Difference in regulation mechanisms of ENaC by aldosterone and glucocorticoids

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ABSTRACT

Na⁺ transport occurs across many epithelial surfaces and plays a key role in regulating salt and water absorption. The molecular pathway underlying this Na⁺ transport is the epithelial Na⁺ channel (ENaC), which is strictly determined by a variety of hormones like aldosterone, ADH and glucocorticoids. In this study, we found that stimulation of either aldosterone or dexamethasone (Dex) distributed ENaC channel on the apical membrane of mouse cortical collecting duct cells (M1). In the single channel recordings from excised membrane, high density ENaC was found in the cell with a dome shape by the treatment of either dex or aldosterone. However, low active ENaC was revealed in intact cells treated with dex, when compared to cells treated with aldosterone. Only 5.84% of cells treated with dex containing ENaC exhibited ENaC current transition in the cell-attach recording, whereas 40% of cells treated with aldosterone containing ENaC exhibited ENaC current transition. ENaC currents appeared rapid rundown within 5 minutes since formation of inside-out configuration in cells treated with aldosterone but not with dex. SKF-525A, a general antagonist of CYP, failed to significantly enhance ENaC activity in intact cells treated with dex, but EGTA, which deforming the cells, increased the ENaC activity in the cells treated with dex. PTX, an antagonist of G-protein, reversed the effect of aldosterone on number of active ENaC in intact cells. Based on our observation, we concluded that there are different mechanisms in regulation of ENaC activity between stimulation of aldosterone and glucocorticoids. The activation of G-protein is required to maintain the activity of ENaC in the collecting ducts.

Keywords: ENaC; Aldosterone; Glucocorticoids;

1. INTRODUCTION

Active transport of Na⁺ across epithelial cells plays an important role in maintenance of homeostasis of electrolytes and water in the body. Hormone sensitive Na⁺ re-absorption is mainly mediated by epithelial Na⁺ channels (ENaC), which are distributed in distal and collecting nephron ducts, and alveolar epithelial cells. Disorder of ENaC could lead to Na⁺ and water retention [1], ‘dry lung’ and even death due to pulmonary edema [2]. Nevertheless, ENaC activities are strictly under the control of a variety of hormones like aldosterone, ADH, and glucocorticoids.

Aldosterone is known to enhance the density of ENaC on apical membrane of epithelial cells by modifying gene expression. Several aldosterone sensitive genes including serum and glucocorticoid (GC) inducible kinase (SGK-1) [3,4], PI3-K [5] have implicated the action mechanism of GC. Previously in vivo and in vitro studies in a number of models show that endogenous and exogenous GC could enhance expression of ENaC and increase the Na⁺ uptake in airway [6-13], gut [14] and kidney [6,15]. The regulation of ENaC by GC determines alveolar fluid clearance and lung development [9]. However, it still remains unclear of the mechanism in regulation of ENaC by GC, even by aldosterone. A slight difference in the expression of ENaC subunits has been observed in some studies. The expression of α subunit of ENaC is the most pronounced in lung, correlating with the high GC level, whereas expression of α, β and γ subunits of ENaC is gradually increased in kidney and colon [9]. Aldosterone is the major regulator in kidney, although GC could also exert the effects on ENaC expression in collecting ducts [6,15]. Till today it is still less clear whether GC and aldosterone exert the same effects on ENaC. In this study, we employed the electrophysiology methods to examine ENaC activity in cells stimulated with either dexamethasone (dex) or aldosterone.
2. MATERIALS AND METHODS

2.1. Cell Culture

M1 cells (mouse kidney cortical collecting duct cells) were purchased from European Collection of Cell Cultures at 21\textsuperscript{st} passages. Cells were grown in the medium containing DME: Ham’s F12 medium (1:1) (Sigma), 2 mM glutamine (Gibco), 5 µM dexamethasone (Sigma), 5% FBS (Sigma) in a 5% CO\textsubscript{2} and 37 ºC incubator. 1.5 µM aldosterone (Sigma) was added into the culture medium instead of 5 µM dexamethasone. When cells in the culture flasks reached 70% confluence, cells were seeded in low density to either coverslips or culture inserts (BD). SKF-525A was added in the culture medium containing dex 8 or 24 hours prior to experiments. Pertussis toxin (PTX) was prepared according to previous description \cite{16} and manufacture instruction. Pertussis toxin was added in the culture medium containing aldosterone in final concentration 300 ng/ml 18 hours prior to experiments.

2.2. Single Channel Patch Clamp Recording

Cell-attached single channel recording was performed as previously described \cite{17}. Briefly, M1 cells on the coverslip or insert were transferred into the recording chamber mounted on a Nikon inverted microscope (Nikon TE 2000U). Patch pipettes with a tip resistance of 7 M\textOmega were fabricated from borosilicate glass capillary (1.5 od, 0.86 id) (Warner) on a Sutter Puller (P97). Bath solutions contained (in mM): 110 NaCl, 4.5 KCl, 1 MgCl\textsubscript{2}, 1 CaCl\textsubscript{2}, 5 HEPEs, 5 Na-HEPEs, pH 7.2. Pipette solution contained (in mM): 110 NaCl, 4.5 KCl, 0.1 EGTA, 5 HEPEs, 5 Na-HEPEs, pH 7.2. Single channel currents were recorded with an Axon 1D amplifier and Axon clampex 9.0. The recording was analyzed by axon clampfit 9.0. Due to the variance of channel open probability, the first 2-5 minute single channel recording in normal bath medium was used as the control and NPo of ENaC during the period of applying chemicals was directly compared with NPo of the control. Data is presented as means ± S.E.M., and statistical differences were compared using ANOVA, taking P < 0.05 as significant and represented as *.

2.3. Chemicals

The following chemicals were used: dexamethasone(Sigma), aldosterone (Sigma), SKF-525A (Biomol), pertussis toxin (PTX) (Sigma). Most chemicals which dissolved in ethanol were made up as 1000 to 5000 times stock. Preparation of PTX was following the previous description and manufacture instruction. All chemical solutions were made as required on the day of experiments. The solvent, ethanol and DMSO at the same dilution, was tested alone in controls and had no effect.

3. RESULTS

3.1. Identification of ENaC Currents

ENaC currents were identified in either inside-out or outside-out recording by the unique ENaC channel conductance, sensitivity to amiloride and channel kinetics. Figure 1a represented the example currents recorded in

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**Figure 1.** ENaC currents obtained in single channel recordings from excised membrane. a. In an inside-out recording, small conductance inward currents were recorded, when pipette voltage was held at +100 mV and +60 mV, respectively. b. In an outside-out recording, small conductance inward currents were monitored, when pipette voltage was held at -60 mV. Bath application of 5 µM amiloride almost abolished these currents. The currents were reversed, when bath amiloride was washed off. c. Currents corresponding to the voltages from inside-out and outside-out recordings were plotted. The points were fitted by a linear line.

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**Figure 2.** ENaC currents obtained in cell-attach recordings. a. In a cell-attach recording, inward currents were recorded, when pipette voltage was held at 0 mV and +40 mV, respectively. b. The figure represented multiple step currents recorded in cell-attach recording. c. The conventional histogram shows the current events against the current size. d. The I-V curve is obtained from currents in cell-attach recordings.
an inside-out recording. Currents were elicited when pipette voltages (Vp) were held at +100 mV and +60 mV, respectively. In an outside-out recording, the small conductance currents possessed conductance similar to those obtained in inside-out recording. These currents were reversely sensitive to amiloride in the range of micromolar (Figure 1b). I-V curves constructed by currents obtained in either inside-out recording (n=50) or outside-out recording (n=20) were almost overlapped (Figure 1c) and possessed the same conductance ($5.1 \text{ pS}$, fitted by linear line, $r^2=0.99$). Figure 2a and Figure 2b represented the currents obtained in the cell-attach recordings. Currents in Figure 2b contained a multiple transition status. I-V curve (Figure 2d) constructed by currents in cell-attach recordings possessed the similar conductance ($5.0 \text{ pS}$, fitted by linear line, $r^2=0.99$). Therefore, we concluded that currents detected in single channel recording in M1 cells were mediated by ENaC channel.

3.2. Different ENaC Activities in M1 Cells Treated with Either Dexmethasone (Dex) or Aldosterone

There was no significant difference in cell morphology of M1 cells by different treatments with either dex or aldosterone. Electrophysiology experiments were performed, when cells formed a monolayer. As a standard protocol, cell-attach recording was carried out for 5-10 minutes in each cell prior to single channel recording from the excised membrane including inside-out recording and outside-out recording. In cells treated with dex, 325 cells exhibited ENaC currents in either inside-out or outside-out recording, but only 19 of 325 cells possessed the detectable current transition of ENaC channel during 5-minute of cell-attach recording (Table 1). However, in cells treated with aldosterone, 150 cells exhibited ENaC currents in either inside-out or outside-out recording, and 60 of 150 cells possessed detectable current transition of ENaC channel during 5-minute of cell-attach recording. Based on our observation, a large number of cells contained ENaC channels but they did not mediate Na+ currents, when cells were in the intact condition. About 5.84% of cells treated with dex could mediate Na+ currents, whereas 40% of cells treated with aldosterone could mediate Na+ currents, although these cells contained the ENaC in apical side of membrane. However, cells possessed low active ENaC ($p<0.05$, in comparing with aldosterone treatment group) on the apical membrane when cells were incubated with aldosterone and PTX (300 ng/ml) 18 hours prior to experiments.

3.3. Rundown of ENaC Currents in Cells Treated with Either Dex or Aldosterone

Previous studies have demonstrated that ENaC currents in A6 [18,19] were stable for the initial 4 minutes and exhibited a sudden rundown during the period from the 5th to 10th minute. In cells treated with dex, ENaC currents
Table 1. Active ENaC detected in different single channel configurations and cells with different treatment. In the cells treated with dex, 325 cells possessed ENaC activity in inside-out recordings but only 19 out of 325 cells possessed ENaC activity in cell-attach recordings. In cells treated with aldosterone, 150 cells possessed ENaC activity in inside-out recordings and 60 out of 150 cells possessed ENaC activity in cell-attach recordings. Deforming cells by EGTA enhanced the ENaC activity in cells treated with dex. SKF-525A did not significantly affect ENaC activity in cells treated with dex. PTX significantly reduced the number of active ENaC in intact cells.

<table>
<thead>
<tr>
<th>Number of cells detected ENaC</th>
<th>Dex</th>
<th>Aldosterone</th>
<th>Dex+EGTA</th>
<th>Dex+SKF 525A</th>
<th>Aldosterone+PTX</th>
</tr>
</thead>
<tbody>
<tr>
<td>Inside out recording Or outside recording (EC)</td>
<td>325</td>
<td>150</td>
<td>80</td>
<td>62</td>
<td>75</td>
</tr>
<tr>
<td>Cell-attach recording (CA)</td>
<td>19</td>
<td>60</td>
<td>15</td>
<td>4</td>
<td>1</td>
</tr>
<tr>
<td>Percentage % (CA/EC)</td>
<td>5.84%</td>
<td>40%</td>
<td>18.8%</td>
<td>6.45%</td>
<td>1.3%</td>
</tr>
</tbody>
</table>

from most cells maintained fairly constant open probability (Po) for over 10 minutes, since formation of inside-out configuration (Figure 3a). However, in cells treated with aldosterone, ENaC currents were stable for the initial 3-5 minutes and possessed a sudden rundown from then on (Figure 3b and Figure 3c).

3.4. Enhancement of Na\(^+\) Currents in Dex Treated Cells

Although the apical side membrane of M1 cells contained ENaC channels, only 5.84% of cells treated with Dex could mediate Na\(^+\) currents. SKF-525A, a general antagonist of CYP, was then used to test whether the low activity of ENaC was due to 11,12-EET [20,21]. M1 cells were incubated with SKF-525A (60 µM) for 8 hours and even 24 hours. Activity of ENaC was then determined in the inside-out recording from 62 cells. About 4 out of 62 cells possessed the ENaC currents in the cell-attach recording (Table 1). There was no significant difference, when compared to cells treated with dex. Bath EGTA was then used to stretch the membrane and rearrange the cytoskeleton (Figure 4e and Figure 4f), which may affect ENaC activities. After the treatment of EGTA for 10 minutes, the cell morphology displayed change within monolayer. The change always occurred in a patch. The patch was in the shape of a circle. Cells in the patch lost the flat structure and became swollen (Figure 4e) with dome. Cells in the right side of Figure 4f were still flat and tight, but cells on the left side were pulled by EGTA. After the wash off EGTA, single channel recordings were then performed. ENaC currents were detected in the inside-out or outside-out recordings from 80 cells. About 15 out of 80 cells possessed the ENaC currents in the cell-attach recording.

3.5. The Linkage Between Cell Morphology and Enac Detection

The common morphology of M1 cells from a monolayer could be summarized into 7 shapes as shown in Figures 4a-4d. Type 1 was flat and could be found in a very flat part of the monolayer. Types 2, 3, 5 and 6 had recognizable surface domes. Type 4 also had a reasonable dome and formed a circle with other similar shape cells. Type 7 was constituted by round cells found on the joint of flat patches. The percentage to detect ENaC currents in Type 1 was 3% (3 out of 100), as compared to over 60% (90 out of 150) in Type 4. In addition, before cells formed the monolayer, there were the patterns as shown in Figure 5a and Figure 5b with high probability to detect ENaC currents over 60% (12 out of 20).

4. DISCUSSION

Dex and aldosterone were suggested to stimulate the activity and expression of ENaC in renal and pulmonary systems. In this study, we showed that both dex and aldosterone could allocate ENaC on the apical membrane. However, these ENaC channels mediating Na\(^+\) entry depended on the treatment on the cells. Only 5.84% of them in cells treated with dex could permeate Na\(^+\) entry, as compared to over 40% of them in cells treated with aldosterone. ENaC in cells treated with aldosterone possessed a quick rundown. The low percentage of ENaC permeating Na\(^+\) entry in cells treated by dex was not related to inhibition by 11,12-EET and likely due to activation of G-protein and how channels were anchored on the membrane. Deformation of cytoskeleton of cells could enhance the percentage of ENaC to permeate Na\(^+\) entry in intact cells.

In addition to the regulation on the ENaC transcription, hormones like dex, aldosterone and epidermal growth factor [22] exert the acute mediation on ENaC activity. These acute effects were revealed to link with the cellular level of PIP2. Anionic phospholipids such as PIP2 [18,19,23] and PIP3 [24-27], located in the inner leaflet of plasma membrane, were suggested to regulate the activity of ENaC [26]. The negatively charged head group of PIP2 or PIP3 could directly interact with the positive charged cytoplasmic parts in β and/or γ subunits [19,24,26,28,29] of ENaC to exert the regulation. Evidences further indi-
activated that aldosterone elevates the cellular concentration of PIP3 by activating PI-3-kinase to lock the cytoplasmic termini of ENaC to the inner surface of plasma membrane [3] and/or pull the channel to activate ENaC [25]. Aldosterone-induced protein K-Ras localizes PI3-K near ENaC. The spatial organization of phosphatidylinositolides and specific phospholipid precursors [30,31] within cellular-membrane might determine the turnover of ENaC to allow Na⁺ currents [32]. Therefore, the activation of PI3-K and spatial organization of phosphatidylinositolides on the apical membrane by aldosterone might explain the active ENaC in the cells treated with aldosterone, but not dex, suggesting the presence of another mechanism.

Degeneration of PIP2 by PLC [33] could lead to unlocking of cytoplasmic termini of ENaC [25,33], resulting in decreased ENaC activity. It was manifested as the rundown of ENaC in excise membrane recording. Rundown of ENaC currents in excise membrane recordings reflected the simultaneous/continuous activation of PLC in a platform interacting with PIP2 and ENaC. Such platforms were revealed in G protein activated inward rectifying K⁺ channel [34], trp channel [35] and suggested to locate in β and γ subunits of ENaC [19]. Moreover, PLC-γ via receptor tyrosine kinase [22] and PLC-β via Gα11 [33,36] were demonstrated to mediate the degeneration of PIP2 in epithelial cells. However, the effects of PIP2 on ENaC activity are suggested to be permissive rather than regulatory [26]. Declines in PIP2 levels are parallel to the loss of ENaC activity, whereas addition of exogenous PIP2 rarely causes ENaC open probability to exceed the control level in excise membrane recording [36]. Increase in ENaC open probability by addition of PIP2 with GTP in inside-out recordings [19] implicates that the activation of G-protein plays an important role in regulating ENaC activity. Decrease in ENaC open probability by dialysis of intracellular GTP, which is not dependent on PI-3 K [37], further supported this hypothesis. We, therefore, proposed that activation of G-protein in cells treated with aldosterone might be the key to keep active ENaC on the membrane. Activation of G-protein could activate PLC, resulting in PIP2 degeneration. The rapid rundown of ENaC in cells treated with aldosterone might reflect activation of G-protein. In intact cells, the degeneration and resynthesis of phosphatidylinositolides reach a balance to maintain the lipid component of the membrane structure [38,39], but the lack of resynthesis of PIP2 pathway in excised membrane recording causes the decline of PIP2, resulting in rundown of ENaC. Like aldosterone, dex too distributes ENaC on the apical membrane via a similar mechanism as aldosterone via SGK-1 [3,4,7,40,41]. It is demonstrated by the ENaC activity detected in inside-out recordings in cells treated with dex. The low active ENaC in intact cells treated with dex and constant ENaC open probability in inside-out recording might be attributed to the low activity of G-protein. The linkage between aldosterone and activation of G-protein could be implicated by experiments that prevention of ATP release due to stimulation of aldosterone abolishes the aldosterone action [17]. Our observation that PTX incubation significantly reverses the effect of aldosterone on active ENaC in intact cells is consistent to previous report [42] and further supports our hypothesis.

11,12 - EET, a metabolite of AA via CYP epoxygenase, was demonstrated to mediate the direct inhibition of ENaC [20]. The evidence that EET mediates the inhibition effect of adenosine [21] on ENaC indicates a profound pathway to regulate the cellular EET level. The other rational explanation to low active ENaC in cells treated with dex is due to presence of cellular inhibitor of ENaC. We, therefore, applied SKF-525A, a general antagonist for CYP, to reduce the cellular EET level. There was no significant difference in the ENaC activity in intact cells, suggesting that EET is not the case to respond to low active ENaC.

It is widely accepted that ENaC activity is strongly regulated by cytoskeleton elements [43-45]. Recent evidence that activation of P2 receptors in epithelial basolateral membrane could deform the cells and eventually activate ENaC addressed the classic observation that aldosterone action on Na⁺ reabsorption relies on ATP production. Cell cytoskeletons could be deformed by many methods like ATP stimulation and EGTA. Our previous work demonstrated that elevation in [Ca²⁺], could reduce ENaC open probability, but there was no significant difference in ENaC activity, when 500 nM and 0 nM Ca²⁺ were in cytoplasmic medium. EGTA was, therefore, used to deform the cell. A significant increase in ENaC activity in cells treated with dex demonstrated that elements of cytoskeleton regulate ENaC activity. It also explains that ENaC activity is commonly observed in the excised membrane recording. However, the mechanism remains unclear. Consistent to previous observation [11], ENaC channels are in high density and distributed in the cells containing splitting pattern, suggesting that ENaC might involve cell differentiation and proliferation. The recognized dome in some cells, which contain ENaC on apical membrane, might be formed due to Na⁺ entry via ENaC with subsequent water entry.

In conclusion, aldosterone and glucocorticoids could increase the expression of ENaC and distribute ENaC on the apical membrane of cortical collecting duct cells. Low active ENaC is found in cells treated with dex. Activation of G-protein is required to keep the channel active. Aldosterone could lead to activation of G-protein to therefore possess high ENaC activity in intact cells.

REFERENCES


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