The epithelial sodium channel mediates the directionality of galvanotaxis in human keratinocytes

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Summary

Cellular directional migration in an electric field (galvanotaxis) is one of the mechanisms guiding cell movement in embryogenesis and in skin epidermal repair. The epithelial sodium channel (ENaC), in addition to its function of regulating sodium transport in kidney, has recently been found to modulate cell locomotory speed. Here we tested whether ENaC has an additional function of mediating the directional migration of galvanotaxis in keratinocytes. Genetic depletion of ENaC completely blocks only galvanotaxis and does not decrease migration speed. Overexpression of ENaC is sufficient to drive galvanotaxis in otherwise unresponsive cells. Pharmacologic blockade or maintenance of the open state of ENaC also decreases or increases, respectively, galvanotaxis, suggesting that the channel open state is responsible for the response. Stable lamellipodial extensions formed at the cathodal sides of wild-type cells at the start of galvanotaxis; these were absent in the ENaC knockout keratinocytes, suggesting that ENaC mediates galvanotaxis by generating stable lamellipodia that steer cell migration. We provide evidence that ENaC is required for directional migration of keratinocytes in an electric field, supporting a role for ENaC in skin wound healing.

Key words: Keratinocyte, Directional migration, ENaC, Lamellipodia

Introduction

Skin functions as an important protective organ, shielding the body from infections, and fluid or thermal loss. The presence of a transepithelial electrical potential is an interesting physiological feature of healthy intact skin, observed to be present in human skin as early as the 19th century (reviewed by McCaig et al., 2005). A 20–65 mV potential difference is measured between the surface of the epithelium and dermis (Barker et al., 1982; Foulds and Barker, 1983) with the inside positive relative to the surface. The non-conductive stratum corneum and polar distribution of higher concentrations of cations including calcium, potassium and sodium within the upper level of the epidermis (Denda et al., 2000; Leinonen et al., 2009) are hypothesized to maintain this transepithelial potential.

When the skin is wounded, the transepithelial potential collapses to zero at the cut site, while the persistence of the higher potential within the adjacent intact epidermis at the wound edge generates a weak electric field (EF) which then drives current flow toward the center of the wound (the cathode) (Barker et al., 1982; Ojingwa and Isseroff, 2003; Reid et al., 2007; Nuccitelli et al., 2008). The wound EF can be measured within several millimeters from the wound edge in newt (Borgens et al., 1977), guinea pig (Barker et al., 1982), mouse (Reid et al., 2007; Nuccitelli et al., 2008) and human (Mukerjee et al., 2006; Reid et al., 2007) skin and cornea, and ranges from 90 mV/mm in human to 180 mV/mm in rodent skin wounds. The EF persists until the wound is fully resurfaced with new epithelium (Borgens et al., 1977; Nuccitelli et al., 2008), and thus has been proposed to be an important component of the wound repair process.

The directionality of galvanotaxis is one of the mechanisms guiding cell movement in embryogenesis and in skin epidermal repair. The epithelial sodium channel (ENaC), in addition to its function of regulating sodium transport in kidney, has recently been found to modulate cell locomotory speed. Here we tested whether ENaC has an additional function of mediating the directional migration of galvanotaxis in keratinocytes. Genetic depletion of ENaC completely blocks only galvanotaxis and does not decrease migration speed. Overexpression of ENaC is sufficient to drive galvanotaxis in otherwise unresponsive cells. Pharmacologic blockade or maintenance of the open state of ENaC also decreases or increases, respectively, galvanotaxis, suggesting that the channel open state is responsible for the response. Stable lamellipodial extensions formed at the cathodal sides of wild-type cells at the start of galvanotaxis; these were absent in the ENaC knockout keratinocytes, suggesting that ENaC mediates galvanotaxis by generating stable lamellipodia that steer cell migration. We provide evidence that ENaC is required for directional migration of keratinocytes in an electric field, supporting a role for ENaC in skin wound healing.
human primary keratinocytes (Brouard et al., 1999) and murine epithelium (Mauro et al., 2002). Although the genetic ablation of the αENaC gene in mice results in a postnatal lethal phenotype due to an inability to clear lung fluids (Hummler et al., 1996), the role of this channel in skin epithelial biology is not yet elucidated. αENaC knockout mice do have abnormal lipid composition in their stratum corneum (Mauro et al., 2002; Charles et al., 2008), but these animals do not survive long enough to perform in vivo skin investigations. In humans, loss-of-function mutations of αENaC genes cause a rare autosomal, recessive disease: pseudohypoaldosteronism type 1 (Chang et al., 1996; Strautnieks et al., 1996), resulting in salt loss, hyperkalemia and metabolic acidosis in newborns. Although cutaneous lesions have been described in case reports of the pseudohypoaldosteronism type 1 patients, these lesions are possibly related to the high concentration of salt in sweat glands (Urbatsch and Paller, 2002; Martin et al., 2005).

Several recent studies suggest that αENaC is involved in cell migration in vitro. Inhibition of αENaC activity in bovine corneal endothelial (BCE), rat aortic vascular smooth muscle cells, and human BeWo cells decreases the rate of scratch wound closure compared to the control (Chifflet et al., 2005; Grifoni et al., 2006; Grifoni et al., 2008; Del Mónaco et al., 2009). Additionally, an ENaC-dependent Na⁺-influx at the wound edge of BCE cells was observed upon the generation of the scratch wound (Chifflet et al., 2005), suggesting that opening of ENaC channels may be the initial step in migration and wound closure for these cells. However, little is known about how, or if, αENaC mediates cell motility or directional migration such as galvanotaxis.

In the current study, we examined the hypotheses that ENaC mediates the directional migratory galvanotaxis response in skin-derived keratinocytes, thus contributing to cutaneous wound repair. We found that ENaC expression is required for the galvanotaxis directional response: knocking down ENaC or blocking ENaC channels does not alter overall cell motility yet remarkably inhibits the directional response of keratinocyte galvanotaxis. Overexpressing ENaC could drive galvanotaxis in cells that are ordinarily non-responsive to an applied EF. Additionally, preferential lamellipodial protrusion at the cathodal side of the cell is dependent on ENaC expression, and in ENaC-deficient cells, no cathodally dominant lamellipodia are established. Our findings suggest a novel function for ENaC in mediating the directional migration of keratinocyte, by stabilizing protrusions at the cathodal side during galvanotaxis. These findings help explain the elusive mechanism of cellular directional responses to physiological electrical fields.

**Results**

**ENaC-depleted keratinocytes lose directionally in galvanotaxis**

To investigate the role for ENaC in keratinocyte galvanotaxis, we examined the directionality of the migratory response in an EF of physiologic magnitude in murine and human keratinocytes that had been genetically depleted of αENaC. The morphology of αENaC knockout mouse epithelial keratinocytes (αENaC-KO-MEK) (+/−) is similar to that of the wild-type MEK (+/+) (Fig. 1A; supplementary material Movies 1, 2) (Hummler et al., 1996) (αENaC depletion in the knockout MEK is shown in supplementary material Fig. S1A). Although αENaC-KO-MEK migrated significantly faster than the wild-type MEK (0.86±0.04 μm/minute versus 0.73±0.03 μm/minute for wild-type keratinocytes, P<0.05), the directional response was essentially absent (decreased by 94%, P<0.05) in the αENaC-KO-MEK, with resultant random, rather than cathodally directed migration in the EF (Fig. 1B, right). Similar results were obtained from siRNA knockdown of αENaC in human keratinocytes. The total αENaC expressed in NHK is decreased by 50% after 72 hours of siRNA treatment (supplementary material Fig. S1B), at which time the migratory response to the EF was examined (Fig. 1B, left; supplementary material Movies 3, 4). As in the αENaC-KO-MEK, αENaC siRNA-treated NHK demonstrate a significant decrease in cathodally-directed galvanotaxis (Fig. 1B, left) compared to the control cells, indicating the loss of a directional migratory response to the applied EF.
Overexpression of active αENaC promotes galvanotaxis

To further examine whether αENaC is sufficient to drive the cellular galvanotaxis response, we tested galvanotaxis in cells overexpressing αENaC. A human lung epithelial cell H441 line (parental), its EGFP-tagged αENaC overexpressing cell line derivative (αC3-3-GFP) as well as a control EGFP expressing cell line (GFP7) were examined for their directional migration in response to an applied EF. The αC3-3-EGFP ENaC overexpressing line demonstrates increased levels of total αENaC relative to the parental cell line H441 or the control EGFP expressing line (the endogenous and the GFP-αENaC, supplementary material Fig. S1C) (Woollhead and Barnes, 2006). All three cell lines H441, GFP7 and αC3-3-GFP migrated with similar speeds of 0.55±0.04, 0.57±0.02, and 0.60±0.02 μm/minute (insignificantly different) in the EF. However, the cosine values for the H441 and GFP7 lines are close to zero, indicating no directedness of migration (Fig. 1C), demonstrating that these cells migrated randomly without a directional response to the EF. On the other hand, the αC3-3-GFP cells overexpressing active αENaC showed a significant increase in cosine value (cosine=0.35, P<0.05) compared to control GFP7 and the parental line (cosine=0), indicating a directional cathodal response to the EF.

The open state of ENaC is involved in NHK directionality in galvanotaxis

In addition to genetic manipulations of levels of expression of αENaC, we investigated whether ENaC in the open state is required to drive galvanotaxis, using pharmacologic modulation of its channel activity. Previously we showed that amiloride, a blocker of the open state of the ENaC channel at a low concentration of 10 μM has no effect on galvanotaxis (Trollinger et al., 2002); therefore, we treated the cells with a more potent and ENaC-specific amiloride analog, phenamil (IC50 is 284 nM for amiloride and 75 nM for phenamil tested on human bronchial epithelial cells; Hirsh et al., 2004) and monitored the NHK migration in the EF. Migration speeds of control and 20 μM phenamil-treated NHK are similar at 1.02±0.43 μm/minute. While control cells demonstrate robust cathodal directional migration, in the presence of phenamil, the cosine value of NHK migrating toward the cathode is decreased by 30% (Fig. 2A, P<0.05), suggesting that the sodium influx via ENaC channel is involved in, but is not the major signal for the NHK directional migration in an electric field.

The inhibitory peptide derived from αENaC cleavage impairs NHK galvanotaxis

Activation of the ENaC channel by proteases releases a short self-inhibitory sequence from the mouse αENaC of amino acids 182–190, LPHPLQRL (Carattino et al., 2006; Carattino et al., 2008). The cleaved ENaC channel is constitutively active and stays in the open state; treating mouse cortical collecting duct cells and human airway epithelial cells with synthetic LPHPLQRL peptide blocks the opening of ENaC (Carattino et al., 2006; Carattino et al., 2008). We used this inhibitory peptide to further confirm the role of open ENaC channels in galvanotaxis. Addition of the peptide to the cells did not alter migration speed, since NHK cultured with either the inhibitory peptide or the scrambled control peptide migrated at the same speed. Similar to the phenamil inhibition, with the pre-treatment of 20 μM αENaC inhibitory peptide, the net cosine value of NHK galvanotaxis decreased by 30% (Fig. 2B, P<0.05) from control values. These results provide additional support to the notion that the ENaC channel in open state is required for galvanotaxis.

Opening ENaC channels promotes NHK directional migration in an EF

On the other hand, if ENaC in the open state is required for galvanotaxis, it is reasonable to propose that constitutively active ENaC should promote NHK directional migration. An ENaC activating agent, S3969, increases the open probability of the channel by activating the beta subunit (Lu et al., 2008). When normal human keratinocytes are pre-treated with S3969 prior to the EF exposure, they demonstrate a 17% increase in the net cosine value from 0.65 to 0.76 (P<0.05) at the end of the 60 minutes galvanotaxis assay (Fig. 2C) compared to the control, while maintaining similar migratory speeds. Opening ENaC by S3969 also increases the slope of the track cosine in the first 20 minutes during galvanotaxis (0.019 in DMSO versus 0.027 in S3969, Fig. 2D), indicating that the S3969 treated keratinocytes migrate on a more persistent, directional route toward cathode in the EF than control cells. These results indicate that ENaC channel at the open state could promote galvanotaxis.

Since the accumulated data support a role for ENaC in keratinocyte galvanotaxis in vitro, and skin wounds generate an
EF of the magnitude used in these studies, we examined how wound healing is affected by the ENaC inhibitor phenamil using an organotypic, ex vivo human wound assay, in which the early stage of wound re-epithelialization, dependent on keratinocyte migration, can be quantitated (Kratz, 1998; Lu and Rollman, 2004). Wound re-epithelialization of control wounds was 90–95% complete after 7 days of ex vivo cultivation and a single layer of NHK covered the original wound area (supplementary material Fig. S2A,B). However, in the presence of 10 μM phenamil, the re-epithelialization is blocked by 70% compared to the control solvent DMSO treatment at day 7 (87.8% in DMSO versus 28.2% in phenamil, P<0.05, supplementary material Fig. S2B–D) and the NHK migrated only minimally into wound. The results suggest that inhibiting ENaC activity by phenamil inhibited keratinocyte migration and re-epithelialization of the wound in the first 4 days of healing, and at later times, impaired viability may have contributed to the decrease in wound re-epithelialization (PCNA staining and extracellular lactate dehydrogenase activity in supplementary material Fig. S3).

NHE1 is not involved in keratinocyte galvanotaxis

Another sodium transporter, NHE-1, present in cultured NHK and skin epithelium mediates the pH homeostasis and the epithelial barrier function (Hachem et al., 2005). It is also required for establishment of fibroblast polarity during directional migration (Denker et al., 2000; Frantz et al., 2007). Amiloride or phenamil, inhibitors of ENaC activity and galvanotaxis, can also inhibit NHE-1 at high concentration (Frelin et al., 1988) and theoretically could impair galvanotaxis by this mechanism. Therefore, we examined whether NHE1 is involved in galvanotaxis by using the NHE1-specific inhibitor cariporide to treat keratinocytes and also by examining galvanotaxis of PS120 fibroblast strains expressing wild-type NHE1 (PSN) or the dead NHE1 transporter (E266f, a mutant NHE1 containing an isoleucine substitution to glutamate 266, Denker et al., 2000).

NHK migration speeds were not inhibited in a high concentration of 50 μM cariporide (cariporide K_D=0.05 μM for NHE-1, Scholz et al., 1995) (10 μM cariporide inhibits a melanoma cell line MV3 migration, Stock et al., 2005) (Fig. 3A, left), and the directionality in response to an applied EF was also not altered (Fig. 3A, right). Similar results were observed when we compared the fibroblasts expressing the defective NHE1 (E266f) to wild-type NHE1 (PSN). Galvanotaxis of the fibroblasts expressing either wild-type NHE1 or the deficient NHE1 show the same degree of directionality in the EF (Fig. 3B, right), suggesting that unlike ENaC, the sodium transporter NHE1 is not required for galvanotaxis.

ENaC translocates to the leading edge of NHK after 60 minutes exposure of the EF

One mechanism by which EF may lead directional migration is to induce asymmetric relocalization of pro-migratory signals to the leading edge of the cell, as has been shown for the EGF receptor (Fang et al., 1999; Zhao et al., 1999) and PI(3)K gamma (Zhao et al., 2006). To study if ENaC translocates to the cathodal edge of the cell during galvanotaxis, αENaC was immunolocalized in both NHK and MEK at different time points after exposure to the EF, using antibodies that detect all forms of αENaC: the full-length 80–85 kDa and the cleaved, active 60–65 kDa (Fig. 4B) (Carattino et al., 2006; Carattino et al., 2008). At time zero, αENaC was randomly located at the cell periphery and in perinuclear areas (Fig. 4A). When keratinocytes are exposed to the EF, αENaC is not detected at the leading edge of NHK until 60 minutes after exposure (Fig. 4A,C). At this time, the staining intensity is 14% higher at the cathodal side than at the anodal side in NHK (Fig. 4C, P<0.05). Since the directional migratory response to the applied EF is detected as early as 15 minutes after exposure (Nishimura et al., 1996), the temporal lag in relocalization of ENaC to the lamellipodium suggests that polarization of ENaC to the leading edge of the lamellipodium may not mediate the galvanotaxis response.

ENaC stabilizes lamellipodial protrusion at the cathodal side during galvanotaxis

The role of lamellipodial dynamics in mediating the directional migratory response in galvanotaxis has not yet been investigated. However, in the directional response in chemotaxis, analysis of lamellipodial and pseudopodial dynamics has provided two models for directional cell steering. A chemotactant could induce (1) de novo formation of cell protrusions toward the gradient or (2) the differentiation of existing protrusions and stabilization of the ones facing the gradient (Andrew and Insall, 2007).

Similar to chemotaxis, keratinocytes could orient their cell bodies during galvanotaxis using lamellipodial steering. NHK in the control galvanotaxis groups, were not always polarized to the...
In wild-type MEK, the protrusion distance is 50–65% further at the cathodal as compared to anodal-facing side of the cell (Fig. 5A, 12.5±1.9 μm at line 9 versus 7.8±1.4 μm at line 1; similar results at line 8 versus line 2, the distance panel in Fig. 5C, ρ<0.05) and the protrusion speed of cathodal-facing lamellipodia is faster within the first 10 minutes of EF exposure than at the anodal-facing side (the speed panel in Fig. 5C, line 9 versus line 1), which results in a further extension of lamellipodia at the cathodal side. The switching frequency between protrusion and retraction is similar on both sides. However, in the αENaC-KO keratinocytes the distances of protrusion or retraction are very similar at the distal end of the cells, preventing an asymmetric extension of the lamellipodia during application of the EF, and thus, no directional migratory response develops. These results suggest that the cathodal lamellipodia are more stable and extend faster and further in the wild type, galvanotactically responsive keratinocytes, as compared to the cells in which αENaC is genetically depleted. Based on this analysis, we propose that αENaC in wild-type keratinocytes is required for establishing stable lamellipodia at the cathodal side during galvanotaxis and that the directional stabilization of the lamellipodia is an important mechanism to orient and steer keratinocyte migration in galvanotaxis.

A difference between the overall protrusion speed of the lamellipodia in the wild type and in the αENaC-KO-MEK was observed as well. There is an 18% significant increase of the migration speed of the KO cells (0.86 μm/minute in αENaC-KO-MEK versus 0.73 μm/minute in wild-type keratinocytes), and 24% and 45% increases of the protrusion and retraction distance in the αENaC-KO-MEK compared to the wild-type cells (Table 1, rows 1 and 4). The overall protrusion speed is also increased by 47%, the duration of retraction increased by 17%, and the switching between protrusion and retraction is more frequent in the αENaC-KO-MEK (Table 1, rows 2, 6 and 7). These results suggest that the function of αENaC in wild-type keratinocytes may be not only to stabilize a dominant, directionality-determining lamellipodium, but also to inhibit, rather than promote, multidirectional lamellipodial extension, and, thus, in the absence of αENaC this control mechanism for lamellipodial outgrowth is impaired and cells move faster, but with less directionality.

Discussion

Summary

Although we have known for over a century that EFs exist within wounds, as well as during limb regeneration and embryogenesis, (Borgens et al., 1977; Burr, 1941) it is only recent work that has demonstrated that these EFs provide directional cues to guide migration of cells to accomplish tissue remodeling required for these physiologic events (Nishimura et al., 1996; Sheridan et al., 1996; Zhao et al., 1996; Zhao et al., 1997; Fang et al., 1998; Fang et al., 1999; Zhao et al., 1999; Farboud et al., 2000). While intracellular signaling mediators of the directional response have been identified, including the kinase domain of the EGF receptor, PI3K, PTEN, phosphorylated ERK, phosphorylated Src, Rac-1, and cAMP (Fang et al., 1999; Zhao et al., 2002; Pullar and Isseroff, 2005; Zhao et al., 2006; Pullar et al., 2006), the proximal
cellular sensor that detects the EF and initiates the signaling cascade to initiate directional migration has remained elusive. Our study demonstrates that ENaC plays a novel role in keratinocyte galvanotaxis and may indeed function as the proximal EF sensor, initiating directional keratinocyte migration ultimately required for wound healing. Keratinocytes in which αENaC is genetically depleted completely lose their ability to respond to an applied EF with directional cathodal migration, although their random locomotory speed is unaffected. ENaC is sufficient to establish galvanotaxis when expressed in cells that do not respond with directional migration to EFs. Mechanistically, αENaC regulates directional migration by establishing stable lamellipodial protrusions at the cathodally facing pole of cells exposed to an EF, maintaining cellular polarity that directs the migratory response toward the cathode. Although ENaC has been shown to mediate migratory speed in other types of cells (Chifflet et al., 2005; Grifoni, et al., 2006; Grifoni, et al., 2008), this is the first study to show that ENaC is required for the directional cell migration in an EF and provides an important mechanistic insight as to how electrotaxis is accomplished.

ENaC protein mediates galvanotaxis
Depleting αENaC from keratinocytes abolished galvanotaxis, suggesting that αENaC or αENaC-associated proteins mediate the directionality of the cell migration in the EF. We also tested the hypothesis that the open state of ENaC is required for keratinocyte galvanotaxis. Blocking the open state of ENaC with either phenamil or the ENaC-derived inhibitory peptide decreases the keratinocyte galvanotaxis response. The high dosage of 10, 20 and 40 mM phenamil treatments did show a dose-dependent inhibitory response of galvanotaxis of 20–50%, but the ENaC control of sodium ion transport may not be the major signal for guiding directional migratory galvanotaxis responses in keratinocytes. Another interesting observation is that opening the ENaC channel on the cell surface enhances the directionality, but
it is not the sole requirement to steer the keratinocyte in galvanotaxis.

In the Trollinger et al. (Trollinger et al., 2002) paper, we previously proposed that an unidentified calcium channel is required for galvanotaxis. Calcium channel blockers such as Sr$^{2+}$ and Gd$^{3+}$ inhibited the directionality of keratinocytes, but the localization or the timing of the activation of the calcium channel remains unclear. In the current study, we found ENaC also mediates galvanotaxis. There could be several pathways involved in galvanotaxis, including sensing of the electric field, determining the direction to migrate, maintaining a persistent direction of migration, and the controlling of cell adhesion and migration speed. ENaC appears to be important in sensing of the electric field and determining the direction of migration (see below). Calcium and the postulated calcium channel might be involved in cell adhesion and migration rate. Future studies will be required to test if a calcium channel is downstream of ENaC in guiding the directional response, or whether they are parallel pathways that control different cellular responses in galvanotaxis.

Although the Na$^+/H^+$ exchanger NHE1 has previously been shown to be involved in fibroblast galvanotaxis, this was detected only when cells were exposed to supra-physiological EF strengths (400 mV/mm, Zhao et al., 2006). Under the physiological strength of the 100 mV/mm EF used in the current study, NHE1 does not mediate galvanotaxis in either normal human fibroblasts or keratinocytes. Our finding of ENaC-mediated galvanotaxis in normal human keratinocytes implies a physiologic role for this mechanism in guiding directional migration of these cells in vivo, especially since it is well-documented that wounding generates an endogenous EF to which the wound edge keratinocytes are immediately exposed. Thus, we propose that ENaC-mediated sensing of the EF plays a fundamental physiologic role in directing cell migration and contributing to wound repair. Additionally this work describes a novel role for ENaC in the epidermis, beyond the previously described function of modulating epidermal barrier formation (Charles et al., 2008).

**ENaC localization during galvanotaxis**

Because re-localization of some downstream mediators of the galvanotaxis response has been reported upon exposure of cells to an EF, we investigated whether there is a polarized localization of ENaC in the cells. During galvanotaxis, the directional response (cosine value) of keratinocytes starts increasing as soon as 15 minutes after exposure to the EF (Nishimura et al., 1996). Unlike the rapid polarization of EGFR and PI3K to the leading edge of galvanotactic keratinocytes (Fang et al., 1999; Zhao et al., 2006) or the Par-3 and Tiam-1 at the leading edge of migrating keratinocytes (Pegtel et al., 2007), ENaC does not polarize at the cathodal side of keratinocytes until a later time point (60 minutes) in the galvanotaxis response, suggesting that the polarization of ENaC does not initiate the directional migration of keratinocytes in response to the EF. It is possible that the late ENaC polarization we observe could be secondary to its known association with actin (Mazzochi et al., 2006), since actin reorganizes at the leading edge of migrating cells, in this case, the cathodal-facing edge of the cells. Similar results are found in the chemotactic Dicyostelium and neutrophils (Xiao et al., 1997; Servant et al., 1999). By time-lapsed imaging of the GFP-tagged chemoattractant receptor, the distribution of the chemoattractant receptors does not polarize at the leading edge when cells migrate and turn toward the attractant source, which suggests the polarization of the chemoattractant receptors is not required for the directional response of the cells.

**ENaC and lamellipodial protrusion**

We also examined if EFs induce the formation of dominant protrusions at the cathodal sides of exposed cells to polarize them, or if EF exposure can result in the asymmetric stabilization of protrusions thus promoting directional migration. The keratinocytes do not polarize immediately toward the cathode in an EF. Instead, kymographic analysis of the leading edge of galvanotactic cells shows that wild-type keratinocyte cells stabilize their cathodal protrusions, whereas ENaC-deficient cells cannot differentiate the cathodal and anodal sides of the cells in an EF in terms of lamellipodial protrusion. Lamellipodial protrusion depends on actin polymerization and depolymerization at the leading edge (Watanabe and Mitchison, 2002; Pollard and Borisy, 2003; Ponti et al., 2004). Asymmetric protrusion rate and distance between the cathodal side and anodal side could be an important mechanism to re-orient keratinocytes towards the cathode in the EF. At the present time there is not sufficient information to fully understand the mechanism by which ENaC may mediate lamellipodial dynamics. However, several studies suggest that ENaC associates with actin filaments and actin regulators. The c-terminus of ENaC has been shown to bind to F-actin in MDCK cells (Mazzochi et al., 2006) and the association is proposed to anchor ENaC at the plasma membrane. ENaC also binds to cortactin, an actin regulator to promote stable branching of F-actin (Ilatrovskaya et al., 2011). Thus, we speculate that ENaC could also regulate actin dynamics via its binding to newly assembling actin filaments in the lamellipodium and thereby regulate lamellipodial protrusions. Further investigations will be required to elucidate these potential interactions.

**ENaC could be a cell sensor of an electric field**

We propose the overall hypothesis that ENaC functions as a proximal, cell membrane sensor of the EF. Several lines of evidence from the current study suggest that ENaC is a good candidate of the EF sensor on keratinocytes. First, ENaC is a membrane channel penetrating the well-insulated plasma membrane. The extracellular domain of ENaC is exposed to the medium where the EF is carried and the intracellular c-terminus is associated with actin (Mazzochi et al., 2006). Second, depleting ENaC from keratinocytes abolishes the galvanotaxis responses in keratinocytes (90% diminution in directedness). Inhibiting or deletion of other molecules known to be involved in the galvanotaxis response, such as P3K-γ/p110γ and p105 or β4-integrin only result in a loss of 50% of the directionality (Pullar et al., 2006; Zhao et al., 2006). Partial loss of directionality when these signaling pathways are impaired suggests that parallel pathways converge to mediate galvanotaxis. However, the almost complete loss of directionality of ENaC knockout or knockdown cells implies that ENaC might be the most proximal signal of galvanotaxis signaling cascade initiating the directional response, and thus in the absence of ENaC, none of the downstream signaling pathways can be activated to induce galvanotaxis. Also, unlike the other mediators noted above, ENaC is not involved in the migratory machinery. Knocking out ENaC from keratinocytes does not impair cell motility and the cells maintain lamellipodial extension, albeit in a non-directional, random migration pattern in
the EF, by an ENaC-independent mechanism. ENaC in the wild-type cells may function to orient the persistent migration much like a steering molecule, which then directs the ENaC-independent mechanism of locomotion. ENaC can also be opened by shear stress from stretch activation or fluid flow (Awadya et al., 1995; Carattino et al., 2004; Carattino et al., 2005; Wang et al., 2009; Fronius et al., 2010). Therefore, one may speculate that ENaC on the keratinocyte cell surface is susceptible to other mechanical forces and that voltage change generated by the EF may induce conformational changes in ENaC, resulting in a galvanotaxis response.

In summary, we found that ENaC plays an important, and until now undiscovered, role in mediating the directional migration of keratinocyte in an EF. Here we propose a model for ENaC in mediating galvanotaxis. When an EF such as those endogenously generated in wounded skin is applied to keratinocytes, ENaC in the open state, located at the surface of migrating cells senses the field and promotes galvanotaxis by stabilizing the lamellipodial protrusions of the cathodal side and inhibiting the extensions at the anodal side. Galvanotactic cells then orient and turn slowly toward the cathode by means of their stable cathodal protrusions. Further studies are required to fully elucidate the mechanism by which ENaC mediates the protrusion process in response to an EF. Nevertheless, this novel function for the channel in the epidermis has broad implications for epithelial wound repair and the broad range of physiological responses of cells to endogenous EFs.

Materials and Methods

Cell culture and eNAC knockout mice

Human and mouse keratinocytes

Isolation of human foreskin keratinocytes was as described (Pullar and Isseroff, 2005) using a protocol approved by the UC Davis IRB. Neonatal human keratinocytes were cultured in EpiLife medium with 1:

Chemicals, siRNA and antibodies

**Chemicals**

Phenamil (Sigma–Aldrich), Cariporide (a NHE1 specific inhibitor, gift from Dr Juegen Punter, Sanofi–Aventis, Germany), S3969, (an ENaC activator, gift from Dr Bryan Moyer, Menominy, CA) and DMSO (final dilution of 1:2500 to 1:5000) were added to NHK for 30 minutes before the electric field was applied.

**siRNA**

Silencer® Pre-designed siRNA, C3ˈ3ˈlabeled (Cat#AM16811, siRNA ID# 6972, human SCNN1A, 5ˈ-C3 GGUACC CGGAAUUUAGATT-3ˈ) against εENaC and Silencer® C3ˈ3ˈlabeled Negative Control #1 siRNA (Cat# 1621) were purchased from Ambion (Austin, TX). The siRNA was applied to NHK using the protocol described and an additional treatment of the siRNA was followed 24 hours after the first treatment. Protein levels of ENaC and galvanotaxis were monitored 72 hours after the second siRNA treatment.

**ENaC inhibitory peptides**

Scrambled control peptide (QLHLPPPR) and ENaC inhibitory peptide (LPHEPLQLR) (Carattino et al., 2006; Carattino et al., 2008) with acetylated and amidated ends were ordered (HPLC purified, purity >90%) from GenScript (Piscataway, NJ). NHKs were pre-treated with 20 μM peptides for 1 hour before the galvanotaxis assay.

**Antibodies**

Anti-εENaC aa627-643 rabbit antibodies (Calbiochem/EMD Chemicals, Gibbstown, NJ, 1:1000 for Western blotting and 1:200 for immunostaining), anti-rabbit secondary antibodies, HRP-linked (Cell signaling Tech., Danvers, MA, 1:1000 for western blotting) or Alexa Fluor 488 donkey anti-rabbit IgG (HˈL) secondary antibodies (Invitrogen, 1:500 for immunostaining) were used.

**Galvanotaxis**

Galvanotaxis was performed on passage 2–5 cells from the primary cultures as described (Pullar et al., 2006). The galvanotaxis chambers were pre-coated with 100 μg/ml collagen I solution (Sigma–Aldrich) for cell migration. Cells were detached by 0.25% trypsin–EDTA (Invitrogen), neutralized with 10% FBS, centrifuged, seeded on the pre-coated galvanotaxis chambers at 6–8×10³ cell/ml until cells are attached (2–3 hours for human and mouse keratinocytes; 6 hours for H441, GFPP, and εC3-3-GFP, and 12 hours for PSN and E266I). Chemical treatments were applied to the cells for 30–60 minutes prior to the exposure of a 100 mV/mm DC electric field (EF), comparable to the physiological range at the wound field. Cell migration in the electric field was monitored for 60 minutes except in the experiments with NHE1 expressing fibroblasts PSN and E266I, where galvanotaxis was monitored for 3 hours in an electric field of 200 mV/mm. In each treatment group, 80–120 cells from 3–4 primary cell culture strains (n = 3–4) or from three repeats of a cell line (n = 3) were tracked.

**Western blotting and immunostaining**

**Western blotting**

NHK were rinsed with PBS and scrapped from the culture plates with Costar Cell Lifters (Corning, Corning, NY) in 100–200 μl of lysis solution containing protease inhibitor cocktails (RIPA buffer (Sigma–Aldrich), 50 mM NaF, Sigma–Aldrich, 1 mM Na3VO4 (Sigma–Aldrich), 1X Complete Protease Inhibitor Cocktail (Roche, Basel, Switzerland), 0.2 mg/ml PMSF (Sigma–Aldrich), and 0.5% Triton X-100 (Sigma–Aldrich). Cell lysates were spun at 12,000 rpm at 4°C for 10 minutes and the supernatants were stored at −20°C. Protein concentration in the lysates was determined by Bio-Rad protein assay (Bio-Rad, Hercules, CA) and 30 μg protein from each sample was loaded. SDS-PAGE and western blotting were performed with NuPAGE® Novex Gel Electrophoresis system (4–12% gels, Invitrogen) and iBlot® Dry Blotting System (Invitrogen). The blot was then blocked with 5% skim milk, incubated with primary antibodies for overnight at 4°C, washed and incubated with secondary antibodies for 2 hours at room temperature. ECL Western Blotting Detection Reagents (GE Healthcare, Little Chalfont, UK) was used to develop signals on X-ray films (Fujifilm, Tokyo, Japan). The protein bands on the blots were scanned and quantified by ImageJ program.

**Immunostaining**

After the exposure to the EF for different time periods, NHK attached to coverslips were fixed in 10% formaldehyde for 15 minutes. Cells were blocked by 4% BSA (BSA, Sigma–Aldrich), incubated with primary antibodies overnight at 4°C, washed, and incubated with secondary antibodies conjugated with fluorochromes for 2 hours at room temperature. The coverslips were mounted with ProLong® Gold antifade reagent with DAPI (Invitrogen).

**Image acquisition and analysis**

**Image acquisition**

Images were acquired on a Nikon TE-2000 microscope with a motorized stage, an environmental chamber to maintain at 37°C, 20× (Nikon Plan Fluor, NA=0.45, for
time-lapse imaging) and 60x (Nikon Plan Apo VC, NA=1.40 oil, for immunofluorescence imaging) objectives, a Retiga EX camera (Qmaging, Canada) and OpenLab and Volocity imaging software (PerkinElmer, Waltham, MA).

Quantification of galvanotaxis
For the fast moving cells, such as keratinocytes and H441 cells, time-lapse sequences of cell locomotion were obtained at 5-minute intervals for 1 hour. For slow moving cells, such as the NHE1 cells, the time-lapse movies were acquired at 15-minute interval for 3 hours to allow the cells to move a distance of at least 40–50 μm. Dividing cells in the movies were excluded from analysis since they were not motile until the end of mitosis. Cell tracking was performed with OpenLab software and the cell migration rate and directionality of galvanotaxis were calculated and shown as mean ± s.e.m. In galvanotaxis, cell directionality was represented by the cosine value of trajectory of cell migration relative to the orientation of the EF. If the cells migrate directly toward the cathode or anode (galvanotaxis), the cosine value will be +1 (for cathodal migration) or −1 (for anodal migration). If cells migrate with random directionality, the average of net cosine will be close to zero. The final data were calculated and plotted with FileMaker Pro 8 Advanced (FileMaker, Santa Clara, CA) and Excel (Microsoft, Redmond, WA). The range of the keratinocyte migration speed is 1.0–2.0 μm/minute and the range of the cosine values for keratinocyte galvanotaxis is 0.6–0.9 depending on the cell strains from different donors.

Quantification of immunostaining
To quantify the immunostaining, fan-shaped or bipolar-shaped cells extending lamellipodia to both the cathodal and anodal sides symmetrically in the EF were selected for comparison. Cathodal and anodal sides of the keratinocytes were defined by a line bisecting the cell relative to its orientation in the electric field. For the peripheral staining of αENaC, the fluorescence intensity at the keratinocyte edge was measured and averaged.

Kymograph analysis
To track lamellipodial dynamics during galvanotaxis, fan-shaped cells facing perpendicularly to the axis of the electric field were filmed under a 60x oil objective. The change of the leading edge of the cells was monitored every 2 seconds for 10 minutes in the electric field. To analyze the images of the leading edges, the cell length was divided to 10 portions and 11 of the 1-pixel wide lines were drawn parallel to the migration direction at the cell periphery. Each cell is divided in half from the middle line to delineate the right, cathodal side and the left, anodal facing side. The kymographs were generated from the images of the stacked lines from the time-lapse movies with NIH ImageJ software (Fig.3A). Protrusions are defined as extensions of the cell periphery and the retractions as shrinking of the cell periphery (Libbette et al., 2001; Borm, et al., 2005; Tabouhi et al., 2007). The distance and slope (the speed) of protrusions and retractions in each kymograph were calculated with the kymograph tools in ImageJ. To perform the statistical analysis, the kymograph data from the anodal side were compared to the counterpart from the cathodal side. For example, the kymograph data from line 1 is compared with lines 2, line 4 with line 5 and line 8 with line 7 etc. Nine to thirteen cells were analyzed in each group.

Statistics
All the experiments were repeated 3–4 times and the bar graphs compared using the Student’s t-test in Excel. If the P value is smaller than 0.05, the results between the two treatment groups are considered significantly different.

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**Fig. S1. Alpha-ENaC expressed in cells.** Immunoblots of cell lysates for alpha-ENaC expression. (A) αENaC in MEK. Alpha-ENaC protein (labeled with *) is depleted in αENaC-KO cells (the density of the bands on the right panel). The middle bands about 65kDa appearing in both wild type and αENaC-KO cells are non-specific. (B) αENaC in siRNA-treated NHK. The densitometric analysis showed that the αENaC-siRNA treatment knocked down the protein level by 50% in NHK (right lane) when it is compared to the scrambled-siRNA treatment. (C) αENaC in H441, GFP7 and C3-3 cells. Endogenous αENaC is visualized as the lower molecular weight bands (labeled with *) present in the lysates of the H441, GFP7 and C3-3 cells, and a higher molecular weight band of the GFP-tagged-αENaC band (labeled with #) is expressed in the C3-3 cells. The total ENaC in C3-3 increased by 5-fold compare to the parental H441 cells (right panel). GAPDH and tubulin were used as loading controls.
Fig. S2. Inhibiting ENaC blocks re-epithelialization of ex vivo skin wound. Normal adult human skin tissue obtained from elective plastic surgical abdominal reduction procedures, with a protocol approved by the UC Davis IRB, was trimmed to remove subdermal and adipose tissue. A 3 mm wide linear lesion was created on the epidermis and the wounded skin tissue was incubated in DMEM containing 10% FBS for 7 days. The tissue was fixed, embedded in paraffin for sectioning and staining with H&E. Wound images were taken, the length of the whole wound and re-epithelialized areas were measured to calculate the percentage of re-epithelialization. The arrows indicate the original wound edge. (A) In the control wound, re-epithelialization occurred and keratinocytes resurfaced the wound (the fine line of stained epithelium between 2 arrows). (B) In the DMSO solvent control wound, keratinocytes also migrated and resurfaced the wound. (C) In the presence of phenamil, re-epithelialization (the dotted line) was limited and keratinocytes migrate only a short distance into the wound. (D) Re-epithelialization of the ex vivo wound is blocked by phenamil by 70% on day 7 ($P<0.05$, tissue from 4 patients) when compared with the DMSO control treatment. Scale bar: 250 μm (panels A,B,C).
Fig. S3. Viability of keratinocytes in the ex vivo wound and in culture with phenamil. (A) Human wounded skin tissue that was treated with either DMSO or 10 μM phenamil, was paraffin-embedded and sectioned on day 7. Immunohistochemistry with anti-PCNA to label proliferating cells was performed. The migrating epithelial tongue was observed at the wound edge of the tissue. In both groups, PCNA stained keratinocyte nuclei at the leading edge of migrating epidermis (arrows), demonstrating that cells proliferated during the period of wound re-epithelialization, whether in control solvent DMSO or 10 μM phenamil. The nuclear staining is shown in the DAPI images and the wound edge is shown in the phase contrast images. Scale bar: 100 μm. (B) Using the activity of extracellular lactate dehydrogenase (LDH) as an indicator of cell damage, the viability of human cultured keratinocytes exposed to 10 μM phenamil in DMSO was compared to control cells exposed to the same concentration of DMSO alone. LDH levels are similar in treated and control cells through day 4 of exposure, and then rise on day 6. Thus, there is cytotoxicity to the cultured keratinocytes from the prolonged phenamil exposure at this dose, that may be limiting migratory ability of the cells in the ex vivo model at later times of incubation.
Movie 1. Galvanotaxis of mouse keratinocytes. MEK were exposed to an EF of 100 mV/mm. Cell migration was filmed for 60 min by time-lapse capture of images using phase contrast microscopy. The direction of the EF is from the bottom (anode) to the top (cathode) of the frames and galvanotaxic keratinocytes migrated directionally to the cathode. Track lines and direction of cell migration are shown in the end of the movie. Scale bar: 100 μm.

Movie 2. Galvanotaxis of αENaC-KO-MEK. Alpha-ENaC-KO-MEKs were exposed to an EF of 100 mV/mm. Cell migration was filmed for 60 min by time-lapse capture of images using phase contrast microscopy. The direction of the EF is from the bottom (anode) to the top (cathode) of the frames and keratinocytes lost the directionality and migrated randomly. Track lines and direction of cell migration are shown in the end of the movie. Scale bar: 100 μm.
Movie 3. Galvanotaxis of human keratinocytes with scrambled siRNA treatment. Scrambled siRNA treated NHKs were exposed to an EF of 100 mV/mm. Cell migration was filmed for 60 min by Hoffman modulation contrast microscopy. The direction of the EF is from the bottom (anode) to the top (cathode) of the frames and galvanotaxic keratinocytes migrated directionally to the cathode. Track lines and direction of cell migration are shown in the end of the movie. Scale bar: 100 μm.

Movie 4. Galvanotaxis of human keratinocytes with αENaC siRNA treatment. Alpha-ENaC siRNA treated NHKs were exposed to an EF of 100 mV/mm. Cell migration was filmed for 60 min by Hoffman modulation contrast microscopy. The direction of the EF is from the bottom (anode) to the top (cathode) of the frames and keratinocytes lost the directionality and migrated randomly. Track lines and direction of cell migration are shown in the end of the movie. Scale bar: 100 μm.