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G α_s directly drives PDZ-RhoGEF signaling to Cdc42

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G α proteins promote dynamic adjustments of cell shape directed by actin-cytoskeleton reorganization via their respective RhoGEF effectors. For example, G α_{13} binding to the RGS-homology (RH) domains of several RH-RhoGEFs allosterically activates these proteins, causing them to expose their catalytic Dbl-homology (DH)/pleckstrin-homology (PH) regions, which triggers downstream signals. However, whether additional G α proteins might directly regulate the RH-RhoGEFs was not known. To explore this question, we first examined the morphological effects of expressing shortened RH-RhoGEF DH/PH constructs of p115RhoGEF/ARHGEF1, PDZ-RhoGEF (PRG)/ARHGEF11, and LARG/ARHGEF12. As expected, the three constructs promoted cell contraction and activated RhoA, known to be downstream of G α_{13} . Intriguingly, PRG DH/PH also induced filopodia-like cell protrusions and activated Cdc42. This pathway was stimulated by constitutively active G α_s (G α_s Q227L), which enabled endogenous PRG to gain affinity for Cdc42. A chemogenetic approach revealed that signaling by G α_s -coupled receptors, but not by those coupled to G α_i or G α_q , enabled PRG to bind Cdc42. This receptor-dependent effect, as well as CREB phosphorylation, was blocked by a construct derived from the PRG:G α_s -binding region, PRG-linker. Active G α_s interacted with isolated PRG DH and PH domains and their linker. In addition, this construct interfered with G α_s Q227L's ability to guide PRG's interaction with Cdc42. Endogenous G α_s -coupled prostaglandin receptors stimulated PRG binding to membrane fractions and activated signaling to PKA, and this canonical endogenous pathway was attenuated by PRG-linker. Altogether, our results demonstrate that active G α_s can recognize PRG as a novel effector directing its DH/PH catalytic module to gain affinity for Cdc42.

Migrating cells follow extracellular cues that guide dynamic protrusions and contractions (1, 2). At the plasma membrane, phosphoinositides and signaling proteins allosterically activate Rho guanine nucleotide exchange factors (RhoGEFs) exposing their catalytic DH/PH modules, composed of Dbl-homology and Pleckstrin-homology domains in tandem (3–5). RhoGEFs stimulate their cognate GTPases to exchange GDP for GTP, orchestrating cytoskeleton remodeling pathways (6, 7). RhoA promotes the assembly of stress fibers and contractile actomyosin structures, whereas Rac and Cdc42 lead the extension of

actin-driven plasma membrane protrusions known as lamellipodia and filopodia, respectively (8). Although these Rho GTPases exhibit contrasting effects, they can be alternatively activated at edges of moving cells (9). Therefore, fine-tuning mechanisms are likely involved.

Several RhoGEFs are effectors of heterotrimeric G proteins (3, 10, 11). GTP-bound G α_{13} and G α_q proteins stimulate RhoA, whereas G $\beta\gamma$ activates Rac and Cdc42 (3, 10, 12). G $\beta\gamma$ signaling to Rac is itself directly regulated by G α_{13} and G α_q (13). GTP-G α_{13} proteins allosterically activate RH-RhoGEFs (p115RhoGEF, PDZ-RhoGEF (PRG), and LARG) (3, 14, 15). The mechanism involves direct interaction of G α_{13} with the RhoGEF RGS-homology (RH) domain, which in consequence exposes the catalytic DH/PH cassette (16, 17). The DH domain activates RhoA, and it is positively modulated by the PH domain (6, 18). Here, we investigated whether PRG DH/PH catalytic module is directly targeted by additional G α proteins.

Results

Membrane-anchored PRG-DH/PH promotes cell contraction and formation of filopodia-like protrusions

RH-RhoGEFs are potent activators of RhoA, which generally counteracts cell protruding processes led by Rac and Cdc42 (3, 9). To characterize potential regulatory mechanisms directly targeting RH-RhoGEF DH/PH domains, we first analyzed the cellular effects of membrane-anchored, EGFP-tagged, p115RhoGEF-DH/PH, PRG-DH/PH, and LARG-DH/PH constructs (Fig. 1A). Their morphological effects, assessed by confocal microscopy of transfected endothelial cells (Fig. 1B), revealed the assembly of actin stress fibers, as expected for RhoA activity (8). Intriguingly, ~50% of cells expressing PRG-DH/PH also induced actin-based thin protrusions (Fig. 1, B and C), reminiscent of Cdc42-induced filopodia (19).

PRG-DH/PH catalytic module stimulates Cdc42

Consistent with its morphological effects, PRG-DH/PH significantly stimulated Cdc42 (Fig. 1E). In contrast, all three RH-RhoGEF EGFP-DH/PH-CAAX constructs stimulated RhoA (Fig. 1F). Furthermore, recombinant nucleotide-free Cdc42-G15A, used to isolate active Cdc42-GEFs (20), pulled down PRG-DH/PH, and to a lesser extent p115RhoGEF-DH/PH (Fig. 1G, left panel), whereas RhoA-G17A pulled down the three constructs (Fig. 1G, middle panel). Furthermore, cotransfected

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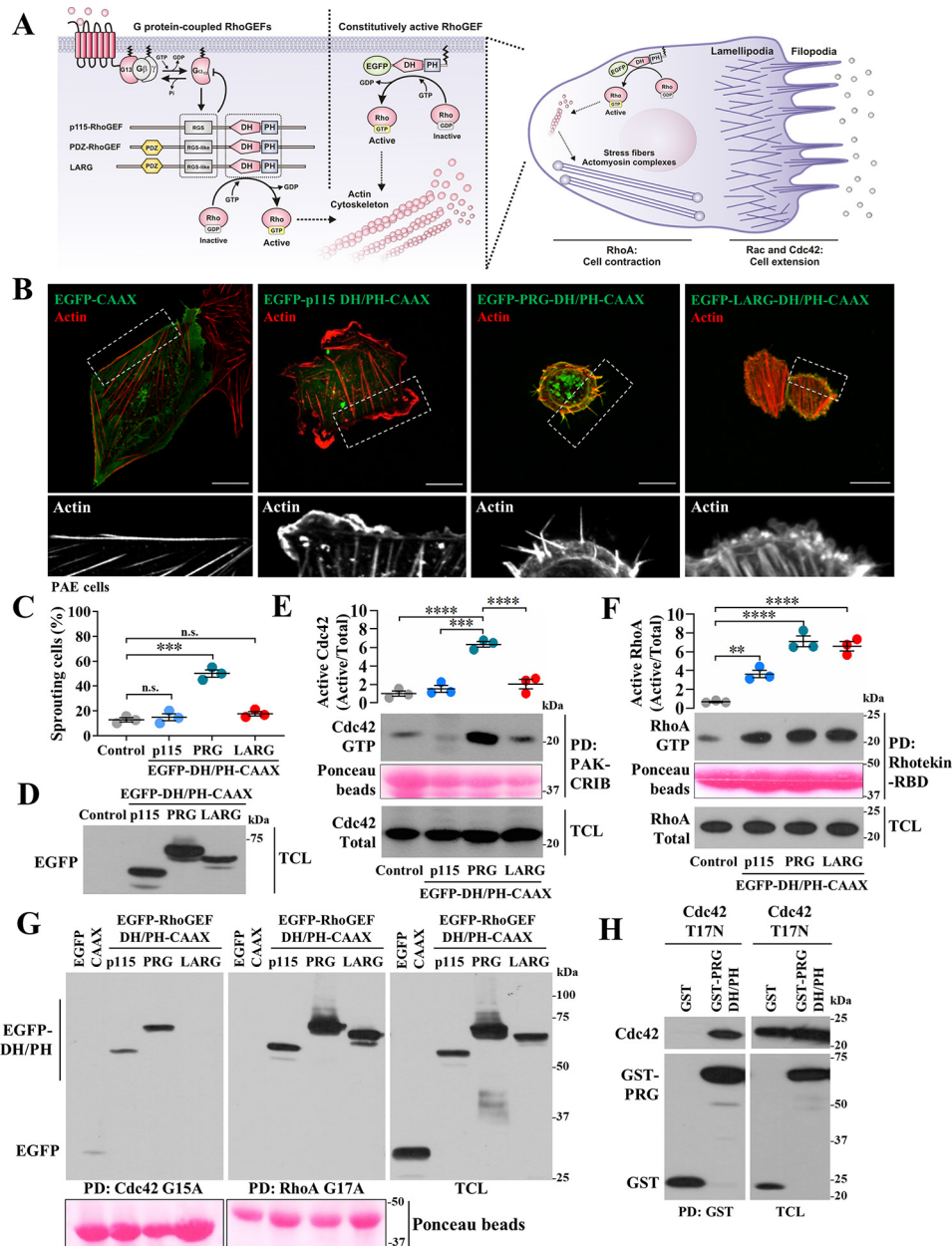


Figure 1. In addition to its canonical effect on RhoA, PRG DH/PH activates Cdc42 and promotes filopodia formation. *A*, hypothetical effects of membrane-anchored RH-RhoGEF DH/PH catalytic modules. *B*, confocal images of PAE cells showing EGFP-RH-RhoGEF DH/PH-CAAX constructs (green) and their effects on F-actin (red). Zoomed-in areas are shown in the bottom row. Scale bar, 20 μ m. *C*, graph shows the percentage of cells exhibiting filopodia-like structures. Each dot represents the means \pm S.E. of at least 30 cells per experiment ($n = 3$). ***, $p < 0.0001$; n.s., no significance, one-way ANOVA followed Tukey. *D*, representative blot shows expression of EGFP-RH-RhoGEF DH/PH-CAAX constructs transfected into HEK293T cells was detected by PAK-CRIB and Rhotekin-RBD pull-down (PD), respectively. The graphs represent the means \pm S.E. densitometric values ($n = 3$). **, $p = 0.005$; ***, $p < 0.001$; ****, $p < 0.0001$, one-way ANOVA followed Tukey. *G*, pull-down of active EGFP-RhoGEF DH/PH-CAAX constructs based on their affinity for nucleotide-free recombinant Cdc42-G15A (left panel) and RhoA-G17A (middle panel). Total cell lysates (TCL) are shown in the right panel. *H*, interaction between PRG DH/PH and Cdc42-T17N was assayed in transfected HEK293T cells subjected to pull-down assays (PD: GST). *D–H*, protein expression is confirmed in total cell lysates.

into HEK293T cells, GST-PRG-DH/PH interacted with Cdc42-T17N, a dominant-negative mutant (Fig. 1H).

Constitutively active $G\alpha_s$ -Q227L increases PRG-DH/PH activity toward Cdc42

The mechanistic basis of p63RhoGEF and TRIO activation by $G\alpha_q$ (21, 22) inspired us to assess whether GTPase-

deficient $G\alpha$ -QL mutants bind PRG DH/PH to stimulate Cdc42 (Fig. 2A). With pull-down assays, we revealed that $G\alpha_s$ -Q227L significantly increased PRG-DH/PH interaction with Cdc42-G15A (Fig. 2B) without affecting its interaction with RhoA-G17A (Fig. 2C). $G\alpha_s$ -Q227L was also detected in the pull-down assay (Fig. 2B), depending on the presence of PRG-DH/PH (Fig. 2D). In HEK293T cells, $G\alpha_s$ -QL interacted with PRG-DH/PH (Fig. 2E), as well as with other RH-

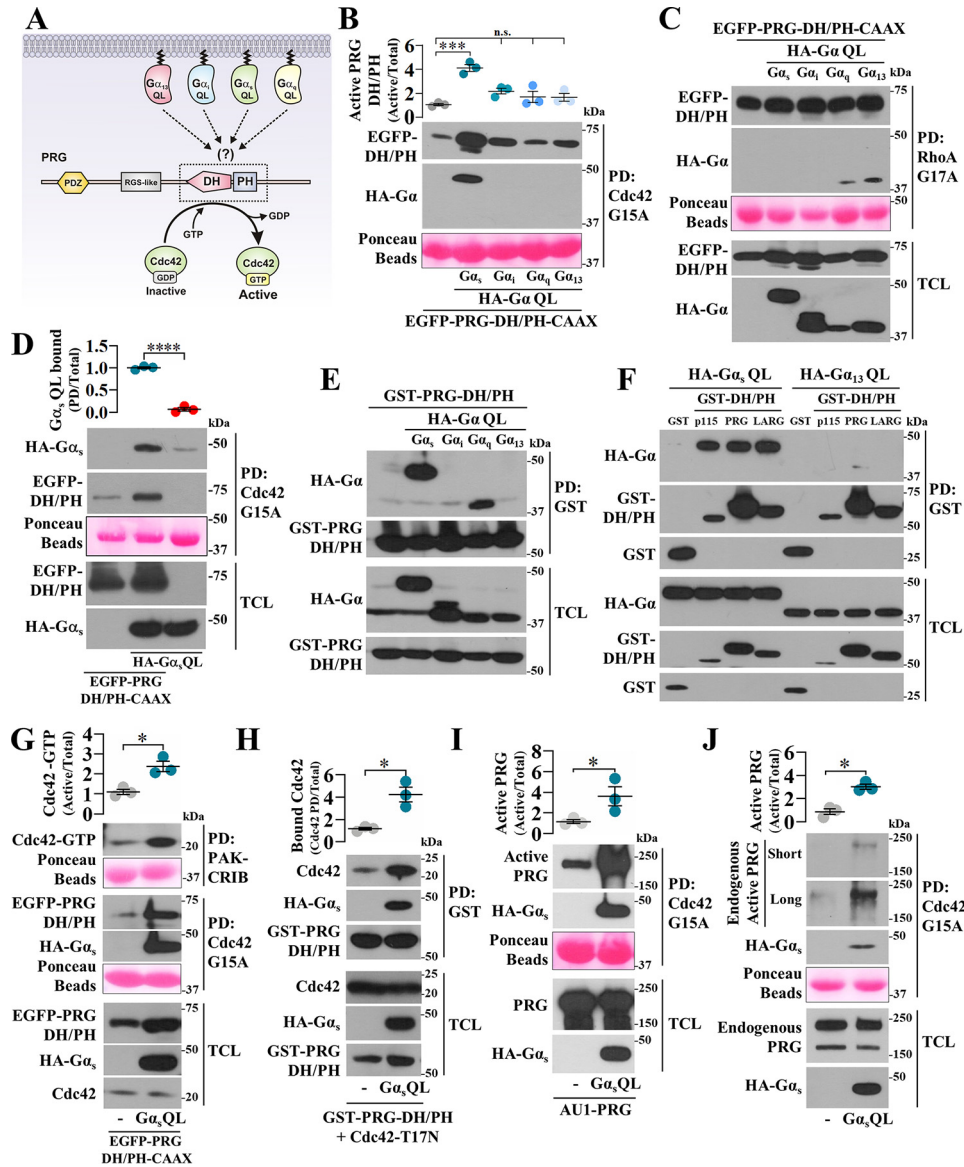


Figure 2. $G\alpha_s$ -Q227L binds PRG DH/PH enabling this prototypical RhoA-specific GEF to directly activate Cdc42. *A*, hypothetic model postulating $G\alpha$ subunits as potential regulators of PRG DH/PH catalytic module. *B* and *C*, the effect of GTPase-deficient $G\alpha$ subunits on the interaction of EGFP-PRG-DH/PH-CAAX with Cdc42-G15A (*B*) and RhoA-G17A (*C*) was analyzed by pull-down (PD) using lysates from HEK293T cells transfected with HA-tagged $G\alpha_s$, $G\alpha_i$, $G\alpha_q$, or $G\alpha_{13}$ QL mutants and EGFP-PRG-DH/PH-CAAX. The graph in *B* represents the means \pm S.E. ($n = 3$). ***, $p < 0.001$; n.s., no significance, one-way ANOVA followed Tukey. *D*, to address whether $G\alpha_s$ -QL detected in the PRG-DH/PH:Cdc42-G15A pull-down was part of a ternary complex, pull-down experiments were done in the presence or absence of PRG-DH/PH. The graph represents the means \pm S.E. ($n = 3$). ***, $p < 0.0001$, *t* test. *E*, the potential interaction between active $G\alpha$ subunits and PRG DH/PH was analyzed in HEK293T cells transfected with GST-PRG-DH/PH and HA-tagged GTPase-deficient $G\alpha$ subunits subjected to pull-down assays. *F*, interaction between $G\alpha_s$ -QL and the catalytic domain of the three RH-RhoGEFs was assayed by pull-down using HEK293T cells transfected with HA- $G\alpha_s$ -QL and GST-p115-DH/PH, GST-PRG-DH/PH, or GST-LARG-DH/PH. GST and HA- $G\alpha_{13}$ -QL served as negative controls. *G*, the effect of $G\alpha_s$ -QL on the activation of Cdc42 by PRG-DH/PH was assessed by pull-down using lysates of transfected HEK293T cells. The graph represents the means \pm S.E. ($n = 3$). **, $p = 0.01$, *t* test. Representative blots show the fraction of active Cdc42 (top panel) and the active fraction of PRG-DH/PH with affinity for Cdc42-G15A (middle panel). *H*, the effect of $G\alpha_s$ -QL on the interaction between PRG-DH/PH and Cdc42-T17N was analyzed by pull-down using lysates of transfected HEK293T cells. The graph represents the means \pm S.E. ($n = 3$). *, $p = 0.01$, *t* test. *I* and *J*, the effect of $G\alpha_s$ -QL on full-length PRG affinity for Cdc42 was analyzed in HEK293T cells that were transfected with full-length AU1-PRG (*I*) without or with HA- $G\alpha_s$ -QL or only with HA- $G\alpha_s$ -QL to address its effect on endogenous PRG (*J*). The active fraction of full-length PRG with affinity for Cdc42-G15A was isolated by pull-down and revealed by immunoblotting with anti-PRG antibodies. The graphs represents the means \pm S.E. ($n = 3$). *, $p = 0.01$ in *H* and 0.04 in *I*, *t* test.

RhoGEFs (Fig. 2*F*), as revealed by pull-down assays using GST-DH/PH constructs. In contrast, $G\alpha_{13}$ -QL, known to interact with RGS-like domain of RH-RhoGEFs (14, 23), was absent in the GST-DH/PH pull-downs (Fig. 2*F*). We also found that $G\alpha_q$ -QL interacted with PRG-DH/PH (Fig. 2*E*); however, it did not affect PRG-DH/PH:Cdc42 binding (Fig. 2*B*).

Consistent with a functional effect, $G\alpha_s$ -QL stimulated PRG-DH/PH to activate Cdc42 (Fig. 2*G*, top panel and graph) and remained bound to the PRG-DH/PH construct pulled down with nucleotide-free Cdc42 (Fig. 2, *B*, *D*, and *G*, middle panel). Furthermore, the interaction of PRG DH/PH with Cdc42-T17N was further stimulated by $G\alpha_s$ -QL (Fig. 2*H*).

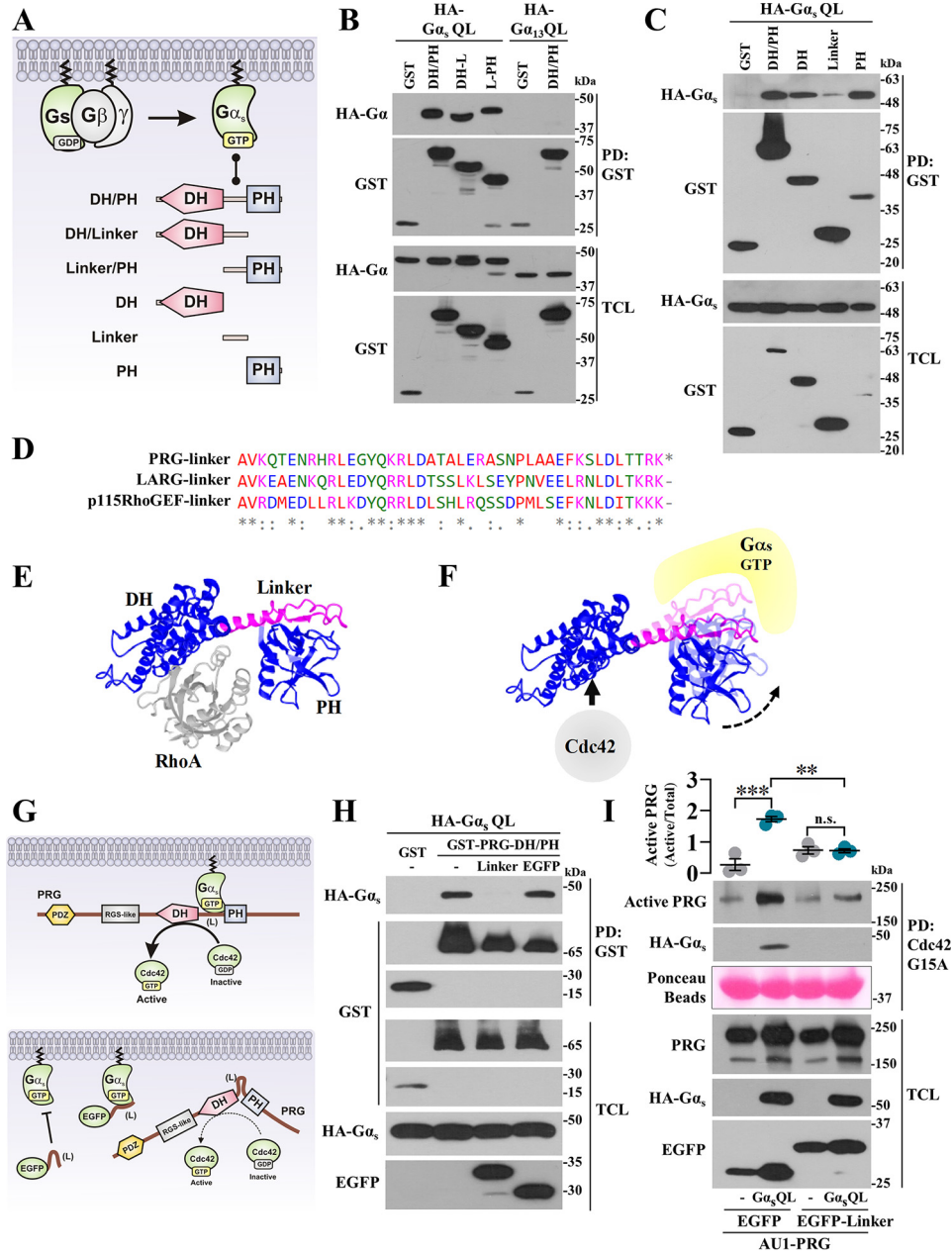


Figure 3. $G\alpha_s$ -Q227L binds PRG DH and PH domains and the linker region joining them. *A*, model showing GST-tagged PRG-DH/PH constructs used to map $G\alpha_s$ -PRG-DH/PH interaction. *B* and *C*, interaction between HA- $G\alpha_s$ -QL and the indicated GST-PRG-DH/PH constructs was analyzed by pull-down using lysates of transfected HEK293T cells. HA- $G\alpha_s$ -QL and HA- $G\alpha_{13}$ -QL (used as control) were revealed with anti-HA antibodies. *D*, multiple alignment of p115RhoGEF, LARG, and PRG-linker regions. *E*, structure of PRG-DH/PH-RhoA complex (24). *F*, model showing $G\alpha_s$ -GTP-PRG-DH/PH complex; hypothetically, active $G\alpha_s$ constrains PRGDH/PH to bind *Cdc42*. *G*, model showing the potential inhibitory effect of the EGFP-PRG-linker construct on PRG activation by $G\alpha_s$ -QL. *H*, HEK293T cells transfected with EGFP-tagged PRG-linker construct (or EGFP) together with HA- $G\alpha_s$ -QL and GST-PRG-DH/PH (or GST) were subjected to GST pull-down assays. *I*, $G\alpha_s$ -dependent PRG-*Cdc42* interaction was analyzed in HEK293T cells transfected with HA- $G\alpha_s$ -QL (or control plasmid) and AU1-PRG together with EGFP-PRG-linker or EGFP and subjected to *Cdc42*-G15A pull-down. The graph represents the means \pm S.E. ($n = 3$). **, $p = 0.001$; ***, $p = 0.0001$; *ns*, no significance, one-way ANOVA followed Tukey.

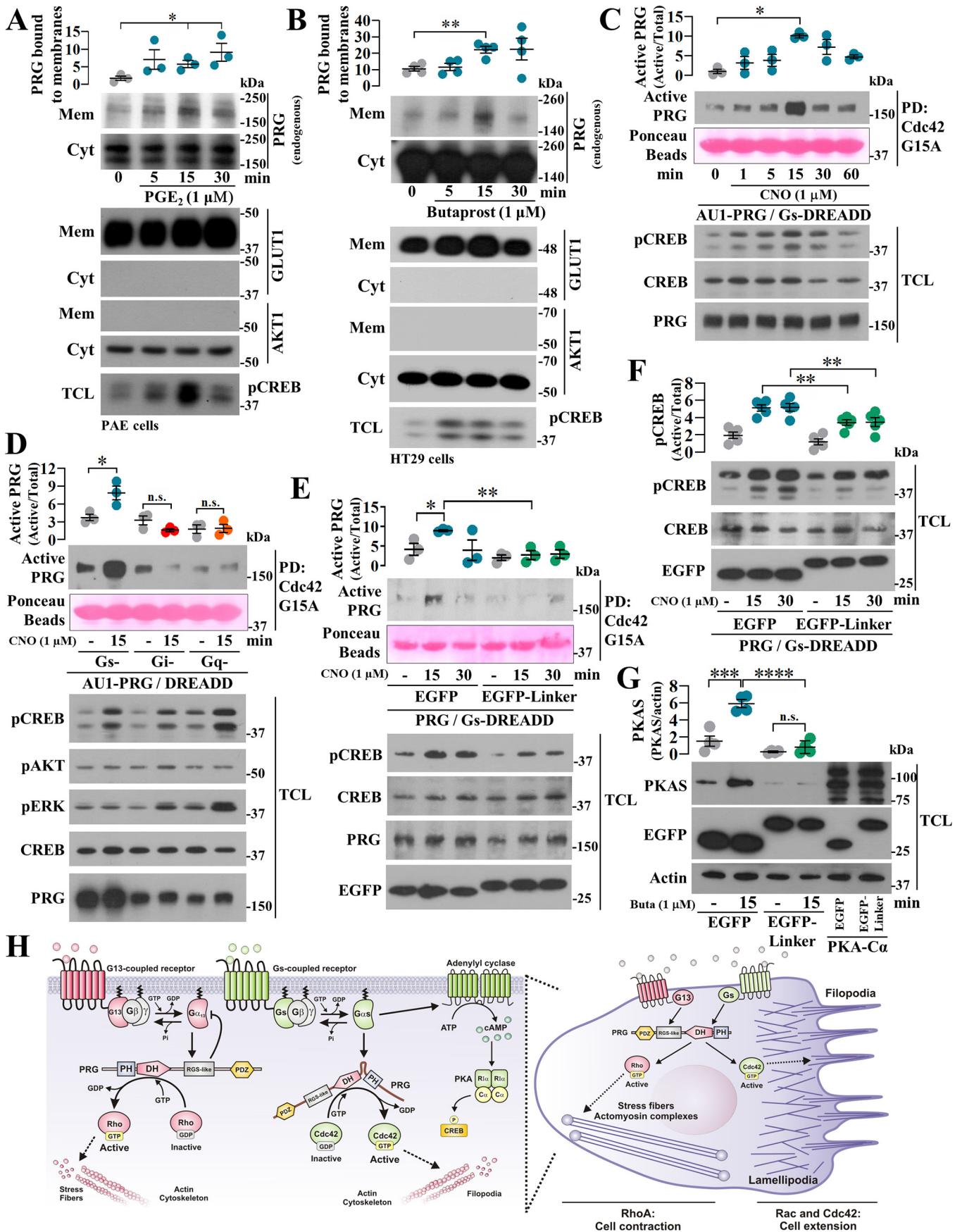
$G\alpha_s$ -Q227L drives full-length PRG to interact with *Cdc42*

To address whether full-length PRG is sensitive to be driven by $G\alpha_s$ -QL to gain affinity for *Cdc42*, we used lysates from transfected HEK293T cells. We found that $G\alpha_s$ -QL stimulated full-length PRG, either transfected (Fig. 2*I*) or endogenous (Fig. 2*J*), to bind *Cdc42*. Furthermore, $G\alpha_s$ -QL remained bound to PRG pulled down with nucleotide-free *Cdc42*. Expression of transfected and endogenous proteins was confirmed in total cell lysates (Fig. 2, *C–J*, *TCL*).

$G\alpha_s$ -Q227L interaction interface at PRG involves the DH and PH domains and the linker region joining them

To characterize how $G\alpha_s$ -Q227L binds PRG-DH/PH guiding this prototypic RhoA-specific GEF to interact with *Cdc42*, we cotransfected $G\alpha_s$ -Q227L with different PRG constructs spanning the DH/PH module, fused to GST (Fig. 3*A*), and addressed by pull-down their potential interaction. As shown in Fig. 3*B*, $G\alpha_s$ -QL interacted with the three PRG-DH/PH constructs that had in common the linker region that joins the DH and PH

ACCELERATED COMMUNICATION: $G\alpha_s$ activates *Cdc42* via PDZ-RhoGEF



domains. When PRG-DH, PRG-linker, and PRG-PH were used as independent constructs, the three of them interacted with $G\alpha_s$ -QL, indicating that PRG-linker is critical to strengthen the interaction (Fig. 3C). Consistent with previous results, $G\alpha_{13}$ -QL served as negative control (Fig. 3B). Because the PRG-linker sequence is conserved among the three RH-RhoGEFs (Fig. 3D) but lacks homology with any other protein, we used an EGFP-tagged PRG-linker construct as a potential inhibitor of $G\alpha_s$ -dependent PRG·Cdc42 interaction. Based on previous results and the structure of the PRG-DH/PH·RhoA complex (Fig. 3E) (24), we postulated that $G\alpha_s$ -Q227L interacts with PRG-DH/PH, forming a complex with affinity for Cdc42 (Fig. 3F). Then we tested the potential inhibitory effect of PRG-linker on the interaction between $G\alpha_s$ -Q227L and PRG (Fig. 3G). As predicted, the PRG-linker construct not only inhibited $G\alpha_s$ -QL·PRG-DH/PH interaction (Fig. 3H) but also interfered on the effect of $G\alpha_s$ -QL to drive full-length PRG to be pulled down as an active Cdc42-GEF (Fig. 3I); EGFP served as negative control.

Agonist-dependent stimulation of G_s -coupled receptors drives PRG to gain affinity for Cdc42

To investigate whether G_s -coupled receptors stimulate PRG to acquire affinity for Cdc42, we first used PAE and HT29 cells as models of endogenous prostaglandin-dependent G_s signaling. As an initial readout of agonist-driven G_s -dependent effect on PRG, we stimulated PAE and HT29 cells with PGE₂ and butaprost, respectively, and assessed PRG recruitment to membrane fractions. In both cases, PRG exhibited a significant time-dependent association to membrane fractions (Fig. 4, A and B, respectively). We then used COS7 cells expressing Gs-DREADDs to test the effect of endogenous G_s on PRG·Cdc42 interaction. In these cells, clozapine *N*-oxide (CNO), the agonist of G_s -DREADDs, enabled PRG to bind nucleotide-free Cdc42 in a time-dependent manner (Fig. 4C). This effect was elicited by G_s -coupled, but not by G_i - or G_q -coupled DREADDs (Fig. 4D), and was inhibited by the PRG-linker peptide (Fig. 4E), which also interfered on CREB phosphorylation (Fig. 4F). Endogenous G_s -coupled endothelial EP2 receptors signaling to cAMP/PKA pathway was also inhibited by the PRG-linker construct, as indicated by a decrease on butaprost-dependent phosphorylation of PKA substrates (Fig. 4G).

Discussion

RH-RhoGEFs link heterotrimeric G proteins to Rho GTPases (25–27). They are activated by $G\alpha_{12/13}$ proteins, which bind the RH domains unleashing the catalytic DH/PH region, known as specific for RhoA (14, 23). Here we demonstrate that active $G\alpha_s$ directly constrains the PRG DH/PH catalytic module to activate Cdc42, whereas its effect on RhoA is unaltered. Although future work using purified proteins is guaranteed, our results suggest that RhoGEF DH/PH domains can be allosterically controlled to expand their specificity.

Direct activation of RhoGEF DH/PH domains by active $G\alpha$ subunits of heterotrimeric G proteins has been described. Specifically, $G\alpha_q$ stimulates p63RhoGEF and TRIO (21, 28). Physiological control of this system is lost by *GNAQ* mutation, causing the $G\alpha_q$ ·TRIO signaling system to drive uveal melanoma progression (28, 29). Similarly, mutant *GNAS* is a driving oncogene in neuroendocrine cancers (30); however, a pathological link to Rho GTPases has not been established. Our results are reminiscent of the regulation of p63RhoGEF and TRIO by $G\alpha_q$ (21, 22, 28) but differ in the fact that $G\alpha_s$ expands PRG specificity directing the DH/PH module to gain affinity for Cdc42 without an apparent effect on RhoA. We speculate that $G\alpha_s$ pulls the PRG DH/PH module to accommodate Cdc42. Consistent with this possibility, conserved residues that directly bind the GTPase are more distant in intersectin-1, a Cdc42-specific GEF compared with the PRG DH/PH module in complex with RhoA (31–33).

Consistent with their reported effects (25, 27, 32, 34–36), RH-RhoGEF DH/PH catalytic modules strongly activated RhoA and promoted the assembly of actin stress fibers and cell contraction. These results confirmed that DH/PH constructs maintain catalysis and specificity (36–44). However, we found that PRG DH/PH also exhibited a previously unrecognized ability to stimulate Cdc42 and filopodia formation. Thus, we addressed the possibility that PRG directly activates Cdc42. We used pulldown assays to isolate active RhoGEFs based on their affinity for nucleotide-free GTPases (20, 45) and revealed that $G\alpha_s$ stimulates PRG to gain affinity for Cdc42, pointing to a direct effect (attenuated by PKA; Fig. S1). We demonstrated that GTPase-deficient $G\alpha_s$ binds the DH/PH module. The linker region joining these domains strengthen their interaction with active $G\alpha_s$. Our evidence arguing for a functional relevance of this interaction derives from the inhibitory effect of the PRG-linker construct, which prevented PRG response to

Figure 4. Agonist-dependent stimulation of G_s -coupled receptors enables PRG to bind Cdc42. A and B, membrane recruitment of endogenous PRG promoted by G_s -coupled GPCR signaling was assessed in PAE (A) and HT29 (B) cells stimulated with 1 μ M PGE₂ or butaprost, respectively. PRG in membrane fractions was revealed by Western blotting. GLUT1 and AKT1 were used as membrane and cytosolic markers, respectively. The graphs represent the means \pm S.E. ($n = 3$, * $p < 0.05$ in A; and $n = 4$, ** $p < 0.01$ in B; *t* test). C, time course of PRG·Cdc42 interaction was assessed in COS7 expressing G_s -DREADD receptors. The cells were stimulated with 1 μ M CNO and subjected to Cdc42-G15A pulldown. The graph represents the means \pm S.E. ($n = 3$). * $p < 0.05$, one-way ANOVA followed Tukey. D, the effect of different endogenous heterotrimeric G proteins on PRG affinity for Cdc42 was studied in COS7 cells transfected with AU1-PRG and G_s , G_i , or G_q -DREADDs. The cells were stimulated with CNO for 15 min and subjected to Cdc42-G15A pulldown assays. The graph represents the means \pm S.E. ($n = 3$). * $p < 0.05$, one-way ANOVA followed Tukey. E, effect of the PRG-linker construct on agonist-stimulated interaction between PRG and Cdc42 was assessed in COS7 cells transfected with Gs-DREADD, AU1-PRG, and EGFP-PRG-linker or EGFP. The cells were stimulated with CNO for 15 and 30 min and subjected to Cdc42-G15A pulldown. The graph represents the means \pm S.E. ($n = 3$). * $p = 0.0342$; ** $p = 0.0056$, *t* test. F, effect of PRG-linker on agonist-dependent phosphorylation of CREB was assessed in COS7 cells expressing G_s -DREADDs and stimulated with CNO. The graph represents the means \pm S.E. ($n = 5$). ** $p < 0.01$, *t* test. G, agonist-dependent phosphorylation of PKA substrates was assessed using PAE cells expressing EGFP or EGFP-PRG-linker and stimulated with butaprost. Lysates from EGFP-PKA- α -transfected cells served as control to detect PKA substrates. The graph represents the means \pm S.E. ($n = 4$). *** $p = 0.0009$; **** $p < 0.0001$; *n.s.*, no significance, one-way ANOVA followed Tukey. H, model depicts the canonical G_{13} -PRG signaling axis to Rho and the emerging GPCR- $G\alpha_s$ -PRG-Cdc42 pathway based on the current findings. In cells, both systems putatively guide dynamic adjustments on actin-cytoskeleton reorganization.

agonist-dependent stimulation of Gs-DREADDs and to GTPase-deficient $G\alpha_s$ coexpression. Our results not only indicate that $G\alpha_s$ guides PRG to bind Cdc42 but also suggest that this effector competes with other $G\alpha_s$ -dependent effectors. The Gs/PKA pathway activates Rho GTPases and regulates cytoskeletal dynamics at multiple levels. Recent evidence documented a role for PKA R1 α subunit as a cAMP-dependent activator of P-REX1, a RacGEF (46), whereas kinase activity of PKA is linked to cytoskeletal dynamics at cell edges and is reciprocally regulated during cell migration (47–49). Our findings showing that $G\alpha_s$ activates a PRG/Cdc42 pathway expand the mechanisms of Gs signaling to Rho GTPases.

Although further experiments are needed to define the spatiotemporal conditions in which PRG is guided to activate Cdc42, our current model (Fig. 4H) illustrates the potential of Gs-coupled receptors to activate this pathway. Our work raises new questions and research avenues on how Gs and G13 signaling pathways are integrated to fine-tune Cdc42 activity in the context of strong RhoA activation to regulate cytoskeletal dynamics and set the basis to further investigate how the Gs/PRG/Cdc42 pathway guides polarized cell migration and its potential pathological implications, particularly in cancers in which mutant *GNAS* is a driving oncogene.

Experimental procedures

Plasmids and cDNA constructs

RH-RhoGEF DH/PH catalytic modules and PRG DH-PH fragments were amplified by PCR and cloned into pCEFL-EGFP-CAAX, pCEFL-EGFP, and pCEFL-GST. Primer sequences are available upon request. Other constructs have been previously described (13, 46).

Cell culture, transfection, immunoblotting, and GST pulldown

HEK293T, PAE, HT29, and COS7 cells were maintained and transfected as described (46). The cells were serum-starved for 16 h before experiments and were all done 48 h after transfection. GST fusion proteins and their interactors were detected by pulldown (13, 46). The cell lysates and pulldowns were analyzed by Western blotting using the following antibodies: EGFP SC-9996, GST B-14 SC-138, Cdc42 SC-8401, RhoA SC-418, p-AKT1/2/3 Ser473 SC-7985-R, and ERK2 SC-154 from Santa Cruz Biotechnology; HA from Covance; p-CREB Ser-133 9191, p-ERK1/2 T202/Y204 9191, CREB 9197S, and PKA 9624 from Cell Signaling Technology; PRG/ARHGEF11 and HPA014658 from Atlas Antibodies; and AKT1 and P2482 from Sigma. The secondary antibodies were goat anti-mouse (Zymed Laboratories Inc., Invitrogen, or KPL) or goat anti-rabbit (Rockland Immunochemicals or KPL).

GTPase (Cdc42 and RhoA) activation and active GEF capture assays

Activation of RhoA and Cdc42, using cells grown in 10-cm dishes, was assessed by pulldown using recombinant Rhotekin and PAK effector domains (13, 46). Active RhoGEFs were detected by pulldown with nucleotide-free GTPases (RhoA G17A and Cdc42 G15A) fused to GST (20, 45). Before lysis, the

cells were washed with PBS containing 10 mM MgCl₂ and lysed with 1 ml of ice-cold lysis buffer (50 mM Tris, 150 mM NaCl, pH 7.5, containing 1% Triton X-100, 5 mM EDTA, and protease and phosphatase inhibitors (13)). Lysates were subjected to GST pulldowns as described (13, 46). COS7 cells were transfected with DREADDs exclusively coupled to Gs, Gi, or Gq and stimulated with CNO (Tocris) (13).

Membrane and cytoplasmic fractionation of PAE and HT29 cells

Serum-starved PAE and HT29 cells, grown in 10-cm Petri dishes, were stimulated with 1 μ M prostaglandin E2 or butaprost, as indicated in Fig. 4. The cells were washed with cold PBS, scraped into 1 ml of cold PBS containing protease and phosphatase inhibitors, and subjected to three freeze/thaw cycles. The lysates were centrifuged at low speed (1,400 rpm for 10 min at 4 °C). Supernatants were centrifuged at 13,000 rpm for 10 min at 4 °C. Cytosol-enriched supernatants were prepared with Laemmli buffer. The pellets were washed once with cold PBS, centrifuged again, incubated with 250 μ l of lysis buffer containing 1% Triton X-100 for 20 min, and centrifuged at 13,000 rpm for 10 min at 4 °C. Supernatants containing solubilized membranes were prepared with Laemmli buffer. PRG was analyzed by Western blotting, together with GLUT1 and AKT1, as membrane and cytosol markers, respectively.

Cytoskeletal effects of RH-RhoGEF DH/PH constructs

PAE cells were seeded at low density on gelatin-coated coverslips. Transfected cells were starved for 16 h with serum-free medium. Subsequently, the cells were fixed in 4% paraformaldehyde in PBS for 20 min, washed twice with PBS, and prepared for conventional phalloidin staining. The cell images were visualized in a Leica confocal laser scanning microscope TCS SP8 using a 63 \times 1.4 oil immersion objective. The images were analyzed with FIJI-ImageJ software. The cells were counted as having filopodia-like structures when they had at least nine of these finger-like protrusions containing F-actin (19).

Statistical analysis

The data are presented as means \pm S.E. of at least three independent experiments. Densitometric quantitation of Western blots was done with ImageJ. Active proteins and interactions in pulldowns were normalized respect to total proteins and pull-down efficiency. Statistical analysis was performed using Sigma Plot 11.0, and graphs were prepared with Prism software V8.0. Statistical tests are indicated at the figure legends.

Data availability

All the described data are contained within this article.

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Abbreviations—The abbreviations used are: RH, RGS homology; DH, Dbl homology; PH, pleckstrin homology; GEF, guanine nucleotide exchange factor; PRG, PDZ-RhoGEF; EGFP, enhanced GFP; CNO, clozapine *N*-oxide; GST, glutathione *S*-transferase; HA, hemagglutinin; ANOVA, analysis of variance; GPCR, G protein-coupled receptor.

References

- Ridley, A. J., Schwartz, M. A., Burridge, K., Firtel, R. A., Ginsberg, M. H., Borisy, G., Parsons, J. T., and Horwitz, A. R. (2003) Cell migration: integrating signals from front to back. *Science* **302**, 1704–1709 [CrossRef Medline](#)
- Pollard, T. D., and Borisy, G. G. (2003) Cellular motility driven by assembly and disassembly of actin filaments. *Cell* **112**, 453–465 [CrossRef Medline](#)
- Aittaleb, M., Boguth, C. A., and Tesmer, J. J. (2010) Structure and function of heterotrimeric G protein-regulated Rho guanine nucleotide exchange factors. *Mol. Pharmacol.* **77**, 111–125 [CrossRef Medline](#)
- Cook, D. R., Rossman, K. L., and Der, C. J. (2014) Rho guanine nucleotide exchange factors: regulators of Rho GTPase activity in development and disease. *Oncogene* **33**, 4021–4035 [CrossRef Medline](#)
- Lawson, C. D., and Ridley, A. J. (2018) Rho GTPase signaling complexes in cell migration and invasion. *J. Cell Biol.* **217**, 447–457 [CrossRef Medline](#)
- Rossman, K. L., Der, C. J., and Sondek, J. (2005) GEF means go: turning on RHO GTPases with guanine nucleotide-exchange factors. *Nat. Rev. Mol. Cell Biol.* **6**, 167–180 [CrossRef Medline](#)
- Etienne-Manneville, S., and Hall, A. (2002) Rho GTPases in cell biology. *Nature* **420**, 629–635 [CrossRef Medline](#)
- Ridley, A. J. (2015) Rho GTPase signalling in cell migration. *Curr. Opin. Cell Biol.* **36**, 103–112 [CrossRef Medline](#)
- Machacek, M., Hodgson, L., Welch, C., Elliott, H., Pertz, O., Nalbant, P., Abell, A., Johnson, G. L., Hahn, K. M., and Danuser, G. (2009) Coordination of Rho GTPase activities during cell protrusion. *Nature* **461**, 99–103 [CrossRef Medline](#)
- Vázquez-Prado, J., Bracho-Valdes, I., Cervantes-Villagrana, R. D., and Reyes-Cruz, G. (2016) $G\beta\gamma$ pathways in cell polarity and migration linked to oncogenic GPCR signaling: potential relevance in tumor microenvironment. *Mol. Pharmacol.* **90**, 573–586 [CrossRef Medline](#)
- Hernández-Vásquez, M. N., Adame-García, S. R., Hamoud, N., Chidiac, R., Reyes-Cruz, G., Gratton, J. P., Côté, J. F., and Vázquez-Prado, J. (2017) Cell adhesion controlled by adhesion G protein-coupled receptor GPR124/ADGRA2 is mediated by a protein complex comprising intersectins and Elmo-Dock. *J. Biol. Chem.* **292**, 12178–12191 [CrossRef Medline](#)
- Welch, H. C. (2015) Regulation and function of P-Rex family Rac-GEFs. *Small GTPases* **6**, 49–70 [CrossRef Medline](#)
- Cervantes-Villagrana, R. D., Adame-García, S. R., García-Jiménez, I., Color-Aparicio, V. M., Beltrán-Navarro, Y. M., König, G. M., Kostenis, E., Reyes-Cruz, G., Gutkind, J. S., and Vázquez-Prado, J. (2019) $G\beta\gamma$ signaling to the chemotactic effector P-REX1 and mammalian cell migration is directly regulated by $G\alpha_q$ and $G\alpha_{13}$ proteins. *J. Biol. Chem.* **294**, 531–546 [CrossRef Medline](#)
- Kozasa, T., Hajicek, N., Chow, C. R., and Suzuki, N. (2011) Signalling mechanisms of RhoGTPase regulation by the heterotrimeric G proteins G12 and G13. *J. Biochem.* **150**, 357–369 [CrossRef Medline](#)
- Vázquez-Prado, J., Miyazaki, H., Castellone, M. D., Teramoto, H., and Gutkind, J. S. (2004) Chimeric $G\alpha_{12}/G\alpha_{13}$ proteins reveal the structural requirements for the binding and activation of the RGS-like (RGL)-containing Rho guanine nucleotide exchange factors (GEFs) by $G\alpha_{13}$. *J. Biol. Chem.* **279**, 54283–54290 [CrossRef Medline](#)
- Bodmann, E. L., Krett, A. L., and Bünemann, M. (2017) Potentiation of receptor responses induced by prolonged binding of $G\alpha_{13}$ and leukemia-associated RhoGEF. *FASEB J.* **31**, 3663–3676 [CrossRef Medline](#)
- Meyer, B. H., Freuler, F., Guerini, D., and Siehler, S. (2008) Reversible translocation of p115-RhoGEF by $G_{12/13}$ -coupled receptors. *J. Cell. Biochem.* **104**, 1660–1670 [CrossRef Medline](#)
- Chen, Z., Medina, F., Liu, M. Y., Thomas, C., Sprang, S. R., and Sternweis, P. C. (2010) Activated RhoA binds to the pleckstrin homology (PH) domain of PDZ-RhoGEF, a potential site for autoregulation. *J. Biol. Chem.* **285**, 21070–21081 [CrossRef Medline](#)
- Nobes, C. D., and Hall, A. (1995) Rho, rac, and cdc42 GTPases regulate the assembly of multimolecular focal complexes associated with actin stress fibers, lamellipodia, and filopodia. *Cell* **81**, 53–62 [CrossRef Medline](#)
- García-Mata, R., Wennerberg, K., Arthur, W. T., Noren, N. K., Ellerbroek, S. M., and Burridge, K. (2006) Analysis of activated GAPs and GEFs in cell lysates. *Methods Enzymol.* **406**, 425–437 [CrossRef Medline](#)
- Lutz, S., Shankaranarayanan, A., Coco, C., Ridilla, M., Nance, M. R., Vettel, C., Baltus, D., Evelyn, C. R., Neubig, R. R., Wieland, T., and Tesmer, J. J. (2007) Structure of $G\alpha_q$ -p63RhoGEF-RhoA complex reveals a pathway for the activation of RhoA by GPCRs. *Science* **318**, 1923–1927 [CrossRef Medline](#)
- Rojas, R. J., Yohe, M. E., Gershburg, S., Kawano, T., Kozasa, T., and Sondek, J. (2007) $G\alpha_q$ directly activates p63RhoGEF and Trio via a conserved extension of the Dbl homology-associated pleckstrin homology domain. *J. Biol. Chem.* **282**, 29201–29210 [CrossRef Medline](#)
- Fukuhara, S., Chikumi, H., and Gutkind, J. S. (2001) RGS-containing RhoGEFs: the missing link between transforming G proteins and Rho? *Oncogene* **20**, 1661–1668 [CrossRef Medline](#)
- Bielnicki, J. A., Shkumatov, A. V., Derewenda, U., Somlyo, A. V., Svergun, D. I., and Derewenda, Z. S. (2011) Insights into the molecular activation mechanism of the RhoA-specific guanine nucleotide exchange factor, PDZ-RhoGEF. *J. Biol. Chem.* **286**, 35163–35175 [CrossRef Medline](#)
- Hart, M. J., Jiang, X., Kozasa, T., Roscoe, W., Singer, W. D., Gilman, A. G., Sternweis, P. C., and Bollag, G. (1998) Direct stimulation of the guanine nucleotide exchange activity of p115 RhoGEF by $G\alpha_{13}$. *Science* **280**, 2112–2114 [CrossRef Medline](#)
- Fukuhara, S., Murga, C., Zohar, M., Igishi, T., and Gutkind, J. S. (1999) A novel PDZ domain containing guanine nucleotide exchange factor links heterotrimeric G proteins to Rho. *J. Biol. Chem.* **274**, 5868–5879 [CrossRef Medline](#)
- Fukuhara, S., Chikumi, H., and Gutkind, J. S. (2000) Leukemia-associated Rho guanine nucleotide exchange factor (LARG) links heterotrimeric G proteins of the G_{12} family to Rho. *FEBS Lett.* **485**, 183–188 [CrossRef Medline](#)
- Bandekar, S. J., Arang, N., Tully, E. S., Tang, B. A., Barton, B. L., Li, S., Gutkind, J. S., and Tesmer, J. J. G. (2019) Structure of the C-terminal guanine nucleotide exchange factor module of Trio in an autoinhibited conformation reveals its oncogenic potential. *Sci. Signal.* **12**, eaav2449 [CrossRef Medline](#)
- Feng, X., Degese, M. S., Iglesias-Bartolome, R., Vaque, J. P., Molinolo, A. A., Rodrigues, M., Zaidi, M. R., Ksander, B. R., Merlino, G., Sodhi, A., Chen, Q., and Gutkind, J. S. (2014) Hippo-independent activation of YAP by the GNAQ uveal melanoma oncogene through a trio-regulated rho GTPase signaling circuitry. *Cancer Cell* **25**, 831–845 [CrossRef Medline](#)
- O'Hayre, M., Vázquez-Prado, J., Kufareva, I., Stawiski, E. W., Handel, T. M., Seshagiri, S., and Gutkind, J. S. (2013) The emerging mutational

- landscape of G proteins and G-protein-coupled receptors in cancer. *Nat. Rev. Cancer* **13**, 412–424 [CrossRef Medline](#)
31. Kapp, G. T., Liu, S., Stein, A., Wong, D. T., Remenyi, A., Yeh, B. J., Fraser, J. S., Taunton, J., Lim, W. A., and Kortemme, T. (2012) Control of protein signaling using a computationally designed GTPase/GEF orthogonal pair. *Proc. Natl. Acad. Sci. U.S.A.* **109**, 5277–5282 [CrossRef Medline](#)
 32. Derewenda, U., Oleksy, A., Stevenson, A. S., Korczynska, J., Dauter, Z., Somlyo, A. P., Otlewski, J., Somlyo, A. V., and Derewenda, Z. S. (2004) The crystal structure of RhoA in complex with the DH/PH fragment of PDZRhoGEF, an activator of the Ca(2+) sensitization pathway in smooth muscle. *Structure* **12**, 1955–1965 [CrossRef Medline](#)
 33. Hernández-García, R., Iruela-Arispe, M. L., Reyes-Cruz, G., and Vázquez-Prado, J. (2015) Endothelial RhoGEFs: a systematic analysis of their expression profiles in VEGF-stimulated and tumor endothelial cells. *Vascul. Pharmacol.* **74**, 60–72 [CrossRef Medline](#)
 34. Rügenapp, U., Blomquist, A., Schwörer, G., Schabrowski, H., Psoma, A., and Jakobs, K. H. (1999) Rho-specific binding and guanine nucleotide exchange catalysis by KIAA0380, a dbl family member. *FEBS Lett.* **459**, 313–318 [CrossRef Medline](#)
 35. Dubash, A. D., Wennerberg, K., García-Mata, R., Menold, M. M., Arthur, W. T., and Burridge, K. (2007) A novel role for Lsc/p115 RhoGEF and LARG in regulating RhoA activity downstream of adhesion to fibronectin. *J. Cell Sci.* **120**, 3989–3998 [CrossRef Medline](#)
 36. Jaiswal, M., Gremer, L., Dvorsky, R., Haeusler, L. C., Cirstea, I. C., Uhlenbrock, K., and Ahmadian, M. R. (2011) Mechanistic insights into specificity, activity, and regulatory elements of the regulator of G-protein signaling (RGS)-containing Rho-specific guanine nucleotide exchange factors (GEFs) p115, PDZ-RhoGEF (PRG), and leukemia-associated RhoGEF (LARG). *J. Biol. Chem.* **286**, 18202–18212 [CrossRef Medline](#)
 37. Solski, P. A., Wilder, R. S., Rossman, K. L., Sondek, J., Cox, A. D., Campbell, S. L., and Der, C. J. (2004) Requirement for C-terminal sequences in regulation of Ect2 guanine nucleotide exchange specificity and transformation. *J. Biol. Chem.* **279**, 25226–25233 [CrossRef Medline](#)
 38. Worthylake, D. K., Rossman, K. L., and Sondek, J. (2000) Crystal structure of Rac1 in complex with the guanine nucleotide exchange region of Tiam1. *Nature* **408**, 682–688 [CrossRef Medline](#)
 39. Reuther, G. W., Lambert, Q. T., Booden, M. A., Wennerberg, K., Becknell, B., Marcucci, G., Sondek, J., Caligiuri, M. A., and Der, C. J. (2001) Leukemia-associated Rho guanine nucleotide exchange factor, a Dbl family protein found mutated in leukemia, causes transformation by activation of RhoA. *J. Biol. Chem.* **276**, 27145–27151 [CrossRef Medline](#)
 40. Rossman, K. L., Worthylake, D. K., Snyder, J. T., Siderovski, D. P., Campbell, S. L., and Sondek, J. (2002) A crystallographic view of interactions between Dbs and Cdc42: PH domain-assisted guanine nucleotide exchange. *EMBO J.* **21**, 1315–1326 [CrossRef Medline](#)
 41. Snyder, J. T., Worthylake, D. K., Rossman, K. L., Betts, L., Pruitt, W. M., Siderovski, D. P., Der, C. J., and Sondek, J. (2002) Structural basis for the selective activation of Rho GTPases by Dbl exchange factors. *Nat. Struct. Biol.* **9**, 468–475 [CrossRef Medline](#)
 42. Kristelly, R., Gao, G., and Tesmer, J. J. (2004) Structural determinants of RhoA binding and nucleotide exchange in leukemia-associated Rho guanine-nucleotide exchange factor. *J. Biol. Chem.* **279**, 47352–47362 [CrossRef Medline](#)
 43. Cash, J. N., Davis, E. M., and Tesmer, J. J. (2016) Structural and biochemical characterization of the catalytic core of the metastatic factor P-Rex1 and its regulation by PtdIns(3,4,5)P₃. *Structure* **24**, 730–740 [CrossRef Medline](#)
 44. Lucato, C. M., Halls, M. L., Ooms, L. M., Liu, H. J., Mitchell, C. A., Whistock, J. C., and Ellisdon, A. M. (2015) The phosphatidylinositol (3,4,5)-trisphosphate-dependent Rac exchanger 1-Ras-related C3 botulinum toxin substrate 1 (P-Rex1-Rac1) complex reveals the basis of Rac1 activation in breast cancer cells. *J. Biol. Chem.* **290**, 20827–20840 [CrossRef Medline](#)
 45. Guilluy, C., Dubash, A. D., and García-Mata, R. (2011) Analysis of RhoA and Rho GEF activity in whole cells and the cell nucleus. *Nat. Protoc.* **6**, 2050–2060 [CrossRef Medline](#)
 46. Adame-García, S. R., Cervantes-Villagrana, R. D., Orduña-Castillo, L. B., Del Rio, J. C., Gutkind, J. S., Reyes-Cruz, G., Taylor, S. S., and Vázquez-Prado, J. (2019) cAMP-dependent activation of the Rac guanine exchange factor P-REX1 by type I protein kinase A (PKA) regulatory subunits. *J. Biol. Chem.* **294**, 2232–2246 [CrossRef Medline](#)
 47. McKenzie, A. J., Svec, K. V., Williams, T. F., and Howe, A. K. (2020) Protein kinase A activity is regulated by actomyosin contractility during cell migration and is required for durotaxis. *Mol. Biol. Cell* **31**, 45–58 [CrossRef Medline](#)
 48. Howe, A. K., Baldor, L. C., and Hogan, B. P. (2005) Spatial regulation of the cAMP-dependent protein kinase during chemotactic cell migration. *Proc. Natl. Acad. Sci. U.S.A.* **102**, 14320–14325 [CrossRef Medline](#)
 49. Lim, C. J., Han, J., Yousefi, N., Ma, Y., Amieux, P. S., McKnight, G. S., Taylor, S. S., and Ginsberg, M. H. (2007) $\alpha 4$ integrins are type I cAMP-dependent protein kinase-anchoring proteins. *Nat. Cell Biol.* **9**, 415–421 [CrossRef Medline](#)