Introduction

Chemotaxis is a pivotal response of many cells to spatial cues (Van Haastert and Devreotes, 2004; Affolter and Weijer, 2005; Wu, 2005). It plays important roles in diverse functions, such as finding nutrients in prokaryotes, forming multicellular structures in protozoa, and tracking bacterial infections in neutrophils (Baggiolini, 1998; Campbell and Butcher, 2000; Crone and Lee, 2002). Research on directional movement by external cues in eukaryotes is dominated by chemoattraction, which is the movement toward the chemical compound. Repellents play an important role in morphogenesis, especially during embryonic development (Yang et al., 2002; Schmitt et al., 2005). Cell movement during chick primitive streak formation is controlled by FGF8-mediated chemorepulsion of the cells away from the streak, followed by chemoattraction toward the FGF4 signal produced by the forming notochord (Yang et al., 2002). Axon guidance during spinal chord development away from the roof plate is regulated by multiple repellents, such as BMP (Butler and Dodd, 2003), and by the attractant netrin toward the floor plate (Kennedy et al., 2006).

The mechanism by which repellents work is not well known (Dormann and Weijer, 2006). We envision that a critical step of the signal transduction pathway for cell movement is stimulated by a chemoattractant and inhibited by a repellent. It is essential that this hypothetical step is somehow connected with cell polarity to obtain directional movement. Dictyostelium discoideum cells have been instrumental in resolving the mechanism by which cells sense and respond to chemoattractants. It has been shown that phosphatidylinositol-3,4,5-trisphosphate (PI[3,4,5]P3), which is formed at the side of the cell closest to the source of chemoattractant, is a very strong inducer of pseudopod extensions (Parent et al., 1998; Hirsch et al., 2000; Servant et al., 2000; Funamoto et al., 2002; Iijima and Devreotes, 2002). D. discoideum cells are known to be repelled by unidentified compounds that are secreted by starving cells (Keating and Bonner, 1977; Kakebeeke et al., 1979), indicating that D. discoideum cells have a mechanism to process repellents. Previously, we have shown that several analogues of the attractant cAMP behave as a repellent (Van Haastert et al., 1984). The analogues mediate their effect through binding to the surface cAMP receptor cAR1 (Johnson et al., 1992; Bominaar and Van Haastert, 1993, 1994). We show that these analogues inhibit PLC, contrary to activation of PLC by cAMP. As a consequence, they induce dominant PI(3,4,5)P3 signaling in the rear of the cell, by which cells move away from the repellent.
Results and discussion

*D. discoideum* cells were stimulated with a micropipette containing either the agonist cAMP or the commercially available antagonist 8CPT-cAMP. The cells moved toward the pipette with cAMP, but did not move effectively toward the pipette with 8CPT-cAMP, and actually moved away from the pipette (Fig. S1 and Videos 1 and 2, available at http://www.jcb.org/cgi/content/full/jcb.200611046/DC1). Experiments have been repeated with 3′NH-cAMP, yielding the same results as with 8CPT-cAMP (unpublished data). Fig. 1A shows four frames from a movie in which cells were stimulated with two pipettes containing cAMP and 8CPT-cAMP, respectively (Video 3). In buffer, cells move in random directions (Fig. 1A, 1 min), and cells move away from the pipette with 8CPT-cAMP (Fig. 1A, 16 min). Upon application of the pipette with cAMP (cAMP and 8CPT-cAMP; Fig. 1A, 26 min) cells moved in nearly random directions. However, upon withdrawal of the pipette containing 8CPT-cAMP, cells immediately moved toward the pipette with cAMP (Fig. 1A, 38 min). The trajectories of the cells were analyzed. Data are presented as the chemotaxis index, which is the distance moved in the direction of the gradient (“upgradient”) divided by the total distance moved in 30-s intervals. Data from Video 3 are presented in Fig. 1B, and the means and the SEMs for six independent experiments are presented in Fig. 1C. Wild-type cells show an excellent chemotactic response toward cAMP, with a chemotaxis index of 0.81 ± 0.05. Cells are not attracted to the pipette containing 8CPT-cAMP, but instead exhibit a significant negative chemotaxis index of −0.52 ± 0.04 (P < 0.005). The chemotaxis index of cells stimulated simultaneously with cAMP and 8CPT-cAMP is −0.18 ± 0.11, indicating that 8CPT-cAMP antagonizes the positive chemotaxis toward cAMP and cAMP antagonizes the negative chemotaxis induced by 8CPT-cAMP. Finally, starting with stimulation by the two pipettes, upon withdrawal of the pipette with 8CPT-cAMP the chemotaxis index toward cAMP rapidly increases to 0.72 ± 0.06. The results demonstrate that 8CPT-cAMP is a repellent that can reversibly inhibit the chemotactic response to cAMP.

*D. discoideum* cells move using actin filaments in the front of the cell, which induce the formation of local pseudopodia, and actomyosin filaments in the rear of the cell, which inhibit pseudopod formation and retract the uropod. We coexpressed Myosin-RFP and the filamentous actin-binding protein LimE-GFP from a single plasmid. A pipette with cAMP induces the expected movement of the cells upgradient with LimE-GFP localized in the front and Myosin-RFP in the rear of the cell (Fig. 2A and Video 4, available at http://www.jcb.org/cgi/content/full/jcb.200611046/DC1). Interestingly, the localization of LimE-GFP in the protruding front and Myosin-RFP in the retracting back is identical in cells stimulated with 8CPT-cAMP, except that the front is downgradient and cells move away from the pipette (Fig. 2A and Video 5).

Figure 1. **Antagonism of chemotaxis to cAMP by 8CPT-cAMP.** Starved *D. discoideum* cells were spread on a polystyrene surface in a droplet with 0.5 ml of 10 mM phosphate buffer, pH 6.1, at room temperature, and stimulated by micropipettes filled with 10 mM 8CPT-cAMP (bottom right, open circle) or 0.1 mM cAMP (top left, asterisk); see Materials and methods for further details. (A) Four frames from a movie (Video 3) showing the distribution of cells without stimulus, 8 min after stimulation with 8CPT-cAMP, 5 min after stimulation with 8CPT-cAMP and cAMP, and 12 min after stimulation with cAMP alone (by removing the pipette with 8CPT-cAMP). (B) The chemotaxis index was determined for 20 cells that were shown in Video 3, (top) the filled sections show the simulation with cAMP and/or 8CPT-cAMP. (C) The chemotaxis index was calculated for 85 cells from 6 independent experiments; data shown are the mean ± the SEM. Video 3 is available at http://www.jcb.org/cgi/content/full/jcb.200611046/DC1.
To investigate the mechanism by which 8CPT-cAMP induces negative chemotaxis, wild-type cells expressing the PI(3,4,5)P3 detector PHcracGFP were stimulated with cAMP and 8CPT-cAMP. Similar to previous investigations (Parent et al., 1998; Huang et al., 2003), a pipette with cAMP induces strong localization of PHcracGFP to the plasma membrane at the upgradient side of the cell. Pseudopodia are extended from PHcracGFP-containing areas and cells move upgradient toward the pipette (Fig. 2 B and Video 6, available at http://www.jcb.org/cgi/content/full/jcb.200611046/DC1). 8CPT-cAMP also induces strong localization of PHcracGFP at the plasma membrane, but with opposite polarity compared with cAMP, which is downgradient (Fig. 2 B and Video 7). Cells extend pseudopodia from these PHcracGFP-containing areas, and therefore move away from the pipette with 8CPT-cAMP. The size of the PHcracGFP patches induced by 8CPT-cAMP (9.0 ± 0.43 μm) is only slightly larger than the patches induced by cAMP (6.6 ± 0.17 μm), indicating that 8CPT-cAMP effectively reverses the PI(3,4,5)P3 polarity.

PI(3,4,5)P3 is formed by PI3-kinase (PI3K) and degraded by PTEN that, in cAMP gradients, are localized at the leading edge and the rear of the cell, respectively. In 8CPT-cAMP gradients, the localization of PI3K and PTEN is reversed compared with cAMP gradients (Fig. 2 B). To investigate the role of PI3K activity in polarity and chemotaxis reversal, we investigated the chemotactic activity of pi3k1/2-null cells toward cAMP and 8CPT-cAMP. In pi3k1/2-null cells, two PI3Ks are deleted that, together, mediate the vast majority of cAMP-stimulated PI(3,4,5)P3 production (Zhou et al., 1998; Funamoto et al., 2002; Huang et al., 2003). These experiments are possible because PI3K is not essential for chemotaxis, and directional sensing can be mediated by other pathways (Hirsch et al., 2000;...
Funamoto et al., 2002; Iijima and Devreotes, 2002; Huang et al., 2003; Postma et al., 2004; Loovers et al., 2006). Fig. 3 shows that pi3k1/2-null cells exhibit a good chemotactic response toward a pipette with cAMP (chemotaxis index is 0.80 ± 0.13). In contrast to the negative chemotaxis induced by 8CPT-cAMP in wild-type cells, pi3k1/2-null cells do not exhibit a significant negative or positive response to 8CPT-cAMP (chemotaxis index is 0.11 ± 0.12). More importantly, using two pipettes with cAMP and 8CPT-cAMP, respectively, pi3k1/2-null cells effectively move toward cAMP and are not inhibited by 8CPT-cAMP and cAMP. The results show that wild-type cells move away from 8CPT-cAMP, whereas pi3k1/2-null and plc-null are not repelled from 8CPT-cAMP. Moreover, chemotaxis toward cAMP is antagonized by 8CPT-cAMP in wild-type cells, but not in pi3k1/2-null and plc-null cells. Videos 3, 8, and 9 are available at http://www.jcb.org/cgi/content/full/jcb.200611046/DC1.

The molecular mechanism by which cAMP mediates PI(3,4,5)P3 accumulation upgradient in D. discoideum cells has been well described. PI3K is activated and enriched upgradient in the cell, whereas the PI(3,4,5)P3-degrading enzyme PTEN strongly localizes downgradient in the cell (Funamoto et al., 2002; Iijima and Devreotes, 2002). PTEN has been demonstrated to bind to phosphatidylinositol-3,4,5-trisphosphate (PI(4,5)P3), suggesting that PI(4,5)P3 is depleted upgradient in the cell (Iijima et al., 2004). This depletion of PI(4,5)P3 could be induced by several nonexclusive methods, such as the observed conversion of PI(4,5)P2 to PI(3,4,5)P3 upgradient by PI3K (Funamoto et al., 2002; Huang et al., 2003), but also by the conversion of PI(4,5)P2 to Ins(1,4,5)P3 and DAG by PLC, which is known to be activated by cAMP (Drayer and van Haastert, 1992; Boininaar et al., 1994). We propose a mechanism by which 8CPT-cAMP could revert the polarity of chemotactic sensing that is based on the observation that cAMP stimulates PLC, whereas 8CPT-cAMP inhibits this enzyme (Peters et al., 1991; Boininaar and Van Haastert, 1993; Boininaar and Van Haastert, 1994; supporting biochemical data are presented in Fig. S2, available at http://www.jcb.org/cgi/content/full/jcb.200611046/DC1). Upgradient stimulation of PLC by cAMP will lead to local depletion of PI(4,5)P2, and thereby prevent PTEN binding, by which the upgradient PI(3,4,5)P3 accumulation is stabilized. In contrast, the upgradient inhibition of PLC by 8CPT-cAMP will lead to the local accumulation of PI(4,5)P2, thereby inducing PTEN binding and upgradient PI(3,4,5)P3 degradation (Fig. 4). This relatively simple model for polarity reversal predicts that 8CPT-cAMP does not induce polarity switching in plc-null cells. D. discoideum cells contain a single plc gene encoding a PLCδ isoform (Drayer and van Haastert, 1992), which, like PI3K, is instrumental but not essential for chemotaxis (Drayer et al., 1994). Expression of GFP-tagged reporter proteins in plc-null cells reveal, as predicted, cytosolic...
localization of PH-cracGFP and enhanced PTEN-GFP expression at the membrane in cAMP and 8CPT-cAMP gradients (Fig. S3). As presented in Fig. 3 B and Video 9, plc-null cells show a similar chemotactic response toward 8CPT-cAMP as pi3k-null cells: they move in random directions in the presence of 8CPT-cAMP alone and, subsequently, move effectively toward an additional pipette with cAMP. This indicates that PLC is also essential for mediating the inhibitory effect of 8CPT-cAMP, as is PI3K. Finally, pien-null cells were investigated, showing that these cells are attracted toward cAMP, but are not repelled by 8CPT-cAMP (unpublished data).

A scheme for PI(3,4,5)P3-mediated chemotaxis reversal by 8CPT-cAMP consists of three parts (Fig. 4). The basis is a PLC/PI(4,5)P2 polarity switch. In D. discoideum, PLC is regulated by the activating Gα2 and inhibitory Gα1, which, in a gradient of attractant or repellent, will determine the polarity of the PI(4,5)P2 gradient. The attractant cAMP shows predominant activation of PLC, leading to lower PI(4,5)P2 levels upgradient, while the repellent 8CPT-cAMP inhibits PLC, leading to higher PI(4,5)P2 levels upgradient. The resulting gradients of PI(4,5)P2 and colocalized PTEN mediate opposite gradients of PI(3,4,5)P3, leading to the localized polymerization of actin. The gradients of localized PTEN and PI3K are stabilized because PTEN accumulates at the site of its product PI(4,5)P2, whereas PI3K accumulates at sites of its effector, PI(3,4,5)P3-induced F-actin. This mutually spatial exclusion of PI3K and PTEN will result in symmetry breaking, by which small spatial differences in the underlying polarity gradient can be amplified to the observed strong PI(3,4,5)P3 gradient. Although PI3K and PLC are not essential for chemotaxis, the results clearly demonstrate that local formation of PI(3,4,5)P3 is a very strong inducer of pseudopod formation, such that the cells can even move downgradient, overruling any upgradient signaling that 8CPT-cAMP may induce.

In our model, a compound is a repellent because it binds to a receptor that is preferentially coupled to PLC via an inhibitory G protein, whereas it is an attractant when the receptor is coupled to a stimulatory G protein. The regulation of D. discoideum PLC by the stimulatory G2 and inhibitory G1 forms the basis for the polarity switch, and it allows the cell to respond to chemical gradient with repulsion or attraction. This polarity switch may be used by the cell during development. D. discoideum cells grow on bacteria. Cells starved for <1 h secrete unidentified compounds that induce repulsion of the cells, by which cells may find bacteria in a larger area (Keating and Bonner, 1977; Kakebeeke et al., 1979). Cells starved for ~5 h secrete cAMP, to which they are attracted and which allows the cells to form a multicellular structure. Interestingly, G1 is expressed throughout development, whereas G2 is nearly absent during early starvation and expressed only after ~4 h (Pupillo et al., 1989). Thus, in early starved cells with the predominant inhibitory G1, the PLC–PI3K system is pruned for repulsion, whereas it becomes a system for attraction by expression of the stimulatory G2 during late starvation.

The mechanism of polarity reversal of PLC–PI3K signaling could be instrumental in mammalian cells to navigate in complex chemotactic gradients. During development, many cells, such as neurons and gonads, are projected in the body by mixtures of attractants and repellents (Yang et al., 2002; Schmitt et al., 2005). Observations on the action of Slit2 may be instrumental. Slit2 is a repellent for neuronal cells (Niclou et al., 2000; Ringstedt et al., 2000). In contrast, Slit2 does not affect the direction of movement of vascular smooth muscle cells, but strongly inhibits PDGF-stimulated chemotaxis by inhibition of PDGF stimulation of Rac1 (Wu et al., 2001; Chen et al., 2004). It is possible that, in neuronal cells, Slit2 induces a polar inhibition of Rac1, thereby inducing repulsion, whereas in vascular smooth muscle cells Slit2 induces uniform inhibition of Rac1 and is therefore not a repellent, but only an inhibitor of chemoattractants. Rac1 is known to be regulated by PIP3 in mammalian (Srinivasan et al., 2003; Kunisaki et al., 2006) and D. discoideum (Park et al., 2004) cells. The observed simplicity by which PLC-mediated polarity inversion of PI3K signaling in D. discoideum converts attraction to repulsion may provide a single mechanism to integrate complex positive and negative chemotactic signals during development.
Materials and methods

Plasmids and cells
The plasmid pWF38 (PHcracGFP) expressing the 700-bp Ntermonal PH domain of CRAC fused to GFP (Parent et al., 1998), and plasmids expressing PI3K2-GFP (Funamoto et al., 2002; Iijima and Devreotes, 2002) and PTEN-GFP (Iijima and Devreotes, 2002) were provided by P. Devreotes (Johns Hopkins University School of Medicine, Baltimore, MD). Plasmid 339 expressing mRFPmars (Fischer et al., 2004) was provided by A. Muller-Taubenberger (Ludwig Maximilians University Munich, Munich, Germany). Plasmid pBIG-GFP-myo expressing a GFP fusion with myosin heavy chain II (Levi et al., 2002) was a gift from T. Egelhoff (Case Western Reserve University, Cleveland, OH). p3K1/2-null cells were provided by R. Firtel (University of California, San Diego, La Jolla, CA).

Plasmid LB15 expressing Lim-GFP and Myo-RFP was constructed as follows. The neomycin resistance gene of MB74 was exchanged for the HPH hygromycin resistance gene that was preceded by an actin 15 promoter and terminated with a cab 1 terminator. The DNA coding for the actin-binding domain of Lim (aa 1–145) was cloned behind an actin 15 promoter and 5 adenosines, which serve as the Kozak sequence. This was followed by a Spel site (coding for Thr and Ser) and the complete open reading frame of GFP (S65T variant), followed by a stop codon and an actin 8 terminator. This yielded the plasmid MB74Hyg-Lim-GFP. The gene encoding the monomeric red fluorescent protein mRFPmars (Fischer et al., 2004) was amplified by PCR on plasmid DNA. The gene was preceded by a NgoMIV site, an actin 15 promoter, and 5 adenosines, and was followed by a BamHI site (encoding Gly and Ser), the sequence encoding 2–211 of myosin heavy chain, the myosin terminator from the vector pBIG-GFP-myo (Levi et al., 2002), and a NgoMIV site. Finally, the gene encoding the mRFPmars-myosin fusion was released using the NgoMIV site and cloned into the single NgoMIV site of MB74Hyg-Lim-GFP.

The D. discoideum strain AX3 was used as wild-type control in all experiments. The mutants strains used are the p3K-null strain 1.19 (Drayer et al., 1995), the p3K-null p3K1/p3K2 strain GMPI (Funamoto et al., 2001), and p3K-null cells (Iijima and Devreotes, 2002). Cells were grown in shaking culture in HGM 5 medium (containing per liter: 14.3 g oxoid peptone, 7.15 g bacto yeast extract, 1.36 g Na2HPO4 × 12H2O, 0.49 g KH2PO4, 10.0 g glucose) at a density of 5 × 106 and 6 × 105 cells/ml. Cells were centrifuged for 3 min at 300 g, washed in PB (10 mM KH2PO4/Na2HPO4, pH 6.5), and starved in PB in 6-well plates (Nunc) for 5 h. Cells were then resuspended in PB, centrifuged, and washed once in PB, and resuspended in PB at a density of 6 × 106 cells/ml.

Recording of movies
Unless otherwise mentioned, digital images of cells in PB at room temperature were captured at 10-s time intervals over 45 min. Videos 1, 2, 4, and 5–7 were captured using a confocal laser scanning microscope (LSM 510 META-NLO; Carl Zeiss Microimaging, Inc.) equipped with a 63×/NA 0.4 objective (Plan-Apochromatic; Carl Zeiss Microimaging, Inc.). For excitation of the fluorochromes, GFP (S65T variant), and mRFPmars, a 488-nm argon/krypton laser was used, respectively. The fluorescence was filtered through a BPF500-530 IR and a LP560 filter, and was detected by a photomultiplier tube. The field of observation was 206 × 206 μm; Videos 1 and 2 present the phase-contrast channel, whereas the fluorescent channel is shown in Videos 4–7. For Videos 3 and 9, an inverted light microscope (Type CK40 with a LWD A240 20×/NA 0.4 objective; Olympus) fitted with a charge-coupled device camera (TK-C1381; JVC) was used. Digital images were captured on a PC using VirtualDub software and Indeo video 5.10 compression. The field of observation is 358 × 269 μm. Video 8 was captured using a 10× numerical aperture 0.25 objective, and presents a selected field of the same size, namely 358 × 269 μm. For all individual videos, specific time periods were selected that start at the moment the pipette was lowered to the plane just above the cells. In the phase-contrast videos, the pipette tips are visible as dark triangular shadows. In the fluorescence videos (Videos 4–7), the place of the pipette tip is indicated with an asterisk.

Analysis of chemotaxis
The chemotaxis index, which is defined as the ratio of the cell displacement in the direction of the gradient and its total traveled distance, was determined for ~25 cells in a video, as follows. First, the position of the centroid of a cell was determined with ImageJ [National Institutes of Health; rsb.info.nih.gov/ji] for frames at 30-s intervals, yielding a series of coordinates for that cell. Using these coordinates, the chemotaxis index of each 30-s step was calculated and was averaged, yielding the chemotaxis index for that cell in the movie. The data shown are the average and SEM of the chemotaxis indices from three independent experiments, with ~25 cells per experiment.

Online supplemental material
Fig. S1 shows cell trajectories of wild-type cells in a gradient of cAMP and BCF7-cAMP, revealing that cells are attracted toward cAMP, but repelled from BCF7-cAMP. Fig. S2 shows inhibition of PLC signaling by the antagonist 3′NH-cAMP. BCF7-cAMP has similar properties to 3′NH-cAMP. Video S3 shows the localization of PHcrac-GFP, PTEN-GFP, and PI3K-GFP in plc-null cells in a gradient of cAMP or BCF7-cAMP. Video 1 shows chemotaxis toward a pipette with cAMP. Video 2 shows chemotaxis away from a pipette with BCF7-cAMP. Video 3 shows cells moving in gradients of BCF7-cAMP and cAMP + BCF7-cAMP, followed by movement in only cAMP. Video 4 shows the localization of F-actin at the leading edge and myosin in the back of cells chemotaxing toward cAMP. Video 5 shows the localization of F-actin at the leading edge and myosin in the back of cells chemotaxing away from BCF7-cAMP. Video 6 shows the localization of PHcrac-GFP (detecting PI3K1/2) at the leading edge of cells chemotaxing toward cAMP. Video 7 shows the localization of PHcrac-GFP (detecting PI3K1/2) at the leading edge of cells chemotaxing away from BCF7-cAMP. Video 8 shows chemotaxis of p3K1/2-null cells toward cAMP in the presence of BCF7-cAMP. Video 9 shows chemotaxis of plc-null cells toward cAMP in the presence of BCF7-cAMP. The online version of this article is available at http://www.jcb.org/cgi/content/full/jcb.200611046/DC1.

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