Nhe1 Is Essential for Potassium but Not Calcium Facilitation of Cell Motility and the Monovalent Cation Requirement for Chemotactic Orientation in Dictyostelium discoideum

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In Dictyostelium discoideum, extracellular K⁺ or Ca²⁺ at a concentration of 40 or 20 mM, respectively, facilitates motility in the absence or presence of a spatial gradient of chemoattractant. Facilitation results in maximum velocity, cellular elongation, persistent translocation, suppression of lateral pseudopod formation, and myosin II localization in the posterior cortex. A lower threshold concentration of 15 mM K⁺ or Na⁺ or 5 mM Ca²⁺ is required for chemotactic orientation. Although the common buffer solutions used by D. discoideum researchers to study chemotaxis contain sufficient concentrations of cations for chemotactic orientation, the majority contain insufficient levels to facilitate motility. Here it has been demonstrated that Nhe1, a plasma membrane protein, is required for K⁺ but not Ca²⁺ facilitation of cell motility and for the lower K⁺ but not Ca²⁺ requirement for chemotactic orientation.

Extracellular Ca²⁺ and K⁺, ubiquitous in the soluble environment of cells, both free-living and in multicellular organisms (5, 15, 21, 36, 56, 79, 80, 84, 91), have long been known to affect the motility of cells ranging in complexity from bacterial motility (5, 15, 21, 36, 56, 79, 80, 84, 91), have long been known to affect the motility of cells ranging in complexity from bacterial to human (7, 8, 19, 25, 31, 60, 62, 65, 98). In Dictyostelium discoideum, which serves as a powerful model for studying animal cell motility (4, 6, 20, 34, 41, 42, 43, 45, 50, 82, 87, 86, 92, 98), either extracellular K⁺ or Ca²⁺ at an optimum concentration of 40 mM or 20 mM, respectively, enhances most aspects of basic cell motility in the absence of chemoattractant, including cell elongation, uropod formation, the suppression of lateral pseudopod formation, velocity, directional persistence, and localization of myosin II in the posterior cell cortex (51, 78). These facilitating concentrations of K⁺ or Ca²⁺ cannot be replaced with other monovalent or divalent cations (51) and are within the range of the soluble concentrations of the two cations found in soil (1, 5, 32, 44, 49, 56) and manure (9, 66, 83), two common niches in which soil amoebae thrive (12, 27, 33, 111). These concentrations of K⁺ and Ca²⁺ are close to those found to be optimum for enhancing the motility of a variety of other animal cells. For instance, 9 mM Ca²⁺ has been found to be optimum for flagellated sea urchin sperm motility (99), and Ca²⁺ ranging in concentration from 1 to 10 mM has been found optimum for vertebrate cell motility (28, 58, 63). Concentrations of K⁺ greater than 100 mM have been found to be optimum for the polarization and motility of human polymorphonuclear leukocytes (52, 69).

The surface molecules and mechanisms regulating K⁺ and Ca²⁺ facilitation in D. discoideum have not been elucidated. However, Patel and Barber (64) reported a defective behavioral phenotype for cells of the nhe1 null mutant in a facilitating concentration of K⁺ that was strikingly similar to the behavior of wild-type cells in a nonfacilitating concentration of K⁺ (51). Nhe1 is a plasma membrane protein related to cation/H⁺ exchangers (64). We therefore tested the possibility that Nhe1 mediated K⁺ facilitation. First, to define and illustrate the process of cationic facilitation, we compared cell motility and chemotaxis among buffer solutions containing a facilitating concentration of K⁺ or Ca²⁺ and buffer solutions commonly used by D. discoideum researchers to study motility and chemotaxis. The majority of the latter solutions contain nonfacilitating cation concentrations. The comparative study revealed that the majority of common buffer solutions support chemotactic orientation but are not optimum for motility, thus illustrating cationic facilitation. This comparison also revealed for the first time that a threshold concentration of 15 mM K⁺ or Na⁺ or a threshold concentration of 5 mM Ca²⁺ was required for chemotactic orientation. We then tested the role of Nhe1 by analyzing the nhe1 null mutant. Our results revealed that Nhe1 mediates both K⁺ facilitation and the K⁺/Na⁺ requirement for chemotactic orientation, but plays no role in Ca²⁺ facilitation or the Ca²⁺ requirement for chemotactic orientation. The possible mechanisms through which Nhe1 might mediate these monovalent cation effects, including the possibility that Nhe1 acts as a monovalent cation sensor, are discussed.

MATERIALS AND METHODS

Strain maintenance and development. The parent strain Ax2, the nhe1⁻ deletion strain, and the nhe1⁻/nhe1⁺ (nhe1⁻/act15::nhe1::HA) complemented strain were obtained from the Dictybase stock center (http://dictybase.org/StockCenter/StockCenter.html) and subcloned. The mutant strains were originally generated by Patel and Barber (64). To select for mutant strains, the nhe1⁻ cells were grown in the presence of 10 µg/ml of blasticidin S (Sigma-Aldrich, St. Louis, MO), and nhe1⁻/nhe1⁺ cells were grown in blasticidin S plus 10 µg/ml G418 (Sigma-Aldrich, St. Louis, MO). To obtain aggregation-competent amoebae, cells were developed on filters according to methods described previously (76, 94, 96). Developmental filters were saturated with a buffer containing 20 mM KCl, 2.5 mM MgCl₂, 20 mM KH₂PO₄, and 5 mM Na₂HPO₄, pH 6.4 (81). For

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experimental purposes, cells were washed four times in the above buffer solution or in Tricine buffer (TB) (pH 7.0) (50).

**Analysis of basic motile behavior and chemotaxis.** Cells were perfused with test solution in a Sykes-Moore chamber according to the methods previously described (8). For chemotaxis, cells were distributed on the bridge of a Plexiglas gradient chamber according to methods previously described (51, 89). Gradients were generated by filling one trough bordering the bridge with buffer solution lacking Ca\(^{2+}\) and in the other trough with buffer solution containing 10 mM cAMP. Cell were incubated for 5 min, and then images were digitally acquired every 4 s for a 10-min period.

**DIAS analysis.** Cells images were digitally acquired and analyzed with two-dimensional (2D)-DIAS software (DIAS) as previously described (77, 86). Movements were exported to a QuickTime format and imported into DIAS. Instantaneous velocity, percent cells moving at >9 \(\mu\)m per min, directional persistence, chemotactic index (CI), and percent positive chemotaxis were computed from centroid positions (75, 77). “Instantaneous velocity” was computed as the average velocity (distance between two consecutive centroids divided by interval time) for two consecutive intervals (54). It is computed for the two intervals \(a\) to \(b\) and \(b\) to \(c\) and then \(b\) to \(c\) and \(c\) to \(d\). It then prescribes the average of each pair to frame \(b\) and \(c\), respectively. It is a method for smoothing the velocities computed for consecutive intervals (77). The parameter \% cells \(\geq 9 \mu\)m/min was computed as the proportion of cells in the classified population moving with an instantaneous velocity equal to or greater the 9 \(\mu\)m per minute. “Directional persistence” was computed as the net distance between the first and last centroids of a centroid track divided by the summed distances between all intervals. The intervals between centroids for calculating motility and chemotaxis parameters were 4 s and are based on the stability of persistence measurements, which was consistent with a subsequent methods paper (D. R. Soll, D. Wessels, and S. Kuhl, unpublished data). The “chemotactic index” was computed as the net distance moved up a gradient divided by the total distance (the sum of the distances at 4-s intervals). A CI of \(-1.00\) represented direct movement down the gradient, +1.00 represented direct movement up the gradient, 0.00 represented random movement, and +0.20 to +0.99 represented positive chemotaxis. The \% positive chemotaxis was computed as the proportion of a cell in a population with a chemotactic index greater than 0.00. “Positive flow” was computed by calculating the region in the second of two consecutive cell parameter images not overlapping the first and converting that area to the percentage of the area of the first image. The interval between centroids for calculating motility and chemotaxis parameters was 4 s and is based on the stability of persistence measurements, which was consistent with a subsequent methods paper. (Soll et al., unpublished).

The percent suppression of lateral pseudopods was determined from the expansion zones present in different pictures generated by DIAS according to the methods of Wessels et al. (96).

**Strains expressing MhcA-GFP.** To generate a nhe1– derivative expressing MhcA-green fluorescent protein (GFP), the plasmid containing mhca-GFP (57) was isolated from the nhe1A/mhca-GFP strain, a generous gift from Thomas Egelhoff of Case Western University, Cleveland, OH, using the Rapid-Plasmid kit (Marligen Biosciences, Ijamsville, MD) according to the manufacturer’s instructions. The plasmid, containing a neomycin resistance marker (61), was transformed into DH5a cells (Invitrogen, Carlsbad, CA) and subcloned, and the isolated plasmid was sequenced. The plasmid was then transformed into both the parental strain, Ax2, and the nhe1– deletion mutant according to the methods of Eichinger and Rivero (30). The concentration of G418, a substitute for neomycin, was subsequently increased over a period of 2 to 3 weeks from 2 to 20 mg/ml to select for transformants.

**Fluorescence imaging of GFP-labeled cells.** Live imaging of GFP-labeled cells was performed as previously described (94) using a Bio-Rad Radiance 2100 MP laser scanning confocal microscope and a 40× Nikon 5 Fluor 1.30 oil objective. Simultaneous fluorescent and differential interference contrast (DIC) images were generated using the 488-nm and 476-nm argon lasers, respectively. The LaserSharp 2000 (release 5.2) software program was used for image acquisition and for measuring the pixel intensities of line profiles across labeled cells as described by Lusche et al. (51).

**Nucleotide sequence accession numbers.** The GenBank accession numbers for Nhe1 to -4 cDNAs from *D. discoideum* strain Ax2 are as follows: for Nhe2 (NhekB), HQ730876; for Nhe3 (NhekC), HQ730877; and for Nhe4 (NhekD), HQ730878.

**RESULTS**

**Cationic facilitation of motility.** To obtain optimum motility, either the Ca\(^{2+}\) concentration must be 10 to 20 mM or the K\(^+\) concentration 40 mM (51). Combining 20 mM Ca\(^{2+}\) and 40 mM K\(^+\) does not increase motility parameters above that achieved by either alone (see Table S1 in the supplemental material). We have referred to the optimization of motility by 20 mM Ca\(^{2+}\) or 40 mM K\(^+\) as “the cationic facilitation of cell motility.” Since the objective here was to test whether Nhe1 specifically mediated K\(^{+}\) facilitation, it was paramount that cationic facilitation was clearly defined and illustrated. One way to do this was to compare cell behavior in solutions containing a facilitating concentration of K\(^{+}\) or Ca\(^{2+}\) with cell behavior in buffers commonly used by researchers of *D. discoideum*, the majority of which do not contain facilitating concentrations of K\(^{+}\) or Ca\(^{2+}\). This comparison could not be performed retrospectively from the published literature since the conditions and strains used in different studies varied dramatically. Moreover, a variety of methods have been used to calculate motility parameters, which can compromise comparisons. Differences in the interval time between data points (see Fig. S1A and B) and the velocity thresholds for subtracting nonmotile cells (see Fig. S1C and D) can have a significant impact on computed motility parameter.

Therefore, to compare cell behavior in different buffer solutions, we used the single laboratory strain Ax2, uniform culture conditions, the same perfusion chamber for assessing behavior in the absence of a chemoattractant (88), the same gradient chamber for assessing chemotaxis (89), the same interval time (4 s) between data points, and the same instantaneous velocity threshold (≥3 \(\mu\)m per min) for identifying motile cells in a population (77). The three motility parameters (average instantaneous velocity, percent cells with speeds of ≥9 \(\mu\)m per min, and average percent positive flow) are defined in Materials and Methods. In the absence of cAMP, these parameters were maximal in 20 mM Ca\(^{2+}\) solution, as they were in 20 mM Ca\(^{2+}\) in a cAMP gradient (Fig. 1A and B, respectively). We therefore used the values in 20 mM Ca\(^{2+}\) solution as the references for comparison with cells translocating in the other solutions. In the absence of a chemoattractant, the three motility parameters in 40 mM K\(^{+}\) solution and in the buffer solution (BB) used by Patel and Barber (61), were below that in 20 mM Ca\(^{2+}\) solution but still on average higher than that in the six other buffer solutions (Fig. 1A). Of the six other buffer solutions, the mean instantaneous velocity was highest in KK buffer (Fig. 1A) (45), which contained the highest concentration of K\(^{+}\) (24.3 mM) (Table 1). The motility parameters in the five test buffers BS, DB, SS, PB, and DB–DC were no better on average than that in Tricine buffer (TB) alone (Fig. 1A), which contains a nonfacilitating K\(^{+}\) concentration of 5 mM (Table 1). Cell shape also reflected facilitation. Cells translocating in 20 mM Ca\(^{2+}\) solution or 40 mM K\(^{+}\) solution were more elongate on average than cells in the other test solutions (Fig. 1C).

When a spatial gradient of cAMP was generated in the tested buffer solutions, the three measured motility parameters were maximal in 40 mM K\(^{+}\) solution and 20 mM Ca\(^{2+}\) solution (Fig. 1B). Two of the three parameters were close to maximal in solution BB (64) (Fig. 1B). All three parameters were reduced in the remaining six buffer solutions and TB (Fig. 1B). In marked contrast, the chemotactic index (CI) was on average as high in the seven test solutions as it was in 20 mM Ca\(^{2+}\) or 40 mM K\(^{+}\) solution (Fig. 1B). These results indicate that although
the majority of commonly used buffer solutions do not contain high enough cation concentrations to facilitate motility, they all support chemotactic orientation maximally and similarly.

The low CI in TB compared to that in the other test buffers (Fig. 1B) also suggested that a monovalent cation concentration threshold also existed for maximum chemotactic orientation and that this threshold was lower than that for the facilitation of motility. To identify the threshold for orientation, we compared the CIs of cells in TB containing 5, 10, and 15 mM Na\(^+\) (Table 1). The CI in 15 mM Na\(^+\) was significantly different from the lower CIs in 5 and 10 mM Na\(^+\) (\(P = 0.05\) and 0.002, respectively). Together, our results indicate that a monovalent cation (K\(^+\) and/or Na\(^+\)) concentration threshold of 15 mM is necessary for maximum chemotactic orientation. Interestingly, all of the buffer solutions empirically formulated by various researchers to study *D. discoideum* chemotaxis contained concentrations of K\(^+\) and/or Na\(^+\) greater than or equal to 15 mM (Table 1).
We previously demonstrated that in a low concentration of monovalent cations, a Ca\(^{2+}\) threshold also existed for optimum chemotactic orientation (51). This threshold was 5 mM, again below the threshold concentration for Ca\(^{2+}\) facilitation of motility. These results, therefore, illustrate the cationic facilitation of cell motility and reveal that most researchers use buffers containing concentrations of cations that are suboptimal for cell motility. These results also reveal these buffers do contain concentrations of Na\(^+\) and/or K\(^+\) equal to or above the threshold for maximum chemotactic orientation.

**Nhe1, a candidate for mediating K\(^+\) facilitation.** We considered the possibility that the putative monovalent cation/H\(^+\) exchanger Nhe1 was the surface molecule mediating the facilitation of motility and the monovalent cation requirement for chemotactic orientation, because the behavioral phenotype of the nhe1\(^-\) null mutant in a facilitating concentration of K\(^+\) (64) was similar to that of wild-type cells in a nonfacilitating concentration of K\(^+\) (51). A BLAST search (http://blast.ncbi.nlm.nih.gov/Blast.cgi) of the *D. discoideum* genome revealed that Nhe1 was a member of a family of four related proteins, Nhe1 (NheA), Nhe2 (NheB), Nhe3 (NheC), and Nhe4 (NheD). Patel and Barber (64) previously noted that Nhe1 was a member of a family of chemosensory receptors in *Caenorhabditis elegans*, according to a motif scan at http://myhits.isb-sib.ch (40) and confirmed with the Worm Database (http://www.wormbase.org) (70, 85) (Fig. 2B, blue box). We have used the same Nhe1, rather than NheA, the correct name according to *D. discoideum* nomenclature (http://dictybase.org), because Nhe1 has been used by Barber and coworkers in previous papers (24, 64).

**K\(^+\) facilitation is lost in nhe1\(^-\) cells in the absence of cAMP.** If Nhe1 is the plasma membrane protein that mediates K\(^+\) facilitation, then deleting it should result in the loss of K\(^+\) facilitation. Cells of the parental strain Ax2 and a complemented nhe1 strain, the nhe1\(^-\)/nhe1\(^+\) strain, translocated with mean instantaneous velocities of 8.0 \pm 3.0 and 8.2 \pm 3.6 \mu m per minute, respectively, in 40 mM K\(^+\) solution (Fig. 3A). In marked contrast, nhe1\(^-\) cells translocated with a mean instantaneous velocity of 3.9 \pm 0.8 \mu m per minute, less than half that of control cells and only 0.9 \mu m per minute above the threshold of 3 \mu m per minute used to discriminate motile cells (Fig. 3A).

The velocity defect in the velocity of nhe1\(^-\) cells was most evident in comparisons of high-end velocities (i.e., the proportion of cells with velocities of \(\geq 9 \mu m\) per minute). While the proportions of cells of the parental and complemented control strains were 36 and 46%, respectively, the proportion of nhe1\(^-\) cells was 0% (Fig. 3A). Finally, the directional persistence of nhe1\(^-\) cells, defined in Materials and Methods, was 33% lower than that of Ax2 cells and 52% lower than that of nhe1\(^+\)/nhe1\(^+\) cells (Fig. 3A). These behavioral defects were evident in perimetre tracks of nhe1\(^-\) cells in 40 mM K\(^+\) solution in the absence of chemoattractant (Fig. 3B). The motile phenotype of nhe1\(^+\) cells translocating on a substrate in 40 mM K\(^+\) solution in the absence of a chemoattractant (Fig. 3A and B) was therefore highly similar to that of parental control cells in TB alone containing K\(^+\) at the nonfacilitating concentration of 5

### TABLE 1. Compositions of tested buffer solutions

| Buffer solution (abbr\(^a\)) | Composition | [Na\(^+\)] (mM) | [K\(^+\)] (mM) | [Total monovalent cations] (mM) | [Ca\(^{2+}\)] (mM) | [Mg\(^{2+}\)] (mM) | Reference |
|-----------------------------|-------------|----------------|---------------|--------------------------------|------------------|----------------|-----------|
| Tricine buffer (TB) | 5 mM Tricine, 5 mM KCl | 5 | 5 |  |  |  | 51 |
| 20 mM Ca\(^{2+}\) solution (20 Ca\(^{2+}\)) | 5 mM Tricine, 5 mM KCl, 20 mM CaCl\(_2\) | 5 | 5 | 20 |  |  | 51 |
| 40 mM K\(^+\) solution (40 K\(^+\)) | 2.5 mM MgCl\(_2\), 20 mM KH\(_2\)PO\(_4\), 20 mM KCl, 5 mM Na\(_2\)HPO\(_4\) | 5 | 40 | 45 | 2.5 |  | 51 |
| Bonner solution (BS) | 2.7 mM CaCl\(_2\), 10 mM KCl, 10 mM NaCl | 10 | 10 | 20 | 2.7 |  | 11 |
| Developmental buffer (DB) | 5 mM Na\(_2\)HPO\(_4\), 5 mM NaH\(_2\)PO\(_4\), 2 mM MgSO\(_4\), 200 \mu M CaCl\(_2\) | 15 | 15 | 0.2 | 2.5 |  | 23 |
| Sorensen buffer (SB) | 2.7 mM Na\(_2\)HPO\(_4\), 14 mM KH\(_2\)PO\(_4\), 3.9 mM K\(_2\)HPO\(_4\), 16.5 mM KH\(_2\)PO\(_4\), 2 mM MgSO\(_4\), 0.1 mM CaCl\(_2\) | 5.4 | 14 | 19.4 |  | 100 |
| Phosphate buffer (PB) | 4 mM Na\(_2\)HPO\(_4\), 7.4 mM KH\(_2\)PO\(_4\), 8 | 7.4 | 15.4 |  | 14 |
| DB minus divalent cations (DB-DC) | 5 mM Na\(_2\)HPO\(_4\), 5 mM NaH\(_2\)PO\(_4\) | 15 | 15 |  | 55 |
| Barber buffer (BB) | 20 mM KH\(_2\)PO\(_4\), 20 mM K\(_2\)HPO\(_4\) | 60 | 60 |  | 64 |

\(a\) abbr, abbreviation.
These results demonstrate that in the absence of cAMP, the motile behavior of \textit{nhe1}/H11002 cells was not facilitated by 40 mM K\textsuperscript{+}/H11001 (51). Ca\textsuperscript{2+} facilitation is intact in \textit{nhe1}/H11546 cells in the absence of chemoattractant. If Nhe1 specifically mediates K\textsuperscript{+} facilitation, then deletion of the gene \textit{nhe1} should not affect Ca\textsuperscript{2+} facilitation. We tested this prediction in 20 mM Ca\textsuperscript{2+}/H11001 solution in the absence of a chemoattractant. The mean velocities of parental and complemented control cells were 10.5\pm 4.3 and 8.5\pm 2.9 m per min, respectively (Fig. 3A). The mean velocity of \textit{nhe1}/H11002 cells was 9.2\pm 5.0 m per min (Fig. 3A), which was statistically indistinguishable from that of the two control strains. The velocity in 20 mM Ca\textsuperscript{2+}/H11001 solution was more than twice that in 40 mM K\textsuperscript{+}/H11001 solution (Fig. 3A). The percentage of \textit{nhe1}/H11002 cells with velocities of \textless 9 m per min in 20 mM Ca\textsuperscript{2+} solution was high (50%), like that of control strains (67 and 25%, respectively) (Fig. 3A). It differed markedly from that of \textit{nhe1}/H11002 cells in 40 mM K\textsuperscript{+} solution, which was 0% (Fig. 3A). The directional persistence parameter of \textit{nhe1}/H11002 cells in 20 mM Ca\textsuperscript{2+} solution was also similar to that of control cells (Fig. 3A). The similarity between the behavior of \textit{nhe1}/H11002 cells and that of control cells in 20 mM Ca\textsuperscript{2+} solution was evident in a comparison of perimeter tracks (Fig. 3C). A repeat of this experiment in 10 mM Ca\textsuperscript{2+} provided similar results (data not shown). Together, these results demonstrate that in the absence of a chemoattractant, deletion of \textit{nhe1} results in the selective loss of K\textsuperscript{+}, but not Ca\textsuperscript{2+}, facilitation.

**K\textsuperscript{+} facilitation and chemotactic orientation are defective in \textit{nhe1}/H11002 cells in a cAMP gradient.** In a spatial gradient of the chemoattractant cAMP generated in 40 mM K\textsuperscript{+} solution in a chamber designed based on that of Zigmond (89, 101), the velocities of \textit{nhe1}/H11002 cells were 39% and 31% lower than those of parental and complemented control cells, respectively (Fig. 4A). The proportion of cells with velocities of \textless 9 m per min was 0%, compared to 50 and 28% for the two respective control strains (Fig. 4A). Directional persistence was also far lower than that of either of the control strains (Fig. 4A). The same motility defects were observed in a cAMP gradient generated in 40 mM K\textsuperscript{+} in TB (Fig. 4A).

To assess chemotactic orientation, the chemotactic index and the percentage of cells exhibiting a positive chemotactic index (defined in Materials and Methods) were compared between the mutant and the two control strains. Cells of the \textit{nhe1}/H11002 mutant strain exhibited weak to negligible chemotactic orientation in a cAMP gradient generated in 40 mM K\textsuperscript{+} solution, formulated in phosphate buffer (Table 1) or TB. The mean chemotactic index of \textit{nhe1}/H11002 cells in the two buffers was 0.11\pm 0.28 and 0.04\pm 0.27, respectively, compared to 0.69\pm 0.14 and 0.73\pm 0.30, respectively, for parental control cells and 0.89\pm 0.11 and 0.62\pm 0.21 for complemented control cells (Fig. 4A). The percent positive chemotaxis of \textit{nhe1}/H11002 cells in 40 mM K\textsuperscript{+} in the two buffers was 42 and 55%, respectively, compared to 100 and 94%, respectively, for parental control cells and 100 and 93%, respectively, for com-
implemented control cells (Fig. 4A). Perimeter tracks reflected these results (Fig. 4B). These results demonstrate first that 40 mM K\(^+\)/H\(_{11001}\) does not facilitate motility in \(nhe1\)/H\(_{11002}\) cells in a spatial gradient of cAMP and second that \(nhe1\)/H\(_{11002}\) cells have lost not only K\(^+\)/H\(_{11001}\) facilitation but also K\(^+\)/H\(_{11001}\)-dependent chemotactic orientation in a cAMP gradient. These quantitative results obtained in gradients generated in a chemotaxis chamber are in agreement with the decrease in chemotactic efficiency observed earlier by Patel and Barber (64).

We also tested whether \(nhe1\) cells also lost chemotactic orientation in a simple phosphate buffer solution, DB–DC, used by McCann et al. (55), which contained 15 mM Na\(^+\)/H\(_{11001}\) (Table 1). The chemotactic index of \(nhe1\^-\) cells in this solution was negligible (+0.04 ± 0.29; \(n = 63\)), and the percent positive chemotaxis was 61%. Hence, neither 40 mM K\(^+\)/H\(_{11001}\) solution nor a phosphate buffer solution containing 15 mM Na\(^+\) supported chemotactic orientation in \(nhe1\^-\) cells. These results demonstrate that Nhe1 is essential not only for K\(^+\) facilitation but also for the K\(^+\) or Na\(^+\) requirement for chemotactic orientation in a cAMP gradient.

Ca\(^{2+}\) facilitation is intact in \(nhe1\^-\) cells in spatial gradient of cAMP. In contrast to the aberrant behavior observed in a spatial gradient of cAMP generated in 40 mM K\(^+\)/H\(_{11001}\) solution or in the Na\(^+\)-based buffer solution DB–DC (Table 1), both the motility and chemotactic orientation of \(nhe1\^-\) cells were indistinguishable from those of control cells in a cAMP gradient generated in 20 mM Ca\(^{2+}\)/H\(_{11001}\) solution (Fig. 4A and C). The instantaneous velocity, percent cells with velocities > 9 \(\mu\)m/min, the proportion of cells moving with average velocities > 9 \(\mu\)m/min; Dir. persist., directional persistence; ns, not significant.

![FIG. 3. Cells of the \(nhe1\^-\) mutant have lost K\(^+\) facilitation but have retained Ca\(^{2+}\) facilitation. (A) Motility parameters for parental control strain Ax2, the \(nhe1\^-\) mutant, and the \(nhe1^-/nhe1^+\) complemented control in the absence of chemoattractant either in 40 mM K\(^+\)/H\(_{11001}\) solution or 20 mM Ca\(^{2+}\)/H\(_{11001}\) solution. \(P\) values were obtained by Student’s \(t\) test. Parameters are defined in Materials and Methods. (B) 2D-DIAS-generated stacked perimeter tracks of three representative Ax2, \(nhe1^-\), and \(nhe1^-/nhe1^+\) cells in 40 mM K\(^+\)/H\(_{11001}\) solution. Perimeters were drawn at 12-s intervals over a 10-min period. The final outline of the cell is color coded blue. (C) 2D-DIAS-generated stacked perimeter tracks of three representative Ax2, \(nhe1^-\), and \(nhe1^-/nhe1^+\) cells in 20 mM Ca\(^{2+}\)/H\(_{11001}\) solution with perimeters drawn as described for panel B. No. cells, the number of cells analyzed; Inst. vel., instantaneous velocity; % cells > 9 \(\mu\)m/min, the proportion of cells moving with average velocities > 9 \(\mu\)m/min; Dir. persist., directional persistence; ns, not significant.](325)
shown to enhance cell elongation and suppress lateral pseudopod formation during wild-type cell migration \((51)\). These facilitated changes are important in attaining maximum velocity \((51, 78)\). The effects of \(40\, \text{mM} \ K^+\) and \(20\, \text{mM} \ Ca^{2+}\) on cell elongation and lateral pseudopod formation were therefore compared between \(nhe1^-\) and control cells in a cAMP gradient. Differential interference contrast microscopy revealed that either \(40\, \text{mM} \ K^+\) or \(20\, \text{mM} \ Ca^{2+}\) induced parental and complemented control cells to elongate, move in the direction of the single anterior pseudopod, suppress lateral pseudopod formation, extend new pseudopods primarily at the anterior end, and form a tapered uropod (Fig. 5A). However, while \(20\, \text{mM} \ Ca^{2+}\) induced these same changes in \(nhe1^-\) cells, \(40\, \text{mM} \ K^+\) did not (Fig. 5A).

Lateral pseudopods have previously been defined as new projections emanating from the flank of the cell body \((78, 93, 94)\) or new anterior pseudopods resulting from anterior pseudopod bifurcation, a process also referred to as “splitting” \((3)\). For a measure of the unsuppressed frequency, pseudopod formation was assessed in a cAMP gradient generated in \(1\, \text{mM} \ Ca^{2+}\) solution \((51)\). The “frequency of lateral pseudopod formation” was computed as the frequency in a test solution divided by the frequency in the reference solution, multiplied by 100. In \(40\, \text{mM} \ K^+\) solution, the frequencies of lateral pseudopod formation of parental control \((Ax2)\) and complemented control \((nhe1^-/nhe1^-)\) cells were 55 and 30%, respectively, compared to those in \(1\, \text{mM} \ Ca^{2+}\) solution (Fig. 5B). The frequency of lateral pseudopod formation of \(nhe1^-\) cells in \(40\, \text{mM} \ K^+\),
A. 

![Images of cell patterns](image)

**FIG. 5.** Cells of the mutant nhe1− do not respond to 40 mM K⁺ by elongating and suppressing lateral pseudopod formation, but they do respond to 20 mM Ca²⁺. (A) Differential interference microscopy images of three representative Ax2 (nhe1+), nhe1−, and nhe1−/nhe1− cells in spatial gradients of cAMP generated in solutions containing either 40 mM K⁺ (top panels) or 20 mM Ca²⁺ (bottom panels). Arrows at the bottom of each panel indicate the direction of the cAMP gradient. (B) Percent lateral pseudopod formation was determined by dividing the number obtained in 40 mM K⁺ or 20 mM Ca²⁺ solution for Ax2, nhe1−/nhe1−, and nhe1− cells by the number obtained for Ax2 cells in a nonfacilitating 1 mM Ca solution and multiplying the fraction by 100. Light-gray bars, Ax2 control (nhe1+); dark-gray bars, nhe1−/nhe1− complemented control; black bars, nhe1− strain.

However, was almost the same as that of control cells in 1 mM Ca²⁺ solution (Fig. 5B). These results indicate that 40 mM K⁺ does not suppress lateral pseudopod formation in nhe1− cells as it does in control cells. In 20 mM Ca²⁺ solution, however, lateral pseudopod formation in Ax2, nhe1−/nhe1−, and nhe1− cells was suppressed by 97%, 80%, and 85%, respectively (Fig. 5B). These results support the conclusion that the motile behavior of nhe1− cells is facilitated by 20 mM Ca²⁺ but not 40 mM K⁺.

**Induced cortical localization of myosin II.** In the process of cationic facilitation, myosin II localizes to the posterior cell cortex (51). Posterior localization is believed to be important for cell elongation, the suppression of lateral pseudopod formation, uropod formation, efficient translocation, and efficient chemotaxis (29, 39, 48, 73, 78, 95). One would therefore expect 20 mM Ca²⁺ solution but not 40 mM K⁺ solution to induce posterior localization of myosin II in nhe1− cells. To test this prediction, parental control and nhe1− cells were transformed with a plasmid containing GFP-tagged myosin II (57) and the pattern of GFP fluorescence was analyzed by confocal microscopy. Projections of the center five images of a Z-series of 16 scans of each cell were stacked and analyzed for combined pixel intensity through a zigzag track protocol diagrammed in the upper portion of each panel in Fig. 6 (51). Intensity plots of representative control cells in a spatial gradient of cAMP generated in 40 mM K⁺ solution (Fig. 6A) or 20 mM Ca²⁺ solution (Fig. 6C) revealed broad peaks and troughs, the former representing the scans through the cell body and the latter the extracellular regions. In both cases, the intensity was highest in the cortex of the posterior region of the cell body (Fig. 6A and C). Intensity plots of representative nhe1− cells in a spatial gradient of cAMP generated in 40 mM K⁺ solution (Fig. 6B) or 20 mM Ca²⁺ solution (Fig. 6D) also revealed broad peaks and troughs. In 20 mM Ca²⁺, the intensity was higher in the posterior cortex of the cell body of nhe1− cells (Fig. 6D), just as it was in control cells (Fig. 6C). However, in 40 mM K⁺, the intensities along a nhe1− cell were similar in the anterior and posterior cell cortex (Fig. 6B). In these cells, the posterior end of apolar nhe1− cells was identified by the position of tail fibers (38), not shown here. These results demonstrate that 20 mM Ca²⁺ but not 40 mM K⁺ induces the localization of myosin II to the posterior cortex of nhe1− cells.

**K⁺ concentration.** The preceding results demonstrating that nhe1− cells have selectively lost K⁺ facilitation were based on tests performed with 40 mM K⁺, the optimum concentration for the facilitation of wild-type cell motility (51). To test the possibility that K⁺ might facilitate motility and polarity in nhe1− cells but at a different concentration, we analyzed the effects of higher and lower K⁺ concentrations. We found that in the range of 5 to 80 mM, K⁺ did not facilitate cell motility or mediate chemotactic orientation in nhe1− cells (Table 2).

**DISCUSSION**

Previously we demonstrated that 40 mM K⁺ or 20 mM Ca²⁺ facilitated cell motility (51). To illustrate the process of facilitation, we compared motility in solutions containing facilitating concentrations of K⁺ and Ca²⁺ with that in buffer solutions commonly employed by researchers studying motility and chemotaxis in *D. discoideum*. Six of the seven contained nonfacilitating concentrations of Na⁺ and/or K⁺, ranging from 15 to 24 mM. None contained a facilitating concentration of Ca²⁺. Our results first demonstrated that the six buffer solutions were suboptimal for cell motility, but all performed as well as 40 mM K⁺ or 20 mM Ca²⁺ solution in supporting chemotactic orientation. However, chemotactic orientation in TB alone, which contains 5 mM K⁺, was reduced dramatically. This suggested that there was a monovalent cationic requirement for chemotactic orientation that was above 5 mM but below 40 mM. By varying the Na⁺ concentration, we found this cationic requirement to be 15 mM, the minimum concentration in the seven tested buffer solutions, suggesting that in the independent development of the seven buffer solutions used to study *D. discoideum* motility and chemotaxis, an unrecognized consensus had emerged that a minimum monovalent cation concentration of 15 mM was necessary for efficient chemotactic orientation. While 15 mM Na⁺ or K⁺ will fulfill the requirement for orientation, only 40 mM K⁺ will facilitate motility. Na⁺ at 40
mM is inhibitory (51). Furthermore, 5 mM Ca\(^{2+}\) also fulfills the requirement for orientation (51).

There are three possible ways in which Nhe1 may mediate the effects of extracellular monovalent cations: through changes in cytosolic pH, through changes in the concentration of cytosolic monovalent cations, or as a coupled sensor. Whatever the mechanism, Nhe1 must function by assessing differences in the steady-state concentration of extracellular monovalent cations.

Nhe1 has been classified as a Na\(^{+}/H^{+}\) exchanger (64, 90). However, the steady-state intracellular pH (pHi) of wild-type, complemented mutant, and mutant cells in the absence of cAMP was measured by Patel and Barber (64), using the fluorescent pH indicator 2',7'-bis(2-carboxyethyl)-5(6)-carboxyfluorescein (BCECF). They reported them to be 6.58 ± 0.06, 6.63 ± 0.07, and 6.48 ± 0.06, respectively. Since the pHi of complemented mutant cells differed from that of wild-type cells by half the difference between those of wild-type and nhe1\(^{-}\) cells, it was not clear if the latter difference was functionally meaningful. Moreover, the cytosolic specificity of BCECF fluorescence is in question, because it has been demonstrated that BCECF stains organelles and vesicles, as well as the cytosol (10, 18, 59, 72). Therefore, it is not clear if Nhe1

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FIG. 6. Myosin II does not localize normally in the posterior cortex of nhe1\(^{-}\) cells in chemotaxis when K\(^{+}\) is the facilitating cation, but it does when Ca\(^{2+}\) is the facilitating cation. The maximum intensity for a scan was normalized to 100%. (A) Ax2 control cells in 40 mM K\(^{+}\) solution. (B) nhe1\(^{-}\) cells in 40 mM K\(^{+}\) solution. (C) Ax2 control cells in 20 mM Ca\(^{2+}\) solution. (D) nhe1\(^{-}\) cells in 20 mM Ca\(^{2+}\) solution. a, anterior; p, posterior. Horizontal bars in each graph indicate the average peak intensities for the posterior and anterior regions of the representative cell images. Insets show the zigzag path of intensity measurements along the cell that is plotted in the corresponding graph below.
regulates intracellular pH. Alternately, Nhe1 may function by increasing intracellular monovalent cations. It has been demonstrated that incremental increases in the concentration of an extracellular monovalent cation result in increases in the cytosolic concentration of that cation (74). Therefore, Nhe1 might function by regulating the cytosolic concentration of monovalent cations. Finally, and perhaps more interestingly, Nhe1 may function as a coupled monovalent cation sensor. Nhe1 belongs to the group CPA1, members of which do not cause changes in intracellular monovalent cations. Finally, and perhaps more interestingly, Nhe1 may function by sensing concentration of that cation (74). Therefore, Nhe1 might function by increasing intracellular monovalent cations. It has been demonstrated in Tricine buffer, and each solution was equilibrated with the NIH.

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TABLE 2. The nhe1− mutant is insensitive to K+ at concentrations ranging from 5 to 80 mM

| [K+] (mM) | Strain name or description | No. of cells | Instantaneous velocity (μm/min) | Directional persistence | Chemotactic index |
|----------|-----------------------------|--------------|---------------------------------|------------------------|------------------|
| 5        | Ax2                         | 28           | 6.8 ± 1.8                       | 0.47 ± 0.22            | +0.34 ± 0.35     |
|          | nhe1                        | 19           | 3.7 ± 1.3                       | 0.15 ± 0.11            | +0.03 ± 0.12     |
| 20       | Ax2                         | 27           | 7.4 ± 2.5                       | 0.63 ± 0.17            | +0.59 ± 0.21     |
|          | nhe1                        | 36           | 4.7 ± 1.3                       | 0.42 ± 0.21            | +0.03 ± 0.34     |
| 40       | Ax2                         | 17           | 9.8 ± 3.5                       | 0.79 ± 0.23            | +0.77 ± 0.30     |
|          | nhe1                        | 43           | 4.2 ± 1.7                       | 0.35 ± 0.21            | +0.04 ± 0.27     |
| 60       | Ax2                         | 21           | 6.7 ± 2.6                       | 0.68 ± 0.21            | +0.63 ± 0.27     |
|          | nhe1                        | 45           | 3.2 ± 0.2                       | 0.37 ± 0.14            | +0.01 ± 0.28     |
| 80       | Ax2                         | 54           | 6.4 ± 3.0                       | 0.66 ± 0.24            | +0.56 ± 0.24     |
|          | nhe1                        | 40           | 3.5 ± 0.7                       | 0.37 ± 0.19            | +0.02 ± 0.25     |

* The K+ concentration was varied in Tricine buffer, and each solution was used as the buffer for generating a cAMP gradient.

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