an active constituent of Chinese medicine for treatment of asthma symptoms may partially be attributed to inhibition of the reversed mode of NCX\(_1\), and that these compounds may stimulate the search for new anti-asthmatic drugs.

Keywords

Sodium-Calcium Exchanger, Asthma, Voltage Clamp, Glycyrrhizic Acid, Cyclophylin A
1. Introduction

Asthma is a disease characterised by reversible contraction of airway smooth muscle (ASM). Several signalling pathways are now known to be related to the process of ASM contraction, and almost all of them involve Ca\(^{2+}\) handling [1]. Cytoplasmic Ca\(^{2+}\) activity (Ca\(^{2+}\)) homeostasis is controlled by several ionic signalling mechanisms, one of which is the reversed mode of the Na\(^+\)/Ca\(^{2+}\) exchanger (NCX) of the plasma membrane [2] [3]. Although little is currently known about NCX in the airways and its involvement in asthma, some investigators have shown that expression and function of the isoform 1 of the exchanger (named NCX) in ASM is up-regulated in asthmatic animals and inhibition of NCX may ameliorate the symptoms of asthma. This observation makes NCX a potential target for asthma treatment [2] [4]. The aim of the present study was to examine effects of various chemicals that might act as inhibitors of NCX.

In traditional medicine, herbal extracts are often applied in treatment of asthma, and the search for natural components has become promising to discover new anti-asthmatic drugs. Extracts of roots of licorice (Glycyrrhiza glabra) have been applied in treatment of a large variety of diseases (see e.g. [5]). The triterpene glycoside glycyrrhizic acid (GA) is one of the major active constituents of licorice:

![GA molecule](image)

GA has been used as a hepatoprotective drug [5], and recent studies also revealed its anti-asthmatic effects [6] [7]. Though modulation of various pathways has been discussed (see [8] [9]), it is unclear whether GA can interfere with NCX, and hence exert its anti-asthmatic effects.

In a recent investigation on rats, we found significantly elevated cyclophilin A (CyPA) level in the serum of acupuncture-treated asthmatic rats compare to that of untreated rats [10]. CyPA is a member of the cyclophilin (CyP) family, which possesses peptidyl-prolyl isomerase (PPIase) activity. CYPs are involved in diverse cellular processes including cell-cycle regulation, receptor signalling, protein folding, and they form cellular targets for immune-suppressant drugs such
as cyclosporine A (CsA) [11]. CyPA has multiple intracellular functions (see e.g. [12] [13]), but can also be secreted [14] and act extracellularly as an inflammatory mediator that may be involved in inflammatory diseases such as atherosclerosis [15] [16] and rheumatoid arthritis [17]. In our investigation we consider extracellular CyPA as a drug for treatment of asthma by inhibiting NCX, which might be a molecular mechanism of asthma therapy by acupuncture. In this study, we choose the human cyclophilin A (hCyPA), one of 7 major cyclophilins in humans [18].

To monitor changes of transport activity of the exchanger, we used the *Xenopus* oocyte for heterologous expression of NCX. NCX operates at a 3:1 or 4:1 Na⁺:Ca²⁺ stoichiometry [19] [20], the transporter is electrogenic, therefore, the activities can be monitored by measuring current using two-electrode voltage clamp (TEVC). Since the oocytes have functionally expressed only a limited number of endogenous membrane proteins, the application of this model system allows investigating effects on NCX with low background signals and restricted functional interference from other membrane proteins.

2. Materials and Methods

2.1. NCX₁-cRNA and hCyPA Preparation

The construct with NCX₁ of dog was kindly provided by Dr. Luis Beauge (Laboratorio de Biophisica, Cordoba, Argentina) and linearised with *XbaI*, then transcribed into cRNA *in vitro* using mMESSAGEmMACHINE SP6 kit (Ambion, USA). The final concentration of cRNA was adjusted to 0.2 ng/nL.

The plasmid pQE30-CyPA was kindly provided by Dr. Xu Sheng (Drug Discovery and Design Center and State Key Laboratory of Drug Research, Shanghai Institute of Materia Medica, Chinese Academy of Sciences). Expression and purification of the hCyPA protein was performed as described elsewhere [21] [22].

Gels and buffers used for native PAGE were made according to the standard Laemmli SDS protocol omitting the SDS. Native gels (12% polyacrylamide) were run at 12 mA and 45 min and stained with Coomassie Brilliant Blue R-250. The molecular weight of hCyPA was slightly less than 20 kDa (compare Figure 1) in line with the reported weight of 18 kDa. Unstained protein-molecular-weight marker was from Fermentas Life Science (USA).

2.2. *Xenopus* Oocytes Preparation and Microinjection

*Xenopus* oocytes were used as expression for NCX, and as a model system to test the effects of herbal extracts and CyPA. This expression system is particularly suited because endogenous ion channels and transporters are functionally expressed only to a low extend, and hence, exogenous current components can easily be extracted. Females of the clawed toad *Xenopus laevis* (purchased from Maosheng Bio-Technology Com., Shanghai, China) were anaesthetised in a bath medium containing 1 g/L tricaine (Sandoz, Basel, Switzerland) and kept on ice. Parts of ovary were removed and treated with 0.5 or 0.25 mg/mL collagenase (Sigma) for 2 - 4 h, or overnight, respectively. Full-grown prophase-arrested
oocytes of Dumont stages V and VI [23] were selected for cRNA injection, and cultured with daily changed G-ORi solution or C-ORi solution (ORi plus 0.07 μg/mL gentamycin (Sigma) or ciprofloxacin hydrochloride (Sigma), respectively) at 20˚C. For expression of NCX, about 1.5 ng NCX1-cRNA was microinjected into an oocyte at a flow-rate of 8 nL/s. Uninjected oocytes served as controls.

For investigating the effect of intracellular hCyPA, oocytes were first microinjected with NCX1-cRNA and divided into 4 groups. The cells of each group were additionally microinjected with hCyPA (5, 10, 20 or 40 nl per oocytes, respectively, at a concentration of 4.27 µg/µL). For investigating extracellular effect of hCyPA, oocytes were first microinjected with NCX1-cRNA and divided into 4 groups. The cells of each group were then incubated with different amounts of hCyPA; 200 µL of incubation medium contained 1, 2, 4 or 8 µL of hCyPA, respectively (4.27 µg/µL). Thereafter, oocytes were cultured in G-ORi or C-ORi at 20˚C for up to 2 days.

2.3. Electrophysiological Recording

Since NCX is electrogenic, NCX-mediated current is a measure for transporter activities. To investigate the function of NCX1, membrane currents were measured by conventional two-electrode voltage clamp (TEVC) using Turbo TEC-03 with Cell Works software (NPI electronic, Tamm, Germany). Glass microelectrodes were filled with 3 M KCl, and balanced in ORi solution for at least 30 min before recording. Before measurements, oocytes were loaded with Na⁺ by incubating the cells in Na⁺-loading solution for 30 min [24]. Thereafter, cells were kept for at least 30 min in post-loading solution (see solutions). Steady-state current-voltage dependencies were determined by averaging membrane currents during the last 20 ms of 200-ms rectangular voltage pulses from −150 to +30 mV in 10 mV increments that were applied from a holding potential of −60 mV. NCX-dependent current was determined as the difference of total membrane

![Figure 1. SDS-PAGE analysis of hCyPA. Lane 1: hCyPA-antibody marked band at slightly less than 20 kDa. Lane 2: marker.](image-url)
current in the absence and presence of 2 mM NiCl₂ as a specific inhibitor of NCX. The data were collected after analogue filtering at 300 kHz and analysed by Origin software (OriginLab Corp., USA). All experiments were performed at room temperature (about 25°C).

2.4. Solutions and Drugs

The composition of ORi was (in mM): 90 NaCl, 2 KCl, 2 CaCl₂ and 5 MOPS (adjusted to pH 7.4 with Tris). To elevate intracellular Na⁺, cells were incubated for 30 min in “Na-loading solution” consisting of (in mM): 110 NaCl, 2.5 Na-citrate, 5 MOPS (adjusted to pH 7.4 with Tris) and stored thereafter for at least another 30 min in “Post-loading solution” consisting of (in mM): 100 NaCl, 5 BaCl₂, 20 TEA, 5 MOPS (adjusted to pH 7.4 with Tris). Standard test solutions contained (in mM): 100 Na-gluconate, 2 CaCl₂, 0.1 niflumic acid, 5 MOPS, and 0 or 2 mM NiCl₂ (adjusted to pH 7.4 with Tris).

Herbal extracts were kindly provided from Shanghai Institute Materia Medica (CAS) by Drs. CG Huang and CH Ma (supercritical fluid extraction of root of Acorus tatarinowii Schott) and by Dr. LJ Xuan (dried ethanol extracts of Ilex pubescence and Gossampinus malabarica), and were dissolved in DMSO. Final concentration of the herbal extracts in test solution was 40 mg/L.

GA (CAS 1405-86-3, purity ≥95%) and α-asarone (CAS 2883-98-9, purity 98%) were purchased from SIGMA. Stock solutions of 1 or 100 mM were prepared in DMSO and diluted to the final concentration in the test solution. DMSO concentrations in all test solutions were below 1%, which was without effect on the membrane currents.

2.5. Data Analysis

Analysed data were represented as means (±SEM) from N experiments. Means were considered as significantly different by Student’s t test on the basis of p < 0.05.

3. Results

The Na⁺/Ca²⁺ exchanger is considered to transport 3 or 4 Na⁺ against 1 Ca²⁺, and hence generating in its forward mode (Ca²⁺ extrusion) an inward-directed current, and in its reversed mode (Ca²⁺ uptake) an outward-directed current (for a review see [25]). To determine this current, we used 2 mM Ni²⁺ as an inhibitor of NCX-dependent current. Total membrane current was measured under voltage clamp, and the difference of steady-state current in the absence and presence of Ni²⁺ was taken as a measure for NCX-dependent current.

3.1. Ni²⁺-Sensitive Currents Represent only in Part NCX₁-Mediated Current

In control oocytes not injected with cRNA of NCX₁, Ni²⁺-sensitive could never be detected (Figure 2(a)), confirming that the cells do not express significant endogenous Na⁺/Ca²⁺ exchanger on the plasma membrane [26]. On the contrary,
Figure 2. Ni\textsuperscript{2+}-sensitive current in un-injected and NCX\textsubscript{1}-cRNA-injected oocytes. (a) Only injected oocytes show Ni\textsuperscript{2+}-sensitive current. (b) Addition of 100 nM niflumic acid strongly blocked the outward-directed current component, while inward current became enhanced. (c) Reduction of external Cl\textsuperscript{-} from 100 to 6 mM strongly blocked the outward-directed component. The data represent averages of N = 5 to 7 oocytes (±SEM).
oocytes being injected with cRNA of NCX₁ exhibited huge currents, in particular at positive potentials in outward direction. Interestingly, the NCX-dependent current could even exceed 10 µA, which can hardly be mediated by a carrier protein even at high density in the cell membrane. Nevertheless, such large currents had been considered to be mediated by NCX (see e.g. [27]). On the other hand, *Xenopus* oocytes exhibit Ca²⁺-activated Cl⁻ current (see [28]). Therefore, an alternative interpretation would be that the transporter operates at these potentials in reversed mode accumulating Ca²⁺ at the intracellular membrane surface. This accumulated Ca²⁺ would activate the Cl⁻ channels. After blocking NCX₁ by Ni²⁺ also the Cl⁻ channels will no longer be activated.

To reduce this Ca²⁺-dependent background current, we used as an inhibitor of the Cl⁻ channels 100 nM niflumic acid [29] in the test solutions, which indeed led to a considerable reduction of the Ni²⁺-sensitive outward-directed current (Figure 2(b)); in addition the inward-directed current was enhanced. Figure 2(c) illustrates that reduction of external Cl⁻ from 100 mM to 6 mM, also led to strong inhibition of the outward-directed current. Therefore, in most experiments, if not state otherwise, we used niflumic acid and the Cl⁻-reduced test solution as given in the Materials & Methods Section. The still remaining Ni²⁺-sensitive current in this standard test solution was considered to be mediated by NCX₁. In the following we will differentiate between “NCX₁-dependent” (or total Ni²⁺-sensitive) and “NCX₁-mediated” current.

### 3.2. Drug Effects on NCX₁-Dependent Current

For a first rough screening we looked for drug effects on total Ni²⁺-sensitive current. To determine this current, we used standard external oocyte-Ringer’s-like solution (ORi) in the absence of niflumic acid without and with 2 mM NiCl₂. The extracts of *Ilex* and *Gossampinus* showed slight, but statistically significant inhibition at 40 mg/L by about 15% or 25%, respectively; the *Acorus* extract showed slight stimulation by about 15% of the Ni²⁺-sensitive outwardly directed current at +10 mV (Table 1). We also tested several pure compounds in addition to the *Acorus* extract α-asarone, which showed slight inhibition by 10% (Table 1). Out of several tested drugs only the GA from *Glycyrrhiza galabra* exhibited significant and clear inhibition by about 60% at 40 mg/L (Table 1). In the following, therefore, our focus was on the effect of GA. Since the scatter of current measurements under voltage clamp to positive potentials in general is pretty large, we will concentrate on the analysis at negative potentials.

### 3.3. GA Inhibits NCX₁-Mediated Current

In the standard experiment, membrane currents were measured in different solutions with low Cl⁻ and 100 nM niflumic acid that were applied usually in the sequence:

\[
0\text{Ni}^{2+} \rightarrow 2\text{Ni}^{2+} \rightarrow 0\text{Ni}^{2+} \rightarrow 0\text{Ni}^{2+} + \text{drug} \\
\rightarrow 2\text{Ni}^{2+} + \text{drug} \rightarrow 0\text{Ni}^{2+} + \text{drug} \rightarrow 0\text{Ni}^{2+} \rightarrow 2\text{Ni}^{2+} ightarrow 0\text{Ni}^{2+}
\]

The currents measured in the respective 0Ni²⁺ solution before and after the
Table 1. Effect of selected drugs on Ni²⁺-sensitive current measured under voltage clamp at +10 mV. Data represent averages of N measurements (±SEM). p values refer to difference to 1 (one-sample t-test).

<table>
<thead>
<tr>
<th>Drug</th>
<th>Relative change in current at +10 mV</th>
<th>Significance p</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Ilex pubescens</em> extract (40 mg/L)</td>
<td>0.83 ± 0.04 (N = 5)</td>
<td>0.01</td>
</tr>
<tr>
<td><em>Gossampinus malabrica</em> extract (40 mg/L)</td>
<td>0.74 ± 0.05 (N = 4)</td>
<td>0.007</td>
</tr>
<tr>
<td><em>Acorus tatarinowii</em> extract (40 mg/L)</td>
<td>1.15 ± 0.01 (N = 8)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>α-asarone (–40 mg/L = 200 µM)</td>
<td>0.90 ± 0.03 (N = 4)</td>
<td>0.04</td>
</tr>
<tr>
<td>Glycyrrhic acid (GA) (–40 mg/L = 85 µM)</td>
<td>0.39 ± 0.13 (N = 5)</td>
<td>0.04</td>
</tr>
</tbody>
</table>

application of 2 mM Ni²⁺ were averaged to partially compensate for small drift with time. The currents in 2 mM Ni²⁺ were then subtracted to obtain the NCX₁-mediated current. In stable experiments, the ⁰Ni²⁺ → ²Ni²⁺ → ⁰Ni²⁺ sequence could be repeated with another drug concentration. In some experiments an abbreviated protocol was applied with the solution sequence

⁰Ni²⁺ → ²Ni²⁺ → ⁰Ni²⁺ → ²Ni²⁺ + drug → ⁰Ni²⁺ → ²Ni²⁺ → ⁰Ni²⁺

The result of the effect of 20 µM GA on the current-voltage dependence of NCX-mediated current is illustrated in Figure 3(a) showing a significant inhibition of the activity of the exchanger over the entire potential range. A more detailed analysis of the concentration dependency yielded an IC₅₀ value of about 40 µM at −100 mV (Figure 3(b)). The inhibition did not significantly depend on membrane potential.

3.4. The Effect of hCyPA on NCX-Mediated Current

Microinjection of hCyPA up to 86 ng/µL (corresponding to 4.8 µM within the cytoplasm, calculated by assuming an oocyte volume of 1 µL) hardly affected the NCX₁-mediated current (Figure 4(a)). Interestingly, extremely high concentration (9.6 µM) obviously stimulated the current by about 30%. Despite the large error bars, the current increase is statistically significant.

Incubation of NCX₁-expressing oocytes in 1.2 µM hCyTA resulted in significant inhibition of NCX-mediated current (p < 0.05 compared to untreated cells); higher concentration 4.8 µM produced only insignificantly more inhibition (Figure 4(b)). Despite the considerable scatter of data an IC₅₀ value for 50% inhibition of less than 1 µM could be estimated (see inset of Figure 4(b)). For the experiments described above, the oocytes were incubated in the respective hCyPA solution for 2 days. One hour of incubation already showed some tendency of inhibition (about 10%), but only after one day maximum inhibition could be detected. From 2 batches of oocytes we found that the current at −100 mV was after 1 h of incubation reduced to 0.87 ± 0.16 and after one day to 0.34 ±
Figure 3. Inhibition of NCX-mediated current by GA. (a) Effect of 20 μM GA on current-voltage dependency. The data are normalised to the current at −60 mV in the absence of drug, and represent averages of N = 12 oocytes (±SEM). (b) Dependence of NCX-mediated current at −100 mV on GA concentration. Data represent averages of N = 5 to 12 oocytes (±SEM). The solid line is a fit of

\[ I_{IC_{50}} - \frac{IC_{50}}{IC_{50} + [GA]} \]

to the data with \( IC_{50} = 40 \mu M \) (n = 0.5). \( I_{IC_{50}} \)
is the normalised current at the respective drug concentration [GA].

0.06 compared to untreated cells.

4. Discussion

Asthma is a reversible airway restriction based on ASM contraction, which is governed by intracellular calcium. A key role plays the release of \( Ca^{2+} \) from intracellular stores, and the refilling of the stores involves the reversed mode of NCX (for a brief review see [3]). The aim of the present study was to examine effects of various drugs that inhibit the NCX, and hence might act as antiasthmatic drugs. We used Xenopus oocytes with heterologously expressed NCX, as a
Figure 4. The effect of hCyPA on NCX₁-mediated current. (a) Effect of intracellular hCyPA. Oocytes injected with NCX₁-cRNA were additionally microinjected with hCyPA to gain the respective intracellular concentration. Data represent averages of N = 5 to 11 oocytes ± SEM; (b) The effect of extracellular hCyPA. Oocytes were incubated in the respective amounts of hCyPA for two days. The inset shows the dependence of NCX₁-mediated normalised current at −100 mV on hCyPA concentration. Data represent averages of N = 7 to 10 oocytes ± SEM. The inhibition of current at all concentrations is significant on the basis of p < 0.05.

model system, and Ni²⁺ as specific inhibitor of the exchanger. Effects on NCX₁-mediated current could only be investigated as Ni²⁺-sensitive current in the presence of niflumic acid to block Ca²⁺-activated Cl⁻-currents and at lowered external Cl⁻ activity. Otherwise the Ni²⁺-dependent current also included a large component of Ca²⁺-activated Cl⁻ current (compare [27]). Since the oocytes have functionally expressed only a limited number of endogenous membrane proteins, the application of this model system allows investigating effects on NCX with low background signals and restricted interference with other membrane proteins.
Our drug screening revealed that GA is a potent inhibitor of NCX₁. GA has particularly been used in the treatment of liver diseases [5], but also seems to have anti-asthmatic effects [6] [7]. Since inhibition of NCX₁ in ASM will lead to reduced Ca²⁺ influx and reduced refilling of the intracellular Ca²⁺ stores, reduced muscle tone can be expected [2] [3] [4]. Though our measurements with GA were performed on the forward mode of NCX₁, the screening experiments with effects on the outward-directed current at +10 mV indicate that also the reversed mode can be inhibited.

Several lines of evidence implicate that intracellular CyPA plays a critical multifunctional role, and interaction with cyclosporine A (CsA) has been shown to be an important step [18] [30]. Our functional analysis revealed that intracellular CyPA also affected the NCX₁ protein, and up-regulated NCX₁-mediated current was observed though only at very high concentration of about 10 μM. The dissociation constant of CyPA from CsA is in the submicromolar range [31]. The much higher concentration in the micromolar range for intracellular stimulation of NCX₁ by CyPA, therefore, makes physiological relevance unlikely. On the other hand, Ca²⁺ uptake experiments with HEK cells co-transfected with NCX₁ and CyPA suggest involvement of CyPA in the regulation of NCX₁ expression and transport activity [32].

In fact CyPA can be secreted [16] via a vesicular pathway [14]. It had been demonstrated previously that acupuncture treatment on asthmatic rats can reduce airway restrictions [33], and this was associated with the elevation of CyPA in the serum [10]. Our results suggest that extracellular hCyPA down-regulates NCX₁-mediated current, which could account for release of the airway restrictions.

The involvement of extracellular CyPA in allergic lung inflammation had been suggested on the basis of the anti-inflammatory effect of an extracellularly applied membrane-impermeable CsA derivative [34]. Whether the effect of hCyPA found in our experiments results from direct interaction with the NCX₁ protein needs further investigation. Our finding that the extracellular inhibition needed several hours of incubation is in favour of an indirect effect.

5. Conclusion

In conclusion, our data suggest that GA and acupuncture-induced elevation of hCyPA in the serum may both contribute via inhibition of reversed NCX to reduced refilling of Ca²⁺ stores, which can promote reduced contraction of ASM, reduced airway restriction, and thus relieve the asthmatic symptoms. Such effects need to be verified in future animal experiments. Nevertheless, derivatives of GA and CyPA may form the basis for development of a new generation of more potent drugs for asthma therapy.

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