MINIREVIEW

Structure and Function of Heterotrimeric G Protein-Regulated Rho Guanine Nucleotide Exchange Factors

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ABSTRACT

Activation of certain classes of G protein-coupled receptors (GPCRs) can lead to alterations in the actin cytoskeleton, gene transcription, cell transformation, and other processes that are known to be regulated by Rho family small-molecular-weight GTPases. Although these responses can occur indirectly via cross-talk from canonical heterotrimeric G protein cascades, it has recently been demonstrated that Dbl family Rho guanine nucleotide exchange factors (RhoGEFs) can serve as the direct downstream effectors of heterotrimeric G proteins. Heterotrimeric \( \alpha_{12/13}, \alpha_q, \) and \( \beta\gamma \) subunits are each now known to directly bind and regulate RhoGEFs. Atomic structures have recently been determined for several of these RhoGEFs and their G protein complexes, providing fresh insight into the molecular mechanisms of signal transduction between GPCRs and small molecular weight G proteins. This review covers what is currently known about the structure, function, and regulation of these recently recognized effectors of heterotrimeric G proteins.

Heterotrimeric G proteins are master regulators of cell homeostasis. By coordinating signaling between the ~800 G protein-coupled receptors (GPCRs) in the human genome and a relatively small handful of effector enzymes and channels in the cell, they control processes such as muscle contractility, glycogen metabolism, neurotransmission, and the concentration of intracellular ions. Their profound impact on nearly all cellular processes and their therapeutic potential have rendered them one of the most intensely studied signal transduction paradigms at the biochemical and molecular level (Sprang et al., 2007).

When heterotrimeric G proteins are in their inactive, GDP-bound state, they exist as an inert complex composed of \( \alpha, \beta, \) and \( \gamma \) subunits (Ga\( \beta\gamma \)). In this state, they are substrates for activated GPCRs, which catalyze nucleotide exchange on the \( \alpha \) subunit (G\( \alpha \)). When bound to GTP, G\( \alpha \) releases the effector binding surface of the \( \beta \) and \( \gamma \) heterodimer (G\( \beta\gamma \)) so that both G\( \alpha \) and G\( \beta\gamma \) can interact with and modulate the activity of specific downstream enzymes and channels. The G\( \alpha \) subunit has weak guanine nucleotide triphosphatase (GTPase) activity that slowly returns the G protein to its GDP-bound state. G\( \alpha \)GDP then becomes resequestered by G\( \beta\gamma \).

Beyond serving as conduits for extracellular signals, heterotrimeric G proteins contribute to the fidelity, duration, and amplitude of GPCR signaling. A given class of heterotrimeric G protein can typically recognize only a subset of GPCRs, and can only interact with one or a few downstream effector targets, ensuring the specificity of signaling from receptor to effector. The rate of GTP hydrolysis on G\( \alpha \) dictates the length of time that its signal is in play, and this rate

ABBREVIATIONS: Dbs, Dbl’s big sister; DEP, Disheveled/EGL10/Pleckstrin; DH, Dbl homology; GAP, GTPase-activating protein; GEF, guanine nucleotide exchange factor; GPCR, G protein-coupled receptor; HEK, human embryonic kidney; IP4P, inositol polyphosphate-4-phosphatase; LARG, leukemia-associated RhoGEF; Lbc, lymphoid blast crisis; LPA, lysophosphatidic acid; PDZ, postsynaptic density 95, disk large, zona occludens-1; PH, pleckstrin homology; PIP3, phosphatidylinositol 3'-kinase; PIP2, phosphatidylinositol 4,5-bisphosphate; PKA, cAMP-dependent protein kinase; PLC, phospholipase C; P-Rex, PIP3-dependent Rac exchanger; rgrGS, RhoGEF fragment containing both N-terminal GAP motif and RH domain; RGS, regulators of G protein signaling; RhoGEF, Rho guanine nucleotide exchange factor; RH, RGS homology; SH3, Src homology 3; SRF, serum response factor.
Fig. 1. Structural studies of heterotrimeric G protein-regulated RhoGEFs. a, structure of Go13 in its active, GDP-AlF4-bound conformation. The Ras-like domain is colored cyan, and the α-helical domain is gray. The three nucleotide-dependent switch regions (switch I–III) are red. The canonical effector docking site, a shallow canyon formed between switch II and the α3 helix, is indicated by the transparent yellow ellipse. b, the p115RhoGEF rgRGS domain in complex with Go13/i1GDP-AlF4. The Go subunit is shown as a molecular surface and is colored as in a. The p115RhoGEF RH domain (orange) binds to the effector docking site of the Go13/i1 chimera with its α8-α9 loop (yellow). Noncanonical helices that extend the C terminus of the RH domain are colored in gold. The N-terminal motif responsible for GAP activity (green) binds to the α-helical domain with its IIIG motif, followed by residues that stabilize catalytic residues in the GTPase active site. c, the rgRGS domain of PDZ-RhoGEF in complex with Go13GDP-AlF4. A short α helix (αE, colored yellow) binds in the effector docking site, which is expanded relative to the effector docking site in the Go13/i1-p115RhoGEF complex. The N-terminal motif follows the same path as in the complex with p115RhoGEF, but there is no productive interaction with catalytic...
can be dramatically accelerated by either the effector target or regulators of G protein signaling (RGS) proteins, which serve as GAPs for some classes of Gα subunits (Ross and Wilkie, 2000). Finally, if the activated receptor is not uncoupled from the G protein, such as via phosphorylation by GPCR kinases and binding of arrestins, then Gβγ can undergo multiple rounds of activation. Thus, regulating the relative rates of G protein activation and deactivation can control the amplitude of signals initiated by GPCRs.

Crystallographic studies have uncovered the molecular mechanism by which activated Ga and Gβγ subunits engage their effector targets. In the case of Ga, the binding of GTP induces a conformational change in structural elements that interact with the γ-phosphate of GTP, known as switch I, II, and III (Fig. 1a). This conformational change lowers the affinity of G for Gβγ but increases its affinity for effectors. Upon binding GTP, a shallow hydrophobic canyon is created between switch II and the α3 helix of Ga that is used for binding all the effectors that have been structurally characterized thus far (Tesmer et al., 1997b, 2005; Slep et al., 2001; Chen et al., 2005, 2008; Lutz et al., 2007). There is sufficient variability in the residues that line the perimeter of this effector docking site to explain the exquisite specificity of Ga for its downstream targets. In the case of Gβγ, its ability to bind effectors depends on the unmasking of a broad, relatively hydrophobic surface that is sequestered in the Gaβγ complex. Conformational changes between the Ga- and effector-bound states of Gβγ have thus far proved relatively minor. Although the binding surface for the effectors of Gβγ is believed to overlap with that of Ga, they are believed to involve distinct regions in each case (Ford et al., 1998; Davis et al., 2005).

A Role for GPCRs in Cell Migration and Transformation

GPCRs are traditionally thought of as regulators of cellular metabolism or as mediators of sensory perception, such as vision, smell, and taste. The canonical effector enzymes involved in these processes include adenyl cyclase, cGMP phosphodiesterase, and phospholipase Cβ (PLCβ). The Ga subunits that couple receptors to these effectors represent three of the four subfamilies of human heterotrimeric Ga subunits: Gaαs, Gaα14, and Gaαq, respectively. However, GPCRs can also serve as potent oncogenes in a manner more typical of tyrosine kinase growth factor receptors (Whitehead et al., 2001). Indeed, constitutively activated Gaα subunits, in particular members of the Ga12/13 subfamily, can lead to strong cell proliferative responses (Dhanasekaran et al., 1998). The mechanism by which this occurs began to be understood by the observation that lysophosphatidic acid (LPA) and bombesin, known ligands for GPCRs, can activate the small-molecular-weight G protein RhoA and induce the formation of stress fibers and focal adhesions (Ridley and Hall, 1992). In subsequent years, numerous studies lent additional support to the idea that Rho GTPases were activated downstream of many GPCRs (Seasholtz et al., 1999; Sah et al., 2000).

The Rho GTPases are a family of peripheral membrane proteins that regulate essential cellular processes, including cell shape, cell migration, cell cycle progression, and gene transcription (Etienne-Manneville and Hall, 2002; Wennerberg and Der, 2004). Like their larger Ga subunit homologs, they cycle between an inactive GDP-bound and an active GTP-bound state that can interact with specific downstream effector targets, depending on the Rho GTPase. In vivo, Rho GTPases require an upstream guanine nucleotide exchange factor (GEF) for activation. The largest and best characterized RhoGEF family is characterized by a catalytic Dbl homology (DH) domain, an extended helical domain of ~200 amino acids that forms the principal binding site for the nucleotide-free GTPase. The DH domain is almost always immediately followed in the primary sequence by a pleckstrin homology (PH) domain (Schmidt and Hall, 2002; Rossman et al., 2005). In some DH-family RhoGEFs, the PH domain inhibits the intrinsic GEF activity of the DH domain (Das et al., 2000; Welch et al., 2002; Bellanger et al., 2003; Lutz et al., 2007), a constraint that is presumably released upon the interaction of the PH domain with phospholipids or other proteins. In other Dbl family members, the PH domain can play a positive signaling role (Liu et al., 1998; Reuther et al., 2001; Rossman et al., 2002).

GPCRs could regulate Rho GTPase signaling either directly via the interaction of activated heterotrimeric G proteins with a RhoGEF, or indirectly via, for example, activation of protein kinase A (PKA) or PKC, which can then phosphorylate RhoGEFs, their upstream regulators, or their downstream targets. In 1998, it was shown that Ga13 subunits bind directly to p115RhoGEF and stimulate its activity (Hart et al., 1998), thereby revealing a simple, direct path from LPA or thrombin receptors to RhoA (Wang et al., 2004). In 2005, it was shown that Ga11 could activate the activity of another RhoA-selective RhoGEF, p63RhoGEF, providing another direct pathway, this time initiated by bombesin, muscarinic, and angiotensin receptors (Lutz et al., 2005). Gβγ is also now known to directly bind and activate RhoGEFs, such as phosphatidylinositol (3,4,5)-triphosphate (PIP3)-dependent Rac exchanger 1 (P-Rex1), which links Gα12/13-coupled chemokine receptors in neutrophils to the activation of Rac2 (Welch et al., 2002).
In this review, we examine what is currently known about the structure and function of Dbl-family RhoGEFs that are directly regulated by heterotrimeric G proteins (Tables 1 and 2, Fig. 2). We focus primarily on three aspects relevant to molecular pharmacology: 1) the structures of RhoGEF domains involved in signal transduction, 2) the specific details of how heterotrimeric G proteins are known to interact with these domains, and 3) the potential mechanisms by which these interactions lead to RhoGEF activation.

### TABLE 1

<table>
<thead>
<tr>
<th>Heterotrimeric G Protein Family &amp; RhoGEF</th>
<th>Known Physiological Roles</th>
<th>Rho Family Substrate</th>
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<tr>
<td>Gα12/13p115RhoGEF/Lsc</td>
<td>Neutrophil migration, leukocyte homeostasis (Girkontaitė et al., 2001; Francis et al., 2006); regulation of apoptosis in thymocytes (Harenberg et al., 2005)</td>
<td>Rho</td>
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<tr>
<td>LARG</td>
<td>Signal transduction downstream of thrombin receptors (Wang et al., 2004); growth cone collapse (Swiercz et al., 2002; Hata et al., 2009); induction of salt-induced hypertension in vascular smooth muscle (Ying et al., 2006; Wirth et al., 2008)</td>
<td>Rho</td>
</tr>
<tr>
<td>PDZ-RhoGEF</td>
<td>Signal transduction downstream of LPA receptors (Wang et al., 2004); Ca2+ sensitization in smooth muscle (Derewenda et al., 2004); neutrophil polarization (Wong et al., 2007); neurite retraction (Togashi et al., 2000); angiotensin II induced contraction in vascular smooth muscle cells (Hilgers et al., 2007; Ying et al., 2009)</td>
<td>Rho</td>
</tr>
<tr>
<td>Lbc</td>
<td>Promotion of cardiac hypertrophy (Appert-Collin et al., 2007; Carnegie et al., 2008)</td>
<td>Rho</td>
</tr>
<tr>
<td>Gα13p63RhoGEF</td>
<td>Gα13-dependent activation of RhoA in smooth muscle cells (S. Lutz, personal communication)</td>
<td>Rho</td>
</tr>
<tr>
<td>TriC</td>
<td>In <em>C. elegans</em>, the C-terminal DH/PH domain of UNC-73 promotes acetylcholine vesicle release at neuromuscular junctions (Williams et al., 2007) and regulation of pharynx pumping, speed of locomotion, and egg-laying (Steven et al., 2005)</td>
<td>Rho</td>
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<tr>
<td>KalirinC</td>
<td>Neurite extension and neuronal neuronal morphology (Penzes et al., 2001)</td>
<td>Rho</td>
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<tr>
<td>Gβγ</td>
<td>P-Rex1: ROS generation and rate of chemotaxis in neutrophils (Dong et al., 2005; Welch et al., 2005); Purkinje cell morphology and cerebellar function (Donald et al., 2008)</td>
<td>Rac</td>
</tr>
<tr>
<td>P-Rex2</td>
<td>Purkinje cell morphology and cerebellar function (Donald et al., 2008)</td>
<td>Rho/Rac</td>
</tr>
<tr>
<td>Cig/PLEKHG2</td>
<td>Unknown</td>
<td>Rac/Cdc42</td>
</tr>
<tr>
<td>Tim/Arhgef5</td>
<td>Dendritic cell chemotaxis, allergic airway inflammation (Wang et al., 2009)</td>
<td>Rho</td>
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*GTPase specificity has not been demonstrated in vitro.*

### TABLE 2

<table>
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<tr>
<th>Structural analysis of heterotrimeric g protein-regulated RhoGEFs</th>
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<tr>
<td><strong>RhoGEF &amp; Domain(s)</strong></td>
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<td>------------------------</td>
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<tr>
<td><strong>p115RhoGEF</strong></td>
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<tr>
<td><strong>Gα13-rGDS domain complex</strong></td>
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<td><strong>LARG</strong></td>
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<td><strong>DH/PH-RhoA complex</strong></td>
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<tr>
<td><strong>PDZ domain</strong></td>
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<td><strong>PDZ-RhoGEF</strong></td>
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<td><strong>Gα13-DH/PH-RhoA complex</strong></td>
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whether their homology correlates with an ability to be regulated by heterotrimeric G proteins. The supporting data are weakest for Lbc, which has not yet been shown to bind or to be activated by \( \text{G}_{\alpha_{12/13}} \) subunits using purified proteins in vitro.

The RH-RhoGEFs

**Background.** Members of the \( \text{G}_{\alpha_{12/13}} \) subfamily are potent regulators of the actin cytoskeleton (Buhl et al., 1995) and are important for vascular development and chemotaxis (Offermanns et al., 1997). Given that Rho GTPases were also well established regulators of these responses, the hunt was on for RhoGEFs that could operate directly downstream of \( \text{G}_{\alpha_{12/13}} \). One candidate was p115RhoGEF, a RhoA-selective GEF containing a region with very weak homology to the catalytic RGS homology (RH) domain characteristic of RGS proteins (Tesmer, 2009). Indeed, this proved to be the binding site for activated \( \text{G}_{\alpha_{12/13}} \) subunits, although only \( \text{G}_{\alpha_{12}} \) could stimulate GEF activity (Hart et al., 1998; Kozasa et al., 1998; Mao et al., 1998). Soon thereafter, PDZ-RhoGEF (Fukuhara et al., 1999), also known as GTRAP48 (Jackson et al., 2001), and LARG (Fukuhara et al., 2000) were identified as RH-domain containing RhoGEFs. These three proteins form the RH-RhoGEF subfamily, a small subfamily of \( \text{G}_{\alpha_{13}} \)-regulated RhoGEFs. LARG is the only one currently known to be activated by \( \text{G}_{\alpha_{12}} \), although this seems to require phosphorylation of the RhoGEF by Tec tyrosine kinase (Suzuki et al., 2003). All three RH-RhoGEFs are widely expressed in mammals (Hart et al., 1996; Fukuhara et al., 1999; Kourlas et al., 2000; Kuner et al., 2002), but p115RhoGEF and PDZ-RhoGEF are most abundantly expressed in hematopoietic cells and the central nervous system, respectively.

In resting cells, RH-RhoGEFs are predominantly localized to the cytosol, with some presence at the cell membrane or bound to actin. Upon activation of \( \text{G}_{\alpha_{12/13}} \), coupled receptors or coexpression of constitutively active mutants of \( \text{G}_{\alpha_{12}} \) or \( \text{G}_{\alpha_{13}} \), RH-RhoGEFs translocate to the membrane surface (Siehler, 2009; Suzuki et al., 2009a). RH-RhoGEFs also exhibit some receptor selectivity. In HEK293 and PC3 cells, RhoA is activated by PDZ-RhoGEF via LPA receptors and by LARG via thrombin receptors (Wang et al., 2004). The mechanism by which this selectivity occurs is not known, but it is not apparently due to differences in their PDZ domains, because both PDZ-RhoGEF and LARG can bind to the C termini of LPA receptors (Yamada et al., 2005).

The RH RhoGEFs are complex proteins with multiple signaling domains and functional motifs hidden within the low-complexity regions that bracket them. There is considerable evidence that the mechanism of activation is also complex, involving not only recruitment to specific sites at the membrane surface, but also multiple intra- and intermolecular interactions that mediate inhibition of basal activity as well as allosteric activation (Bhattacharyya and Wedegaertner, 2003; Bhattacharyya et al., 2009; Suzuki et al., 2009b; Zheng et al., 2009). The specific details vary among subfamily members (e.g., p115RhoGEF lacks a PDZ domain) and context

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**Fig. 2.** Domain structure of RhoGEFs reported to be directly regulated by heterotrimeric G proteins. The amino acid numbers shown above each protein correspond to the human ortholog. For domains that have not been structurally characterized, domain boundaries were assigned based on sequence alignment with their closest homologs of known structure unless otherwise indicated. a, RhoGEFs regulated by \( \text{G}_{\alpha_{12/13}} \) subunits. p115RhoGEF, LARG, and PDZ-RhoGEF constitute the RH-RhoGEF subfamily and are characterized by an N-terminal RH domain. Short sequence motifs that seem to play functional roles are shown as small bars and are color-coded according to the key at the bottom of the panel. b, RhoGEFs regulated by \( \text{G}_{\alpha_{q}} \) subunits. All three RhoGEFs have closely related DH/PH domains that constitute the TrioC subfamily. c, RhoGEFs regulated by \( \text{G}_{\alpha_{13}} \) subunits. None of these RhoGEFs are closely related, although p114RhoGEF belongs to the Lbc subfamily of RhoGEFs, hence, its DH/PH domains are closely related to those of the RH-RhoGEF subfamily and Lbc. The boundaries for the PDZ domains and the domains shown in gray were assigned based on the GenPept entry for each protein.
(e.g., LPA versus thrombin receptors). In the following sections, we summarize what is known about the structure and function of the major domains and other functional motifs of RH-RhoGEFs, highlighting their potential roles in regulating the activity of the DH/PH domains. We end by describing recent solution-based studies of LARG and PDZ-RhoGEF that provide new insights into the molecular mechanism of RH-RhoGEF activation.

The RhoGEF RH Domain. As the principal binding site for \(G_{\alpha_{12/13}}\) subunits, structural analysis of the RhoGEF RH domain was anticipated to explain the molecular basis for selectivity and GAP activity and to shed some light on the mechanism of regulation by heterotrimeric G proteins. The canonical RH domain is a bundle of nine helices that forms a relatively flat domain with two lobes (Tesmer, 2009). In RGS proteins, the \(\alpha5-\alpha6\) loop of the domain interacts with catalytic residues in switch I and II of \(G_{\alpha}\) to accelerate GAP activity (Tesmer et al., 1997a). In contrast, the RH domains of p115RhoGEF and LARG do not have GAP activity. Instead, a short sequence motif positioned immediately N-terminal to the RH domain is required for stimulating GTP hydrolysis in p115RhoGEF and LARG (Chen et al., 2003). The region spanning this N-terminal motif and the RH domain has been termed the “rgRGS” domain (Chen et al., 2001) (Fig. 2a).

Crystal structures of the p115RhoGEF (Chen et al., 2001) and the PDZ-RhoGEF (Longenecker et al., 2001) RH domains confirmed that they indeed retain most of the canonical RH domain fold, albeit with a \(\sim65\)-residue extension that contributes three extra \(\alpha\) helices to the C terminus (Fig. 1, b and c). However, these initial structures did not reveal an obvious mode of interaction with \(G_{\alpha_{12/13}}\) subunits, in part because the \(G_{\alpha}\)-interacting residues of RGS proteins are not conserved in the RhoGEF RH domain (Tesmer, 2009). Although the entire rgRGS protein of p115RhoGEF was crystallized, the N-terminal GAP motif was not ordered.

The structure of the p115RhoGEF rgRGS domain in complex with a chimeric \(G_{\alpha_{13/i1}}\) Protein (Chen et al., 2005) revealed a surprising bipartite interaction mediated by the N-terminal GAP motif and the RH domain (Fig. 1b). The \(G_{\alpha_{13/i1}}\) chimera used in this study was created to circumvent the \(G_{\alpha}\)-interacting residues of RGS proteins that are not conserved in the RHGEF RH domain (Tesmer, 2009). Although the entire rgRGS protein of p115RhoGEF was crystallized, the N-terminal GAP motif was not ordered.

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The PDZ Domain. PDZ-RhoGEF and LARG can also be activated by receptors other than GPCRs via interactions mediated by the PDZ domains found at their N termini. PDZ domains are common structural domains that typically bind the C termini of proteins found at the cell membrane (Nourry et al., 2003). Compared with other PDZ domains, the RH-RhoGEF PDZ domain binds to surprisingly diverse biological targets (for review, see Smietana et al., 2008). Perhaps the best characterized is plexin B1, a semaphorin 4D receptor that mediates RhoA activation and axonal growth cone collapse (Aurandt et al., 2002; Hirotani et al., 2002; Perrot et al., 2002; Swierz et al., 2002). Both RhoGEFs also bind to LPA-1 and -2 receptors, which couple primarily with \(G_{\alpha_{12/13}}\) subunits. Interaction between the C termini of these receptors and the PDZ domain was reported to be required for RhoA activation (Yamada et al., 2005).

NMR structures of the LARG PDZ domain alone and in complex with a plexin-B1 C-terminal octapeptide have been reported (Liu et al., 2008) (Fig. 1d). A solution structure of the PDZ-RhoGEF PDZ domain is also available (Protein Data Bank entry 2dls). They retain the typical PDZ domain fold, which consists of six antiparallel strands (\(\betaA-\betaP\)) and two helices (\(\alphaA\) and \(\alphaB\)). Peptide ligands dock in an extended conformation into a groove located between \(\betaB\) and \(\alphaB\), where they form a \(\beta\)-sheet like interaction with the \(\betaB\) strand. The LARG PDZ domain undergoes an atypical con-
formational change upon ligand binding relative to other characterized PDZ domains (Fig. 1d), and the apo form exhibits a high degree of conformational flexibility (Liu et al., 2008). These characteristics may facilitate the binding of these PDZ domains to a wide variety of targets.

**DH/PH Domain Structures.** The canonical DH domain is composed of 10 or more helices that assemble into an ovoid shape reminiscent of a chaise lounge (Worthylake et al., 2000). The last helix (called α6) transverses the longest dimension of the domain and often forms a continuous helix with αN, the first helix of the PH domain (Fig. 1e). The PH domain consists of a flattened seven-stranded antiparallel β-barrel, capped at one end by αC, the hallmark C-terminal helix found in all PH domains. In some PH domains, loops on the opposite end of the barrel from αC are used to bind specific phosphohelipid head groups. The αN helix is characteristic of PH domains found in DH/PH tandems.

Structures have been determined for the LARG DH/PH domains, both alone and in complex with RhoA (Kristelly et al., 2004) and for the PDZ-RhoGEF DH/PH-RhoA complex (Derewenda et al., 2004). Both RhoA complexes exhibit similar domain organization and intersubunit contacts (Fig. 1f). As in DBs, another RhoGEF in which the PH domain contributes positively to GEF activity (Rossman et al., 2002, 2003), direct contacts are formed between RhoA and the PH domain. Although this interface accounts for only ~8% of the total buried surface area in the LARG DH/PH-RhoA complex, mutation of residues in the PH domain-RhoA interface diminished GEF activity to the level of the DH domain alone (Kristelly et al., 2004). Mutation of residues in the PDZ-RhoGEF interface did not lead to significant differences in vitro (Oleksy et al., 2006), and hence the functional importance of the RhoA-PH domain interface seems to vary among RH-RhoGEFs.

Examination of the DH/PH structures reveals several structural elements that could play a role in regulation. The first is a flexible, helical extension at the N terminus of the DH domain (Fig. 1e). Either truncation of the extension or mutation of a tryptophan (LARG-Trp769) buried between the extension and the DH domain reduced GEF activity (Kristelly et al., 2004). However, the LARG-W769D mutation exhibited only a minor defect in serum response element transcription assays in cells (Aittaleb et al., 2009). Sequence analysis predicts that the extension exists in all Lbc subfamily RhoGEFs, suggesting that it is not involved in regulation specific to the RH-RhoGEFs. Instead, recent data suggest that the N-terminal extension could be more important for autoinhibition of basal activity (see **Solution Studies**).

The second notable feature is a relatively long helical linker between the DH and PH domains that allows for considerable conformational flexibility between the two domains. For example, there is a ~30° difference in orientation of the PH domain relative to the DH domain between the LARG DH/PH and DH/PH-RhoA structures, the hinge occurring at the junction of the α6 and αN helices. The four unique DH/PH domains in the LARG DH/PH-RhoA structure also exhibit conformational differences in the α6-αN linker (Kristelly et al., 2004). Because the mutation of residues in the PH domain that directly interact with RhoA inhibits GEF activity (Kristelly et al., 2004), these contacts were proposed to alter the conformation of the α6-αN linker region in a manner that facilitates the binding of RhoA. Indeed, residues at the end of α6 make critical interactions with switch II of the bound GTPase (Aghazadeh et al., 1998). Interactions with other proteins or phospholipids that likewise influence the relative orientation of the DH and PH domains could likewise affect LARG activity. Solution studies of the PDZ-RhoGEF DH/PH domains found no evidence for an analogous large conformational change in the linker region (Cierpicki et al., 2009), suggesting that in PDZ-RhoGEF the PH domain may simply serve to stabilize the α6 helix of the DH domain.

Finally, in all three DH/PH structures, a solvent-exposed patch of conserved hydrophobic residues is observed in the PH domain (Fig. 1e,f). Depending on the crystal structure, the patch forms either a 2-fold crystallographic or a quasi-2-fold molecular interface that buries ~800 Å² of surface area. Located on the side of the PH domain β barrel, the patch would be in position to bind adjacent peripheral membrane proteins or domains at the cell membrane. Mutation of residues in the hydrophobic patch of LARG had no effect on nucleotide exchange activity in vitro but abrogated the ability of LARG to induce RhoA activation and stress fiber formation in cells (Aittaleb et al., 2009). Because the activity of these mutants could be rescued by fusion with nonspecific membrane targeting motifs, the hydrophobic patch seems to contribute to the proper localization of LARG by interacting with unknown target(s) at the cell membrane. However, membrane targeting by constitutively activated Gα13 did not rescue activity, suggesting that the hydrophobic patch also plays an essential role in GPCR-mediated regulation, although it does not yet seem to involve direct interactions with activated Gα13 (Aittaleb et al., 2009). The hydrophobic patch is conserved among all Lbc subfamily members, once again indicating a common functional role. Indeed, mutation of the hydrophobic patch of Lbc likewise abolished its ability to promote RhoA-mediated transcriptional activation in cells (Aittaleb et al., 2009).

**Other Functional Motifs.** p115RhoGEF, LARG, and PDZ-RhoGEF undergo homo- and heterodimerization mediated by their C-terminal tails, and specific coiled-coil motifs have been identified in p115RhoGEF and LARG (Eisenhaure et al., 2003; Chikumi et al., 2004; Grabocka and Wedegaertner, 2007) (Fig. 2a). Deletion of the C terminus augments the ability of RH-RhoGEFs to induce RhoA activation in cells, suggesting a negative regulatory role for dimerization, presumably via the recruitment of unknown inhibitory factors (Eisenhaure et al., 2003; Chikumi et al., 2004). A proline-rich motif positioned immediately C-terminal to the PDZ-RhoGEF PH domain (residues 1081-1119) has been shown to be essential for plasma membrane localization of PDZ-RhoGEF and is required for PDZ-RhoGEF induced cortical actin reorganization and cell rounding (Togashi et al., 2000). The RH-DH/PH linker region also contains functional elements. In addition to an acidic autoinhibitory motif (Zheng et al., 2009), the RH-DH/PH linker of PDZ-RhoGEF contains a unique actin-binding motif (Banerjee and Wedegaertner, 2004; Banerjee et al., 2009).

**Solution Studies.** Two recent solution-based studies have provided new insights into RH-RhoGEF regulation (Fig. 3a). Using surface plasmon resonance, the first study showed that Gα13 could interact not only with the RH but also with the DH/PH and C-terminal regions of LARG (Suzuki et al., 2009b), consistent with earlier observations using coimmunoprecipitation (Wells et al., 2002; Bhattacharyya and
The presence of all three domains (RH, DH, and PH) was required for formation of the highest affinity complex, and the K204A mutation of Gα13, which abrogates binding to the RH domain (Nakamura et al., 2004; Grabocka and Wedegaertner, 2005), did not eliminate binding to the DH/PH domain. Thus, Gα13 seems to exhibit at least two LARG binding surfaces: one for the rgRGS domain that is dependent on the activation status of Gα13 and another for the DH/PH domains that uses activation-independent structural elements, potentially the C-terminal region of the Ras-like domain (Kreutz et al., 2007).

Truncation studies of PDZ-RhoGEF identified a conserved acidic motif (residues 706-712) a few amino acids upstream of the N-terminal helical extension of the DH domain that

**Fig. 3.** Models of RhoGEF activation. a, LARG/PDZ-RhoGEF activation. Membrane recruitment and RhoA activation is mediated by receptors or other proteins that interact with the PDZ domain, such as the pleck B1 receptor, or by activated Gα13 subunits, which interact with the RH domain. Plexin B1 signaling to RhoA does not seem to require Gα13 (Perrot et al., 2002). Although membrane recruitment is one aspect of signal transduction to RH-RhoGEFs, additional interactions at the cell membrane involving the various structural domains and sequence motifs of the RH-RhoGEF are probably required for full activation (Bhattacharyya and Wedegaertner, 2003; Bhattacharyya et al., 2009). Indeed, there is evidence that Gα13 interacts with other regions in addition to the RH domain (indicated by red arrows) to stabilize a high-affinity complex and stimulate GEF activity (Suzuki et al., 2009b).

Autoinhibition of basal activity is believed to be imposed by the cooperative actions of the N-terminal region and C-terminal tail of the RhoGEF, as well as by an acidic sequence in the RH-DH linker region (Zheng et al., 2009). Not included in this schematic is the hydrophobic patch of the PH domain, which is required for RhoA activation in cells and probably mediates protein-protein interactions at the membrane surface (Aittaleb et al., 2009). Unlike in RH-RhoGEFs, p63RhoGEF/TrioC subfamily activation does not seem to require Gα13 subunits, which interact with the RH-DH-Arklik regions of p63RhoGEF. The bridging interactions of the Gα subunit are proposed not only to fix the PH domain in a less inhibitory conformation but also to activate the DH domain allosterically. c, Activation of P-Rex1 by Gβγ subunits in neutrophils. In neutrophils, P-Rex1 is a Rac2-specific GEF (Dong et al., 2005; Welch et al., 2005) that can be fully activated by GPCRs because of the presence of class IB PI3K (Donald et al., 2004), which are likewise activated by Gβγ subunits. The mechanism of activation shown is based on that proposed by Urano et al. (2008) with Gβγ binding to an intramolecular complex formed by the second DEP, first PDZ, and IP4P domains. Binding to Gβγ is expected to relieve the inhibition imposed by the DEP and PDZ domains (Hill et al., 2005) and recruit the enzyme to the cell surface.
serves an autoinhibitory role (Zheng et al., 2009). NMR spectroscopy comparing the chemical shifts in DH/PH fragments with or without this acidic motif suggests that it interacts with basic residues on the DH domain, in particular Arg867 and Arg868. Because these residues also interact with the bound GTPase, their interaction with the acidic motif would form an autoinhibited state reminiscent of that of Vav (Aghazadeh et al., 2000). Indeed, charge-reversing mutations of four of the acidic residues in the motif (the “activation box”; Fig. 2a) enhanced GEF activity ~20-fold in a DH/PH fragment of PDZ-RhoGEF. The enhancement was curiously only ~2-fold for the same mutants in the context of a larger fragment that also contained the PDZ and RH domain. This may indicate that the acidic motif and the PDZ and RH domains act synergistically to autoinhibit membrane activity. Acidic motifs also occur in the RH-DH linker regions of p115RhoGEF and LARG, although it remains to be seen if they function similarly. If the activation box is used in all Lbc subfamily members to regulate basal activity, it may provide a functional explanation for the N-terminal extension of the DH domain, which folds in such a manner that it would juxtapose the acidic motif of PDZ-RhoGEF with Arg867 and Arg868 on the surface of the DH domain.

### Lbc-RhoGEF

Lbc RhoGEF was originally isolated as a transforming oncogene (onco-Lbc) from patients with myeloid leukemia (Toksoz and Williams, 1994) and is the founding member of the Lbc subfamily of RhoGEFs. No atomic structures are currently available for this enzyme. Beyond the DH/PH domains, proto-Lbc contains a predicted α-helical domain and a proline rich motif in its C terminus that have been shown to act as negative regulators of proto-Lbc transforming activity and localize the enzyme to the membrane fraction of cells (Sterpetti et al., 1999). Onco-Lbc lacks this C-terminal region. The Lbc-RhoGEF splice variant AKAP-Lbc, or AKAP13, is highly expressed in the heart and functions both as a cAMP-dependent protein kinase (PKA)-scaffolding protein and a GEF for RhoA (Diviani et al., 2001). Similar to the RH-RhoGEFs, there is evidence that AKAP-Lbc can undergo homo-oligomerization through a leucine zipper motif in the C-terminal region. Deletion of this C-terminal region in AKAP-Lbc enhances its GEF activity, suggesting a negative regulatory role for oligomerization (Baisamy et al., 2005).

Proto-Lbc and AKAP-Lbc have been shown to be specifically stimulated by Gα12 as well as via activation of the LPA receptor (Majumdar et al., 1999; Diviani et al., 2001; Dutt et al., 2004). The activation seems direct, because Lbc can be coimmunoprecipitated from HEK293 cells with constitutively active Gα12 (Diviani et al., 2001; Dutt et al., 2004). However, it should be noted that Gαq can also be precipitated with Lbc, although with no obvious effect on RhoA activation (Sagi et al., 2001; Pi et al., 2002). The binding site for Gα12 is not known, but a region C-terminal to the PH domain with debatable homology to RH domains has been identified (Dutt et al., 2004). This region overlaps with the leucine zipper motif shown to mediate homodimerization (Baisamy et al., 2005).

### Gαq-Regulated RhoGEFs

The involvement of Gαq in Gq-mediated signaling has been recognized for more than a decade. Dominant-negative RhoA and C3 endotoxin were used to demonstrate the existence of a unique Rho-dependent pathway in response to the activation of Gαq-coupled receptors in cardiac myocytes (Sah et al., 1996). Later, the development of cell-based, RhoA pulldown assays enabled a direct demonstration of an increase in RhoA-GTP in response to the activation of Gαq-coupled receptors in HEK293 cells, establishing that Gαq operated upstream of RhoA, at least in some signaling contexts (Chikumi et al., 2002). Studies of mouse embryonic fibroblasts deficient in either Gα12/Gα13 or Gα12/Gα11 subunits confirmed that these G protein families activate RhoA via independent pathways, although RhoA activation by Gα12/Gα13 subunits was more robust (Vogt et al., 2003).

A number of RhoGEFs have been reported to interact with, and are thus potentially regulated by, Gαq. These include LARG (Booden et al., 2002; Chikumi et al., 2002; Vogt et al., 2003) and Lbc (Sagi et al., 2001; Pi et al., 2002). In neither case has a direct interaction between the RhoGEF and Gαq been demonstrated with purified components, and no Gαq-mediated stimulation of GEF activity has been reconstituted in vitro. Furthermore, reports conflict on whether or not the RH domain of LARG is responsible for binding Gαq.

### p63RhoGEF

p63RhoGEF, which is expressed predominantly in the brain and heart, has been shown to be activated by Gα12-coupled receptors in a manner that directly competes with the activation of PLCβ, the canonical effector of Gαq (Lutz et al., 2005). A splice variant lacking the first 106 amino acids, known as GEFT, was reported to activate Cdc42 and Rac1 in cell-based assays and with purified proteins in vitro (Guo et al., 2003). Although there is the possibility of cross-talk in cells, GEF activity toward Cdc42 and Rac1 has not been demonstrated by other groups (Souchet et al., 2002; Lutz et al., 2004; Rojas et al., 2007). Primary sequence analysis also strongly suggests that p63RhoGEF and its splice variants are selective for RhoA.

The primary sequence of p63RhoGEF lacked an obvious RH domain that could serve as a binding site for Gαq. Initial truncation studies instead localized the binding site to a fragment of p63RhoGEF that includes the PH domain (Lutz et al., 2005). Subsequently, more detailed truncation studies, site-directed mutagenesis, and structure determination revealed that the C-terminal helix of the PH domain formed the principal binding site for activated Gα12 (Lutz et al., 2007; Rojas et al., 2007). The closely related, C-terminal set of DH/PH domains in Trio and Kalirin were also shown to bind and be activated by Gαq in vitro (Lutz et al., 2007; Rojas et al., 2007) and in cells (Lutz et al., 2007). Although the physiological role of p63RhoGEF in mammals probably involves smooth muscle function (S. Lutz, personal communication), UNC-73E, the p63RhoGEF homolog in Cenorhabditis elegans, is known to serve as a bona fide downstream effector of Gαq and works in conjunction with PLCβ to regulate the release of acetylcholine vesicles at neuromuscular synapses (Williams et al., 2007). Its knockout, in combination with a PLCβ-null allele, generates the same phenotype as knockouts of Gαq, confirming the necessary involvement of both pathways...
downstream of Goq (Williams et al., 2007). Finally, the kinase domain of mixed lineage kinase 3 was reported to bind to p63RhoGEF and inhibit its interaction with Goq, thereby regulating directed cell migration (Swenson-Fields et al., 2008). Although it is not yet known what moiety of p63RhoGEF could be involved in binding mixed lineage kinase 3, the PH domain would be an obvious candidate. It is noteworthy that Go16, a divergent member of the Goq subfamily, binds but fails to activate p63RhoGEF, suggesting that the domain-bridging interactions of Goq strain the DH and PH domains in this conformation and that the DH/PH domains adopt a more Dbs-like conformation in the basal state.

**Mechanism of Activation.** Unlike the RH-RhoGEFs, p63RhoGEF is predominantly found localized to actin-rich structures (Souchet et al., 2002), and overexpressed p63RhoGEF is localized to the cell surface (A. Shankaranarayanan, C. A. Boguth, S. Lutz, C. Vettel, F. Uhleman, M. Aittaleb, T. Wieland, and Tesmer JJ, submitted). Therefore, membrane translocation is not expected to play a major role in activation. In addition, unlike RH-RhoGEFs and Dbs, the PH domain of p63RhoGEF is autoinhibitory, the isolated DH/PH fragment having far less activity than the isolated DH domain (Lutz et al., 2004, 2007; Rojas et al., 2007). Thus, the PH domain must somehow interfere with the binding of RhoA. Indeed, the binding of Goq greatly enhances the affinity of RhoA for the PH/DH domains (A. Shankaranarayanan, C. A. Boguth, S. Lutz, C. Vettel, F. Uhleman, M. Aittaleb, T. Wieland, and Tesmer JJ, submitted). Because the PH domain does not inhibit GEF activity when added in trans (A. Shankaranarayanan, C. A. Boguth, S. Lutz, C. Vettel, F. Uhleman, M. Aittaleb, T. Wieland, and Tesmer JJ, submitted), the inhibition mediated by the PH domain seems to require a covalent linkage between the DH and PH domains, once again implicating the α6-αN linker as a region of regulatory importance.

Inspection of residues in the α6-αN linker region indicate that Arg341 of p63RhoGEF might sterically and electrostatically repel Arg68 in switch II of RhoA if the PH domain were in a more Dbs-like orientation relative to the DH domain. An R341A mutation indeed enhances the basal activity of p63RhoGEF but, surprisingly, does not eliminate activation by Goq (A. Shankaranarayanan, C. A. Boguth, S. Lutz, C. Vettel, F. Uhleman, M. Aittaleb, T. Wieland, and Tesmer JJ, submitted). In fact, the R341A mutation allows Goq to enhance activity to levels greater than those of the DH domain alone, indicating that Goq can activate p63RhoGEF not only by removing an autoinhibitory restraint but also by activating the DH domain allosterically. This is presumably carried out via the interactions of the C-terminal region of Goq with the DH domain, as observed in the crystal structure, which in turn alters the conformation of the α6-αN linker (A. Shankaranarayanan, C. A. Boguth, S. Lutz, C. Vettel, F. Uhleman, M. Aittaleb, T. Wieland, and Tesmer JJ, submitted). A schematic model for Goq-mediated activation of p63RhoGEF and related DH/PH domains is shown in Fig. 3b.

**Trio and Kalirin**

Trio and Kalirin are closely related proteins that contain two sets of DH/PH domains and a much more complex domain structure than p63RhoGEF (Fig. 2b). Both proteins play critical roles in neurite growth and neuronal morphology (Table 1). The second set of DH/PH domains in Trio (TrioC) is closely related to that of p63RhoGEF and is likewise selective for RhoA. The first set, TrioN, which is selective for RhoG and Rac, has been structurally characterized (Liu et al., 1998; Skowronek et al., 2004; Chhatrivala et al., 2007). The PH domain of TrioC likewise inhibits GEF activity (Bellanger et al., 2003), and the regulation of the TrioC and KalirinC DH/PH domains by Goq is expected to be similar to that of p63RhoGEF (Lutz et al., 2007; Rojas et al., 2007). It is not known how the other domains in these com-
plex proteins will affect regulation by heterotrimeric G proteins, especially in light of the fact that downstream signaling from each set of DH/PH domains has the potential to antagonize each other. There are, however, isoforms of Trio and Kalirin in mammals (Kawai et al., 1999; McPherson et al., 2005) and of UNC-73 in C. elegans (Steven et al., 2005) that contain only the C-terminal DH/PH domains. These splice variants are thus expected to act in a manner analogous to that of p63RhoGEF in cells.

**Gβγ-Regulated RhoGEFs**

As yet, there are no reported structures of Gβγ in complex with a RhoGEF, and compared with the Gαs and Gαq-regulated RhoGEFs, relatively little is known about the specific residues involved in binding or the molecular mechanisms involved. In two of the four examples covered here, the physiological roles of the enzyme are unknown (Table 1). The example with the strongest experimental support is P-Rex1, in which Gβγ-mediated regulation has been demonstrated by a number of different groups, and a direct interaction was quantitatively measured in vitro using surface plasmon resonance (Urano et al., 2008). Gβγ is also reported to regulate directional sensing mediated by the RhoGEF PIXα via complex formation with p21-associated protein kinase (Li et al., 2003). However, because the interaction of Gβγ with PIXα is thought to be indirect, it is not discussed further in this review.

**The P-Rex Subfamily**

The P-Rex subfamily of RhoGEFs is made up of P-Rex1 (Welch et al., 2002) and P-Rex2 (Donald et al., 2004; Rosenfeldt et al., 2004). P-Rex1 and P-Rex2 are of similar size (~185 kDa) and have high sequence identity (59%) (Donald et al., 2004). Their domain architecture begins with the characteristic DH/PH tandem domains, followed by two DEP domains, two PDZ domains, and a C-terminal domain that exhibits homology to inositol polyphosphate-4-phosphatase (IP4P) (Welch et al., 2002) (Fig. 2c). The IP4P-like domain does not seem to have phosphatase activity (Hill et al., 2005). P-Rex2, a splice variant of P-Rex2, lacks the IP4P domain (Rosenfeldt et al., 2004). P-Rex1, P-Rex2, and P-Rex2b are all directly regulated by Gβγ subunits and PIP3 (Welch et al., 2002; Donald et al., 2004; Li et al., 2005).

P-Rex1 is highly expressed in white blood cells and the brain. In neutrophils, P-Rex1 couples GPCRs to Rac-dependent formation of reactive oxygen species (Welch et al., 2002). Upon binding chemokines, Gαi-coupled GPCRs release Gβγ subunits, which increase the concentration of PIP3 by activating class IB phosphatidylinositol-3 kinases (PI3Ks). Thus, in neutrophils, the activation of Rac can occur independently from activation of tyrosine kinase receptors. P-Rex2 has wide tissue distribution but is most strongly expressed in skeletal muscle, small intestine, and placenta, whereas the splice variant P-Rex2b is expressed highly in the heart (Donald et al., 2004; Rosenfeldt et al., 2004). Neither isoform of P-Rex2 is expressed in neutrophils. Because the cells in which P-Rex2 and P-Rex2b are expressed generally lack class IB PI3Ks, it is believed that P-Rex2 integrates signals generated from both GPCRs and tyrosine kinase receptors, the latter of which activates the more ubiquitously expressed class IA PI3Ks (Donald et al., 2004).

**Mechanism of Activation.** In cells, P-Rex1 is activated when both Gβγ and PI3Ks act synergistically to activate the enzyme (Welch et al., 2002). Not all Gβγ subunit isoforms can activate P-Rex1 equally, although the ones with the most activity seem to be the same ones that activate PI3K the most (Mayeenuddin et al., 2006). Gγ1 and Gγ11, two of the Gγ subunits that were less effective at activating P-Rex1, are farnesylated instead of geranylgeranylated, suggesting that their ability to stably associate with the cell membrane is important for regulation by Gβγ. Phosphorylation of P-Rex1 by PKA inhibits activation by Gβγ by approximately 50-fold (Mayeenuddin and Garrison, 2006), and thus P-Rex1 can be regulated by GPCRs via both direct (Gβγ) and indirect (PIP3 and PKA) pathways.

To delineate the regions that are important for Gβγ and PIP3 binding, truncation studies of P-Rex1 were conducted. The isolated DH/PH tandem domain fragment exhibited higher activity than the full-length protein, indicating that other domains of the protein are involved in autoinhibition of basal activity. Of all the domains, the PH domain was found to be most important for PIP3 activation of P-Rex1, although it lacks obvious sequence signatures for phospholipid binding found in other PIP3-binding PH domains, and there was evidence for at least one other PIP3-binding site in the protein (Hill et al., 2005).

Identification of the Gβγ-binding site has not been as clear-cut. The isolated DH/PH tandem domain of P-Rex1 was reportedly activated by Gβγ, as was a variant with the PH domain deleted, suggesting that the DH domain, being the only domain in common between the constructs, was sufficient for binding and regulation by Gβγ (Hill et al., 2005). However, direct activation of the isolated DH domain was not shown. Coimmunoprecipitation assays using various fragments of P-Rex2b suggested instead that the PH domain contained the binding site for Gβγ (Li et al., 2005). A more recent study of P-Rex1, again using coimmunoprecipitation, indicated that the DH domain does not directly bind Gβγ. Rather, an intramolecular complex formed by the second DEP, first PDZ, and IP4P domains creates a binding site for Gβγ. PKA phosphorylation, presumably at consensus sites found in these domains, weakened these interdomain contacts and thus inhibited Gβγ binding (Urano et al., 2008). This latter result, however, is difficult to reconcile with the fact that P-Rex2b seems to be regulated by Gβγ and altogether lacks an IP4P domain (Li et al., 2005) and that the DH/PH tandem domains of P-Rex1 are sufficient to observe PIP3- and Gβγ-mediated membrane translocation (Barber et al., 2007). Thus, although PIP3-mediated membrane recruitment represents one component of the activation mechanism for P-Rex subfamily RhoGEFs, the molecular basis for allosteric control of enzymatic activity by Gβγ remains obscure, except that it probably results in the release of the inhibition mediated by domains C-terminal to the DH domain (Fig. 3c).

**p114RhoGEF**

p114RhoGEF is another member of the Lbc subfamily of RhoGEFs that can be activated via stimulation of LPA or M2-muscarinic receptors. Beyond the characteristic tandem DH/PH module, the protein contains no other recognized structural domains but does contain a C-terminal proline-rich region (Blomquist et al., 2000). The protein is widely expressed (Blomquist et al., 2000; Niu et al., 2003), and a
shorter variant was detected in placenta and smooth muscle (Niu et al., 2003). There are conflicting reports on whether it is selective for RhoA (Blomquist et al., 2000; Nagata and Inagaki, 2005) or both RhoA and Rac1 (Niu et al., 2003). However, it should be noted that its DH/PH domains are densely related in sequence to those of other Lbc subfamily RhoGEFs, which are all believed to be specific for RhoA. The physiological roles of p114RhoGEF are unknown, but it has been identified as a binding partner for septin 9b, a protein involved in cytokinesis. This interaction is believed to be mediated by regions C-terminal to the DH/PH domains (Nagata and Inagaki, 2005).

Initial studies indicated that p114RhoGEF activity is directly regulated by Gβγ and not by Gα12/13 subunits (Niu et al., 2003). A subsequent study, however, found no evidence that the enzyme could be activated by Gβγ (Ueda et al., 2008). The reason for this discrepancy is not clear. Truncations of p114RhoGEF demonstrated that the DH/PH module has higher basal GEF activity than wild type, suggesting that the C-terminal region is inhibitory (Niu et al., 2003); hence, Gβγ binding would be anticipated to release autoinhibition. To identify the Gβγ binding site, an N-terminal fragment containing the DH/PH domains and C-terminal fragments (with and without the proline-rich region) were coexpressed in the presence or absence of Gβγ in NIH3T3 cells (Niu et al., 2003). Although the N-terminal catalytic fragment (residues 1–647) was as fully responsive to Gβγ activation as WT in driving SRF-mediated transcription, immunoprecipitation from cell lysates suggested that Gβγ could bind to either N- or C-terminal fragments.

Clg

Common-site lymphoma/leukemia GEF (Clg; also known as PLEKHG2 or FLJ00018) is a widely expressed GEF that is selective for Cdc42, although some activity for Rac1 has been demonstrated (Himmel et al., 2002; Ueda et al., 2008). The enzyme was singled out in a screen designed to identify RhoGEF enzymes for which SRF-mediated transcription could be enhanced by the coexpression of Gβγ subunits. The existence of such an enzyme could help explain how Gγ-coupled receptors induce cell-spreading in NIH3T3 fibroblasts (Ueda et al., 2001). In addition to the DH/PH tandem domains, putative formin homology and PDZ domains are predicted in the C terminus of Clg. The importance of these motifs is not known, although the C terminus of the enzyme clearly plays a role in inhibition of basal GEF activity (Ueda et al., 2008). Constructs of Clg that contain only the DH domain do not activate transcription, indicating that the PH domain is necessary for activity in cells, perhaps via the direct interactions of this domain with the Rho GTPase or the cell membrane. Treatment with sortase demonstrated that Clg activity is not dependent on PIP3, which is known to interact with the PH domain of P-Rex1.

To identify the binding site for Gβγ, a series of fragments of Clg with increasing portions of the C terminus removed were expressed, purified, and tested for their ability to bind Gβγ (Ueda et al., 2008). A DH domain fragment was also tested. All fragments except the DH domain fragment could bind Gβγ in a GST-pulldown assay. The smallest fragment that retained nearly full ability to bind Gβγ spans residues 1 to 134, which includes a small portion of the DH domain. This fragment could inhibit the ability of Clg to promote SRF-dependent gene transcription and induce cell spreading in a concentration-dependent manner, supporting its identification as the Gβγ binding site (Ueda et al., 2008).

**Tim/Arhgef5**

Tim, also known as Arhgef5, was first identified in a screen for focus-forming genes in NIH3T3 cells (Chan et al., 1994) and more recently in a small interfering RNA based screen for enzymes essential for macrophage chemotaxis (Wang et al., 2009). The enzyme is selective for RhoA and related GTPases (Snyder et al., 2002; Xie et al., 2005; Wang et al., 2009). Tim is a relatively simple RhoGEF, having in addition to the DH/PH domains a C-terminal SH3 domain that interacts with an N-terminal proline-rich region of the protein (Yohe et al., 2008). In addition, a helix N-terminal to the DH domain is predicted to bind directly to the DH domain to autoinhibit activity (Yohe et al., 2007). Both release of the SH3 domain (e.g., via competition with other proline rich motifs) and tyrosine phosphorylation of the N-terminal helix are postulated to be required for full activation of Tim (Yohe et al., 2008).

Because Gβγ subunits are known to be involved in leukocyte chemotaxis, the ability of Gβγ to bind and regulate the activity of Tim was tested. In coimmunoprecipitation assays, Gβγ was shown to bind to fragments that contained the PH domain and the preceding DH/PH linker region. Pulldown assays using purified proteins demonstrated that the interaction between Tim and Gβγ was direct. Gβγ also activated Tim in a serum response element reporter assay as well as in vitro using purified proteins (Wang et al., 2009). It remains to be seen how regulation by Gβγ is integrated into the layers of autoinhibition already mediated by the SH3 domain and N-terminal helix motif, or if it simply provides a mechanism to bypass them. It also remains to be seen whether other members of the Tim subfamily of RhoGEFs, which includes ephexin/NGEF, neuroblastoma, WGEF, SGEF, and Vsm-RhoGEF (Rossman et al., 2005), are also regulated by Gβγ.

**Conclusions**

Crystal structures and other biophysical studies of RH-RhoGEFs and p63RhoGEF have yielded substantial new insights into the molecular mechanisms governing their regulation by heterotrimeric G proteins. The current data suggest that in most cases, heterotrimeric G proteins interact directly with the catalytic core of the enzyme to stimulate activity. For example, the Gα13-RH domain complex is expected to bind to the DH/PH domain in RH-RhoGEFs, and Gαi forms direct contacts with both DH and PH domains of p63RhoGEF in the Gαi-p63RhoGEF-RhoA crystal structure. The binding of heterotrimeric G proteins does not always have to be stimulatory, as it has been proposed that Gα12 and Gα16 can antagonize signaling by their closely related paralogs. These proteins bind with high affinity to the same sites as Gα12 and Gα16, respectively, yet somehow fail to induce a conformational change that leads to RhoGEF activation. The molecular basis for this failure to activate is not yet understood.

Based on the available structural data from the RH-RhoGEFs and p63RhoGEF, the confocation of the α6-αN linker, and hence the relative orientation of their DH and PH domains (Fig. 1, e–g), is expected to play a role in tuning activity. In RH-RhoGEFs, the PH domain augments the ac-
tivity of the adjacent DH domain, even in the absence of heterotrimeric G proteins, by proteins that link the trimeric domain in a more optimal conformation. In this case, it is tempting to speculate that the role of heterotrimeric G proteins may primarily be to displace autoinhibitory elements from the DH/PH core domain. In TrioC RhoGEFs, the PH domain and linker adopt a conformation that instead inhibits the binding of RhoA in the basal state. Confronted with this scenario, heterotrimeric G proteins must bind directly to the DH/PH core and constrain the DH and PH domains in a manner that relieves inhibition, that does not block the binding of RhoA, and that allosterically activates nucleotide exchange (Fig. 1g).

As in other Gα-effector complexes, the effector docking site of Gα forms the principal binding site for downstream RhoGEFs. The effector docking site of Gα13 seems to be conformationally flexible, not only in response to GTP loading and hydrolysis but also when challenged by the unique docking motifs presented by p115RhoGEF and PDZ-RhoGEF (Chen et al., 2008) (Fig. 1b and c). Thus, even closely related effectors need not necessarily have well conserved Gα-binding motifs. Contacts made with Gα outside of the effector docking site mandate specificity and can also contribute to regulation, as in the case of RH-RhoGEFs and p63RhoGEF.

Unfortunately, the structural data reported thus far are still not definitive with regard to the mechanism of signal transduction for any heterotrimeric G protein-regulated RhoGEF. In no case are structures known for both the inactive and active forms of these enzymes, and future efforts are likely to focus on obtaining other snapshots of these enzymes along their reaction coordinates. Structures of RH-RhoGEFs that include the PH domain, in addition to the DH/PH domains, and of apo-p63RhoGEF would be especially informative. This is a challenging task, especially when considering that the domains are connected by long, unstructured linkers. High-resolution spectroscopic studies, such as those employed by Zheng et al. (2009), hold promise in these situations. Our understanding of the molecular basis for regulation of the Gα-subunits. Structural analysis of these enzymes and their Gββx complexes will be important for rapidly advancing our understanding of the molecular mechanisms that control their activity.

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References


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