Epithelial Na$^+$ channels (ENaCs) control extracellular fluid volume by facilitating Na$^+$ absorption across transporting epithelia. In vitro studies showed that Cys-palmitoylation of the γENaC subunit is a major regulator of channel activity. We tested whether γ subunit palmitoylation sites are necessary for channel function in vivo by generating mice lacking the palmitoylated cysteines (γC33A,C41A) using CRISPR/Cas9 technology. ENaCs in dissected kidney tubules from γC33A,C41A mice had reduced open probability compared with wild-type (WT) littermates maintained on either standard or Na$^+$-deficient diets. Male mutant mice also had higher aldosterone levels than WT littermates following Na$^+$ restriction. However, γC33A,C41A mice did not have reduced amiloride-sensitive Na$^+$ currents in the distal colon or benzamil-induced natriuresis compared to WT mice. We identified a second, larger conductance cation channel in the distal nephron with biophysical properties distinct from ENaC. The activity of this channel was higher in Na$^+$-restricted γC33A,C41A versus WT mice and was blocked by benzamil, providing a possible compensatory mechanism for reduced prototypic ENaC function.

We conclude that γ subunit palmitoylation sites are required for prototypic ENaC activity in vivo but are not necessary for amiloride/benzamil-sensitive Na$^+$ transport in the distal nephron or colon.
Mice lacking γENaC palmitoylation sites maintain benzamil-sensitive Na⁺ transport despite reduced channel activity

Andrew J. Nickerson, Stephanie M. Mutchler, Shaohu Sheng, Natalie A. Cox, Evan C. Ray, Ossama B. Kashlan, Marcelo D. Carattino, Allison L. Marciszyn, Aaliyah Winfrey, Sebastien Gingras, Annet Kirabo, Rebecca P. Hughey, and Thomas R. Kleyman

Department of Medicine, Department of Computational and Systems Biology, Department of Cell Biology, and Department of Pharmacology and Chemical Biology, University of Pittsburgh, Pittsburgh, Pennsylvania, USA.

Epithelial Na⁺ channels (ENaCs) control extracellular fluid volume by facilitating Na⁺ absorption across transporting epithelia. In vitro studies showed that Cys-palmitoylation of the γENaC subunit is a major regulator of channel activity. We tested whether γ subunit palmitoylation sites are necessary for channel function in vivo by generating mice lacking the palmitoylated cysteines (γC33A,C41A) using CRISPR/Cas9 technology. ENaCs in dissected kidney tubules from γC33A,C41A mice had reduced open probability compared with wild-type (WT) littermates maintained on either standard or Na⁺-deficient diets. Male mutant mice also had higher aldosterone levels than WT littermates following Na⁺ restriction. However, γC33A,C41A mice did not have reduced amiloride-sensitive Na⁺ currents in the distal colon or benzamil-induced natriuresis compared to WT mice. We identified a second, larger conductance cation channel in the distal nephron with biophysical properties distinct from ENaC. The activity of this channel was higher in Na⁺-restricted γC33A,C41A versus WT mice and was blocked by benzamil, providing a possible compensatory mechanism for reduced prototypic ENaC function. We conclude that γ subunit palmitoylation sites are required for prototypic ENaC activity in vivo but are not necessary for amiloride/benzamil-sensitive Na⁺ transport in the distal nephron or colon.

Introduction

Na⁺ transport in the aldosterone-sensitive distal nephron (ASDN) is a primary determinant of extracellular fluid volume and blood pressure (1, 2). Epithelial Na⁺ channel–mediated (ENaC-mediated) transport in principal cells of the ASDN is one of the most highly regulated renal Na⁺ handling mechanisms (3–7), controlled mainly by volume-regulatory hormones such as aldosterone (8), vasopressin (9), and angiotensin II (10). ENaC activity also facilitates K⁺ secretion in the ASDN by providing the depolarizing force necessary for K⁺ efflux into the tubule lumen (11). Specific loss-of-function ENaC mutations cause type 1 pseudo-hypoaldosteronism (PHA-1), a disorder of renal salt wasting and hyperkalemia (12–14). Gain-of-function mutations have been described in patients with Liddle syndrome, where reduced rates of ENaC degradation lead to early-onset hypertension and hypokalemia (15, 16). Other ENaC gain-of-function variants may also contribute to salt-sensitive hypertension (17–22).

ENaCs are heterotrimeric (αβγ), amiloride-sensitive channels that are regulated by several mechanisms (23). Synthesis and apical membrane trafficking of channel subunits are both aldosterone-regulated events that increase the number of channels (N) at the cell surface (24, 25). Other factors that increase ENaC open probability (Pₒ) include proteolytic cleavage of the α and γ subunits (26–28), binding of acidic phospholipids such as phosphatidylinositol 4,5-biphosphate (29–31), extracellular Na⁺ and H⁺ (32–34), shear stress (35), and Cys-palmitoylation (36–38).

Cys-palmitoylation is the reversible attachment of a 16-carbon acyl group (palmitate) to intracellular cysteines. ENaC β and γ subunits were shown to be palmitoylated, which increases the channel’s Pₒ (36, 37). Palmitoylation of the γ subunit at 2 residues in the intracellular N-terminus (γC33 and γC41) was necessary for normal
channel activity in vitro (37). Channels lacking either palmitoylation site had substantially reduced $P_o$ but no decrease in surface expression of $\alpha$ or $\gamma$ subunits. Coexpression with a palmitoyltransferase (DHHC2) enhances currents from wild-type (WT) but not palmitoylation site-deficient channels ($\gamma^{C33A,C41A}$) expressed in *Xenopus* oocytes. Accordingly, catalytically dead DHHC2 did not increase WT currents. Inhibition of endogenous palmitoyltransferase activity using 2-bromopalmitate reduces amiloride-sensitive ENaC currents in cultured mouse cortical collecting duct cells (37). Specific DHHC proteins co-immunoprecipitate with ENaC and increase currents in cultured cells, and at least 1 isoform (DHHC3) colocalizes with ENaC in mouse kidney (38). However, the physiological importance of ENaC regulation by palmitoylation in vivo remains unknown.

We developed a mouse model lacking $\gamma$ subunit palmitoylation sites ($\gamma^{C33A,C41A}$) using CRISPR/Cas9 technology to study the role of this mechanism in regulating ENaC in vivo (Figure 1). Using electrophysiological approaches, we assessed ENaC function in the kidneys and distal colons of WT and $\gamma^{C33A,C41A}$ littermates under normal and Na⁺-depleted conditions. We also assessed ENaC subunit expression and processing, as well as systemic electrolyte handling via metabolic cage analyses. Our results demonstrate that $\gamma$ subunit palmitoylation sites, $\gamma$C33 and $\gamma$C41, are required for channel activity in vivo. However, salt and fluid balance can still be maintained, likely because of other benzamil-sensitive transport mechanisms.

**Results**

$\gamma^{C33A,C41A}$ mice express ENaCs with reduced activity in the cortical connecting tubules/collection ducts but maintain normal blood electrolytes and basal metabolic parameters. We first assessed whether $\gamma^{C33A,C41A}$ mice exhibited phenotypic abnormalities at baseline due to possible deficits in ENaC-dependent Na⁺ absorption. Age-matched $\gamma^{C33A,C41A}$ mice and WT littermate controls (6–7 weeks) showed no differences in blood electrolytes or plasma aldosterone levels when maintained on a standard chow diet (Table 1). However, cell-attached patch clamp recordings from split-open cortical connecting tubules/collection ducts (CNT/CCDs) revealed that $\gamma^{C33A,C41A}$ mice expressed ENaCs with significantly reduced activity at baseline compared with WT controls. This was detected as a decrease in total channel activity ($NP_o$ measured as number of channels $\times P_o$), driven by a reduced $P_o$ without a change in the number of observed channels per patch (Figure 2, C–E). The very low apparent single-channel $P_o$ from $\gamma^{C33A,C41A}$ mice (Figure 2E) reduced the likelihood of all channels opening simultaneously, leading to a likely underestimation of $N$ and an overestimation of $P_o$. An obvious reduction in $NP_o$ was nonetheless apparent, indicating a decrease in total ENaC activity in the apical membranes of PCs from $\gamma^{C33A,C41A}$ compared with WT mice. This observation is consistent with published data from heterologous expression systems using channels that carry the same $\gamma$ subunit cysteine mutations (38). Estimated single-channel conductance was similar between genotypes (WT vs. $\gamma^{C33A,C41A}$: 7.3 ± 0.9 vs. 8.2 ± 0.7 pS with Li⁺ as the charge carrier; $P = 0.11$).

ENaC protein expression is not altered in $\gamma^{C33A,C41A}$ versus WT mouse kidney. Under basal conditions, we did not observe a change in either $\alpha$ or $\gamma$ subunit expression between WT and $\gamma^{C33A,C41A}$ mouse kidney homogenates (Figure 3; see complete unedited blots in the supplemental material). These results again comport with previous findings in Fischer rat thyroid cells or *Xenopus* oocytes. We did detect a small increase in the N-terminal cleavage fragment of the $\alpha$ subunit in $\gamma^{C33A,C41A}$ versus WT male mice (Figure 3D) but not female mice. These data hint that male mice may be more sensitive to the observed loss of ENaC function.

ENaC activity in the CNT/CCD remains low in $\gamma^{C33A,C41A}$ mice under dietary Na⁺ restriction. ENaC-mediated absorption in the ASDN has a minor role in Na⁺ and fluid homeostasis in mice under basal dietary conditions. However, during dietary Na⁺ restriction, elevated aldosterone levels dramatically enhance ENaC activity, which then becomes a more prominent component to tubular Na⁺ reabsorption (8, 25, 39). We therefore evaluated the effects of Na⁺ restriction on channel activity and systemic salt and fluid balance in these mice.

During 7 days on a low-Na⁺ diet (<0.01%), mutant mice maintained similar body weight and total body water composition compared to WT littermates (Figure 4, A and B), suggesting no global deficits in maintaining fluid balance. Blood electrolytes and metabolic parameters were also similar between groups following treatment (Table 2). However, plasma aldosterone levels were significantly higher in male $\gamma^{C33A,C41A}$ mice than in WT male controls (Figure 4C and Table 2), indicating these animals were more sensitive to Na⁺ restriction and required more aldosterone to maintain homeostasis. As with $\alpha$ subunit cleavage on a Na⁺-replete diet (Figure 3), this effect was not observed in females. Sex differences with respect to salt handling in the nephron have been reported (40) and are likely involved in the differing compensatory responses between male and females in this experiment.
Cell-attached recordings of PC apical membranes from WT mice maintained on low-Na⁺ diet revealed robust ENaC activity, as expected (Figure 5A). Under these conditions, NP was approximately 3 times higher compared with WT mice on a standard diet (1.3 ± 0.9 vs. 0.4 ± 0.4, P = 0.02 by unpaired t test; Figure 2C vs. Figure 5C). However, channel activity in γC33A,C41A mice remained very low, even after Na⁺ restriction (Figure 5, C–E). This suggests that channels lacking γ subunit N-terminal cysteines cannot be activated by elevated aldosterone signaling that is associated with dietary Na⁺ restriction. As mentioned above, the very low PO observed in mutant mice likely results in an underestimation of N in these experiments.

ENaC activity is also observed in the late distal convoluted tubule (DCT2) and early CNT (8). In the DCT2, relatively low levels of 11β-hydroxysteroid dehydrogenase expression permit mineralocorticoid receptor activation by glucocorticoids (i.e., corticosterone/cortisol), rather than mineralocorticoids (i.e., aldosterone) (41). ENaC activity in the DCT2/CNT is therefore robust under both normal and Na⁺-restricted conditions. Whether ENaC activation by palmitoylation is region specific within the ASDN is unclear. Because ENaC regulation differs between the DCT2 and more distal segments, we assessed whether augmented channel activity in the DCT2/CNT might compensate for the loss of function observed in CNT/CCDs from γC33A,C41A mice. Cell-attached recordings revealed that ENaC activity in the DCT2/CNT was also significantly reduced in γC33A,C41A versus WT mice (Figure 6). This indicates that ENaC function in γC33A,C41A mice is reduced throughout the ASDN and is not a region-specific phenomenon.

WT and γC33A,C41A mice have comparable γ subunit cleavage. One major consequence of aldosterone signaling in the distal nephron is ENaC γ subunit cleavage (42–44), which activates the channel by releasing a short inhibitory peptide located in the extracellular domain (28, 45). This event increases Pₚ in part, by rendering the channel insensitive to the inhibitory effect of extracellular Na⁺ (i.e., by removing Na⁺ self-inhibition) (46). We therefore investigated whether our mutation impaired γ subunit proteolysis in mice, as this could explain the reduced channel activity observed under high-aldosterone conditions.

We examined γ subunit cleavage by treating kidney lysates with peptide:N-glycosidase F (PNGase F) to deglycosylate the protein prior to SDS-PAGE, as previously described (47). This approach facilitates visualization of the uncleaved subunit protein (~70 kDa), subunits processed at the proximal furin cleavage site (~55 kDa), and subunits processed by other proteases at a distal cleavage site (~50 kDa). The 50 kDa band likely reflects twice-cleaved γ subunits corresponding to a fully activated form of the channel.

Figure 1. Schematic for the γC33A,C41A mouse genotype. (A) Amino acid (upper) and nucleotide (lower) partial sequences of mouse γ subunit’s N-terminal region. CRISPR-mediated cysteine to alanine mutations are shown in red. (B) Genotyping of WT and γC33A,C41A mice, showing PCR products after digestion with MluI [see Methods for details]. DNA ladder and corresponding molecular weights are shown to the left. (C) Cartoon depicting the locations of putative palmitoylation sites, C33 and C41, within γ subunit’s N-terminal region (α helices within the γ subunit are depicted in red; α and β subunits are translucent).
of γ subunit cleavage were identical between WT and γC33A,C41A mice under Na+-restricted conditions, as were the overall levels of γ subunit expression (Figure 7). This suggests that the decreased ENaC activity observed in γC33A,C41A mice reflects a reduction in P0, that is independent of channel proteolysis.

WT and γC33A,C41A mice have similar expression of upstream Na+ transport proteins, NCC and NKCC2. Given that ENaC activity in cell-attached patch recordings was remarkably low in γC33A,C41A mice even under Na+-restricted conditions, we assessed whether Na+ transport mechanisms in other upstream nephron segments may be altered in order to compensate for loss of ENaC function. We measured the expression of 2 other critical Na+ transporters, NCC and NKCC2, which mediate electroneutral Na+ reabsorption in the distal convoluted tubule and thick ascending limb, respectively. Western blot analysis revealed no differences in the expression of either total NCC protein or its active phosphorylated form (phosphothreonine at position 53) between WT and γC33A,C41A mice of either sex (Figure 8, A–D). NKCC2 expression was also similar between WT and γC33A,C41A mice (Figure 8, E and F). These data indicate that γC33A,C41A mice do not require enhanced Na+ transport in upstream segments to maintain homeostasis, despite substantial loss of ENaC function.

Interestingly, NCC and NKCC2 co-transporters were both more highly expressed in female versus male mice, regardless of genotype (Figure 8, D and F). This suggests that females may rely more heavily on Na+ reabsorption in the distal convoluted tubule and/or the thick ascending limb under Na+-restricted conditions. This finding may also explain, at least in part, the lower aldosterone levels measured in female versus male mice after maintenance on a low-Na+ diet (Figure 4C).

WT and ENaCγC33A,C41A mice display robust benzamil-sensitive natriuresis. To evaluate the consequences of diminished ENaC activity on renal Na+ handling at the whole-organ level, we used metabolic cages to measure urinary volume and electrolyte composition before and during dietary Na+ restriction, as well as in response to selective ENaC blockade. Urinary Na+ excretion (UNaV) was similar between the groups at baseline and progressively diminished at similar rates during Na+ restriction (Figure 10C). Urinary K+ excretion (UKV) was also similar between genotypes throughout the experiment (Figure 10D). Likewise, body weight (Figure 10A) and effects on urine output over time (Figure 10B) were similar between groups. These data are consistent with the strong antidiuretic and antinatriuretic responses typical of dietary Na+ depletion.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>WT</th>
<th>n</th>
<th>γC33A,C41A</th>
<th>n</th>
<th>p</th>
</tr>
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<tbody>
<tr>
<td>Na+ (mM)</td>
<td>143 ± 1.4</td>
<td>12</td>
<td>143 ± 2.2</td>
<td>12</td>
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<tr>
<td>K+ (mM)</td>
<td>4.4 ± 0.3</td>
<td>12</td>
<td>4.7 ± 0.5</td>
<td>12</td>
<td>0.14</td>
</tr>
<tr>
<td>Cl− (mM)</td>
<td>112 ± 1.8</td>
<td>12</td>
<td>111 ± 4.6</td>
<td>12</td>
<td>0.42</td>
</tr>
<tr>
<td>Total CO2 (mM)</td>
<td>228 ± 1.1</td>
<td>12</td>
<td>23.8 ± 2.3</td>
<td>12</td>
<td>0.23</td>
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<tr>
<td>Glucose (mg/dL)</td>
<td>228 ± 17</td>
<td>12</td>
<td>220 ± 19</td>
<td>12</td>
<td>0.26</td>
</tr>
<tr>
<td>Blood urea nitrogen (mg/dL)</td>
<td>29.0 ± 6.8</td>
<td>12</td>
<td>28.5 ± 5.2</td>
<td>12</td>
<td>0.84</td>
</tr>
<tr>
<td>Hematocrit (% packed cell vol)</td>
<td>41.5 ± 1.6</td>
<td>11</td>
<td>41.6 ± 1.5</td>
<td>12</td>
<td>0.84</td>
</tr>
<tr>
<td>Hemoglobin (g/dL)</td>
<td>14.2 ± 0.6</td>
<td>12</td>
<td>14.1 ± 0.5</td>
<td>12</td>
<td>0.86</td>
</tr>
<tr>
<td>Anion gap (mM)</td>
<td>14.3 ± 1.3</td>
<td>12</td>
<td>13.5 ± 1.6</td>
<td>12</td>
<td>0.21</td>
</tr>
<tr>
<td>Aldosterone (pg/mL)</td>
<td>162 ± 54</td>
<td>10</td>
<td>179 ± 71</td>
<td>10</td>
<td>0.57</td>
</tr>
</tbody>
</table>

Metabolic parameters and aldosterone levels measured in whole blood and plasma, respectively, from male and female WT and γC33A,C41A mice. All mice were maintained on a standard chow diet. Data are listed as mean ± SD. Statistical comparisons were made via Student’s unpaired t test, with no significant differences detected.
In contrast with our patch clamp data, which clearly showed impaired ENaC function in γC33A,C41A mice, both groups responded to benzamil administration (1.5 mg/kg i.p.) with robust natriuresis (0.6 ± 0.2 mL for both groups) over a 3-hour period. Even more surprising were the greater natriuretic (Figure 10E) and chloruretic (Figure 10G) responses to benzamil in γC33A,C41A versus WT mice. These data demonstrate that benzamil-sensitive Na+ transport was not only present, but also greater, in γC33A,C41A mice than in WT littermates under these conditions. Kaliuresis after benzamil was similar between groups (Figure 10F).

Following the benzamil experiment, aldosterone levels in γC33A,C41A male mice may have been higher than WT (Figure 10H), although any difference did not reach significance (P = 0.06). However, aldosterone levels in both groups were substantially higher than previous experimental cohorts on a low-Na+ diet (Figure 4C and Figure 9B), which may reflect benzamil-induced diuresis and/or environmental stress induced by housing in the metabolic cages.

WT and γC33A,C41A mice both display robust amiloride-sensitive Na+ absorption in the distal colon. The distal colonic mucosa represents another major aldosterone-sensitive epithelium that participates in global electrolyte homeostasis (49–51). Here, aldosterone signaling induces the expression of ENaC β and γ subunits (α expression is constitutive), resulting in channel activity that can be readily measured as short-circuit current (Isc) in an Ussing chamber (52, 53). We found no differences between Na+-restricted WT and γC33A,C41A mice regarding their responses to another ENaC-selective blocker, amiloride (100 μM), when added to the mucosal bath (Figure 11, A and B). In distal colon, γ subunit expression was also similar between WT and γC33A,C41A mice under these conditions (Figure 11, C and D). Together, these data suggest that ENaC-dependent (amiloride/benzamil-sensitive) Na+ transport is readily observed in the kidneys and colons of γC33A,C41A mice, despite remarkably low ENaC activity in cell-attached recordings from isolated tubules.

WT and γC33A,C41A mice express a benzamil-sensitive 20 pS channel in the CNT/CCD. Our measurements of ENaC function at the single-channel level (split-open cell-attached recordings, Figures 2 and 5) and the whole-organ level (benzamil-induced natriuresis and amiloride-sensitive colonic Isc, Figures 10 and 11) are contradictory. Others have reported apical membrane, Na+-permeable channels in the distal nephron
as well as other ENaC-expressing epithelia (57–59), that display slow gating kinetics typical of the ENaC/Degenerin family. However, these channels are generally nonselective and usually have a unitary conductance of approximately 20 pS, much larger than that of typical ENaCs. In some cases, these nonselective channels were inhibited by amiloride with varying affinity (55).

We also observed a 20 ± 0.3 pS (n = 12) channel in PC apical membrane patches from both WT and γC33A,C41A mice (Figure 12, A, B, and F). Interestingly, the activity (NPO) of this channel was greater in Na+-restricted γC33A,C41A compared with WT mice (Figure 12C). Given the presence of an inward Cl− concentration gradient, current reversal at a holding potential of 0.1 ± 0.7 mV (n = 12) indicates that this is a nonselective cation channel. By comparison, the 8 pS channel displayed current reversal at 14 ± 2 mV holding potential (n = 13; P < 0.01 vs. 20 pS channel). Benzamil (50 μM) in the bath and patch pipette solutions decreased the likelihood of observing the 20 pS channel from 35% to 8% (P = 0.02; Mann-Whitney U test), as well as canonical 8 pS ENaC from 54% to 8% (P < 0.01; Mann-Whitney U test) (Figure 12G). Furthermore, when the 20 pS channel was observed in the presence of benzamil, its activity was low and diminished rapidly during the recording (example trace shown in Figure 12H), indicating that channel blockage increased with time. These data suggest that an apical membrane channel species, distinct from the canonical 8 pS conduc-

Figure 3. Baseline ENaC α and γ subunit expression is similar between WT and γC33A,C41A mice. (A and B) Representative Western blot images showing the detection of ENaC subunits in kidney homogenates from male (A) and female (B) WT and γC33A,C41A mice maintained on a standard chow diet. Standard molecular weight markers are shown in the far-left lane, along with corresponding weights, in kDa. Full-length α subunit, as well as the cleaved N-terminal fragment, are indicated by arrows and arrowheads, respectively. Stain-free gel images showing total protein content for each sample are shown below blots. (C–E) Densitometric quantification of full-length α subunit (C), cleaved α subunit (D), and γ subunit (E) was performed by first normalizing to the total protein signal for each sample, then transforming the data such that WT values were set equal to 1 (WT and γC33A,C41A males, n = 6; WT and γC33A,C41A females, n = 5). Lines and error bars represent mean ± SD. Comparisons were made via Student’s unpaired t test for each sex.
tance attributed to αβγ ENaC, may be involved in Na⁺ handling within the ASDN. This may also explain our finding of benzamil-sensitive natriuresis in γ C33A,C41A mice despite markedly reduced PO of prototypic ENaC.

Discussion

Previous studies established that palmitoylation of the γ subunit at C33 and C41 activates ENaC in vitro. Disrupting palmitoylation at these sites markedly reduces channel activity and prevents palmitoyltransferase-dependent channel activation when expressed in a heterologous system (37). To date, the broader physiological importance of this regulatory mechanism has been unclear. We now demonstrate that mice lacking these N-terminal cysteines (γ C33A,C41A mice) express ENaCs in the ASDN with greatly reduced activity compared with WT littermates.

Cell-attached recordings from split-open CNT/CCDs revealed that single-channel PO, as well as NP O, were significantly reduced in γ C33A,C41A compared with WT mice receiving a control diet (Figure 2). This finding is consistent with data from 2-electrode voltage clamp and patch clamp experiments using Xenopus oocytes expressing channels that carry the same mutations (37). We further observed that channel activity in γ C33A,C41A mice remained quite low under dietary Na⁺-restricted conditions (Figure 5), suggesting that a loss of function was not overcome by the effects of aldosterone.

Na⁺ restriction induced higher aldosterone levels in male γ C33A,C41A versus WT mice (Figure 4), which was likely a compensatory response to impaired ENaC function, although we did not observe a similar effect in females. Sex differences associated with epithelial ion transport regulation are known to play a major role in the adaptive responses to various dietary manipulations (40, 60–63). Increases in aldosterone levels in response to Na⁺ restriction are more modest in female versus male rats (64). We found that NCC and NKCC2 expression were approximately 50% greater in female versus male mice of either genotype (Figure 8), suggesting an enhanced capacity for Na⁺ reabsorption upstream of the CCD. Thus, females may require less aldosterone to maintain homeostasis, which would be consistent with the observed sex-dependent effect of Na⁺ restriction on aldosterone production between γ C33A,C41A and WT mice.

Patch clamp experiments showed that prototypic ENaC function was clearly diminished in γ C33A,C41A mice, despite no apparent changes in channel subunit expression or proteolysis (Figure 7). Because γ subunit cleavage at the distal site is thought to occur after channels traffic to the apical membrane (65), our biochemical findings further suggest that ENaC surface expression was similar in γ C33A,C41A and WT mice. These data accord with results from WT and γ C33A,C41A channels expressed in Xenopus oocytes (37). However, a direct assessment of ENaC surface expression (e.g., in vivo biotinylation of surface proteins in mouse tubules) was not performed. We therefore rest on the assumption that the abundance of double-cleaved γ ENaC approximates that of the surface-localized channel pool.

Surprisingly, when we examined Na⁺ transport at the whole-organ level, we found that WT and γ C33A,C41A mice both exhibited robust benzamil-induced natriuresis (Figure 10), as well as amiloride-sen-
sensitive $I_{SC}$ in the distal colon (Figure 11). Thus, the response to classical ENaC-selective blockers remains intact in $\gamma^{C33A,C41A}$ mice, despite a profound reduction in $P_{O}$ and no apparent increase in channel expression. Further, $K^+$ handling in response to acute or chronic $K^+$ loading was not impaired in $\gamma^{C33A,C41A}$ mice (Figure 9), indicating these animals possess a sufficient capacity for electrogenic Na$^+$/K$^+$ exchange in the ASDN. How can these conflicting observations be reconciled?

One possible explanation is that other benzamil/amiloride-sensitive transport mechanisms, aside from prototypic ENaCs, contribute to Na$^+$ reabsorption in the ASDN. We observed a second cation channel species in PC apical membranes that displays unitary conductance larger than that of ENaC (20 vs. 8 pS with Li$^+$ as the charge carrier) but was blocked by 50 $\mu$M benzamil (Figure 12). This 20 pS channel is likely nonselective, given its reversal potential of near 0 mV in cell-attached patches. However, opening of a nonselective cation channel at the PC apical membrane would still heavily favor Na$^+$ entry from the tubule lumen as a result of the attendant electrochemical driving forces. Blockage of such a channel could significantly impact Na$^+$ transport in the ASDN, especially considering the relatively high $P_{O}$ and unitary conductance, as well as the frequency at which these channels were observed.

Interestingly, the activity of this 20 pS channel was greater in Na$^+$-restricted $\gamma^{C33A,C41A}$ mice compared with WT littermates, raising the possibility that it may have a compensatory role when prototypic ENaC activity is compromised. We cannot determine the molecular identity of the 20 pS nonselective channel observed in our experiments based on the current data. The role of this channel in tubular Na$^+$ reabsorption therefore remains to be defined. However, future studies will be aimed at exploring the composition and physiological importance of this channel in greater detail.

Other ENaC-like, amiloride-sensitive cation channels with diverse biophysical characteristics have been described in transporting epithelia (54–56, 58). For example, alveolar epithelial cells express a nonselective cation channel species that, when compared with typical ENaCs, exhibits shorter open/closed times, is approximately 100 times less sensitive to amiloride, and has a larger unitary conductance of around 20 pS with Na$^+$ as the charge carrier (57). Eaton and colleagues provide evidence that these nonselective ion channels comprise ENaC$\alpha$ and acid-sensing ion channel (ASIC1a) subunits. ENaC$\alpha$/ASIC1a channels were further shown to have an important role in alveolar fluid clearance, which depends on vectorial ion transport.

In the kidney, Fila et al. recently demonstrated that a novel splice variant of ASIC2 is amiloride sensitive and contributes to Na$^+$ retention in a rodent model of nephrotic syndrome (66), a phenomenon that was previously considered to be largely mediated by ENaC (67–69). Other ~20 pS cation channels have also been documented in patch clamp recordings of native collecting duct PCs (70) and cultured collecting duct cell lines (71–73). In mouse CCDs, such a channel was observed in PC apical membrane patches from animals lacking primary cilia (70). Activity of this channel was increased in ciliopathic mice compared with WT littermates. This was concurrent with increased abundance of TRPV4 and TRPP3 mRNA, although the authors did not demonstrate a functional connection. Interestingly, TRPP3 is sensitive to amiloride and benzamil (74). TRPM4, another nonselective, 23 pS cation channel, was also observed in cell-attached patches from PC apical membranes of $\gamma^{C33A,C41A}$ mice. However, its functional significance remains to be determined.

### Table 2. Blood electrolytes and metabolic parameters in WT and $\gamma^{C33A,C41A}$ mice following dietary Na$^+$ restriction

<table>
<thead>
<tr>
<th>Parameter</th>
<th>WT</th>
<th>$n$</th>
<th>ENaC$\gamma^{C33A,C41A}$</th>
<th>$n$</th>
<th>$P$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Na$^+$ (mM)</td>
<td>144 ± 2.4</td>
<td>14</td>
<td>144 ± 0.9</td>
<td>15</td>
<td>0.93</td>
</tr>
<tr>
<td>K$^+$ (mM)</td>
<td>4.7 ± 0.4</td>
<td>14</td>
<td>4.4 ± 0.4</td>
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<td>0.15</td>
</tr>
<tr>
<td>Cl$^-$ (mM)</td>
<td>110 ± 2.0</td>
<td>14</td>
<td>111 ± 3.5</td>
<td>15</td>
<td>0.40</td>
</tr>
<tr>
<td>Total CO$_2$ (mM)</td>
<td>25.3 ± 1.9</td>
<td>12</td>
<td>24.0 ± 2.8</td>
<td>13</td>
<td>0.21</td>
</tr>
<tr>
<td>Glucose (mg/dL)</td>
<td>213 ± 23</td>
<td>9</td>
<td>207 ± 36</td>
<td>9</td>
<td>0.40</td>
</tr>
<tr>
<td>Blood urea nitrogen (mg/dL)</td>
<td>26.2 ± 2.5</td>
<td>9</td>
<td>24.0 ± 4.9</td>
<td>10</td>
<td>0.18</td>
</tr>
<tr>
<td>Hematocrit (% packed cell vol)</td>
<td>43.6 ± 1.7</td>
<td>9</td>
<td>42.0 ± 2.4</td>
<td>10</td>
<td>0.07</td>
</tr>
<tr>
<td>Hemoglobin (g/dL)</td>
<td>14.8 ± 0.6</td>
<td>9</td>
<td>14.3 ± 0.8</td>
<td>10</td>
<td>0.09</td>
</tr>
<tr>
<td>Anion gap (mM)</td>
<td>13.2 ± 2.5</td>
<td>9</td>
<td>12.6 ± 2.0</td>
<td>10</td>
<td>0.55</td>
</tr>
<tr>
<td>Aldosterone (pg/mL)</td>
<td>1,810 ± 690</td>
<td>14</td>
<td>1,985 ± 1,119</td>
<td>13</td>
<td>0.63</td>
</tr>
</tbody>
</table>

Metabolic parameters and aldosterone levels measured in whole blood from male and female WT and $\gamma^{C33A,C41A}$ mice after 8 days of maintenance on low-Na$^+$ diet. Statistical comparisons were made via Student’s unpaired t test, with no significant differences detected.
channel, was also identified in mpkCCD cell apical membrane patches (75). A functional role for the aforementioned TRP channels with respect to Na⁺ transport in the CCD has not yet been determined.

Our experimental model system is a constitutive disruption of known palmitoylation sites, rather than an inducible mechanism. Therefore, γC33A,C41A mice carry these mutations throughout development and postnatal maturation. Nephron development may be affected in these mutants to compensate for diminished ENaC function in the distal tubule. However, our analysis of transporter protein expression in kidney lysates indicates that distal convoluted tubule– or thick ascending limb–localized Na⁺ transport mechanisms are not upregulated in γC33A,C41A mice (Figure 8). If upstream tubule segment remodeling occurs in γC33A,C41A mice (e.g., hyperplasia or hypertrophy), such adaptations were not evident in these studies.

In agreement with in vitro studies, we observed that prototypic ENaCs in the CNT/CCDs of γC33A,C41A exhibited a dramatically reduced $P_O$ compared with WT littermates. While our results suggest that ENaC palmitoylation has an important role in activating the channel in vivo, we cannot provide direct evidence that the γ subunit is modified by palmitoylation in the kidney. We previously used fatty acid exchange chemistry to demonstrate palmitoylation of N- and C-terminally tagged β and γ subunits using heterologous expression systems (36, 37). To date, we have not been successful in using a similar approach to demonstrate ENaC palmitoylation in native (untagged) proteins.

A recently developed technique to detect palmitoylated cysteine residues involves replacing the acyl group with a large polyethylene glycol, resulting in an upward shift in the target protein when assayed by immunoblot (76). Our attempts at using this approach were also unsuccessful, likely because the palmitoylated cysteines in γENaC reside within the N-terminal segment, which is cleaved during channel processing. Due to the location of the epitope sequence targeted by the γENaC antibody used here (residues 629–650 in the C-terminus), the mature N-terminal fragment that contains the palmitoylated cysteines cannot be visualized by immunoblotting. We therefore cannot rule out the possibility that loss of γ subunit N-terminal cysteines confers a reduction in ENaC activity independent of palmitoylation. For example, Kellenberger et. al. suggested that intracellular cysteines regulate ENaC function through redox chemistry (77). Reducing or oxidizing reagents enhanced or inhibited ENaC activity in excised membrane patches, respectively. Collective mutation of all

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**Figure 5.** CNT/CCD ENaC activity in γC33A,C41A mice remains low under dietary Na⁺ restriction. (A and B) Representative recordings from cell-attached patches of PC apical membranes in split-open CNT/CCDs from WT (A) and γC33A,C41A mice (B) maintained on a low-Na⁺ (<0.01%) diet. Channel closed (C) and open (O) states are depicted to the right of each recording. (C–E) Total channel activity ($NPO$; C), number of observed channels per patch ($N$; D), and single-channel open probability ($P_O$; E) in apical membrane patches from WT ($n = 10$) and γC33A,C41A ($n = 11$) mouse collecting ducts. Data from male and female mice are shown as black and white circles, respectively. Lines and error bars represent mean ± SD. $NPO$, $N$, and $P_O$ data were analyzed via Student’s unpaired $t$ test.
intracellular cysteines in the α, β, and γ subunits attenuated these responses. We cannot exclude the possibility that a similar mechanism is at least partially responsible for the loss of function observed in γC33A,C41A mice.

Evaluating the consequences of a total loss of ENaC function is made difficult by the fact that constitutive knockout of ENaC subunits causes perinatal lethality due to impaired fluid clearance in the lung and severe salt wasting (78–80). Hummler and colleagues developed a series of inducible, nephron-specific ENaC-knockout mouse models as a strategy to circumvent this issue (81–84). Mice with induced γ subunit knockout exhibit a PHA-1–like phenotype with salt wasting, weight loss, hyperkalemia, and elevated aldosterone. This may reflect a downregulation on NCC as well as ENaC, given that urinary Na⁺ wasting and weight loss were not observed in mice maintained on a K⁺-deficient (<0.1% K⁺) diet (82, 85). Here, we show that γC33A,C41A mice display a profound loss of prototypic ENaC activity in the ASDN but have no overt phenotype. It is therefore unlikely that the γC33A,C41A mutation described here is functionally similar to a knockout of the γ subunit. Single-channel recordings from inducible γ subunit–knockout mice were not reported but may yield important insights into Na⁺ transport in the ASDN, including the possible involvement of other channels.

In conclusion, our study demonstrates that 2 key cysteine residues in the γ subunit N-terminus, γC33 and γC41, are required for robust ENaC activity in vivo. However, a benzamil-sensitive, nonselective cation channel in the ASDN may compensate for the loss of prototypic ENaC function caused by mutation of these residues (γC33A,C41A).

Methods
Animals. γC33A,C41A mice were generated on a 129SVE background using CRISPR/Cas9 technology, with the assistance of the mouse embryonic services core facility at the University of Pittsburgh Department of Immunology. Alanine substitutions at cysteines 33 and 41 (γC33A,C41A) were induced via homology-directed
repair (HDR). Embryos were injected with single-guide RNA (50 ng/µL; 5′-TAATACGACTCACTATAGG-CACGATGCCGGCCAGCCGTGTTTTAGAGCTAGAAATAGCA-3′), Cas9 mRNA (100 ng/µL), and single-stranded oligo DNA nucleotides for HDR (50 µM; 5′-AAGAATCTGCCAGTTCGAGGCCCCCAAGGCACCGACCATTAGGACCTGATGCTGGTACGCGTTAAACACCCACGGAGCTC-GGCACACGTTTCTTCGAGCCGGCATCGTGGTGTCCCGAGGCCGCCTTCGGCGCCTGTTGTGGATTGCGTTCACGCTG-3′).

Mlu1 restriction sites were introduced into γC33A,C41A mice near A33 such that digestion of PCR-amplified products yielded fragments of 395 bp in WT and 195/205 bp in mutant mice, respectively (Figure 1). Founder mice were genotyped by PCR (forward primer 5′-GACATAGGGCTGACACACCA-3′, reverse primer 5′-CCCACCAGTTTCTCGACTCA-3′) and confirmed by sequencing the amplification products. These mice were backcrossed against WT 129SVE mice for 3 generations before arranging heterozygous mating schemes for producing experimental cohorts. After establishing colonies, mice were moved to Charles River Laboratories for breeding and genotyping. Experimental cohorts were then shipped to the University of Pittsburgh for experiments and genotype confirmation.

**Blood electrolyte and aldosterone measurements.** Mice were anesthetized with isoflurane and blood was collected via left ventricular puncture or via retro-orbital sinus bleed. Whole blood was analyzed with an iStat analyzer (Abbott Laboratories) using chem8+ cartridges to gather electrolyte and metabolic parameters. For left ventricular punctures, remaining blood was centrifuged for 15 minutes at 1,500 g at 4°C to isolate the plasma. Plasma aldosterone was measured via ELISA (IBL-America IB79134) according to the manufacturer's instructions.

**Immunoblotting.** Kidney homogenates were prepared by lysing tissues in CellLytic MT lysis buffer (MilliporeSigma; 5% weight/volume) using a glass homogenizer. After determining protein concentrations for each lysate, 30 mg of total protein was resolved using 4%–15% TGX-Stain-free polyacrylamide

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**Figure 7.** Expression and cleavage pattern of the γ subunit is similar between Na+-restricted WT and γC33A,C41A mice. (A) Western blot image showing cleavage fragments of the γ subunit after treating samples with (+) or without (-) PNGase F to remove N-linked sugar residues. Standard molecular weight markers are shown in the far-left lane, along with corresponding weights, in kDa. Arrow indicates the full, uncleaved fragment. Single and double arrowheads indicate proximally (furin) and distally (presumably double) cleaved fragments, respectively. All samples were from male mice maintained for 8 days on a low-Na+ diet. (B) Stain-free gel image showing total protein content for each sample. Dark bands appearing in the stain-free gel image correspond to PNGase enzyme (~36 kDa) in the sample preparation. Densitometric quantification of full-length (C) and furin (D) distally cleaved γ subunit (E) was performed as described above (n = 4 for WT and γC33A,C41A mice). Lines and error bars represent mean ± SD. Statistical comparisons were made via Student’s unpaired t test.
Figure 8. Expression of upstream Na+ transport proteins is similar between Na+-restricted WT and γC33A,C41A mice but varies by sex. (A, C, and E) Western blot images showing expression of Thr-53 phosphorylated Na+,Cl– co-transporter (p-NCC; A), total NCC (C), and Na+, K+, 2Cl– co-transporter (NKCC2; E) proteins from whole-kidney lysates of WT and γC33A,C41A mice of both sexes (n = 4 for each group). Data from male and female mice are shown as black and white circles, respectively. Standard molecular weight markers are shown in the far-left lane, along with corresponding weights, in kDa. Stain-free gel images showing total protein content for each sample are shown below each blot. All mice were maintained for 8 days on a low-Na+ diet. (B, D, and F) Densitometric quantification of p-NCC (B), total NCC (D), and NKCC2 (E) was performed as described above. Lines and error bars represent mean ± SD. Statistical comparisons were made via 2-way ANOVA with Tukey’s post hoc analysis.

Figure 9. Responses of WT and γC33A,C41A mice to K+ loading. (A and B) Blood K+ (A) and plasma aldosterone levels (B) measured from WT and γC33A,C41A (n = 13 and 15, respectively for A; n = 10 and 11, respectively for B) male mice maintained on a 10% KCl diet for 8 days. Lines and error bars represent mean ± SD. Statistical comparisons were made via Student’s unpaired t test. (C) Blood K+ levels measured following an acute oral gavage of 150 μL 5% KCl solution in WT (n = 6) and γC33A,C41A (n = 6) mice maintained on a standard chow diet. Sequential measurements were taken at 30- and 60-minute time points postgavage. Statistical comparison was made with 2-way ANOVA with repeated measures. No difference between genotypes was observed.
gels (BioRad). For some experiments, N-linked glycosylation products were removed using PNGase F (New England Biolabs P0704) prior to gel loading, as per the manufacturer’s instructions. Uniform protein loading between samples was confirmed after electrophoresis by imaging the stain-free gels using a Chemidoc imager (BioRad), before transferring the resolved proteins to PVDF membranes (BioRad). After transfer, membranes were incubated in blocking buffer (5% milk dissolved in Tris-buffered saline with 0.1% Tween 20; TBST) for 2 hours at room temperature. Membranes were then incubated at 4°C overnight in blocking buffer containing primary antibodies against the ENaC \( \alpha \) subunit (gift from Johannes Loffing, University of Zurich, Zurich, Switzerland; 1:1,000; raised against mouse \( \alpha \) ENaC residues 2–21) (86), ENaC \( \gamma \) subunit (Stressmarq SPC-405; 1:1,000; raised against rat \( \gamma \) ENaC residues 629–650), NCC (David Ellison, Oregon Health & Science University, Portland, Oregon, USA; 1:5,000; raised against mouse NCC residues 1–98) (87), phospho-NCC (Phosphosolutions p1311-53, 1:5,000; raised against residues surrounding pThr53 in mouse NCC), or NKCC2 (Stressmarq SPC-401; 1,1000; raised against rat NKCC2 residues 33–55). The following day, membranes were washed 4 times, 10 minutes each, with TBST and incubated in blocking buffer containing HRP-conjugated secondary antibody (Thermo Fisher Scientific 31460; goat anti-rabbit, 1:10,000) for 1 hour at room temperature. Target proteins were detected by chemiluminescence using Clarity peroxide substrate reagent (BioRad). Blots were imaged with a Chemidoc imager, and protein abundance was quantified by measuring the signal intensity via ImageJ software (NIH) and normalizing to the total protein signal detected in each sample from corresponding stain-free gel images.

**Patch clamp electrophysiology.** Freshly isolated kidneys were sliced into approximately 1 mm sections and rinsed thoroughly in warm L-15 medium (Gibco). Using a stereomicroscope, small portions of the renal cortex were separated from the medulla and incubated in type II collagenase (MilliporeSigma C2-22; 1 mg/mL) for 40 minutes at 37°C. After incubation, tissue segments were rinsed 5 consecutive times in ice-cold L-15 medium to stop enzymatic digestion. CNT/CCDs were isolated using very fine needles under a stereomicroscope. CNT/CCD segments were identified based on the following spatial/morphological features: 1) being located just downstream of distal convolutions, 2) coarse appearance, 3) presence

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**Figure 10.** WT and \( \gamma^{C33A,C41A} \) mice exhibit robust benzamil-sensitive natriuresis following dietary \( Na^+ \) restriction. (A–D) Body weight (A), daily urine output (B) and urinary \( Na^+ \) (C) and \( K^+ \) excretion (D) recorded from male WT (squares; \( n = 6 \)) and \( \gamma^{C33A,C41A} \) mice (triangles; \( n = 6 \)) over the course of 4 days on a low-\( Na^+ \) diet. No statistical differences were detected for any parameter by 2-way ANOVA with repeated measures. (E–G) Urinary \( Na^+ \) (E), \( K^+ \) (F), and \( Cl^- \) (G) excretion from WT (\( n = 5 \)) and \( \gamma^{C33A,C41A} \) (\( n = 5 \)) mice in response to 1.5 mg/kg benzamil injection. Urines were collected over the first 3 hours following benzamil administration. (H) Plasma aldosterone levels measured in WT (\( n = 6 \)) and \( \gamma^{C33A,C41A} \) (\( n = 6 \)) mice following the benzamil injection experiment. Lines and error bars represent mean ± SD. Comparisons were made via 2-way ANOVA with repeated measures (A–D) or Student’s unpaired t test (E–H).
of branched segments, and 4) medium-sized tubule diameter (i.e., between that of the proximal tubule and loop of Henle segments). DCT2/CNTs were identified as the last third of the convoluted segment and the tubule areas upstream of branching points.

Tubules were mounted to poly-l-lysine–coated coverslips and transferred to a perfusion/recording chamber continuously perfused with physiological saline solution (in mM: 140 Na+, 4.5 K+, 2 Ca2+, 1.2 Mg2+, 150.9 Cl–, 5 glucose, 10 HEPES, 5 Tris, pH 7.4) at a rate of ~1 mL/min. Immobilized tubules were split open using sharpened micropipettes controlled by a micromanipulator (Sutter Instruments), to gain access to the cells’ apical membranes. PCs were identified based on the appearance of a small, flat apical membrane surface and a prominent nucleus. Apical membrane seals were made using micropipettes pulled from borosilicate glass (Sutter Instruments), polished to give a series resistance of 5–8 MΩ, and filled with LiCl solution (in mM: 140 Li+, 144 Cl–, 2 Mg2+, 10 HEPES, pH 7.4). High-resistance seals (>2 GΩ) were made by applying gentle suction to the pipette using a glass syringe. Membrane currents were recorded in a cell-attached configuration using a PC-One Patch Clamp amplifier (Dagan Corp.), operated by pClamp 10 software (Molecular Devices). Currents were recorded at a sampling frequency of 5 kHz and filtered at 100 Hz by a built-in 4-pole low-pass Bessel filter. All recordings were manually inspected and corrected for any baseline drift prior to analyzing channel transitions via single-channel event detection in Clampfit software. Channel transition events were then measured over a period of at least 2 minutes. All analyses were again visually inspected prior to inclusion of the data.

Body composition analysis. Total body water content in restrained mice was determined using a quantitative magnetic resonance imager (EchoMRI-100H). Measurements were performed in duplicate for each mouse during the course of dietary Na+ restriction. Total body water mass (including free water) was taken as a percentage of total body weight to yield percentage body water values. All measurements were taken in 24-hour intervals.

K+ gavage experiments. Mice maintained on standard chow were administrated 150 μL of a solution containing 5% KCl and 2% sucrose via gastric gavage. At 30- and 60-minute time points, mice were briefly

Figure 11. WT and γC33A,C41A mice exhibit amiloride-sensitive Na+ transport in the distal colon. (A) Representative short-circuit current (Isc) recordings of distal colons obtained from WT (black) and ENaCγC33A,C41A mice (red) maintained for 8 days on low-Na+ diet. Where indicated 100 μM amiloride was added to the apical chamber bath. (B) Amiloride-sensitive Isc measured in WT (n = 13) and γC33A,C41A (n = 12) mouse distal colons. Data from male and female mice are shown as black and white circles, respectively. (C) Representative Western blot images showing the detection of the γENaC subunit in colon mucosal homogenates from male WT (n = 4) and γC33A,C41A (n = 4) mice maintained on a low-Na+ diet for 8 days. Standard molecular weight markers are shown in the far-left lane, along with corresponding weights, in kDa. Stain-free gel image showing total protein content for each sample is also provided. (D) Densitometric quantitation of γ subunit abundance was performed as described above. Lines and error bars represent mean ± SD. Statistical comparisons were made via Student’s unpaired t test and no significant differences were detected.
anesthetized with 5% isoflurane, and blood samples were collected via retro-orbital bleed. Blood K⁺ values were measured using an iStat as described above.

Ussing chamber electrophysiology. Colons were harvested from anesthetized mice, and mucosa/submucosa preparations were made by blunt dissection. The colon was opened longitudinally along the mesenteric line and pinned down in a sylgard dish with the mucosal surface facing downward. Using a razor blade, a superficial lateral incision was made near the rectum, allowing separation of the mucosa/submucosa and muscularis propria. The tissue layers were then teased apart using fine forceps under a dissecting microscope. Tissues were mounted on custom-made sliders (aperture 0.3 cm²) for use in an Ussing-style recording chamber (Physiologic Instruments) and were bathed in Ringer’s solution containing the following, in mM: 140 Na⁺, 119.8 Cl⁻, 25 HCO₃⁻, 5.2 K⁺, 1.2 Ca²⁺, 1.2 Mg²⁺, 2.4 HPO₄²⁻, 0.4 H₂PO₄⁻, and 10 glucose. The solution was continuously bubbled with 5% CO₂ balanced with oxygen to maintain pH at 7.41 and gradually warmed to 37°C using a circulating heated water bath. Tetrodotoxin (0.5 μM) was added to the serosal bath to suppress residual neurogenic activity. Trans-epithelial voltage (Vᵣₑₚ) was monitored using a

Figure 12. Presence of a benzamil-sensitive, 20 pS apical membrane conductance in CNT/CCDs of both WT and γC₃₃A,C₄₁A mice. (A and B) Representative recordings from cell-attached patches of PC apical membranes in split-open CNT/CCDs from WT (A) and γC₃₃A,C₄₁A mice (B) maintained on a low-Na⁺ diet for 8 days. Channel “closed” (C) and “open” (O) states are depicted to the right of each recording. (C–E) Total channel activity (Nₚₒ, C), number of observed channels per patch (N, D), and single-channel open probability (Pₒ, E) in apical membrane patches from WT (n = 10) and γC₃₃A,C₄₁A (n = 12) mouse CNT/CCDs. Data from male and female mice are shown as black and white circles, respectively. (F) Current/voltage relationship of 8 pS and 20 pS channels observed in CNT/CCD apical membrane patches with Li⁺ in the patch pipette. (G) Frequency of channel appearances for both 8 and 20 pS conductance populations in the absence (left) or presence (right) of 50 μM benzamil in the bath and patch pipette solutions. (H) Representative recording of a 20 pS channel from a WT male mouse in the presence of benzamil, illustrating diminishing channel activity over time.
multichannel voltage clamp/amplifier (VCC MC6; Physiologic Instruments) until the tissues stabilized. $V_{Te}$ was then clamped to 0 mV using computer-operated software (pClamp 10, Molecular Devices) to measure $I_{sc}$. After currents stabilized, ENaC activity was measured as the change in $I_{sc}$ after addition of 100 μM amiloride to the apical bath.

**Metabolic cage experiments.** For baseline urine collections, mice were singly housed in metabolic cages (Techniplast) for an acclimation period of 3 days with free access to food and water. The standard pelleted diet was replaced with a gel mixture (pelleted diet blended with 3% agar) to prevent contamination of the urine by pellet debris. After acclimation, baseline urines were collected over 24 hours, and Na⁺ and K⁺ concentrations were measured with an EasyLyte analyzer (Medica). Total urine output (in milliliters) was also recorded for the 24-hour period, which was used to calculate rates of UNaV and UKV. For benzamil response experiments, mice were maintained on a low-Na⁺ gel diet for 5 days to enhance ENaC expression. This diet was prepared in the same manner as normal gel diet but with <0.01% Na⁺ chow (Teklad) instead of standard chow. Body weight, daily urine output, and UNaV/UKV were monitored over the treatment period. After 5 days of Na⁺ restriction, each mouse was injected i.p. with 1.5 mg/kg benzamil (dissolved in sterile saline + 0.1% DMSO), and urine was collected over the following 3 hours. Following the experiment, mice were euthanized, and blood was collected for aldosterone measurements.

**Statistics.** Methods of statistical analysis are described in detail in the text and/or figure legends. End-point comparisons between genotypes were made via unpaired 2-tailed Student’s $t$ test. Time course comparisons between genotypes were made via 2-way ANOVA with repeated measures. Sex- and genotype-dependent comparisons were made via 2-way ANOVA with Tukey’s post hoc. Comparison of observed channel appearances between treatment groups was performed via Mann-Whitney $U$ test. For all analyses, $P < 0.05$ was considered significant.

**Study approval.** Animal protocols were approved by the University of Pittsburgh Institutional Animal Care and Use Committee.

**Data availability.** All data presented in this manuscript are accessible in the Supporting Data Values (supplemental material available online with this article; https://doi.org/10.1172/jci.insight.172051DS1) XLS file or by request to the corresponding author.

**Author contributions**
AJN designed and performed experiments, analyzed data, and drafted and revised the manuscript. SMM performed experiments. SS assisted with mouse model generation, performed experiments, and edited the manuscript. NAC performed experiments and analyzed data. ECR assisted with mouse model generation and data analysis. OBK assisted with data analysis and edited the manuscript. ALM assisted with experiments and mouse breeding. AW assisted with mouse breeding. SG assisted with mouse model generation. AK assisted with data analysis and edited the manuscript. RPH assisted with data analysis and edited the manuscript. TRK conceived of the study, designed experiments, analyzed data, and revised the manuscript.

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Address correspondence to: Thomas Kleyman, A919 Scaife Hall, 3550 Terrace Street, Pittsburgh, Pennsylvania 15261, USA. Email: kleyman@pitt.edu.