INTRODUCTION

Hypertension is a major public health issue among industrialized nations, and several clinical and epidemiological studies have shown a clear link between dietary Na+ intake and blood pressure. With Na+ and K+ homeostasis being closely interlinked, a low Na+ and high K+ diet has been shown to lower blood pressure (Mente et al., 2014). Studies on monogenic syndromes such as Gordon’s (Familial Hyperkalaemia & Hypertension, http://omim.org/entry/145260) and Gitelman (OMIM http://omim.org/entry/263800) have shown that mutations in ion channels and transporters and/or their regulators such as WNK proteins in the nephron perturb the Na+/K+ homeostasis and cause significant changes in blood pressure. The distribution of these proteins along the nephron reflects the key importance of these nephron segments including the loop of Henlé (LoH), the distal convoluted tubule (DCT), and the contiguous segments of the connecting tubule (CNT) and...
collecting duct (CD) in the homeostatic process. The WNK/SPAK pathway regulates various ion transporters/channels in this segment of the nephron, and thus WNK kinase expression and activity is pivotal. In the DCT, WNKs are activated by sympathetic stimulation, cortisol, (Terker et al., 2014) low extracellular K⁺, (Terker et al., 2015) and low intracellular Cl⁻ (Piala et al., 2014; Terker, Zhang, et al., 2016), which affect its phosphorylation state that in turn leads to an activation of downstream targets. WNK kinases phosphorylate and activate SPAK (STE20/SPS1-related proline-alanine-rich protein kinase) and OSR1 (Oxidative Stress Responsive 1), which in turn bring about the coordinated phosphorylation of NCC and NKCC2 in the DCT and TAL of the nephron, respectively (Richardson & Alessi, 2008; Vitari et al., 2006). In the DCT, the WNKs are degraded by the proteasome pathway when they are targeted by a kelch-like protein 3 (KLHL3) mediated ubiquitylation by Cullin3 (Cul3) (Ohta et al., 2013; Schumacher et al., 2015).

Studies on the WNK/SPAK/NCC signaling pathway have elucidated the molecular link between WNK kinases and the regulation of blood pressure by their effect on renal electrolyte homeostasis, and how the aldosterone regulated the processes of Na⁺ absorption (from the urine to the blood) and K⁺ secretion (from the blood to the urine) work in concert in the distal nephron (Kahle et al., 2003). Although the benefits of a high-K diet in lowering blood pressure has been well-established, only the recent finding that WNK kinases are chloride-sensitive has provided a clue to the molecular basis for this effect (Ellison & Terker, 2015). WNK kinases switch between active and inactive states based on their phosphorylation status, with phosphorylation stabilizing the active state. Cl⁻ ions stabilize the inactive WNK1, and this prevents any kinase autophosphorylation and activation, thus explaining how the changes in intracellular chloride and toxicity affect the WNK kinase activity (Piala et al., 2014). This activation is rapid, occurring in less than 0.5 min, and is due to the phosphorylation of the serine residue in the T-loop which is conserved across all the WNKs. Crystallographic data on WNK1 have shown that it has an “LGL” motif, also called a “chloride sensor,” that confers chloride sensitivity by blocking the autophosphorylation of the T-loop, and thus confirming that WNKs are the key chloride sensing kinases which regulate the behavior of Na⁺ and K⁺ cation cotransporters such as NCC and NKCC2 (Haas & Forbush, 2000). Mutating the leucine residue in the LGL motif (LL to FF) causes an activation of WNK kinase function which is most pronounced in WNK4 (Piala et al., 2014). Recently, Ellison has proposed a model for DCT function in which low extracellular K⁺ ([K⁺]o) causes the DCT cell to hyperpolarize, resulting in the efflux of Cl⁻, a reduction in [Cl⁻]i, and activation of the WNK/SPAK/NCC pathway. With high [K⁺]o, the contrary occurs— the DCT is depolarized, Cl⁻ enters the cell and the rise in [Cl⁻]i, deactivates the WNK4/SPAK/NCC pathway (Terker et al., 2015). Thus the level of Na⁺ reabsorption at the DCT is directly regulated by the [K⁺]o, which determines the phosphorylation state of NCC by controlling the activity of the WNK/SPAK/NCC pathway.

We have tested the Ellison model directly to show that the effect of plasma [K⁺] on NCC levels is mediated by changes in [Cl⁻]i in response to changes in the membrane potential (V_m) of the cell. To do this, we have used an in vitro system, that is, modified HEK-293T cells generated by transient transfection of the necessary ion channels/transporters, including KCNJ10, Clc-kb, Barttin, typically found in DCT cells. A fluorescence-based approach was used to directly show that increasing [K⁺]o depolarizes the cells and causes subsequent inactivation of the WNK/SPAK/NCC cascade.

2 | MATERIALS AND METHODS

2.1 | Cell culture and transfection

Human embryonic kidney HEK-293 T cells were cultured in DMEM high glucose supplemented with 10% (v/v) fetal bovine serum, 2 mM L-glutamine, 100 units/ml penicillin, and 0.1 mg/ml streptomycin at 37°C in 5% CO₂-95% atmosphere. For transfection experiments, cells were grown on 10 mm petri dishes and were incubated for a period of 36 hr with a mixture of 20 μL of 1 mg/ml polyethyleneimine (Polysciences) and 5–10 μg of the respective plasmid DNA (KCNJ10, Clc-kb, and barttin from Origene), as described (Durocher, Perret, & Kamen, 2002). Depending on whether the V_m or [Cl⁻]i was going to be measured in the cells, the plasmid constructs pcDNA3.1 (−) VSFP3.1 (Addgene) or pcDNA3.1 (+) EYFP H148Q/I152L (Addgene) were transfected into the cells. Transfected cells were then transferred to poly-L-lysine coated 96-well, black microplates (Greiner bio-one), and allowed to adhere overnight before use. Simultaneous measurements of both V_m and [Cl⁻]i, in cells by co-expression of the voltage and Cl sensor were not feasible because of the large overlap of the two emission spectra of CFP and EYFP.

2.2 | Measurement of V_m, and [Cl⁻]i using the Flex station microplate reader

HEK cells transfected with the K⁺ channel KCNJ10, and the voltage-sensitive dye VSFP3.1 (encoded by pcDNA3.1 (−), Addgene) were used to measure the change in V_m at an absorption and emission wavelengths of 444 and 535 nm, respectively. HEK cells transfected with the chloride sensor, pcDNA3.1 (+) EYFP H148Q/I152L (Addgene), KCNJ10, Clc-kb and barttin were used to measure the change in [Cl⁻]i, at an absorption and emission wavelengths of 485 and 555 nm, respectively.
Immediately before the Flex station measurements, 96-well plates with the adhered HEK cells were washed with a solution containing 130 mM NaCl, 10 mM NaI, 2 mM KCl, 2 mM CaCl₂, 1 mM MgCl₂, and 20 mM HEPES pH 7.3. The cells were then incubated in 100 µl of the same solution at 37°C for 45 min and the fluorescence was measured in a FlexStation 3 microplate reader (Molecular Devices) at 2 s intervals for 200 s. NaCl was included only in the solutions which were used for [Cl⁻]i measurement. Solutions with different external KCl concentrations were introduced to the cells in each well after 50s of baseline measurement (the concentration of NaCl was adjusted in each case to maintain the osmolality of the solution). Appropriate negative controls were employed, and selective blockers including DIDS (Sigma) (L’Hoste et al., 2013), Tertiapin Q (Abcam) (Doupnik, 2017), and Ba (Sigma) (L’Hoste et al., 2013), Valsartan (Sigma) (Imbriaco, 2013), and diuron were employed, and selective blockers including DIDS (Sigma) (L’Hoste et al., 2013), Tertiapin Q (Abcam) (Doupnik, 2017), and Ba (Sigma) were used as indicated. The change in fluorescence (ΔF) was defined as F(max) (peak fluorescence) – F(min) (baseline fluorescence). EC50 values were obtained by fitting data to four-variable concentration–response curves using Prism 5 (GraphPad Software Inc., San Diego, CA).

2.3 | Biochemical studies

HEK-293 T cells were grown in normal cell culture conditions, and then transfected with mouse NCC (pECFP-C1-NCC) alone, or with WNK4 WT, WNK4 L322F L324F mutant or the WNK4 LLFF kinase-dead mutant form respectively (in pMO-Myc vector, a generous gift from Prof D. Ellison, University of Oregon), and grown in medium with different external KCl concentrations (2–5 mM).

2.4 | Biotinylation

Transfected HEK 293T cells in 10 cm plates were biotinylated following a previously published method (Murthy, Kurz, & O’Shaughnessy, 2016) Briefly, HEK 293T cells in 10 cm plates were washed once in NES buffer and placed on ice. Cells were incubated in 0.2 mg/ml NHS-biotin (www.thermofisher.com) for 45 min on ice before quenching with Tris buffer (25 mM Tris-HCl; 150 mM NaCl; 10 mM EDTA, pH 7.4). Cells were removed from the plate and centrifuged at 200g for 5 min at 4°C. Supernatant was discarded and cells were incubated in solubilization buffer (25 mM Tris-HCl; 150 mM NaCl; 10 mM EDTA; 1% Triton X-100, pH 7.4) with protease inhibitors (www.roche.co.uk) at 4°C with agitation for 1 hr. The resulting lysate was cleared by centrifugation at 14 K rpm 4°C for 1 hr. Thirty microliters of streptavidin beads (www.thermofisher.com) per sample were washed once in wash buffer (25 mM Tris-HCl; 150 mM NaCl; 10 mM EDTA; 1% Triton X-100, pH 7.4), centrifuged at 1,400 rpm for 5 min at 4°C and supernatant removed. The supernatant was added to streptavidin beads and incubated with agitation for 2 hr at 4°C. Beads were then pelleted at 14K rpm for 5 min at 4°C and washed three times in wash buffer before resuspending the pellet in 4x sample buffer (www.lifetechnologies.com) for Western blotting.

2.5 | Western blotting

For blots with nonbiotinylated cell lysates, cells were harvested 36 hr post-plasmid transfection from the 10 cm plates, and lysed in 300 µl lysis buffer (Terker, Zhang, et al., 2016) (supplemented with protease inhibitors, phosphatase inhibitors, and 1 mM DTT). After protein quantification by Bradford’s method, 20 µg equivalent total protein lysate was loaded on to 3%–8% Tris-acetate Nupage gel and transferred to nitrocellulose membranes. Membranes were Ponceau stained to check for transfer efficiency and loading, then blocked with 5% milk for 1 hr and incubated with primary antibody in 5% (w/v) skimmed milk in TBS-T with the indicated primary antibodies: 1 in 500 diluted rabbit phospho-WNK4 Ser575 antibody (a generous gift from Prof Isao Naguro, Japan) or 2 µg/ml sheep phospho-SPAK Thr373 antibody and peptide (to prevent binding to nonphosphorylated SPAK (10 µg/ml) (Division of Signal Transduction Therapy Unit, University of Dundee) or pNCC Thr58 (a generous gift from Prof J Loffing, University of Zurich) (1 in 1,000 dilution) overnight at 4°C. β-actin (mouse antibody from Thermo Fisher, 1 in 2,000 dilution) was stained as a loading control. The blots were then washed three times with TBS-T and incubated in secondary antibodies, donkey anti rabbit 800CW (926–32213 at 1:5,000 dilution; www.licor.com), and goat anti mouse Alexa680 (A-21058 at 1:5,000 dilution; www.licor.com), and goat anti mouse Alexa680 (A-21058 at 1:5,000 dilution; www.lifetechnologies.com) in TBS-Tween 20 for 1 hr at room temperature in the dark and then washed six times in TBS-Tween 20. Membranes were imaged using the LiCor Odyssey system (www.licor.com), and the blots were quantified using the Image studio Lite software www.licor.com).

2.6 | Statistical analysis

All data are shown as mean ± SEM unless indicated otherwise. Significance was determined using unpaired t test or one-way ANOVA followed by post hoc testing as appropriate. p < .05 was considered significant. The stats package in Prism4 (http://www.graphpad.com/scientific-software/prism/) was used.

3 | RESULTS

3.1 | Measurement of change in membrane potential (Vm)

Cells transfected with the voltage-sensitive VSFP3.1 exhibited a typical CFP (cyan fluorescent protein, emission max at
535 nm) emission spectrum that was sensitive to membrane depolarization when the [KCl]₀ was changed progressively from 2 through 5 mM KCl (Figure 1a). This effect of increasing in external K⁺ resulted from increasing quenching of the CFP fluorescence (Figure 1a) over the range of [KCl]₀ used, the normalized change in fluorescence (ΔF/F₀) exhibited a log-linear relationship with increasing external KCl (Figure 1b), showing that the membrane functions as a K⁺ electrode. The fluorescence quenching in 5 mM [KCl]₀ was affected by the presence of the K channel blockers Ba²⁺ or Tertiapin Q (Figure 1c); Ba²⁺ abolished the quenching almost completely, while Tertiapin Q (50nM) reduced it to about 10%. The quenching response to increasing external KCl was not observed in cells, which were not transfected with KCNJ10 (data not shown).

3.2 | I⁻ affects basal YFP quench

YFP, a genetically modified variant of green fluorescent protein (GFP), is halide sensitive, and the H148Q/I152L mutant form of YFP has increased sensitivity to halides with the relative affinity of anions to YFP-H148Q/I152L being I⁻>NO₃⁻>Br⁻>Cl⁻ (Galietta, Haggie, & Verkman, 2001). About 10 mM NaI was used in all our experimental solutions, based on previous work with this fluorophore (Johansson, Norris, & Peilot-Sjogren, 2013) showing that 10 mM NaI was an optimal concentration for assay performance while ensuring that there is no artifact of agonist-independent I⁻ permeability of the cells. EYFP quenching was not observed when cells were assayed in the absence of I⁻ (data not shown).

3.3 | Changing external [KCl] affects EYFP fluorescence in the cells

HEK cells expressing YFP-H148Q/I152L showed a step-wise fluorescence quenching in response to increasing [KCl]₀. A maximal EYFP fluorescence quenching of over 50% was seen when the concentration of KCl was stepped up from 2 to 5 mM, a physiologically relevant range of [K⁺]₀ (Figure 2a). When the normalized change in fluorescence (ΔF/F₀) was plotted against [K⁺]₀, the data fitted a four-variable dose–response curve, with an IC50 of 2.9 mM KCl (Figure 2b). The quenching response to increasing external KCl was not
observed in cells which were not transfected with *Clc*-kb and *barttin* (data not shown).

In the presence of the Cl channel blockers, the fluorescence quenching was substantially reduced. DIDS (4,4'-Diisothiocyanatostilbene-2,2'-disulfonic acid), an inhibitor of Cl⁻ flux (Wulff, 2008), abolished the EYFP quenching almost completely, suggesting that the K effect on Cl⁻ influx and intracellular EYFP depends on the expressed *Clc*-kb channel, whereas Valsartan (Imbrici et al., 2017) was less effective in comparison to DIDS. Tertiapin Q also reduced the EYFP quenching by about 50%, suggesting that K⁺ influx into the cells through KCNJ10 was necessary for the subsequent Cl⁻ influx, and EYFP quenching (Figure 2c).

### 3.4 Extracellular KCl affects phosphorylated WNK4 (pWK4) levels in NCC and WNK4 transfected cells

Having observed the sensitivity of the “model HEK cell” membrane to coupled changes in [KCl]₀ and transmembrane Cl⁻ movement, the consequence for the cellular WNK4/SPAK/NCC signaling pathway was investigated. Altering [KCl]₀ had distinct effects on the phosphorylation status of the different forms of WNK4 in the transfected cells. WNK4 WT was affected inversely with pWK4 levels falling (Figure 3a) when [KCl]₀ was increased from 2 to 3 mM and above. A steep fall in pWK4 levels was observed upon changing the [KCl]₀ from 2 to 3 mM with a further small reduction in its levels between 3 and 5 mM KCl (Figure 3b). The K channel blocker Tertiapin Q, abolished this effect on WNK4 WT levels (Figure 3c), demonstrating the role of the KCNJ10 channel in mediating the effect of changing [K⁺]₀ on WNK4 levels. Increasing [K⁺]₀ did not affect the intensities of the Cl insensitive mutant WNK4 LLFF (Figure 3d), confirming the role of the kinase domain of WNK4. Finally replacing WT WNK4 with the WNK4 LF kinase-dead markedly reduced or abolished p WK4 levels showing the importance of the kinase domain for the changes seen in pWK4.

### 3.5 Increase in external KCl affects the downstream phosphorylation targets of WNK4

The inverse relationship between [KCl]₀ and pWK4 levels was reflected in its downstream phosphorylation targets:
**Figure 1**

**a**

Western blots showing pWNK4 phosphorylation at various concentrations of KCl. Each lane represents a different condition: 2 mmol/L KCl, 3 mmol/L KCl, 4 mmol/L KCl, and 5 mmol/L KCl.

**b**

A bar graph illustrating the pWNK4 intensities at different KCl concentrations. The x-axis represents KCl concentrations (2, 3, 4, and 5 mmol/L), and the y-axis represents pWNK4 intensities (a.u.). The graph shows a significant increase in pWNK4 intensities at 4 mmol/L KCl compared to 2 mmol/L KCl.

**c**

A Western blot similar to **a**, but with an additional treatment of Tertiapin Q at 4 mmol/L KCl.

**d**

A Western blot similar to **a**, but with a variety of WNK4 mutations (WT, WNK4 LE, WNK4 LE KD, WNK4 LE KD) at different KCl concentrations (2, 3, 4, and 5 mmol/L).
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FIGURE 3 Effect of \([\text{K}]_o\) on WNK4 phosphorylation in HEK 293 T cells transfected with NCC alone or different forms of WNK4: WNK4 WT, WNK4 LLFF, and WNK4 LLFF KD, respectively. (a) Total protein lysates were blotted from HEK 293 T cells transfected with NCC and the WT WNK4 at four different external KCl. Rabbit phospho WNK4 Ser 575 antibody detected a 150 kDa WNK4 band. ß-actin was the loading control. This is representative of Western blots replicated three times with cell lysates from three different passage numbers. (b) Quantification of Western blots showing phospho WNK4 levels normalized to its ß-actin loading control. Error bars represent mean ± SEM of Western blots \((n = 3)\). *p < .0001 by two-tailed t test. (c) The effect of K channel blocker, Tertiapin Q on phospho WNK4 levels at 4 mM, and 5 mM KCl, respectively. (d) Phospho WNK4 levels in cells transfected with the three different WNK4 forms, and at four, different \([\text{K}]_o\).

SPAK and NCC. Both pSPAK (Figure 4a) and pNCC (Figure 4c) levels decreased with increasing \([\text{KCl}]_o\) in the model HEK cells expressing WNK4 WT. This fall in pSPAK levels with an increase in \([\text{KCl}]_o\) from 2 to 5 mM (Figure 4b) possibly explains the decline in the phosphorylation of NCC at a similar range of concentrations of \([\text{KCl}]_o\). The levels of pNCC is a linear function of \([\text{KCl}]_o\), within the range of 2 to 5 mM and the slope of the trendline is comparable (Figure 4d) to the one observed when the pNCC levels were plotted against a similar range of plasma \([\text{K}^+]\) (in the mice untreated with aldosterone) in the in vivo study (Terker, Zhang, et al., 2016). With pNCC being a marker for NCC activity (Pacheco-Alvarez et al., 2006), the effect of \([\text{KCl}]_o\) on pNCC suggests that the NaCl entry via NCC would be reduced at physiologically viable higher \([\text{K}^+]_o\) due to its restricted activity.

4 | DISCUSSION

Although \([\text{K}^+]\) is the most abundant exchangeable cation in the human body, with an intracellular concentration of 140–150 mM, the serum \([\text{K}]\) is normally maintained within a smaller physiological range, 3.5–5 mM. Earlier animal studies have shown that a large increase in dietary \([\text{K}^+]\) affects the abundance and activity of the cotransporter NCC (Terker et al., 2015; Vitzhum et al., 2014); however, a recent paper (Terker, Zhang, et al., 2016) has demonstrated that \([\text{K}^+]\) is a pivotal modulator of NCC at a physiologically relevant concentration, and proposed a possible mechanism (“Ellison hypothesis”) to explain this. Our in vitro study is a working model of this hypothesis, where the physiological and molecular effects of changing \([\text{K}^+]_o\) on intracellular Cl and the WNK/SPAK/NCC pathway (involved in regulating salt transport across the DCT) has been shown in a “DCT-like” HEK cell system.

Using a CFP fluorescence-based approach in the model HEK cell system, we have shown that the cell membrane functions as a \([\text{K}^+]\) electrode which responds to small, step-wise changes of 1 mM KCl when the \([\text{K}^+]_o\) is increased from 2 mM through 5 mM in cells expressing KCNJ10. Significant reduction in EYFP fluorescence was detected as \([\text{KCl}]_o\) was increased, indicating an inverse relation of EYFP fluorescence quenching to increased \([\text{K}^+]_o\). Blockers such as Tertiapin Q and BaCl₂ abolished these effects of changing \([\text{K}^+]_o\), demonstrating the role of the K channels on the DCT-like cell membrane (including KCNJ10), whereas the Cl⁻ channel blockers affected EYFP quenching which is a consequence of Cl⁻ influx across the membrane, and binding of these anions to the EYFP protein. Thus our model cells mimics the hypothesized behavior of DCT cells and confirms they can respond to small changes in \([\text{K}^+]_o\) within the physiological range followed by transmembrane anion transport through CICs.

The model HEK cells expressing WNK4 or its mutant forms respectively responded differently to altering \([\text{K}^+]_o\) in the physiologically relevant range of 2 to 5 mM. The \([\text{K}]_o\) affected the activity of WNK4/SPAK/NCC signaling and its downstream targets in this model cell line as determined by measuring the levels of pWNK4, pSPAK, and pNCC level, respectively. The inverse effect of increasing \([\text{K}^+]_o\) on pWNK4 levels in cells expressing wild-type WNK4 was abolished by K channel blockers (tertiapin and Ba²⁺), thus confirming that DCT K channels (such as KCNJ10) are the route for \([\text{K}^+]\) entry into these cells, which facilitates the subsequent effects on the WNK4/SPAK/NCC pathway as observed in in vivo studies (Wang et al., 2018). The absence of a similar \([\text{K}^+]\) sensitivity to the phosphorylation states of the same proteins in cells expressing the Cl⁻ insensitive mutants, WNK4 LLFF and WNK4 LF KD respectively, emphasises the importance of the WNK4 Cl⁻ binding and kinase domains for transducing the membrane potential changes caused by altered \([\text{K}^+]_o\).

WNK4 has been shown to be the most responsive WNK family member to changes in \([\text{Cl}^-]\) (Terker, Zhang, et al., 2016), so it appears to be the key sensor of \([\text{K}^+]\) in DCT cells where \([\text{Cl}^-]\) is typically around 10–20 mM (Weinstein, 2005). Our model HEK cells behave like the DCT cells, with the downstream signaling targets of WNK4, the SPAK and NCC showing a corresponding decrease in their phosphorylation levels at \([\text{K}^+]_o\) of 5 mM. Although WNK1 has been reported to be the \([\text{K}^+]\) sensor in HEK cells (Terker et al., 2015) (due to its higher \([\text{Cl}^-]\) of >40 mM), the exogenously expressed WNK4 in our model cell probably functions as the primary sensor due to its much greater abundance over WNK1.

The abundance of both pSPAK and pNCC in our in vitro study showed an inverse relationship to the increase in \([\text{K}^+]_o\), and interestingly, the pattern of pNCC reduction with \([\text{K}^+]_o\) was comparable to the one observed in aldosterone untreated
Figure 4  Effect of [K]₀ on NCC phosphorylation in HEK 293 T cells transfected with NCC alone or with WNK4 WT. The transfected cells were treated with external KCl concentrations of 2, 3, 4, and 5 mM respectively. Cells were biotinylated. (a) Nonbiotinylated fraction of the cell lysates were blotted. Sheep pSPAK t373 antibody detected an approximately 60 KD band. (b) Quantification of Western blots showing phospho pSPAK levels normalized to its ß‐actin loading control. Error bars represent mean ± SEM of Western blots (n = 3). p < .05 by one‐way ANOVA. (c) Biotinylated protein lysates were blotted and rabbit pNCC Thr 58 antibody detected a 150 KD phospho NCC band. (d) Quantification of Western blots showing biotinylated phospho NCC levels. Error bars represent mean ± SEM of Western blots (n = 3). *p < .05 by two‐tailed t test.
mice in vivo. However, in the same study aldosterone-treated mice showed a much steeper decline in pNCC abundance with an increase in plasma [K+] (Terker, Zhang, et al., 2016), suggesting a role for aldosterone in ameliorating the phosphorylation and thus activation of NCC at low plasma [K+].

Given NCC is the principal transporter for Na+ reabsorption in the DCT, it plays a crucial role in controlling blood pressure, and has been shown to be regulated by the hormone aldosterone indirectly via changes in plasma [K+] (van der Lubbe et al., 2012; Terker, Yarbrough, et al., 2016). Interestingly, a recent study revealed that aldosterone‐treated mice showed a much steeper decline in pNCC abundance at physiologically relevant [K+]o, within the physiological range and also show the changes in intracellular [Cl−] that affect downstream WNK4/SPAK/NCC signaling. This confirms the hypothesized DCT model in which the transcellular flux of NaCl in the DCT is directly regulated by the membrane potential of the DCT cell and its direct coupling to the phosphorylation state of NCC through the Cl− sensor WNK4.

ACKNOWLEDGMENTS

We would like to thank Professor Sarah Lummis (Department of Biochemistry, University of Cambridge) for giving us access to the Flex station Plate reader. We acknowledge Professors J. Loffing and I. Naguro for giving us the antibodies used in this work.

CONFLICT OF INTEREST

None declared.

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How to cite this article: Murthy M, O'Shaughnessy KM. Modified HEK cells simulate DCT cells in their sensitivity and response to changes in extracellular K. *Physiol Rep*. 2019;7:e14280. https://doi.org/10.14814/phy2.14280