

# Long range mutual activation establishes Rho and Rac polarity during cell migration

Henry De Belly<sup>1,2</sup>, Andreu Fernandez Gallen<sup>3</sup>, Evelyn Strickland<sup>1,2</sup>, Dorothy C. Estrada<sup>1,2</sup>, Patrick J. Zager<sup>1,2</sup>, Tamas L. Nagy<sup>1,2</sup>, Janis K Burkhardt<sup>4,5</sup>, Hervé Turlier<sup>3,4</sup>✉, and Orion D. Weiner<sup>1,2</sup>✉

<sup>1</sup>Cardiovascular Research Institute, University of California, San Francisco, San Francisco, CA, USA.

<sup>2</sup>Department of Biochemistry and Biophysics, University of California, San Francisco, San Francisco, CA, USA.

<sup>3</sup>Center for Interdisciplinary Research in Biology (CIRB), College de France, CNRS, INSERM, Université PSL, Paris, France

<sup>4</sup>Department of Pathology and Laboratory Medicine, Children's Hospital of Philadelphia Research Institute, Philadelphia, PA

<sup>5</sup>Perelman School of Medicine, University of Pennsylvania, Philadelphia, PA, Philadelphia

**In migrating cells, the GTPase Rac organizes a protrusive front, whereas Rho organizes a contractile back. How these GTPases are appropriately positioned at the opposite poles of migrating cells is unknown. Here we leverage optogenetics, manipulation of cell mechanics, and mathematical modeling to reveal a surprising mechanochemical long-range mutual activation of the front and back polarity programs that complements their well-known local mutual inhibition. Rac-based protrusion stimulates Rho activation at the opposite side of the cell via membrane tension-based activation of mTORC2. Conversely, Rho-based contraction induces cortical-flow-based regulation of phosphoinositide signaling to trigger Rac activation at the opposite side of the cell. We develop a minimal unifying mechanochemical model of the cell to explain how this long-range facilitation complements local inhibition to enable robust Rho and Rac partitioning. We show that this long-range mutual activation of Rac and Rho is conserved in epithelial cells and is also essential for efficient polarity and migration of primary human T cells, indicating the generality of this circuit. Our findings demonstrate that the actin cortex and plasma membrane function as an integrated mechanochemical system for long-range partitioning of Rac and Rho during cell migration and likely other cellular contexts.**

Correspondence: Orion.Weiner@ucsf.edu, herve.turlier@college-de-france.fr

## Introduction

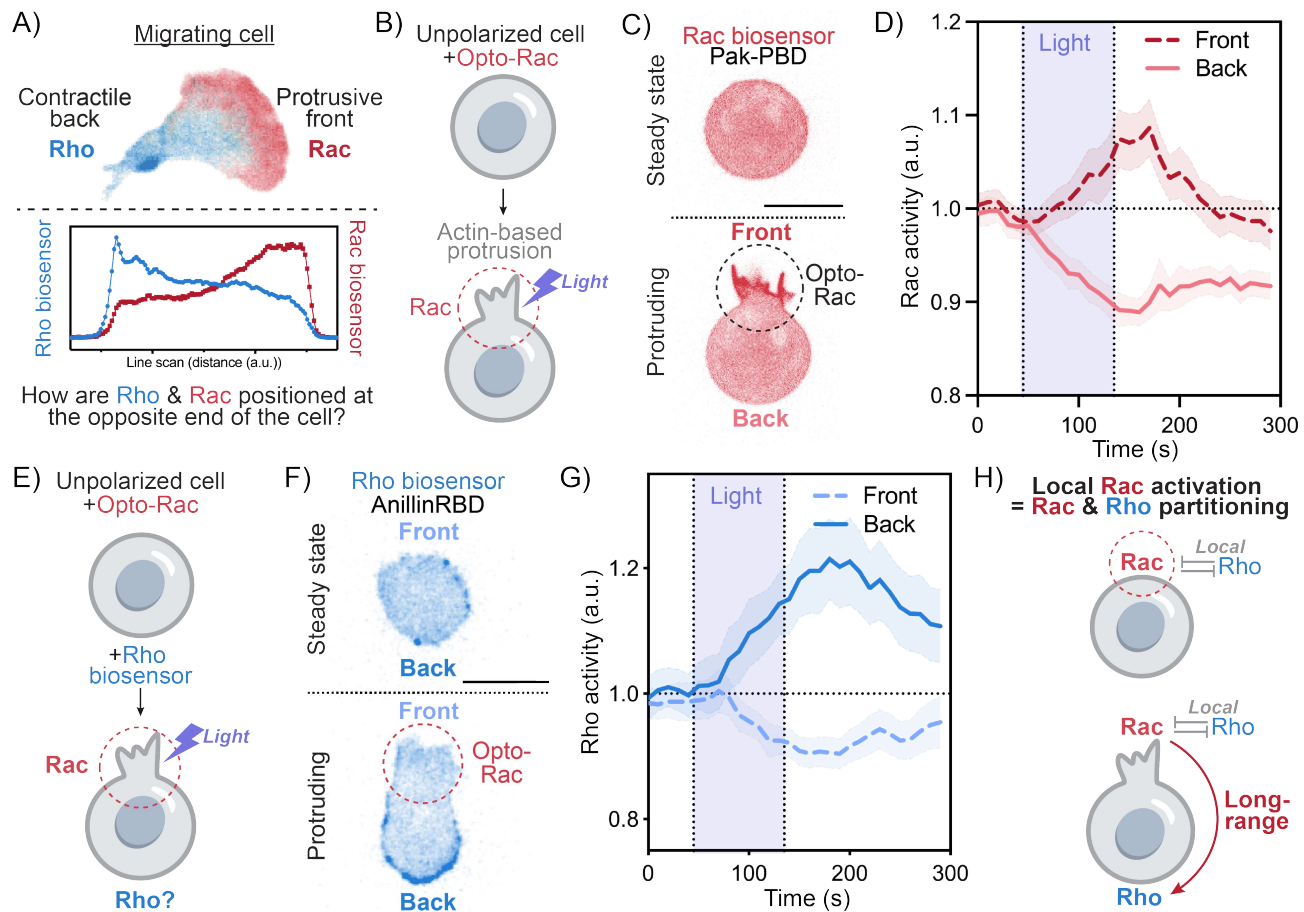
For proper physiology, many cells need to polarize, or restrict different signaling programs to different portions of their cell surface. Rac and Rho GTPases are key regulators of cell polarity and spatially pattern the actin cytoskeleton for mitosis, morphogenesis, migration, and development (1). Rac is localized to the front migrating cells, where it promotes the extension of sheet-like lamellipodia or pressure-driven blebs (Figure 1A) (2, 3). Rho is localized to the back of migrating cells, where it promotes local myosin-2-based contractions and long-range cortical flows (1, 4) (Figure 1A).

While we know that Rho and Rac mutually inhibit each other locally (1, 5–7), how they are properly positioned at opposite ends of the cell during migration remains poorly understood. The majority of models of polarity in migration are based on local biochemical interactions (6, 8–15), with a majority of mathematical models of polarity establishment relying in fact on the so-called wave-pinning mechanism (16–19),

where a travelling front of active Rho GTPase is pinned by a bath of cytosolic inactive form. Such mechanism relies importantly on the fast diffusion of the inactive form and a hypothesis of fixed total protein amount. However, previous studies have challenged these requirements, demonstrating that a diffusion-based inhibitor or depletion mechanism is not sufficient to explain neutrophil polarization (20–23). In contrast, local inhibition is not sufficient either to explain the long-range partitioning of active pools of Rho and Rac and to coordinate front and back at the scale of a migrating cell. The requirement for an additional long-range communication had long been suggested (20–25), but direct evidence and molecular mechanism for this link was unknown.

Cell polarity involves long-range cellular information processing and necessitates information flow at the cellular scale. Forces transmitted via the actin cortex and the plasma membrane have emerged as key conduits for this global coordination. In migration, membrane tension guides shape by relaying actin-based protrusive forces at the front to the contraction of the rear; setting up a global competition that enables the establishment of a single dominant front (26–30). Here we investigate whether the membrane and cortex could act as the mechanical conduit for long-range coordination of the front and back polarity programs.

To probe the long-range spatial coordination of Rac and Rho, we take advantage of optogenetic activators of Rac and Rho in initially unpolarized cells. This enables us to precisely and specifically activate either Rac or Rho in a small region of the cell and observe the global response of the other GTPase. By combining optogenetics with manipulation of cell mechanics and mathematical modeling, we find that both the front and back mutually activate each other at a distance, and this occurs using two distinct pathways. The front stimulates the back via membrane tension, whereas the back stimulates the front via cortical remodeling. We demonstrate the physiological relevance of our findings for immune cell migration using primary human T cells. Our results demonstrate that the actin cortex and plasma membrane act as an integrated mechanochemical system to ensure the proper positioning of the front and back polarity programs during cell migration.



**Fig. 1. Local Rac activation stimulates long-range RhoA activation at the opposite side of the cell**

(A) Confocal image and associated linescan of a migrating neutrophil-like HL-60 cell expressing the polarity biosensors for Rac (PAK-PBD, in red) and Rho (Anillin-RBD, in blue). In migrating cells, Rac and Rho localize to the protruding cell front and contracting cell back, respectively. Here we sought to investigate how Rac and Rho are appropriately positioned at the opposite poles of a migrating cell. (B) We locally activated the front polarity program Rac in an initially unpolarized neutrophil-like HL-60 cell using optogenetics (opto-PI3K, see Methods). (C) Time-lapse confocal images of an unpolarized cell before and during opto-Rac stimulation. Rac activity was monitored via the Rac biosensor Pak-PBD. (D) Average time trace of Rac activity at the plasma membrane at the site of opto-Rac activation (cell front) compared to the opposite side of the cell (cell back). (mean  $\pm$  95%CI;  $n>40$ ,  $N=3$ ). (E) We locally activated Rac in an initially unpolarized cell while simultaneously measuring Rho activity using AnillinRBD. (F) Time-lapse confocal images of an unpolarized cell before and during opto-Rac stimulation. Rho activity was monitored using the biosensor AnillinRBD. (G) Average time trace of Rho activity at the plasma membrane for the front versus back of the cell following opto-Rac activation. (mean  $\pm$  95%CI;  $n>40$ ,  $N=8$ ). (H) Local Rac activation leads to long-range activation of Rho at the opposite side of the cell. Scale bars: 10  $\mu$ m.

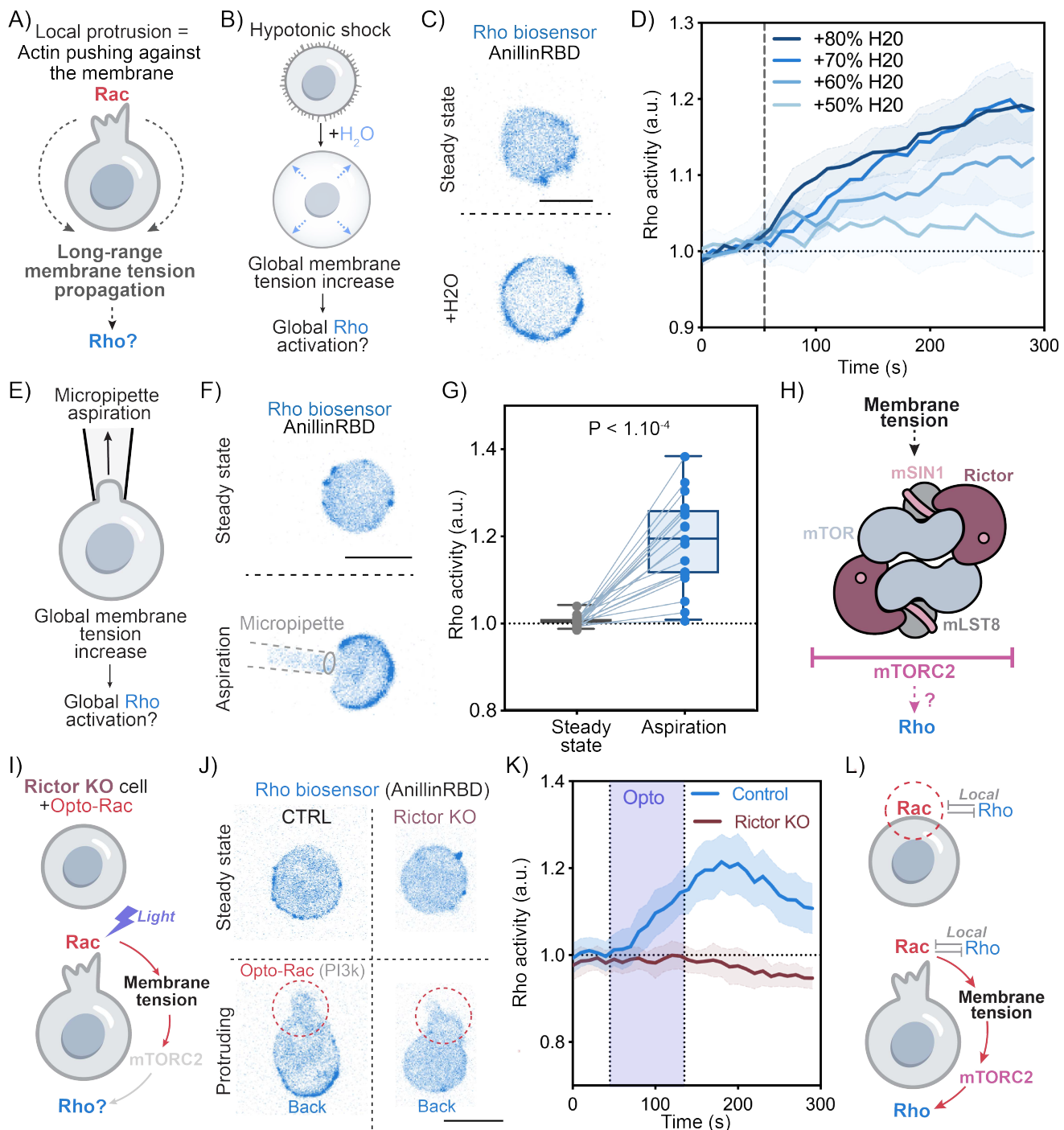
## Results

### Local Rac activation stimulates long-range RhoA activation at the opposite side of the cell.

To investigate how the front and back polarity programs are properly positioned at the opposite ends of the cell, we first analyzed the establishment of cell polarity in a context where we can independently control the spatial and temporal dynamics of each of these programs. Towards this end, we leveraged an optogenetic approach (via local activation of PI3K (31), see Methods), to locally activate the front polarity program Rac in initially unpolarized neutrophil-like HL60 cells (Figure S1A). As previously demonstrated, this local Rac activation leads to actin-driven protrusions like those seen at the cell front during migration (Figure S1) (31, 32). Rac activity (visualized with Pak-PBD) is locally enriched in the zone of activation (cell front), while being depleted at the opposite side of the cell (cell back) (Figure 1B-D, S1B, C

and Video S1). These results confirm that opto-PI3K locally activates Rac. Next, we sought to investigate how this local zone of Rac activity influences the back polarity program. To this end, we used local Rac activation via opto-PI3K while monitoring the activity of Rho using the biosensor Anillin-RBD (Figure 1E and Video S2). Light-induced Rac activation elicits a rapid long-range increase in Rho activity at the opposite side of the cell (Figures 1F,G and S1G-I). Thus, in addition to locally inhibiting Rho, Rac activation also stimulates Rho at the opposite end of the cell.

We next sought to understand how local Rac activation in one part of the cell triggers Rho activity at the opposite end. We considered two possibilities: distal Rho activation could result from local Rac-based biochemical inhibition(1) or indirect long range mechanical signals. One possible means of long-range Rho activation would be Rac-induced protrusions, which generate a membrane tension increase



**Fig. 2. Rac stimulates long-range RhoA activation at the opposite side of the cell via membrane-tension-mediated mTORC2 activation**

(A) Local Rac-mediated cell protrusion leads to a global increase in membrane tension<sup>32</sup>. (B) If protrusions activate Rho by elevating membrane tension, then increasing membrane tension (even in the absence of Rac activation) should suffice to activate Rho. Towards this end, we used hypotonic shock to elevate membrane tension. (C) Time-lapse confocal images of an unpolarized opto-Rac HL-60 cell expressing the Rho biosensor (AnillinRBD) before and after hypotonic shock. (D) Average time trace of Rho activity at the plasma membrane in response for hypotonic shocks of various intensity. (N>3, n>20, means  $\pm$  95%CI). Hypotonic-shock-based elevation of membrane tension suffices to globally increase RhoA activity. (E) As an alternate approach to increase membrane tension, we leveraged micropipette aspiration<sup>(32)</sup>. (F) Time-lapse confocal images of an unpolarized opto-Rac HL-60 cell expressing the Rho biosensor (AnillinRBD) before and after micropipette aspiration (see Methods). (G) Average Rho activity before (steady state) and during aspiration. Micropipette-based elevation of membrane tension significantly stimulates Rho activation. Box and whiskers: median and min to max; p values from Wilcoxon paired Student's t test. (H) We investigate whether mTORC2 is part of the mechanosensory pathway that links increased in membrane tension to the activation of Rho. (I) We used optogenetics to locally activate Rac in control cells versus cells lacking different core components of the mTORC2 complex (Rictor or mSIN1 knockouts). (J) Time-lapse confocal images of an unpolarized control or Rictor KO cell before and during opto-Rac stimulation. Rac activity was monitored via the Rac biosensor Pak-PBD. (K) Local Rac activation potentially stimulates long-range activation of Rho in control cells (blue) but not cells deficient in mTORC2 activity (Rictor KO, magenta) (mean  $\pm$  95%CI; n>20, N= 3). Control curve is same as in Fig. 1D. (L) Local Rac activation stimulates long-range Rho activation via membrane tension mediated mTORC2 activation. Scale bars: 10 $\mu$ m.



that propagates across the cell (32) (**Figure S2A**). To distinguish between these possibilities, we impaired Rac-mediated protrusion formation by applying pharmacological inhibitors of actin assembly (latrunculin) or Arp2/3 complex activation (CK666) to opto-PI3K cells (**Figure S2A-D**). While these perturbations do not impair our ability to optogenetically activate Rac (31), they inhibited Rac-mediated Rho activation at the other end of the cell. To test whether Rho activation depend on the actomyosin network, we impaired actomyosin contractility by treating cells with either the myosin inhibitor blebbistatin or the ROCK inhibitor Y-27632 (**Figure S2E**) and observed no noticeable effect on Rho as a result of local Rac activation (**Figure S2F-H**). These data indicate that protrusion mediated Rho activation is independent of contractility. Altogether, our results support long-range activation of Rho by Rac, operating independently from the well-established local antagonism between these GTPases (**Figure 1H**).

### **Rac stimulates long-range Rho activation via membrane-tension-mediated mTORC2 activation**

If protrusions stimulate Rho activation through an increase in membrane tension (32) (**Figure 2A**), membrane tension increases should suffice to stimulate Rho activation even in the absence of Rac activation or protrusions. To investigate this possibility, we used hypotonic shock to stimulate an increase in membrane tension (33) (**Figure 2B**). Hypotonic shock induced a rapid and global increase in Rho activity across the cell (**Figures 2C, D, S3A-F and Video S3**). To verify that this increase in Rho activity is a function of increased membrane tension and not a secondary consequence of the rebuilding of the actin cortex following hypotonic shock, we combined our hypotonic shock assay with Latrunculin B treatment, a combination that potently depolymerizes the actin cortex (32, 33). Under these conditions, osmotic shock still activates Rho, indicating that the actin cytoskeleton is not necessary for membrane-stretch-based Rho activation (**Figure S3D**). As an alternate approach, we next used a micropipette-based aspiration assay, which we previously demonstrated to increase membrane tension<sup>32</sup>. This mechanical manipulation also simulates global Rho activation (**Figures 2E-G; S3G-J and Video S4**). The activation of Rho by hypotonic shock, micropipette aspiration and opto-induced protrusions likely reflects a role for membrane tension as a Rho activator. To test whether increases in tension also affect Rac activation, we performed measurements of Rac activity in response to both osmotic shock and micropipette aspiration and found that, as previously reported, elevated tension does not activate Rac (**Figure S3E, F and I, J**). Our data show that Rac acts through protrusion-mediated increases in membrane tension to stimulate Rho at a distance.

Next, we sought to identify the mechanosensors that link increases in membrane tension to Rho activation. The mTORC2 complex emerged as a strong candidate, given its responsiveness to membrane tension and its key roles in regulating cell polarity and motility (28, 29, 33–35)(**Figure**

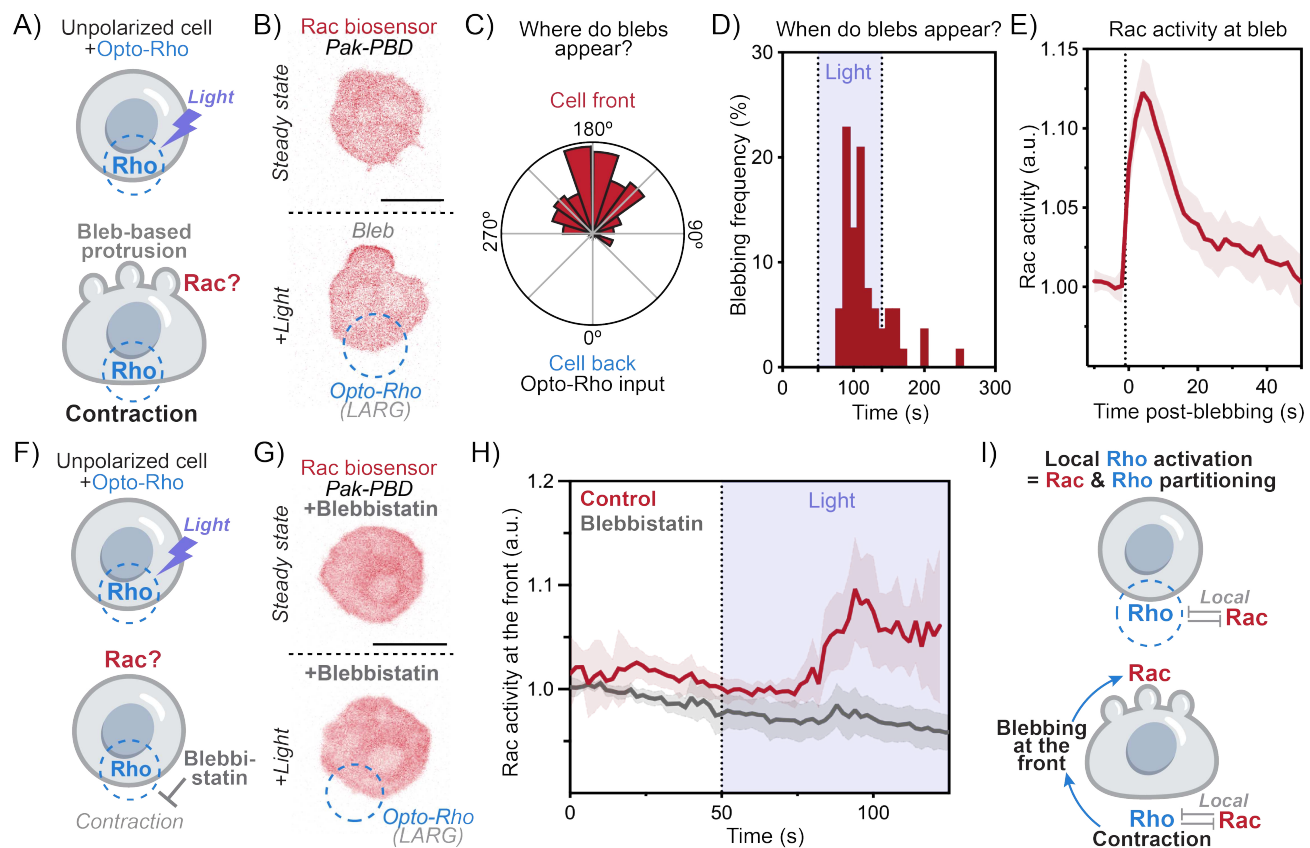
**2H**). To test whether mTORC2 links membrane tension increases to Rho activation, we performed our opto-Rac activation assay in cells lacking different core components of the mTORC2 complex (Rictor or mSIN1 CRISPR KO cell lines) (29) or using a pharmacological inhibition of mTOR. Cells deficient in mTORC2 (Rictor or mSIN1 KO) or treated with mTOR inhibitor KU-0063794 failed to activate Rho activity upon local Rac-activation (**Figures 2I-K; S4A-E and Video S5**) despite retaining their ability to generate protrusions. To further isolate the tension-mediated potentiation of Rho activity, we repeated previous hypotonic and micropipette aspiration assays in mTORC2-impaired cells. Our results confirmed that mTORC2 is essential for Rho activation following both hypotonic shock (**Figures S3F-H**) and micropipette aspiration (**Figures S3I-K**). Our work reveals a molecular pathway in which Rac-induced actin protrusions globally increase membrane tension, activating the mechanosensitive mTORC2, which in turn stimulates Rho.

### **Local Rho activation leads to Rac activation at the opposite side of the cell**

We previously established that local Rac-mediated protrusion is sufficient for long-range Rac and Rho partitioning across the cell. However, not all migrating cells generate branched actin-based protrusions at their front. In bleb-based migration, Rho mediated contraction at the cell back is the primary force generator, producing increased intracellular pressure leading to blebbing on the opposite side (2, 36). Interestingly, while both blebbing programs and actin polymerization-based protrusions use different protrusion engines, both rely on the same Rho-Rac polarity machinery (37, 38). Given the role of Rho activation in promoting bleb-based migration, we next wondered whether local Rho activation is also sufficient to trigger long-range Rac and Rho partitioning.

To investigate the ability of Rho to activate Rac, we used an optogenetic approach to locally activate the back polarity program Rho (via LARG, see Methods) while simultaneously monitoring Rac activity using the biosensor Pak-PBD (**Figure 3A and Video S6**). Our results show that Rho activation leads to long-range Rac activation at the opposite end of the cell, coinciding with another morphological change—blebbing (**Figure 3B**). Given previous studies linking blebbing to Rac activation (38, 39), we hypothesized that Rho's long-range stimulation of Rac might depend on Rho-induced blebbing (**Figures 3C-E and S5A**). To test this hypothesis, we used the actomyosin inhibitor blebbistatin to block cellular contraction and blebbing following optogenetic Rho activation (**Figure 3F**). Indeed, when blebbing was inhibited, Rho could no longer activate Rac at a distance (**Figures 3G, H**). Just as Rac requires its downstream cytoskeletal effects (actin polymerization and protrusion generation) to activate Rho at a distance (**Figure S2**), Rho relies on its downstream cytoskeletal impact (myosin-based contraction and blebbing) to activate Rac at a distance. To further explore whether blebs suffice to activate Rac independent of Rho-mediated stimulation of actomyosin





**Fig. 3. Local Rho activation elicits Rac activation at the opposite side of the cell**

(A) Local Rac activation leads to long-range Rho activation at the opposite end of the cell via membrane tension. Does local Rho activation also lead to long-range Rho and Rac partitioning? To test this, we used optogenetics to locally stimulate Rho (opto-LARG, see Methods) while simultaneously measuring Rac activity using the biosensor Pak-PBD. (B) Time-lapse confocal images of an unpolarized opto-Rac HL-60 cell expressing the Rac biosensor (Pak-PBD) before and during opto-Rho stimulation. Local Rho activation induces blebs at the opposite end of the cell, and these blebs are enriched in active Rac. (C) Polar histogram of the spatial distribution of blebs appearance in relation to opto-Rho (cell back) showing that the majority of blebs are positioned at the opposite side of the cell back (opto-Rho). (N = 5, n > 65). (D) Histogram of bleb distribution in time during opto-Rho stimulation. (N = 3, n > 55). (E) Average time trace of Rac activity at the plasma membrane at the protrusive blebs. Time 0 = blebbing. For times before blebbing, Rac activity is measured at the same membrane spot where the bleb will appear. (see Methods) (mean ± 95%CI; n > 30, N = 3). (F) We hypothesized that Rho's long-range stimulation of Rac might depend on Rho-based contractility which in turns triggers blebbing. (G) We locally activated Rho using optogenetics in the absence or presence of an inhibitor of myosin activation and blebbing (10  $\mu$ M Blebbistatin). (H) Average time trace of Rac activity at the plasma membrane measured at the cell front of control or cells treated with 10M Blebbistatin. These data indicate that myosin-based blebbing is important for Rho-mediated activation of Rac activity (mean ± 95%CI; n>20, N= 3). (I) Local Rho activation leads to long-range activation of Rac at the opposite side of the cell. Scale bars: 10  $\mu$ m.

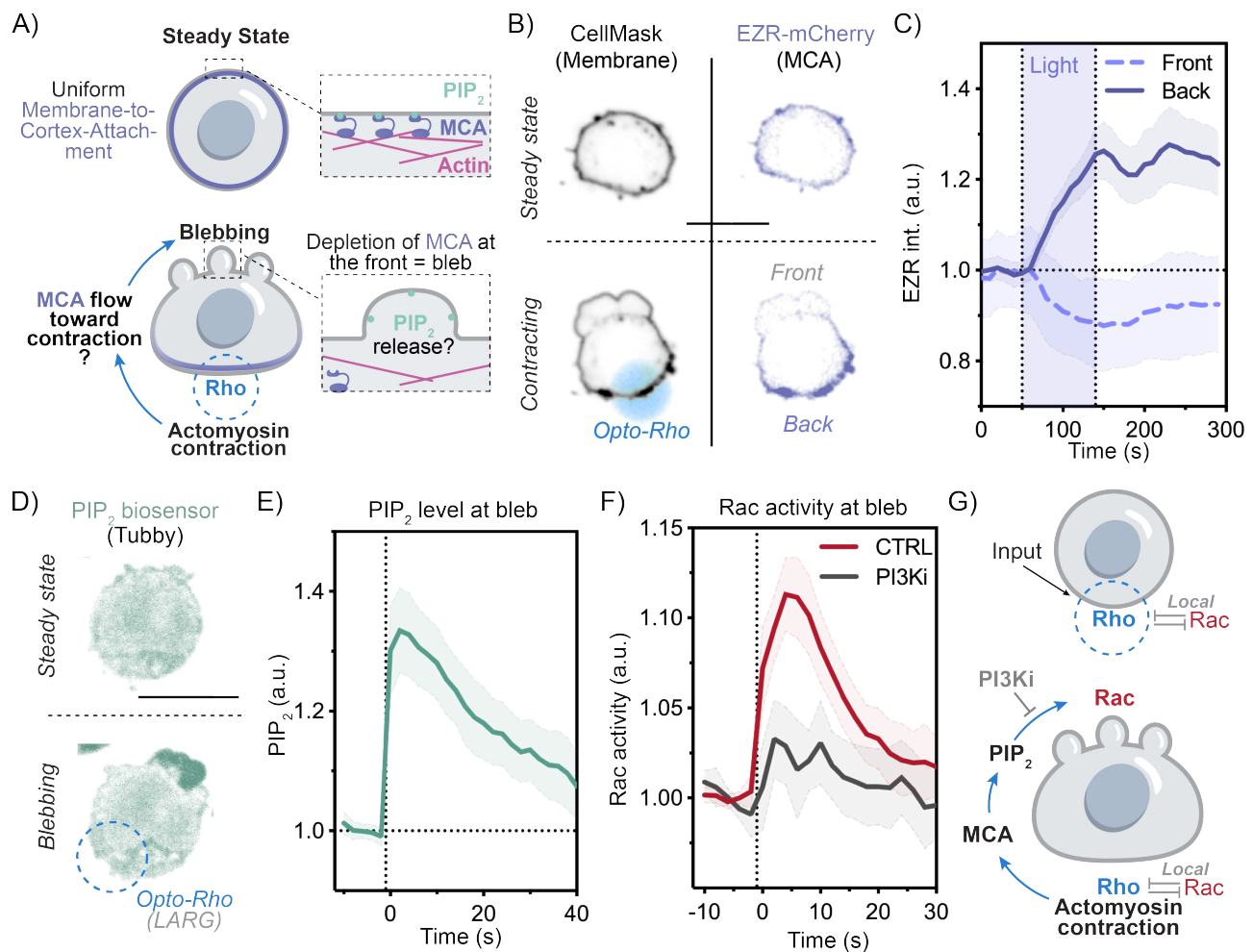
contractility, we used the actin inhibitor Latrunculin B to induce stable blebs (32) (Figure S5B). These stable blebs showed robust Rac enrichment (Figure S5C, D). Our data demonstrate that local Rho activation stimulates Rac activity at the opposite end of the cell through actomyosin-based blebbing.

### Contraction-induced membrane-to-cortex attachment asymmetry leads to PIP2 release and PI3K-dependent Rac activation at the opposite side of the cell

We found that Rac is recruited to blebs opposite the contractile back of the cell, but what determines the position of these blebs relative to the Rho-mediated contraction site? Blebs form when the plasma membrane detaches from the underlying actin cortex, and this process is facilitated by a low local concentration of membrane-to-cortex attachment (MCA) proteins. Localized contraction induces actomyosin flow toward the site of Rho mediated-contraction (32). This

flow could cause MCA proteins to accumulate at the site of contraction while depleting them from the opposite end of the cell (40–42) (Figure 4A). To test this hypothesis, we imaged cells expressing a fluorescent tagged ezrin, a core MCA protein in neutrophils. Upon local Rho activation, ezrin accumulated at the site of contraction and was depleted at the opposite side of the cell (Figure 4B, C; S6A-C and Video S7). To further validate these findings, we used actin-membrane proximity biosensor MPact (43) to assess MCA distribution during opto-contraction. This tool confirmed that local contraction enriches MCA at the contraction site while depleting it from the opposite side of the cell (Figure S6D-F). These results suggest that the depletion of MCA opposite to the contraction site promotes the formation of bleb-based sites of Rac activation.

We next explored the molecular mechanism linking blebbing to Rac activation, hypothesizing the lipid com-



**Fig. 4. Contraction-induced blebbing leads to PIP<sub>2</sub> release and PI3K-dependent Rac activation at the opposite side of the cell**

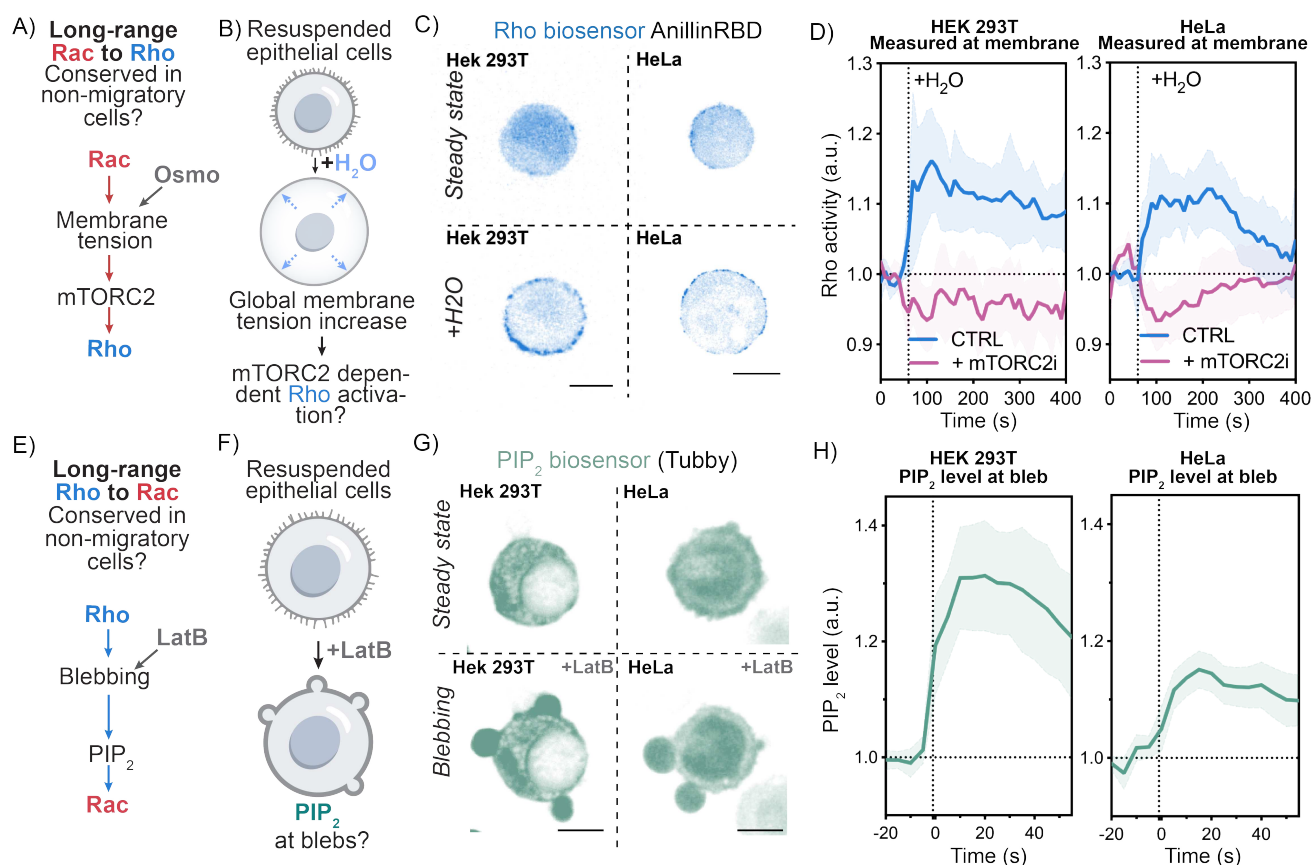
(A) In unpolarized cells at steady state, actin and MCA proteins are uniformly distributed. Upon local contraction (for example induced by opto-Rho), Actin flows toward the site of contraction. We ask whether MCA also flow with the cytoskeleton toward the site of contraction to induce an MCA asymmetry across the cell. (B) Time-lapse confocal images of unpolarized cells expressing the MCA protein Ezrin-mCherry and stained with the membrane marker CellMask before and during opto-Rho stimulation. (C) Average time trace of relative Ezrin intensity at the back (purple) and front (grey) of the cell in response to opto-Rho activation (at the back). (N = 3, n > 40, means ± 95%CI). We next hypothesized that MCA detachment from plasma membrane during blebbing leads to PIP<sub>2</sub> release. (D) Time-lapse confocal images of an unpolarized opto-Rac HL-60 cell expressing the PIP<sub>2</sub> biosensor (Tubby) before and during opto-Rho stimulation demonstrates PIP<sub>2</sub> enrichment in bleb-based protrusions. (E) Average time trace of PIP<sub>2</sub> accumulation at the protrusive blebs (opposite side of the cell from Rho activation) (mean ± 95%CI; n > 30, N = 3). (F) Average time trace of Rac activity at the plasma membrane at the protrusive blebs in control and cells treated with 1 μM PI3Kδ/γ inhibitor, Duvelisib demonstrates a requirement for PI3K for Rac activation in protrusive blebs (mean ± 95%CI; n > 30, N = 3). Control data is the same as in Figure 3E. (G) Local Rho activation leads to contraction-driven localized PIP<sub>2</sub> enrichment at the cell front which act a substrate of the Rac activator PI3K. Scale bars: 10 μm.

position of the blebs to play a critical role. Because MCA proteins such as ERMs (Ezrin, Radixin, Moesin) bind to the membrane through interactions with PIP<sub>2</sub> (44), detachment of MCA proteins from the membrane during blebbing could increase the local concentration of accessible PIP<sub>2</sub>. PI3K could act on this newly-released PIP<sub>2</sub> to produce PIP<sub>3</sub>, a known activator of Rac. To test this hypothesis, we used the PIP<sub>2</sub> biosensor Tubby (45) in conjunction with local opto-Rho activation and observed a marked increase in PIP<sub>2</sub> level at blebs induced by Rho mediated contraction (Figure 4D, E and Video S8). Next, we examined the functional role of this PIP<sub>3</sub> in Rac activation within blebs by inhibiting PI3K using Duvelisib (PI3Ki), a dual PI3Kδ/γ inhibitor (46) (Figures 4F and S6G, H). Cells treated with Duvelisib displayed a significant impairment Rac enrichment

downstream of both opto-Rho-mediated blebs (Figure 4F) as well as Latrunculin-induced stable blebs (Figure S6I, J). These findings provide a molecular mechanism to link local Rho activation to Rac activation at the opposite end of the cell. Local Rho-induced contraction redistributes MCA proteins away from the opposite end of the cell, promoting blebbing. This depletion of MCA proteins releases PIP<sub>2</sub>, which serves as a substrate for PI3K to generate the Rac activator PIP<sub>3</sub> (Figure 4G).

#### Core components of the mutual activation Rho and Rac activation are conserved in non-migratory cells

Our experiments demonstrate that Rho and Rac mutually activate one another at a distance using two distinct mechanisms: (1) membrane tension, which links front generation



**Fig. 5. Core components of Rho and Rac mutual long-range activation are conserved in non-migratory cells**

(A) Are the core components of the long-range communication from Rac to Rho conserved in non-migratory cells? (B) If membrane tension operates through mTORC2 to activate Rho in epithelial cells, then increasing membrane tension should suffice to activate Rho. Towards this end, we used hypotonic shock to elevate membrane tension in resuspended epithelial cells. (C) Confocal images of HeLa and 293T cells expressing the Rho biosensor Anillin-RBD before and during osmotic shock. (D) Average time trace of Rho activity at the plasma membrane in response for hypotonic shocks of control cells (in blue) or cells treated with the 10 $\mu$ M of the mTOR inhibitor KU-0063794 (in pink) (N>2, n>20, means  $\pm$  95%CI). (E) Are the core components of the long-range communication from Rho to Rac conserved in non-migratory cells? (F) To test whether localized PIP<sub>2</sub> release in blebs is conserved in non-migratory cells, we used resuspended epithelial cells treated with the actin inhibitor Latrunculin B to generate persistent cellular blebs. (G) Time-lapse confocal images of an unpolarized HeLa and 293T cells expressing the PIP<sub>2</sub> biosensor (Tubby) before and during Latrunculin B treatment demonstrates that PIP<sub>2</sub> enrichment in bleb-based protrusions is conserved in non-migratory cells. (H) Average time trace of PIP<sub>2</sub> accumulation at the blebs in HeLa and 293T cells treated with 2M of the actin inhibitor Latrunculin B (mean  $\pm$  95%CI; n>30, N= 2).

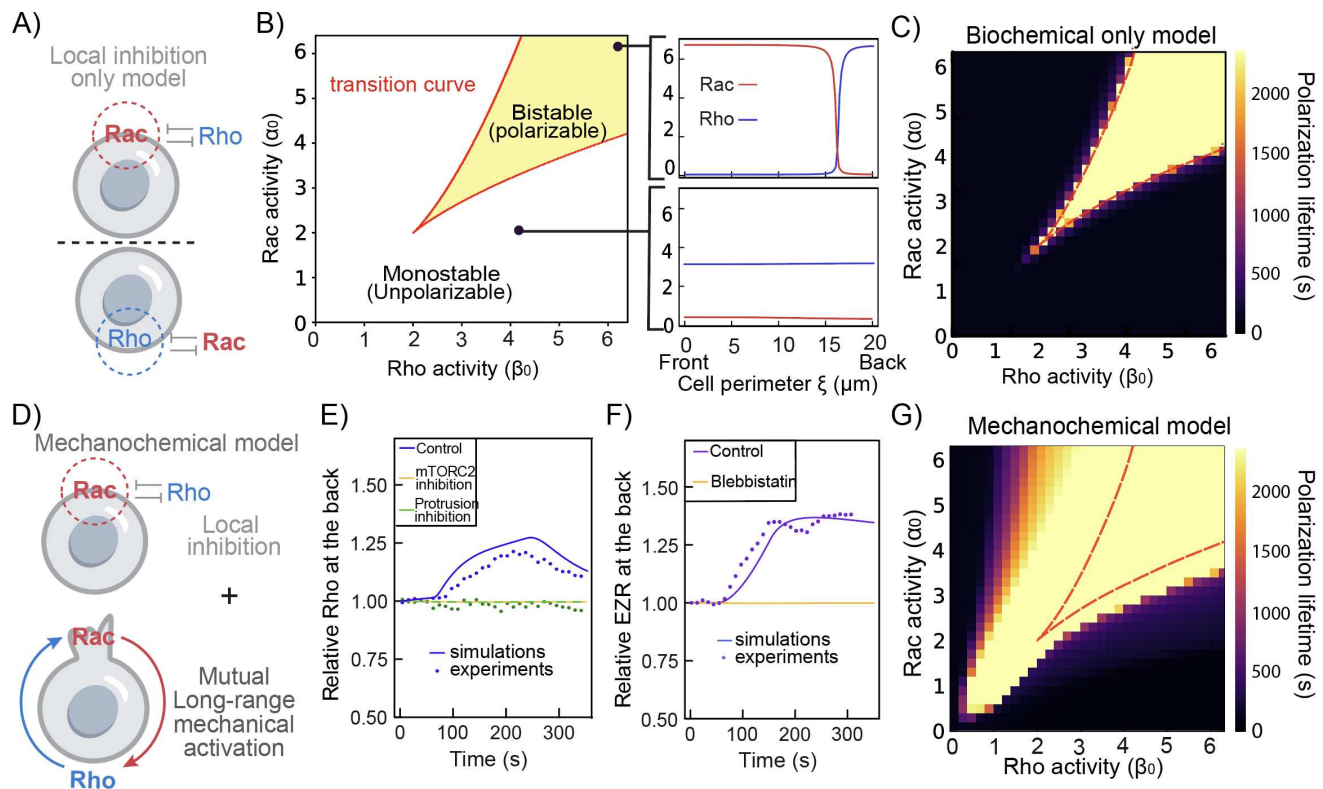
with back activation, and (2) cortical remodeling, which connects back activation with front generation. Given the broad conservation of the components of the pathways we have identified for this long-range coupling (membrane tension propagation, mTORC2 activation, cortical flows, and phospholipid changes), we next investigated the generality of this circuit beyond cell migration. First, we probed whether tension/mTORC2-mediated increases in Rho activity via mTORC2 is conserved in non-migratory cells (**Figure 5A**). To investigate this possibility, we used hypotonic shock in resuspended epithelial cells (Hek-293Ts and HeLa) to stimulate an increase in membrane tension (**Figure 5B**). Consistent with our previous findings, we observed that hypotonic shock induced a rapid and global increase in Rho activity across the cell (**Figures 5C, D**). Cells treated with mTOR inhibitor KU-0063794 failed to activate Rho activity in response to hypotonic shock. These data show a conservation of the pathway linking membrane tension to mTORC2 to Rho activation in non-migratory cells (**Figures 5D and S7**).

Next, we probed whether the MCA mediated release of PIP<sub>2</sub> in blebs is also conserved in non-migratory cells (**Figure 5E**). Towards this end, we used the actin inhibitor Latrunculin B to induce stable blebs in resuspended epithelial cells expressing the PIP<sub>2</sub> biosensor Tubby (**Figure 5F**) and observed a marked increase in PIP<sub>2</sub> in blebs in both HeLa and Hek 293T cells (**Figure 5G, H**). Together, our results show that the core components of the long-range mutual activation between Rac and Rho are conserved in non-migratory cells and are likely to account for the robustness of polarity in a number of other cellular contexts, such as embryo polarization, epithelial polarity, and asymmetric cell division, all of which involve long-range Rac/Rho patterning.

### Mechanochemical model of Rho and Rac partitioning in cells

To understand the key requirements for the long-range Rho and Rac mutual activation, we developed a minimal mechanochemical model of Rho and Rac partitioning. We





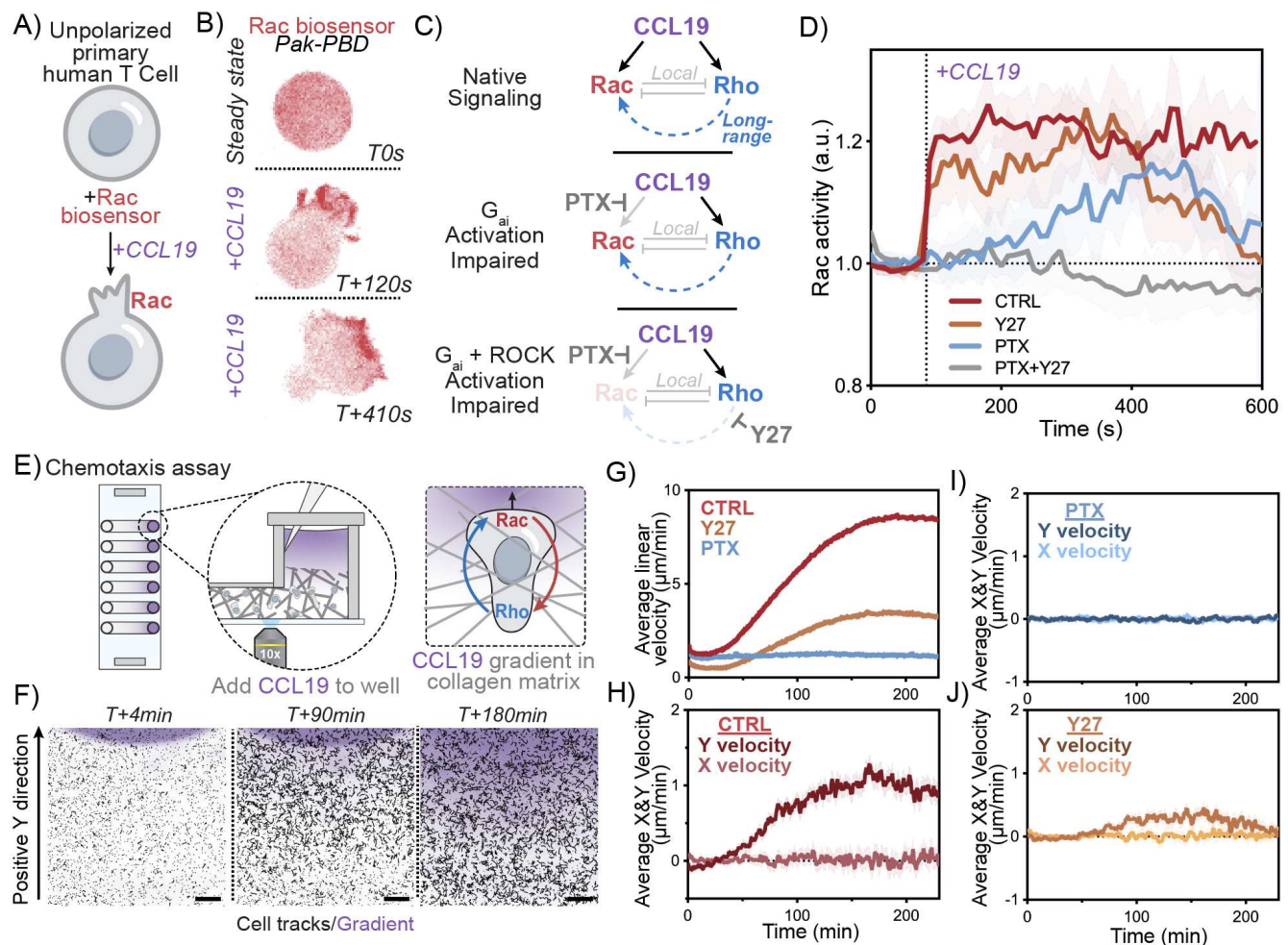
**Fig. 6. Modelling Rac and Rho polarization mechanisms through local inhibition and long-range mechanical feedback**

(A) Simple model of local inhibition between Rac and Rho (B) Phase diagram of polarity obtained from stability analysis using a simple model of local mutual inhibition between Rac and Rho. For any given Rac and Rho basal levels ( $\alpha$  and  $\beta$  respectively), the system is either monostable (unpolarized) or bistable (polarized). On the right, concentration values of Rac and Rho for simulations inside (Top) and outside (Bottom) the bistable region, where (Top) is polarized with a high value of Rac at the front of the cell and low at the back and vice-versa for Rho. (C) Heatmap of the time needed for a polarized cell to relax back to steady state in function of basal Rac and Rho activity ( $\alpha$  and  $\beta$  respectively) using local inhibition only. (D) Mechanochemical model of Rac and Rho polarity, combining local mutual inhibition with long-range mechanical feedback. (E) Rac is locally activated on one end (the 'front') of the cell, while Rho activity is measured at the opposite end (the 'back') of the simulated cell. Simulated (solid lines) and experimental (dots) Rho activity at the back of the cell following Rac activation at the front. Control cells are shown in blue, mTORC2 defective cells (Rictor KO) in orange, cells with inhibited protrusions (CK666) in green. (F) Simulated (solid lines) and experimental (dots) Ezrin levels at the back of the cell show that local Rho activity leads to MCA asymmetry. Control is shown in purple while cells with impaired contraction (blebbistatin) are shown in yellow. (G) Same as (C) but using fully integrated mechanochemical model.

first formulated a mathematical model using two coupled differential equations to describe the local mutual inhibition between Rac and Rho (47) (Figure 6A, Supplementary Text). The model defines two regions of stability for Rac and Rho, depending on their respective levels of antagonism-modulated activation, denoted  $\alpha_0$  and  $\beta_0$ . At low or highly asymmetric production levels, only Rho or Rac will dominate (monostable regime), while at sufficiently high levels of both GTPases, a bistable regime arises where they can coexist (Figure 6B, C). In the bistable regime, one could force the system locally to be either in a high Rac or Rho state, but polarization cannot persist in this simplified biochemical picture only (Supplementary Text, section 2.4). This highlights the limitations of polarity regulation based only on local inhibition, consistent with our previous experimental results (Figure S2A-D and 3F-H).

Next, we combined this mutual local inhibition model with a mechanical model describing the membrane-cortex interplay, building on previous works (32) (see Supplementary Theory, section 4) (Figure 6D). In this model,

the cortex is treated as a viscous-contractile layer attached to the membrane via MCA proteins of varying surface density  $\rho$ . The mechanically relevant membrane tension (also referred to as frame tension (48, 49)) is stored in entropic and cortical-flow-driven membrane ruffles, which are minimally modeled as a linearly elastic component. Rac is assumed to promote protrusion, modeled by a velocity ( $v_p$ ) at the front, while Rho mediates contraction at the back ( $\sigma_a$ ). Based on previous experimental findings, we further introduce two novel couplings: activation of active Rac is negatively modulated by MCA protein density, and Rho is enhanced by membrane tension, each governed by switch-like functions. Using this mechanochemical model, we successfully replicated our previous findings showing where local opto-Rac activation resulted in Rho activation at the cell's opposite (Figure 6E and S7). In line with our experiments, blocking Rac's influence on membrane tension (e.g., via drugs inhibiting protrusion) or disrupting tension's effects on Rho activation (e.g., in mTORC2 mutants) prevented Rac from stimulating Rho activation at the distal end. This integrated mechanochemical model



**Fig. 7. Long range mutual activation establishes robust Rho and Rac polarity during primary T cell migration**

(A) Primary human T cells expressing the Rac biosensor (Pak-PBD) to assay cell polarization following acute stimulation with the chemoattractant CCL19. (B) Time-lapse confocal images of an unpolarized human primary T cell before and during CCL19 stimulation. Rac activity was monitored via the Rac biosensor Pak-PBD. (C) Chemoattractants such as CCL19 are known to activate both Rho and Rac. (D) Average time trace of Rac activity at the cell front following addition of 25nM of CCL19 comparing control cells (red), cells treated with 1  $\mu$ g/ml of  $G_{\alpha i}$  inhibitor PTX (blue), cells treated with 20 $\mu$ M of Y27 (orange) and a combination of PTX and 20 $\mu$ M Y27 (grey). (mean  $\pm$  95%CI;  $n > 20$ ,  $N = 3$ ). (E) Ex-vivo assay for human primary T cell chemotaxis. Cells are premixed with Bovine Dermal Collagen and placed into linear channels. After collagen sets, media is added to one side of the channel (TCM) and media with human CCL19 (100ng total) and 10ug/mL of Dextrans 10k-AF647 (Dex647) is added to the other. Imaging takes place right on the edge of the well containing CCL19 so that T cell responses can be recorded as the CCL19 (as read out by Dex647) diffuses into the channel. (F) Representative control experiment is shown at different timepoints with Dex647 fluorescence (top row) and corresponding tracks of T cells (bottom row); tracks display 3 minutes preceding timepoint listed. (G) Average linear velocity of all tracks at each timepoint recorded. Control volunteer  $N = 3$ , PTX Volunteer  $N = 3$ , Y27 Volunteer  $N = 2$ . (H) Average Y only component of velocity of all tracks at each timepoint recorded. Positive is considered towards the well containing CCL19. Control Volunteer  $N = 3$ , PTX Volunteer  $N = 3$ , Y27 Volunteer  $N = 2$ .

explains how the front positions the back by combining local inhibition and long-range, tension-mediated Rho activation. We next simulated the reverse scenario by locally activating Rho and measuring Rac act the opposite end. The model captured how Rho-driven creates MCA asymmetry (Figure 6F) and induces Rac activation at the distal pole (Figure S7). The minor discrepancy between model and data for Rac activation may be due to variability in blebbing across cells, both in timing (Figure 3D) and quantity (Figure S5A).

Finally, we compared the persistence of polarity generated by the integrated mechanochemical model with that produced by two alternative models: one based solely on local Rac-Rho mutual inhibition, and another using

wave-pinning dynamics (16–19) driven by diffusion and mass conservation (Supplementary Theory, Section 3). Our results indicate that local mutual inhibition alone is insufficient to robustly partition Rho and Rac across the cell (Supplementary Theory, Figure 5). Although wave-pinning models can produce stable polarization, they exhibit hypersensitivity and a binary response to external stimuli or fluctuations in Rho and Rac—features that are inconsistent with the broader spectrum of cellular behaviors observed experimentally. In contrast, the integrated model, which combines local mutual inhibition with long-range mutual activation via mechanochemical coupling, successfully recapitulates experimentally observed behaviors. Specifically, it predicts three distinct regimes of cellular behavior

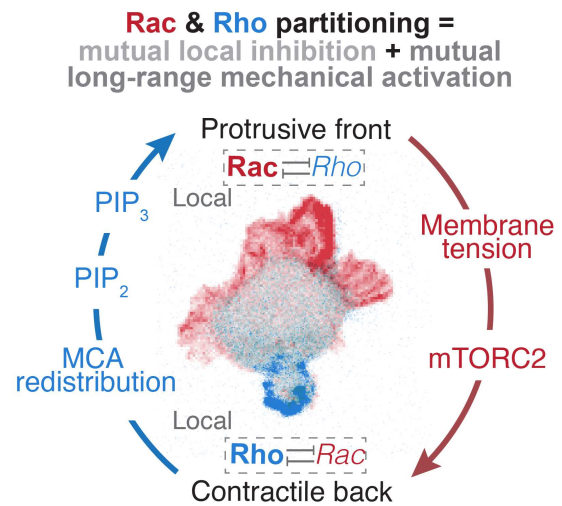
across a wide range of Rac and Rho basal production levels: (1) a responsive but non-polarizable state, characteristic of HL60 cells; (2) a responsive and polarizable state, in which sufficiently strong stimulation can induce persistent polarization through mechanochemical feedback, consistent with the response of primary human T cells (**Figure 7**); (3) a spontaneously polarized state, where the cell maintains a persistent polarized configuration without external cues (**Supplementary Theory, Figure 10**).

### Long range mutual activation establishes robust Rho and Rac polarity during primary T cell migration

Having shown that activating either Rho or Rac leads to a robust establishment of polarity and that Rho and Rac both mutually reinforce each other at a distance, we next investigated the consequences of this long-range cross talk for immune cell polarization and migration. To this end, we used primary human T cells expressing the Rac biosensor (Pak-PBD) to assay cell polarization following acute stimulation with the chemoattractant CCL19 (**Figure 7A**). We found that CCL19 induces rapid cell polarization with sustained Rac activity at the front (**Figure 7B, S8A-C and Video S9**).

CCL19 is known to activate both Rac and Rho (50, 51). We first tested if our observed long-lasting Rac partitioning depends solely on the known local activation of Rac via G $\alpha$ i signaling or if it also depends on the long-range activation via the Rho/myosin pathway defined in this work (**Figure 7C**). To focus on the role of Rho/myosin on Rac regulation, we first treated cell with pertussis toxin (PTX) to inactivate G $\alpha$ i signaling. These cells maintained their ability to stimulate Rac activation and polarization, albeit with a significant delay and more transiently than control cells (**Figure 7D and S8D-F**). The ability of PTX treated cells to polarize in response to chemokines has been previously reported but the mechanism by which this is achieved is unknown (6). To test whether this residual Rac regulation occurred through long-range Rho/myosin activation, we treated cells with both PTX and the actomyosin inhibitor Y27632 (**Figure 7D and S8G-I**). This combination completely abolished even transient Rac polarization, consistent with a long-range back-to-front signaling link that acts in conjunction with G $\alpha$ i-based Rac activation.

We previously showed that long-range Rac activation by Rho is dependent on PI3K which locally converts PIP<sub>2</sub> into the Rac activator PIP<sub>3</sub> at the cell front. We further investigated whether chemoattractant induced back-to-front signaling is also dependent on PI3K activation and found Rac polarization to be blocked when using both PTX (which blocks the G $\alpha$ i-based Rac activation) and the PI3K inhibitor Duvelisib (which blocks the long-range Rho-based Rac activation, **Figure S8J-L**). Finally, we show that preventing the long-range Rho to Rac communication using Y27632 alone elicits a more transient polarization than is seen for control cells (**Figure S8M-O**). These results demonstrate



**Fig. 8. Summary figure**

that chemoattractant signaling communicates with the front program in two different ways, one direct route via G $\alpha$ i and one indirect route via the back-mediated mechanism we define in this work. Cells deficient in either the direct or indirect(long-range) route can only polarize transiently, while WT primary cells combine both direct and indirect routes of Rac activation for robust and sustained polarization in response to chemoattractant.

Finally, we sought to test the role of this robust polarization established by the long-range Rac-Rho communication during chemoattractant induced migration. To stimulate chemotaxis, we embedded T cells in a 3D collagen matrix in a thin channel with CCL19 placed on one side (**Figure 7E**). As CCL19 freely diffuses into the channel, a chemotactic gradient is established, and control cells rapidly polarize and chemotax up the CCL19 gradient (**Figure 7F-H and Video S10**). In contrast, cells treated with PTX or Y27 only briefly polarized and failed to robustly migrate toward the CCL19 gradient (**Figure 7G, I-J and S8Q**).

Our results show that long-range mutual activation of front and back is an essential feature of human lymphocyte polarization and motility. Furthermore, we show that activating either the front or back polarity programs without the other only leads to transient polarization. Dual activation is required to support sustained polarization and efficient chemotactic migration.

## Discussion

Our work demonstrates that the well-appreciated mutual short-range inhibition between Rac and Rho is accompanied by an additional mutual long-range facilitation of the front and back polarity programs in migrating cells (**Figure 8**). The front reinforces the back via membrane tension mediated mTORC2 activation (**Figure 1, 2 S1-4**) The back reinforces the front via cortical remodeling and localized PIP<sub>2</sub> release (**Figure 3, 4 and S5, S6**). Both modes of long-range



facilitation are required for sustained and efficient front organization and chemotaxis in primary human lymphocytes (**Figure 7**). Understanding the partitioning of Rho and Rac in migrating cells has long been a challenge due to the difficulty in establishing the range and directionality of interactions of Rac and Rho. Our use of optogenetics in unpolarized cells enables us to dissect the long-range positive interactions between the front and back programs, and our combined mechanical and biochemical models complement our experimental work by revealing the importance of these links to cell polarization. Our combined multidisciplinary approach is likely to be similarly powerful for other complex interconnected spatiotemporal signaling networks such as axon specification (52), asymmetric cell division (53), epithelial polarity (54), planar cell polarity (55), branching morphogenesis (56), lumen formation (57), and collective cell migration (58).

Robust cell polarization necessitates long-range integration of signals across the cell. A number of currencies have been proposed for this long-range signal relay: diffusion of signals across the cell (14, 17, 47, 59, 60) or force-mediated communication via propagation of forces in the membrane or cortex (26–28, 30, 61) or advection of cellular components via force asymmetries across the cell (62–64). Membrane tension rapidly propagates across cells and is used for long-range integration of cellular processes from cell spreading (65, 66) to phagocytosis (67) to the competition of fronts that enables a dominant protrusion in migrating cells (27, 28). Tension-based communication is particularly powerful for global signal transmission across the cell (37). Our work shows that membrane tension also enables the front to facilitate contraction at the cell rear through mTORC2 activation. Previous studies (34, 68–73), including our own (28, 29), have broadly linked TORC2 and mTORC2 to cell polarity, but how mTORC2 does so is not well-understood. Even its links to myosin regulation have been a point of contention, with some studies showing a positive role (29) and others proposing a negative role (34). Our work provides direct evidence for and a molecular mechanism for this link (**Figure 2**). The evolutionarily-conserved mTORC2 is also the mechanosensitive pathway that enables competition between fronts (28, 29), regulation of plasma membrane homeostasis in budding yeast (74, 75), and chemotaxis in cells from neutrophils (34) to Dictyostelium (70). Given the evolutionarily conserved role of mTORC2, it would be interesting to probe in future work whether the mTORC2 mediated activation of Rho in response to elevated membrane tension is important in other contexts such as cell division.

Force-mediated flow of components is another powerful mechanism of long-range cell communication (62, 76, 77). Our work shows that flow of MCA components secondary to Rho-mediated actin-based contractions enables the cell rear to establish a Rac-based protrusive front at the opposite end of the cell. These flow-mediated modes of communication

are an efficient means for the polarized distribution of signals—in this case the depletion of MCA sites and the facilitation of Rac activation at the opposite end of the cell from Rho-mediated contractions. Flow of cellular signals from the cell front to the cell back along retrograde flows or cortical contractions have been proposed to enable robust polarization (62, 76, 77). Our work provides a molecular mechanism for this long-range integration. It is interesting that the long-range back-to-front and front-to-back communication involve different currencies of membrane tension and cortical flow. These different currencies have different spatial patterns of propagation (relatively global for membrane tension and polarized for cortical flows) and also differ in their molecular requirements, potentially enabling these modes of communication to be independently modulated without interfering with one another.

Rho GTPases are central regulators of the cytoskeleton, and their partitioning underlie many fundamental cellular functions such as cell shape, adhesions and division. In the context of cell locomotion, we find that long-range activation is a key complement to local inhibition to ensure the proper partitioning of Rac and Rho at opposite poles of the cell. Our findings and model highlight the limitations of a system that relies only on local interactions for spatial segregation. We speculate that the coupling between local biochemical interactions and long-range mechanical facilitation is a conserved feature of Rho GTPases partitioning. This is in good agreement with other systems in which Rho GTPase partitioning relies on long-range communication via the actin and the membrane, from polarization in early *C. elegans* embryos by cortical flow (78), to axon branching in neuron via long-range membrane tension propagation (52), to the winner-take-all competition enabling a single front during migration (27, 28). Long-range mechanochemical interactions represent a conserved mechanism to complement local biochemical interactions for the partitioning of Rho GTPases in cell migration and likely other processes.

#### ACKNOWLEDGEMENTS

We thank the Weiner and Turler lab for their critical feedback. We especially want to thank Kate Cavanaugh for kindly sharing the PIP2 biosensor construct and Juan Manuel García-Arcos for his helpful insights. We also thank the blood donors that made the primary human T cell experiment possible. This work was supported by R35GM118167 (ODW, ES), National Science Foundation/Biotechnology and Biological Sciences Research Council grant 2019598 (ODW), K99GM154115 (HDB), the European Research Council (ERC) under the European Union's Horizon 2020 research and innovation program (Grant agreement No. 949267) (HT), the European Union's Horizon 2020 research and innovation program under the Marie Skłodowska-Curie grant agreement No. 101150259 (AFG), the American Heart Association predoctoral fellowship 24PRE1198865 (PJZ) and R21AI1739938 (JKB). HT and AFG are supported by the CNRS and Collège de France.

#### Bibliography

1. William M. Bement, Andrew B. Goryachev, Ann L. Miller, and George von Dassow. Patterning of the cell cortex by rho GTPases. pages 1–19. ISSN 1471-0080. doi: 10.1038/s41580-023-00682-z. Publisher: Nature Publishing Group.
2. Juan Manuel García-Arcos, Ankita Jha, Clare M. Waterman, and Matthieu Piel. Blebology: principles of bleb-based migration. 34(10):838–853. ISSN 1879-3088. doi: 10.1016/j.tcb.2024.02.009.
3. Matteo Parri and Paola Chiarugi. Rac and rho GTPases in cancer cell motility control. 8(1): 23. ISSN 1478-811X. doi: 10.1186/1478-811X-8-23.
4. Olivier Pertz, Louis Hodgson, Richard L. Klemke, and Klaus M. Hahn. Spatiotemporal dynamics of RhoA activity in migrating cells. 440(7087):1069–1072. ISSN 1476-4687. doi: 10.1038/nature04665.

5. K. Rottner, A. Hall, and J. V. Small. Interplay between rac and rho in the control of substrate contact dynamics. 9(12):640–648. ISSN 0960-9822. doi: 10.1016/s0960-9822(99)80286-3.
6. Jingsong Xu, Fei Wang, Alexandra Van Keymeulen, Paul Herzmack, Aaron Straight, Kathleen Kelly, Yoh Takuwa, Naotoshi Sugimoto, Timothy Mitchison, and Henry R. Bourne. Divergent signals and cytoskeletal assemblies regulate self-organizing polarity in neutrophils. 114(2):201–214. ISSN 0092-8674. doi: 10.1016/s0092-8674(03)00555-5.
7. M. Hirose, T. Ishizaki, N. Watanabe, M. Uehata, O. Kranenburg, W. H. Moolenaar, F. Matsumura, M. Maekawa, H. Bito, and S. Narumiya. Molecular dissection of the rho-associated protein kinase (p160rock)-regulated neurite remodeling in neuroblastoma n1e-115 cells. 141(7):1625–1636. ISSN 0021-9525. doi: 10.1083/jcb.141.7.1625.
8. Erik Sahai, Raquel Garcia-Medina, Jacques Pouyssegur, and Emmanuel Vial. Smurf1 regulates tumor cell plasticity and motility through degradation of RhoA leading to localized inhibition of contractility. 176(1):35–42. ISSN 0021-9525. doi: 10.1083/jcb.200605135.
9. Victoria Sanz-Moreno, Gilles Gadea, Jessica Ahn, Hugh Paterson, Pierfrancesco Marra, Sophie Pinner, Erik Sahai, and Christopher J. Marshall. Rac activation and inactivation control plasticity of tumor cell movement. 135(3):510–523. ISSN 0092-8674. doi: 10.1016/j.cell.2008.09.043.
10. Alfonso Bolado-Carrancio, Oleksii S Rukhlenko, Elena Nikonova, Mikhail A Tsyganov, Anne Wheeler, Amaya Garcia-Munoz, Walter Kolch, Alex von Kriegsheim, and Boris N Kholodenko. Periodic propagating waves coordinate RhoGTPase network dynamics at the leading and trailing edges during cell migration. 9:e58165. ISSN 2050-084X. doi: 10.7554/eLife.58165. Publisher: eLife Sciences Publications, Ltd.
11. Rafael Dominik Fritz and Olivier Pertz. The dynamics of spatio-temporal rho GTPase signaling: formation of signaling patterns. 5:F1000 Faculty Rev–749. ISSN 2046-1402. doi: 10.12688/f1000research.7370.1.
12. Hee Won Yang, Sean R. Collins, and Tobias Meyer. Locally excitable cdc42 signals steer cells during chemotaxis. 18(2):191–201. ISSN 1476-4679. doi: 10.1038/ncb3292. Publisher: Nature Publishing Group.
13. Katrin Martin, Andreas Reimann, Rafael D. Fritz, Hyunryul Ryu, Noo Li Jeon, and Olivier Pertz. Spatio-temporal co-ordination of RhoA, rac1 and cdc42 activation during prototypical edge protrusion and retraction dynamics. 6:21901. ISSN 2045-2322. doi: 10.1038/srep21901.
14. Peter N. Devreotes, Sayak Bhattacharya, Marc Edwards, Pablo A. Iglesias, Thomas Lampert, and Yuchuan Miao. Excitable signal transduction networks in directed cell migration. 33:103–125. ISSN 1530-8995. doi: 10.1146/annurev-cellbio-100616-060739.
15. Yu Deng, Tatsat Banerjee, Dhiman Sankar Pal, Parijat Banerjee, Huiwang Zhan, Jane Borleis, Pablo A. Iglesias, and Peter N. Devreotes. PIP5k-ras bistability initiates plasma membrane symmetry breaking to regulate cell polarity and migration. Pages: 2024.09.15.613115 Section: New Results.
16. Alexandra Jilkine, Athanasios F. M. Marée, and Leah Edelstein-Keshet. Mathematical model for spatial segregation of the rho-family GTPases based on inhibitory crosstalk. 69(6):1943–1978. ISSN 0092-8240, 1522-9602. doi: 10.1007/s11538-007-9200-6.
17. Yoichiro Mori, Alexandra Jilkine, and Leah Edelstein-Keshet. Wave-pinning and cell polarity from a bistable reaction-diffusion system. 94(9):3684–3697. ISSN 0006-3495. doi: 10.1529/biophysj.107.120824.
18. Kai H. Kopfer, Willi Jäger, and Franziska Matthäus. A mechanochemical model for rho GTPase mediated cell polarization. 504:110386. ISSN 0022-5193. doi: 10.1016/j.jtbi.2020.110386.
19. Cole Zmurchok, Dhananjay Bhaskar, and Leah Edelstein-Keshet. Coupling mechanical tension and GTPase signaling to generate cell and tissue dynamics. 15(4):046004. ISSN 1478-3975. doi: 10.1088/1478-3975/aab1c0. Publisher: IOP Publishing.
20. Kit Wong, Olivier Pertz, Klaus Hahn, and Henry Bourne. Neutrophil polarization: Spatiotemporal dynamics of RhoA activity support a self-organizing mechanism. 103(10):3639–3644. doi: 10.1073/pnas.0600092103. Publisher: Proceedings of the National Academy of Sciences.
21. Yanqin Wang, Chin-Jen Ku, Elizabeth R. Zhang, Alexander B. Artyukhin, Orion D. Weiner, Lani F. Wu, and Steven J. Altschuler. Identifying network motifs that buffer front-to-back signaling in polarized neutrophils. 3(5):1607–1616. ISSN 2211-1247. doi: 10.1016/j.celrep.2013.04.009.
22. Angela H. Chau, Jessica M. Walter, Jaline Gerardin, Chao Tang, and Wendell A. Lim. Designing synthetic regulatory networks capable of self-organizing cell polarization. 151(2):320–332. ISSN 0092-8674, 1097-4172. doi: 10.1016/j.cell.2012.08.040. Publisher: Elsevier.
23. Wenzhe Ma, Ala Trusina, Hana El-Samad, Wendell A. Lim, and Chao Tang. Defining network topologies that can achieve biochemical adaptation. 138(4):760–773. ISSN 0092-8674, 1097-4172. doi: 10.1016/j.cell.2009.06.013. Publisher: Elsevier.
24. Alexis J. Lomakin, Kun-Chun Lee, Sangyoon J. Han, Duyen A. Bui, Michael Davidson, Alex Mogilner, and Gaudenz Danuser. Competition for actin between two distinct f-actin networks defines a bistable switch for cell polarization. 17(11):1435–1445. ISSN 1476-4679. doi: 10.1038/ncb3246. Publisher: Nature Publishing Group.
25. Calina Copos and Alex Mogilner. A hybrid stochastic-deterministic mechanochemical model of cell polarization. 31(15):1637–1649. ISSN 1059-1524. doi: 10.1091/mbc.E19-09-0549. Publisher: American Society for Cell Biology (mboc).
26. Jan Mueller, Gregory Sze, Maria Nemethova, Ingrid de Vries, Arnon D. Lieber, Christoph Winkler, Karsten Kruse, J. Victor Small, Christian Schmeiser, Kinneret Keren, Robert Hauschild, and Michael Sixt. Load adaptation of lamellipodial actin networks. 171(1):188–200.e16. ISSN 1097-4172. doi: 10.1016/j.cell.2017.07.051.
27. Andrew R. Houk, Alexandra Jilkine, Cecile O. Mejean, Rostislav Boltyskiy, Eric R. Dufresne, Sigurd B. Angenent, Steven J. Altschuler, Lani F. Wu, and Orion D. Weiner. Membrane tension maintains cell polarity by confining signals to the leading edge during neutrophil migration. 148(1):175–188. ISSN 1097-4172. doi: 10.1016/j.cell.2011.10.050.
28. Alba Diz-Muñoz, Kevin Thurlay, Sana Chintamen, Steven J. Altschuler, Lani F. Wu, Daniel A. Fletcher, and Orion D. Weiner. Membrane tension acts through PLD2 and mTORC2 to limit actin network assembly during neutrophil migration. 14(6):e1002474. ISSN 1545-7885. doi: 10.1371/journal.pbio.1002474.
29. Suvrajit Saha, Jason P. Town, and Orion D. Weiner. Mechanosensitive mTORC2 indepen-
- dently coordinates leading and trailing edge polarity programs during neutrophil migration. 34(5):ar35. ISSN 1059-1524. doi: 10.1091/mbc.E22-05-0191. Publisher: American Society for Cell Biology (mboc).
30. Arnon D. Lieber, Shlomit Yehudai-Resheff, Erin L. Barnhart, Julie A. Theriot, and Kinneret Keren. Membrane tension in rapidly moving cells is determined by cytoskeletal forces. 23(15):1409–1417. ISSN 0960-9822. doi: 10.1016/j.cub.2013.05.063.
31. Jason Town and Orion Weiner. Rac negative feedback links local PIP3 rate-of-change to dynamic control of neutrophil guidance.
32. Henry De Belly, Shannon Yan, Hudson Borja da Rocha, Sacha Ichbiah, Jason P. Town, Patrick J. Zager, Dorothy C. Estrada, Kirstin Meyer, Hervé Turlier, Carlos Bustamante, and Orion D. Weiner. Cell protrusions and contractions generate long-range membrane tension propagation. 186(14):3049–3061.e15. ISSN 0092-8674. doi: 10.1016/j.cell.2023.05.014.
33. Chloé Roffay, Guillaume Molinard, Kyoohyun Kim, Marta Urbanska, Virginia Andrade, Victoria Barbarasa, Paulina Nowak, Vincent Mercier, José García-Calvo, Stefan Matile, Robbie Loewith, Arnaud Echard, Jochen Guck, Martin Lenz, and Aurélien Roux. Passive coupling of membrane tension and cell volume during active response of cells to osmosis. 118(47):e2103228118. doi: 10.1073/pnas.2103228118.
34. Lunhua Liu, Satarupa Das, Wolfgang Losert, and Carole A. Parent. mTORC2 regulates neutrophil chemotaxis in a cAMP- and RhoA-dependent fashion. 19(6):845–857. ISSN 1878-1551. doi: 10.1016/j.devcel.2010.11.004.
35. Yuan He, Dong Li, Sara L. Cook, Mee-Sup Yoon, Ashish Kapoor, Christopher V. Rao, Paul J. A. Kenis, Jie Chen, and Fei Wang. Mammalian target of rapamycin and rictor control neutrophil chemotaxis by regulating rac/cdc42 activity and the actin cytoskeleton. 24(21):3369–3380. ISSN 1059-1524. doi: 10.1091/mbc.E13-07-0405.
36. Ewa K Paluch and Erez Raz. The role and regulation of blebs in cell migration. 25(5):582–590. ISSN 0955-0674. doi: 10.1016/j.cel.2013.05.005.
37. Henry De Belly and Orion D. Weiner. Follow the flow: Actin and membrane act as an integrated system to globally coordinate cell shape and movement. 89:102392. ISSN 0955-0674. doi: 10.1016/j.cel.2024.102392.
38. Brian R. Graziano, Jason P. Town, Ewa Sitarska, Tamas L. Nagy, Miha Fošnarič, Samo Penič, Aleš Igljič, Veronika Kralj-Igljič, Nir S. Gov, Alba Diz-Muñoz, and Orion D. Weiner. Cell confinement reveals a branched-actin independent circuit for neutrophil polarity. 17(10):e3000457. ISSN 1545-7885. doi: 10.1371/journal.pbio.3000457.
39. Meghan K. Driscoll, Erik S. Wolf, Andrew Weems, Etai Sapoznik, Felix Zhou, Vasanth S. Murali, Juan Manuel García-Arcos, Minna Roh-Johnson, Matthieu Piel, Kevin M. Dean, Reto Fiolka, and Gaudenz Danuser. Proteolysis-free amoeboid migration of melanoma cells through crowded environments via bleb-driven worrying. pages S1534–S807(24)00342–3. ISSN 1878-1551. doi: 10.1016/j.devcel.2024.05.024.
40. Chantal Roubinet, Barbara Decelle, Gaëtan Chicanne, Jonas F. Dorn, Bernard Payrastre, François Payre, and Sébastien Carreno. Molecular networks linked by moesin drive remodeling of the cell cortex during mitosis. 195(1):99–112. ISSN 0021-9525. doi: 10.1083/jcb.201106048.
41. Shota Hiruma, Tomoko Kamasaki, Kohei Otomo, Tomomi Nemoto, and Ryota Uehara. Dynamics and function of ERM proteins during cytokinesis in human cells. 591(20):3296–3309. ISSN 1873-3468. doi: 10.1002/1873-3468.12844. \_eprint: <https://onlinelibrary.wiley.com/doi/pdf/10.1002/1873-3468.12844>.
42. Elgin Korkmazhan and Alexander R. Dunn. The membrane-actin linker ezrin acts as a sliding anchor. 8(31):eabo2779. doi: 10.1126/sciadv.abo2779. Publisher: American Association for the Advancement of Science.
43. Anjali Bisaria, Arnold Hayer, Damien Garbett, Daniel Cohen, and Tobias Meyer. Membrane proximal f-actin restricts local membrane protrusions and directs cell migration. 368(6496):1205–1210. ISSN 0036-8075. doi: 10.1126/science.aay7794.
44. Andrea I. McClatchey. ERM proteins at a glance. 127(15):3199–3204. ISSN 0021-9533, 1477-9137. doi: 10.1242/jcs.098343.
45. Kathryn V Quinn, Philippe Behe, and Andrew Tinker. Monitoring changes in membrane phosphatidylinositol 4,5-bisphosphate in living cells using a domain from the transcription factor tubby. 586:2855–2871. ISSN 0022-3751. doi: 10.1113/jphysiol.2008.153791.
46. David G. Winkler, Kerrie L. Faia, Jonathan P. DiNitto, Janid A. Ali, Kerry F. White, Erin E. Brophy, Melissa M. Pink, Jennifer L. Proctor, Jennifer Lussier, Christian M. Martin, Jennifer G. Hoyt, Bonnie Tillotson, Erin L. Murphy, Alice R. Lim, Brian D. Thomas, John R. Macdougall, Pingda Ren, Yi Liu, Lian-Sheng Li, Katti A. Jessen, Christian C. Fritz, Jol L. Dunbar, James R. Porter, Christian Rommel, Vito J. Palombella, Paul S. Changelian, and Jeffery L. Kutok. PI3k-δ and PI3k-γ inhibition by IPI-145 abrogates immune responses and suppresses activity in autoimmune and inflammatory disease models. 20(11):1364–1374. ISSN 1879-1301. doi: 10.1016/j.chembiol.2013.09.017.
47. Kate M. Byrne, Naser Monsefi, John C. Dawson, Andrea Degasperi, Jimi-Carlo Bukowski-Wills, Natalia Volinsky, Maciej Dobrzyński, Marc R. Birtwistle, Mikhail A. Tsyganov, Anatoly Kiyatkin, Katarzyna Kida, Andrew J. Finch, Neil O. Carragher, Walter Kolch, Lan K. Nguyen, Alex von Kriegsheim, and Boris N. Kholodenko. Bistability in the rac1, PAK, and RhoA signaling network drives actin cytoskeleton dynamics and cell motility switches. 2(1):38–48. ISSN 2405-4712. doi: 10.1016/j.cels.2016.01.003.
48. Hayato Shiba, Hiroshi Noguchi, and Jean-Baptiste Fournier. Monte carlo study of the frame, fluctuation and internal tensions of fluctuating membranes with fixed area. 12(8):2373–2380. ISSN 1744-6848. doi: 10.1039/C5SM01900A. Publisher: The Royal Society of Chemistry.
49. Marc Durand. Frame tension governs the thermal fluctuations of a fluid membrane: new evidence. 18(20):3891–3901. ISSN 1744-6848. doi: 10.1039/D1SM01765A. Publisher: The Royal Society of Chemistry.
50. José Luis Rodríguez-Fernández and Olga Criado-García. The chemokine receptor CCR7 uses distinct signaling modules with biased functionality to regulate dendritic cells. 11. ISSN 1664-3224. doi: 10.3389/fimmu.2020.00528. Publisher: Frontiers.
51. Lorena Riol-Blanco, Noelia Sánchez-Sánchez, Ana Torres, Alberto Tejedor, Shuh Narumiya, Angel L. Corbi, Paloma Sánchez-Mateos, and José Luis Rodríguez-Fernández. The chemokine receptor CCR7 activates in dendritic cells two signaling modules that independently regulate chemotaxis and migratory speed. 174(7):4070–4080. ISSN 0022-1767. doi: 10.4049/jimmunol.174.7.4070.

52. Zheng Shi, Sarah Innes-Gold, and Adam E. Cohen. Membrane tension propagation couples axon growth and collateral branching.
53. Bharath Sunchu and Clemens Cabernard. Principles and mechanisms of asymmetric cell division. 147(13):dev167650. ISSN 0950-1991. doi: 10.1242/dev.167650.
54. Clare E. Buckley and Daniel St Johnston. Apical-basal polarity and the control of epithelial form and function. 23(8):559–577. ISSN 1471-0080. doi: 10.1038/s41580-022-00465-y. Publisher: Nature Publishing Group.
55. Mitchell T. Butler and John B. Wallingford. Planar cell polarity in development and disease. 18(6):375–388. ISSN 1471-0080. doi: 10.1038/nrm.2017.11. Publisher: Nature Publishing Group.
56. Tsuyoshi Hirashima and Michiyuki Matsuda. ERK-mediated curvature feedback regulates branching morphogenesis in lung epithelial tissue. 34(4):683–696.e6. ISSN 1879-0445. doi: 10.1016/j.cub.2023.12.049.
57. Dhiraj Indana, Andrei Zakharov, Youngbin Lim, Alexander R. Dunn, Nidhi Bhutani, Vivek B. Shenoy, and Ovijit Chaudhuri. Lumen expansion is initially driven by apical actin polymerization followed by osmotic pressure in a human epiblast model. 31(5):640–656.e8. ISSN 1934-5909. doi: 10.1016/j.stem.2024.03.016.
58. Justin Bui, Daniel E. Conway, Rebecca L. Heise, and Seth H. Weinberg. Mechanochemical coupling and junctional forces during collective cell migration. 117(1):170–183. ISSN 0006-3495. doi: 10.1016/j.bpj.2019.05.020.
59. Hans Meinhardt. Orientation of chemotactic cells and growth cones: models and mechanisms. 112(17):2867–2874. ISSN 0021-9533. doi: 10.1242/jcs.112.17.2867.
60. Steven J. Altschuler, Sigurd B. Angenent, Yanqin Wang, and Lani F. Wu. On the spontaneous emergence of cell polarity. 454(7206):886–889. ISSN 1476-4687. doi: 10.1038/nature07119. Publisher: Nature Publishing Group.
61. Juan Manuel García-Arcos, Johannes Ziegler, Silvia Grigolon, Loïc Reymond, Gaurav Shajepal, Cédric J. Cattin, Alexis Lomakin, Daniel J. Müller, Verena Ruprecht, Stefan Wieser, Raphael Voituriez, and Matthieu Piel. Rigidity percolation and active advection synergize in the actomyosin cortex to drive amoeboid cell motility. 0(0), . ISSN 1534-5807. doi: 10.1016/j.devcel.2024.06.023. Publisher: Elsevier.
62. Tony Y.-C. Tsai, Sean R. Collins, Caleb K. Chan, Amalia Hadjitheodorou, Pui-Ying Lam, Sunny S. Lou, Hee Won Yang, Julianne Jorgensen, Felix Ellett, Daniel Irimia, Michael W. Davidson, Robert S. Fischer, Anna Huttenlocher, Tobias Meyer, James E. Ferrell, and Julie A. Theriot. Efficient front-rear coupling in neutrophil chemotaxis by dynamic myosin II localization. 49(2):189–205.e6. ISSN 1878-1551. doi: 10.1016/j.devcel.2019.03.025.
63. Kotryna Vaidzilytė, Anne-Sophie Macé, Aude Battistella, William Beng, Kristine Schauer, and Mathieu Coppey. Persistent cell migration emerges from a coupling between protrusion dynamics and polarized trafficking. 11:e69229. ISSN 2050-084X. doi: 10.7554/eLife.69229. Publisher: eLife Sciences Publications, Ltd.
64. Xiaoguang Li, Dhiman Sankar Pal, Debojyoti Biswas, Pablo A. Iglesias, and Peter N. Devreotes. Reverse fountain flow of phosphatidylinositol-3,4-bisphosphate polarizes migrating cells. 40(4):e105094. ISSN 1460-2075. doi: 10.15252/embj.2020105094.
65. Drazen Raucher and Michael P. Sheetz. Cell spreading and lamellipodial extension rate is regulated by membrane tension. 148(1):127–136. ISSN 0021-9525.
66. Nils C. Gauthier, Marc Antoine Fardin, Pere Roca-Cusachs, and Michael P. Sheetz. Temporary increase in plasma membrane tension coordinates the activation of exocytosis and contraction during cell spreading. 108(35):14467–14472. ISSN 0027-8424. doi: 10.1073/pnas.1105845108.
67. Thomas A. Masters, Bruno Pontes, Virgile Viasnoff, You Li, and Nils C. Gauthier. Plasma membrane tension orchestrates membrane trafficking, cytoskeletal remodeling, and biochemical signaling during phagocytosis. 110(29):11875–11880. ISSN 1091-6490. doi: 10.1073/pnas.1301766110.
68. Pascale G. Charest, Zhouxin Shen, Ashley Lakoduk, Atsuo T. Sasaki, Steven P. Briggs, and Richard A. Firtel. A ras signaling complex controls the RasC-TORC2 pathway and directed cell migration. 18(5):737–749. ISSN 1878-1551. doi: 10.1016/j.devcel.2010.03.017.
69. Avais M. Daulat, François Bertucci, Stéphane Audebert, Arnaud Sérgé, Pascal Finetti, Emmanuelle Josselin, Rémy Castellano, Daniel Birnbaum, Stéphane Angers, and Jean-Paul Borg. PRICKLE1 contributes to cancer cell dissemination through its interaction with mTORC2. 37(4):311–325. ISSN 1878-1551. doi: 10.1016/j.devcel.2016.04.011.
70. Yoichiro Kamimura, Yuan Xiong, Pablo A. Iglesias, Oliver Hoeller, Parvin Bolourani, and Peter N. Devreotes. PIP3-independent activation of TorC2 and PKB at the cell's leading edge mediates chemotaxis. 18(14):1034–1043. ISSN 0960-9822. doi: 10.1016/j.cub.2008.06.068.
71. Susan Lee, Frank I. Comer, Atsuo Sasaki, Ian X. McLeod, Yung Duong, Koichi Okumura, John R. Yates, Carole A. Parent, and Richard A. Firtel. TOR complex 2 integrates cell movement during chemotaxis and signal relay in dictyostelium. 16(10):4572–4583. ISSN 1059-1524. doi: 10.1091/mbc.e05-04-0342.
72. Lunhua Liu, Derek Gritz, and Carole A. Parent. PKC $\beta$  acts downstream of chemoattractant receptors and mTORC2 to regulate cAMP production and myosin II activity in neutrophils. 25(9):1446–1457. ISSN 1939-4586. doi: 10.1091/mbc.E14-01-0037.
73. Fei Zhang, Xiaofang Zhang, Menghui Li, Peng Chen, Bin Zhang, Hua Guo, Wenfeng Cao, Xiying Wei, Xuchen Cao, Xishan Hao, and Ning Zhang. mTOR complex component rictor interacts with PKC $\zeta$  and regulates cancer cell metastasis. 70(22):9360–9370. ISSN 1538-7445. doi: 10.1158/0008-5472.CAN-10-0207.
74. Brad J. Niles, Huzefa Mogri, Andrew Hill, Ariadne Vlahakis, and Ted Powers. Plasma membrane recruitment and activation of the AGC kinase ypk1 is mediated by target of rapamycin complex 2 (TORC2) and its effector proteins slm1 and slm2. 109(5):1536–1541. doi: 10.1073/pnas.1117563109. Publisher: Proceedings of the National Academy of Sciences.
75. Margot Riggi, Beata Kusmider, and Robbie Loewith. The flipside of the TOR coin – TORC2 and plasma membrane homeostasis at a glance. 133(9):jcs242040. ISSN 0021-9533. doi: 10.1242/jcs.242040.
76. Paolo Maiuri, Jean-François Rupprecht, Stefan Wieser, Verena Ruprecht, Olivier Bénichou, Nicolas Carpi, Mathieu Coppey, Simon De Beco, Nir Gov, Carl-Philipp Heisenberg, Carolina Lage Crespo, Franziska Lautenschlaeger, Maël Le Berre, Ana-Maria Lennon-Dumenil, Matthew Raab, Hawa-Racine Thiam, Matthieu Piel, Michael Sixt, and Raphaël Voituriez. Actin flows mediate a universal coupling between cell speed and cell persistence. 161(2):374–386. ISSN 1097-4172. doi: 10.1016/j.cell.2015.01.056.
77. Lawrence Yolland, Mubarik Burki, Stefania Marcotti, Andrei Luchici, Fiona N. Kenny, John Robert Davis, Eduardo Serna-Morales, Jan Müller, Michael Sixt, Andrew Davidson, Will Wood, Linus J. Schumacher, Robert G. Endres, Mark Miodownik, and Brian M. Stramer. Persistent and polarized global actin flow is essential for directionality during cell migration. 21(11):1370–1381. ISSN 1476-4679. doi: 10.1038/s41556-019-0411-5.
78. Edwin Munro, Jeremy Nance, and James R. Priess. Cortical flows powered by asymmetrical contraction transport PAR proteins to establish and maintain anterior-posterior polarity in the early *C. elegans* embryo. 7(3):413–424. ISSN 1534-5807. doi: 10.1016/j.devcel.2004.08.001. Publisher: Elsevier.