PI3K Class II α Controls Spatially Restricted Endosomal PtdIns3P and Rab11 Activation to Promote Primary Cilium Function

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SUMMARY

Multiple phosphatidylinositol (PtdIns) 3-kinases (PI3Ks) can produce PtdIns3P to control endocytic trafficking, but whether enzyme specialization occurs in defined subcellular locations is unclear. Here, we report that PI3K-C2α is enriched in the pericentriolar recycling endocytic compartment (PRE) at the base of the primary cilium, where it regulates production of a specific pool of PtdIns3P. Loss of PI3K-C2α-derived PtdIns3P leads to mislocalization of PRE markers such as TfR and Rab11, reduces Rab11 activation, and blocks accumulation of Rab8 at the primary cilium. These changes in turn cause defects in primary cilium elongation, Smo ciliary translocation, and Sonic Hedgehog (Shh) signaling and ultimately impair embryonic development. Selective reconstitution of PtdIns3P levels in cells lacking PI3K-C2α rescues Rab11 activation, primary cilium length, and Shh pathway induction. Thus, PI3K-C2α regulates the formation of a PtdIns3P pool at the PRE required for Rab11 and Shh pathway activation.

INTRODUCTION

Phosphatidylinositol 3-kinases (PI3Ks) are lipid kinases involved in a large set of biological processes, including membrane receptor signaling, cytoskeletal organization, and endocytic trafficking (Shigo et al., 2010; Vanhaesebroeck et al., 2010). Mammals possess eight PI3K genes, which are divided into three classes on the basis of structural homology and substrate specificity (class I, II, and III). All PI3Ks phosphorylate the D3 position of the inositol ring of phosphatidylinositols (PtdIns), lipids involved in signal transduction as well as in membrane specification and dynamics (Di Paolo and De Camilli, 2006). Of the different 3-phosphorylated PtdIns species, PtdIns3P is the only product that can be directly or indirectly generated by all PI3K classes in vivo (Jean and Kiger, 2012). For example, class I PI3Ks (PI3Kx, PI3Kβ, PI3Kγ, and PI3Kδ) produce PtdIns(3,4,5)P3 that can be converted into PtdIns3P by phosphatases acting on endocytic vesicles (Shin et al., 2005). The unique member of class III, Vps34, is responsible for a major fraction of PtdIns3P produced on endocytic vesicles, where it controls the generation of autophagosomes (Backer, 2008) as well as docking and fusion of endosomes (Christoforidis et al., 1999). Class II PI3Ks (namely, PI3K-C2α, PI3K-C2β, and PI3K-C2γ) produce PtdIns3P as well (Falasca et al., 2007; Maffucci et al., 2003) and are involved in intracellular membrane trafficking, endocytosis, exocytosis (Falasca and Maffucci, 2012), and autophagy (Deveraux et al., 2013). However, the precise function of class II PI3K-produced PtdIns3P remains partially obscure. In flies, the only class II homolog, P3k68D, is required for endosomal sorting from the endocytic compartment to the plasma membrane, likely via regulation of PtdIns3P levels (Jean et al., 2012; Velichkova et al., 2010). Mammalian PI3K-C2α has been proposed to play a similar role in endothelial cells, where it promotes endosomal trafficking via RhoA activation and regulation of PtdIns3P levels. This process is required for the targeting of vascular endothelial (VE)-cadherin to tight junctions and consequent endothelial cell maturation and vessel integrity (Yoshioka et al., 2012). In agreement with PI3K-C2α playing multiple roles in different membrane compartments, PI3K-C2α has been reported to produce PtdIns(3,4)P2 at the plasma membrane. This lipid is crucial for clathrin-coated pit maturation and clathrin-mediated endocytosis (Posor et al., 2013).

Interestingly, vesicular trafficking and metabolism of phosphorylated PtdIns converge in the organization and functional maintenance of the primary cilium (Bielas et al., 2009; Jacoby...
Primary cilia are versatile organelles that, in most mammalian cells, function as motile propellers or sensorial antennas to regulate cell proliferation, polarity, differentiation, and tissue organization. Primary cilia provide a separate highly regulated compartment and, for cell assembly, they require influx of proteins and membranes from the cytosol (Pedersen and Rosenbaum, 2008). Targeting of components to the cilium involves polarized trafficking of vesicles originating from the Golgi and the endocytic recycling compartment (Follit et al., 2006; Hsiao et al., 2012; Pedersen and Rosenbaum, 2008; Wang et al., 2012). Biochemical and genetic approaches indicate that delivery and docking of secretory vesicles at the base of the cilium is regulated by a cascade of events involving the activation of specific small GTPases of the Rab family (Yoshimura et al., 2007). Rab8 and its activator, Rabin8, are essential for the entry of protein cargoes into the ciliary compartment, and impaired Rabin8 localization at the base of the cilium as well as defective Rabin8-dependent Rab8 activation reduces ciliary elongation (Das and Guo, 2011; Nachury et al., 2007). Targeting of Rabin8 to the ciliary base is regulated by another Rab GTPase, Rab11 (Westlake et al., 2011), a known coordinator of endosome recycling to the plasma membrane (Grant and Donaldson, 2009). At the ciliary base, Rab11 directly associates with Rabin8 and stimulates its guanine nucleotide exchange factor activity toward Rab8 (Feng et al., 2012); consequently, loss of Rab11 causes defective Rab8 activation and results in impaired ciliary elongation (Knödler et al., 2010).

Transport of components to and from the cilium is deeply interconnected with primary cilium function in signal transduction. For example, a well-organized and fully functional primary cilium is necessary for the cellular response to Sonic Hedgehog (Shh), a morphogen critically involved in vertebrate embryonic development (Goetz and Anderson, 2010). Consistently, loss of proteins involved in intraflagellar transport prevents correct primary cilium formation and has a dramatic impact on Shh signaling and mouse development (Huangfu and Anderson, 2005; Huangfu et al., 2003; Liem et al., 2012; Ocbina et al., 2011; Tran et al., 2008). Shh effectors, such as the Smoothened transmembrane protein and the Gli transcription factors, accumulate in cilia after pathway activation (Corbit et al., 2005; Ocbina et al., 2011). Translocation of these factors to the ciliary compartment is required for downstream steps of signal transduction, and alterations of this trafficking cause impairment of Shh signaling even in the presence of a fully elongated cilium (Keady et al., 2012). Genetic deletion of Smo in mouse embryos completely disrupts Hedgehog (Hh) signaling, thus impairing turning, left-right axis development, and neural tube patterning and leading to early embryonic lethality (Zhang et al., 2001).

Loss of the Pik3c2a gene in mouse has been reported to cause early embryonic lethality, initially ascribed to defective vasculogenesis (Yoshioka et al., 2012). Here, we show that in addition to abnormal angiogenesis, the lack of Pik3-C2α causes defective primary cilium organization as well as reduced Shh signaling. We report that this phenotype is linked to the ability of Pik3-C2α to control the production of PtdIns3P at the endocytic recycling compartment located at the base of the primary cilium. This specific pool of PtdIns3P was found to be required to activate the Rab11/Rab8 axis and promote Smo translocation to the ciliary shaft. Thus, PI3K-C2α integrates lipid signaling and Rab11 activation necessary for Shh signaling.

RESULTS

PI3K-C2α Localizes at the Ciliary Base

To explore the intracellular distribution of PI3K-C2α, a GFP-tagged form of the protein (GFP-PI3K-C2α) was transfected into mouse embryonic fibroblasts (MEFs) and localization was assessed by fluorescence microscopy. During interphase, GFP-PI3K-C2α was enriched at the perinuclear/pericentriolar area (Figure 1A). In G0 cells, the centrosome constitutes the cilium basal body; thus, endogenous PI3K-C2α was stained together with a centrosomal marker (γ-tubulin) and a ciliary marker (acetyl-tubulin) to analyze relative positioning. This staining failed to show any signal along the ciliary shaft but underlined a specific accumulation of PI3K-C2α in vesicular structures surrounding the ciliary basal body (Figure 1B), a region involved in primary cilium biogenesis (Kim et al., 2010). Enrichment of the protein in proximity of the base of the primary cilium was confirmed in a cell line widely used in cilium studies: the inner medullary collecting duct 3 (IMCD3) cell line. Downmodulation of Pik3c2a through infection with specific small hairpin RNA (shRNA) sequences (Sh1 and Sh2) severely reduced the immunofluorescence signal, thus confirming antibody specificity (Figures S1A and S1B available online).

The pericentriolar enrichment of PI3K-C2α in primary MEFs was found to colocalize with markers of the recycling compartment such as Rab11 and the transferrin receptor (TfR) (Figure 1C, top and bottom panels, respectively). In further agreement, cell fractionation experiments revealed the presence of PI3K-C2α in Rab11+ endosomes and not in Rab7+ late endosomes (Figure 1D). Altogether, immunofluorescence and fractionation experiments reveal enrichment of PI3K-C2α in the pericentriolar recycling endocytic compartment (PRE).

PI3K-C2α Produces a Specific Pool of Perinuclear PtdIns3P

To explore the function of PI3K-C2α in the PRE, PI3K-C2α-deficient (Pik3c2a−/−) MEFs were obtained from animals genetically modified by gene targeting in the mouse germine and showing complete ablation of PI3K-C2α (Figures S2A–S2D). First, localization of PtdIns3P, a phosphoinositide mainly involved in vesicular trafficking and produced by PI3K-C2α (Falasca et al., 2007; Yoshioka et al., 2012), was analyzed in primary MEFs. Interestingly, wild-type cells stained either with a PtdIns3P-selective GFP-FYVE fluorescent probe (Figure 2A) or with anti-PtdIns3P antibodies (Figure S2E) showed abundant labeling of PtdIns3P around the base of the cilium. By contrast, in Pik3c2a−/− MEFs, the pool of PtdIns3P around the ciliary base was reduced, whereas PtdIns3P detected in the rest of the cell did not show significant changes (Figures 2A and 2B; Figure S2E). A small but significant reduction in total PtdIns3P cellular levels (~21%) was also measured by high-pressure liquid chromatography (HPLC) analysis after metabolic labeling of starved Pik3c2a−/− MEFs (Figures 2C and 2D), in line with the notion that a restricted pool of PtdIns3P is specifically produced by PI3K-C2α.
Loss of PI3K-C2α Disrupts Pericentriolar Localization and Activation of Rab11

The reduction of PtdIns3P staining in the pericentriolar area suggested that the lack of PI3K-C2α altered the organization of the PRE compartment around the ciliary base. In agreement with this hypothesis, markers of the PRE such as Rab11 and TfR appeared dispersed and mislocalized in Pik3c2a/C0/C0 MEFs (Figures 3A and 3B, lower panels), whereas they were enriched at the expected pericentriolar location in wild-type cells. This was not due to a reduction of protein levels, because the total amount of Rab11 and TfR in Pik3c2a/C0/C0 MEFs was comparable to wild-type controls (Figures S3A and S3B). This effect was limited to recycling markers, because the early endosomal compartment was unaffected in cells lacking PI3K-C2α (Figure S3C).

To explore whether Rab11 mislocalization was related to its functional state, we analyzed Rab11 activity using a pull-down assay based on the ability of GTP-bound active Rab11 to associate with a fragment of its effector FIP3 (Eathiraj et al., 2006). The specificity of this assay was validated using a constitutively active Rab11 mutant (Rab11 Q70L) as well as Rab5 mutants as negative controls (Figure S3D). Given that Pik3c2a/C0/C0 primary MEFs scarcely proliferated in culture, Rab11 pull-down assays were performed in NIH 3T3 mouse fibroblasts infected with shRNA sequences able to significantly knock down PI3K-C2α (Sh1 and Sh2-3T3; Figure 3C). As shown in Figure 3C, reduction of PI3K-C2α expression levels significantly impaired Rab11 activation. This effect was not dependent on the cell line, because the same result could be repeated in IMCD3 (Sh1 and Sh2-IMCD3) and HeLa cells (Sh1 HeLa; Figure S3E). Overall, these data show a specific involvement of PI3K-C2α upstream of Rab11 localization and activation.

Localization and Activation of Rab11 Require PI3K-C2α-Dependent PtdIns3P Pools

To better characterize the role of PI3K-C2α in the PRE compartment, Pik3c2a/C0/C0 MEFs were engineered to express either a kinase-inactive form of PI3K-C2α (PI3K-C2αKD) or a PI3K-C2α mutant (PI3K-C2αcl) that can produce PtdIns3P, but not PtdIns(3,4)P2 (Posor et al., 2013). Expression of either wild-type or PI3K-C2αcl mutant restored accumulation of Rab11 at the pericentriolar compartment, whereas PI3K-C2αKD did not produce any rescue (Figure 4A). These experiments similarly restored TIR perinuclear localization only in the presence of the PI3K-C2αcl mutant (Figure S4). These data indicate that
PI3K-C2α-derived PtdIns3P is required to properly localize Rab11+ and TfR+ recycling vesicles to the pericentriolar area.

To assess the involvement of PI3K-C2α catalytic activity in triggering Rab11 activation, add-back experiments were performed by transfection of shRNA-resistant PI3K-C2α mutants in Sh1-3T3 cells. Although the wild-type enzyme restored Rab11 activity, a kinase-dead mutant was unable to revert Rab11 inactivation induced by PI3K-C2α downregulation (Figure 4B). Similar to rescue of Rab11 localization, expression of the PI3K-C2αcIII mutant was sufficient to fully restore Rab11 activity (Figure 4B). These results indicate that PI3K-C2α is required and sufficient for production of a pool of PtdIns3P critically needed for Rab11 localization and activation at the PRE.

Loss of PI3K-C2α Impairs Rab8 Activation and Ciliogenesis

At the ciliary base, active Rab11 promotes a cascade of signaling events, including the activation of Rab8 and its localization along the ciliary axoneme, where it controls cilium elongation (Westlake et al., 2011). To test if Rab11 mislocalization and inactivation induced by the loss of PI3K-C2α affected Rab8 function, red fluorescent protein (RFP)-tagged Rab8 was transfected into...
of wild-type cilia, the percentage of Rab8-positive cilia was severely reduced in Pik3c2a \(-/-\) MEFs, thus confirming that PI3K-C2\(\alpha\) promotes Rab8 function (Figures 5A and 5B).

Given the role of the Rab11/Rab8 axis in primary cilium formation (Knödler et al., 2010), the length of primary cilia was measured in wild-type, Pik3c2a \(-/-\), and Pik3c2a \(-/-\) MEFs. Ciliary length was significantly reduced in both Pik3c2a \(-/-\) and Pik3c2a \(-/-\) MEFs (Figures 5C and 5D), with a more pronounced shortening in Pik3c2a \(-/-\) (60%) compared to heterozygous MEFs (23%), indicating a dose-response effect. In agreement, Pik3c2a down-modulation in IMCD3 cells produced a 60% and 70% reduction in Rab8-positive cilia using Sh1 and Sh2, respectively (Figures S5A and S5B), accompanied by a 50% and 60% reduction in ciliary length (Figure S5C), further demonstrating that PI3K-C2\(\alpha\) is involved in Rab11/Rab8-mediated cillum elongation.

Next, rescue of ciliary length was attempted by transfecting Pik3c2a \(-/-\) MEFs with PI3K-C2\(\alpha\) and its mutant variants. Although the add back of a wild-type PI3K-C2\(\alpha\) completely restored ciliary length, the kinase-dead mutant did not (Figure 5E), indicating a crucial role for PI3K-C2\(\alpha\) catalytic activity in cillum elongation. More importantly, reintroduction of the Pik3c2a\(\alpha\)211 mutant fully restored ciliary length (Figure 5E), indicating that the PI3K-C2\(\alpha\)2-dependent production of PtdIns3P is critical for this specific process. Furthermore, to test the involvement of Rab11 in this process, Pik3c2a \(-/-\) MEFs were transfected with either wild-type or constitutively active Rab11. The Q70L constitutively active Rab11 mutant was able to rescue ciliary length in Pik3c2a \(-/-\) MEFs, but neither the wild-type Rab11 nor the constitutively active mutant of a different Rab, Rab5 (Rab5\(^{Q79L}\)), produced any effect (Figure 5F).

### Abnormal Primary Cilia in Pik3c2a-Deficient Embryos

Given the well-established role of primary cilia in embryonic development (Goetz and Anderson, 2010), the embryonic-lethal phenotype induced by the loss of PI3K-C2\(\alpha\) (Yoshioka et al., 2012) was re-evaluated in light of a cillum-related dysfunction. In line with a role in almost ubiquitous primary cilia, detection of the lacZ marker inserted into the mutant allele showed ubiquitous expression in Pik3c2a \(-/-\) embryos at all developmental stages analyzed (Figure S6A). Similarly to what previously reported (Yoshioka et al., 2012), homozygous inactivation of the Pik3c2a locus determined evident growth defects at embryonic day 8.5 (E8.5; Figure S6B) and lethality between E10.5 and E11.5 (Table S1). Consistent with ciliary defects observed in cultured cells, scanning electron microscopy, as well as immunofluorescence, revealed that cilia of the ventral node were frequently bent and swollen and with a 30% and 15% shorter shaft in Pik3c2a \(-/-\) and Pik3c2a \(-/-\) embryos, respectively (Figures 6A and 6B; Figures S6C and S6D). Thus, loss of PI3K-C2\(\alpha\) is compatible with ciliogenesis but impairs primary cilium elongation in vivo.

### PI3K-C2\(\alpha\) Is Required for Hh Pathway Activation

Analysis of Pik3c2a \(-/-\) embryos at E9.5/E10.5 showed that turning and cardiac tube looping were severely disturbed (Figures 6C and 6D; Figures S6E and S6F; Table S1). Furthermore, in situ hybridization failed to detect any expression of Lefty1/2 and Nodal in Pik3c2a \(-/-\) embryos at the 1- to 5-somite stage, whereas these genes were correctly expressed in the ventral...

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**Figure 3. PI3K-C2\(\alpha\) Loss Impairs TIR/Rab11 Localization and Rab11 Activation**

(A) Immunofluorescence with antibody to transferrin receptor (TIR, green) in wild-type and Pik3c2a \(-/-\) MEFs showing that pericentriolar accumulation of this recycling endosome marker is lost in Pik3c2a \(-/-\) MEFs. (B) Immunofluorescence of transfected Rab11-GFP (green) in wild-type and Pik3c2a \(-/-\) MEFs. (C) Pull-down experiment showing the endogenous content of Rab11-GFP in NIH 3T3 cells infected with either a control sequence (empty-pGIPZ; Ctrl) or shRNAs downmodulating PI3K-C2\(\alpha\) (Sh1 and Sh2). Quantification of four independent experiments is provided on the right. Error bars indicate SEM.
node and left lateral plate mesoderm of wild-type embryos at the same developmental stage (Figures 6E and 6F; Table S1), indicating that loss of PI3K-C2α impairs left-right patterning.

Embryonic lethality and laterality defects detected in Pik3c2α−/− embryos resembled the loss of the Hh signal transducer Smo, which depends on membrane traffic to the primary cilium for its function. In the node of 1- to 5-somite Pik3c2α−/− embryos, expression of the Shh mRNA was normally detected (Figure 7A). However, the Shh-dependent translocation of Smo to the shaft of ventral node cilia was severely reduced (Figure 7B; Figure S7A). Similarly, the number of Smo-positive cilia after Hh pathway stimulation was significantly decreased in Pik3c2α-silenced NIH 3T3 cells compared to control cells (Figure 7C; Figure S7B). A reduction of Smo ciliary localization was also obtained by transfecting in wild-type cells a dominant-negative (GFP-Rab11S25N), but not a wild-type, Rab11 (Figure S7C). Conversely, transfection of the constitutively active mutant of Rab11 (GFP-Rab11G70L) (Figures 7D and 7E), as well as the Rab11-activating Pik3c2αS28 mutant (Figure 7E), normalized Smo ciliary localization upon Hh-pathway stimulation in Pik3c2α-deficient cells, demonstrating that the Pik3c2α-Rab11 axis is required for efficient ciliary translocation of Smo.

This suggested that the lack of PI3K-C2α impairs one of the earliest events in Hh signaling, potentially blocking downstream signaling events in the Hh pathway in vivo. In agreement, loss of PI3K-C2α was found to alter the processing of the Gli3 transcription factor, which in 1- to 5-somite Pik3c2α−/− embryos was pre-sent in a significantly higher amount as a repressor (R) short form than as a full-length transcriptional activator form (Figure 7F; Figure S7D). Furthermore, transcripts for Shh downstream targets Ptc1 and Gli1 were reduced by 53% and 57% in Pik3c2α−/− 1- to 5-somite embryos compared to wild-type controls, whereas the Hh unrelated gene FGF8 was equally expressed in both genotypes (Figure 7G). In further agreement with a reduced Hh signaling, Shh-mediated dorsoventral patterning of the neural tube was severely altered in E10.5 Pik3c2α−/− embryos. In particular, expression of Shh-dependent markers of ventral neural cell types, such as Shh, FoxA2, and Nkx2.2, was absent, whereas expression of the Shh-inhibited Pax6 expanded to ventral cells (Figure S7E).

In mutant MEFs, analysis of Shh-induced target gene expression accordingly revealed that the upregulation of Ptc1 and Gli1 after 24 hr stimulation with 100 nM Shh was significantly lower than that of wild-type controls (Figure 7H). A significant reduction in Shh-mediated target gene transcription was also confirmed in Sh1-3T3 cells (Figure 7I). This experimental system was next used to test the role of PI3K-C2α catalytic activity and of Rab11 in Shh pathway activation. Expression of an shRNA-resistant PI3K-C2αS301 mutant showed a significant rescue of both Gli1 and Ptc1 Shh-mediated transcriptional upregulation, indicating that PtdIns3P produced by PI3K-C2α is critically required for Shh signaling pathway activation. Similarly, a constitutively active Rab11Q70L mutant, but not a wild-type Rab11, was able to fully restore sensitivity to Shh (Figure 7I). Collectively, these results establish a critical requirement for PI3K-C2α-mediated production of PtdIns3P for the organization of the perinuclear recycling compartment and for Shh signaling, both in vivo and in cultured cells.

**DISCUSSION**

Four distinct PI3Ks (the three class II α, β, γ enzymes and the class III PI3K) can in principle generate PtdIns3P in vivo (Vanhaesebroeck et al., 2010), but whether each of these enzymes contributes only to a specific pool of PtdIns3P with distinct

**Figure 4. PI3K-C2α-Dependent PtdIns3P Is Necessary for Rab11 Localization and Activity**

(A) Localization of endogenous Rab11 (red) in Pik3c2α−/− MEFs transfected with either wild-type (WT), kinase-inactive (KD), or class III mutated (CIII) PI3K-C2α-GFP (green) showing that only WT and CIII enzymes rescue Rab11 localization. (B) Expression of WT, KD, or CIII PI3K-C2α-GFP in Pik3c2α-silenced NIH 3T3 cells to assess levels of active Rab11 (Rab11-GTP) by pull-down (left) and quantification (right, n = 6). Only PI3K-C2α forms able to produce PtdIns3P can rescue Rab11 activation. Error bars indicate SEM.
intracellular function has long remained obscure. Our results help to elucidate the localization and role of a distinct PtdIns3P pool by demonstrating that PtdIns3P produced by PI3K-C2α is concentrated at the PRE and promotes the correct pericentriolar localization of Rab11+ vesicles as well as the activation of Rab11. This triggers the Rab8-dependent pathway as well as the translocation of Smo to the primary cilium, a process involved in the activation of Shh signal transduction cascade.

Our experiments with fluorescently labeled PtdIns3P probes and anti-PtdIns3P antibodies as well as HPLC analysis showed that in the absence of PI3K-C2α, a small, highly localized PtdIns3P pool was missing. The finding that loss of PI3K-C2α causes reduced levels of PtdIns3P matches what has been reported using an independently generated PI3K-C2α-deficient mouse strain (Yoshioka et al., 2012) as well as silenced cells (Falasca et al., 2007). The small but significant reduction in recycling compartment and suggests that this enzyme plays a specific function at this subcellular location. The finding that PI3K-C2α-derived PtdIns3P controls the Rab11/Rab8 axis further supports this hypothesis. Our results thus indicate that PI3K-C2α has multiple functions at different subcellular locations and that its role at the PRE is distinct from that played, for example, at the plasma membrane (Posor et al., 2013). At the cell surface, PI3K-C2α is located in clathrin-coated pits, where it specifically generates PtdIns(3,4)P2. This lipid then favors Snx9 recruitment, followed by Dynamin association and cleavage of the membrane neck required to form a free vesicle. As a consequence, loss of PI3K-C2α causes a severe impairment of clathrin-dependent endocytosis. A PI3K-C2α mutant (PI3K-C2α CIII) that can only generate PtdIns3P does not rescue this defect (Posor et al., 2013) but does, unexpectedly, fully restore Rab11 localization and activation as well as primary
cilium-dependent Shh signaling. This demonstrates that the activity of PI3K-C2α at the PRE is in principle independent from the endocytic defect. Together with recent work from Posor et al. (2013), our findings suggest that the substrate specificity of PI3K-C2α in vivo may be determined in a compartment-specific manner, e.g., by substrate availability. According to this model, PtdIns serves as a preferential PI3K-C2α substrate at recycling endosomes, which likely contain little PtdIns4P (Krauss and Haucke, 2007), in contrast to the plasma membrane (Hammond et al., 2012). However, we currently cannot rule out the possibility that PI3K-C2α produces a pool of PtdIns(3,4)P2 that is then rapidly converted into PtdIns3P by 4P phosphatases such as INPP4B (Gewinner et al., 2009). The inability to specifically target such phosphatases to the PRE prevented us from examining this possible mechanism of action in detail.

Remarkably, the pool of PtdIns3P generated by PI3K-C2α and associated with the PRE was found to specifically control Rab11 localization and activity, whereas it did not affect localization of the endosomal marker Rab5. PI3K-C2α and Rab11 were found to colocalize but could not be coimmunoprecipitated, suggesting that PI3K-C2α does not directly associate with Rab11. The loss of PI3K-C2α-dependent PtdIns3P generation might thus impair Rab11 activation through other, possibly indirect mechanisms that require further investigation. Nonetheless, our results clearly point to an unexpected epistatic interaction placing PI3K-C2α-dependent PtdIns3P production upstream of Rab11 activation.

In the endosomal compartment, class III PI3K-dependent PtdIns3P synthesis leads to late endosome maturation (Backer, 2008; Stein et al., 2003), whereas our results suggest that PI3K-C2α-dependent PtdIns3P is involved in endosomal recycling. Consistently, the Drosophila homolog of class II PI3Ks, Pi3k68D, produces PtdIns3P required for the exit of vesicles from the endocytic compartment, sorting toward the plasma membrane, and cell protrusion formation (Jean et al., 2012; Velichkova et al., 2010). Furthermore, a previous report indicates that the loss of PI3K-C2α in endothelial cells leads to defective delivery of VE-cadherin to cell junctions, thereby causing impaired assembly of endothelial junctions and disrupted vessel integrity in Pik3c2a mouse mutants (Yoshioka et al., 2012). Considering the established role of Rab11 in the control of cadherin traffic (Lock and Stow, 2005), and in light of the results we present here, this
Figure 7. Defective Hh Signaling in Pik3c2a Mutant MEFs and Embryos

(A) Expression of Shh in the ventral node area (arrows) by in situ hybridization; ventral view. vn, ventral node.

(B) Immunofluorescence analysis of Smoothened (Smo, red) localization at the nodal cilia (acetylated α-tubulin, green) of presomite wild-type and Pik3c2a−/− embryos (n = 5). Bar = 500 nm.

(F) Expression of GLI3 in Pik3c2a−/− and wild-type embryos (n = 5).

Hh signaling and downstream target gene expression in Pik3c2a−/− MEFs and embryos

(A) GLI3 expression in wild-type and Pik3c2a−/− MEFs and embryos (n = 5).

(B) GLI3FL/GIL3R mRNA ratio in wild-type and Pik3c2a−/− MEFs and embryos (n = 5).

(C) GLI3R mRNA expression in wild-type and Pik3c2a−/− MEFs and embryos (n = 5).

(D) Ptc1 mRNA expression in wild-type and Pik3c2a−/− MEFs and embryos (n = 5).

(E) Gli1 mRNA expression in wild-type and Pik3c2a−/− MEFs and embryos (n = 5).

(F) FGF8 mRNA expression in wild-type and Pik3c2a−/− MEFs and embryos (n = 5).

(G) Ptc1 fold induction in wild-type and Pik3c2a−/− MEFs and embryos (n = 5).

(H) Gli1 fold induction in wild-type and Pik3c2a−/− MEFs and embryos (n = 5).

(I) Smo fold induction in wild-type and Pik3c2a−/− MEFs and embryos (n = 5).

(legend continued on next page)
phenotype is conceivably linked to the loss of Rab11 activity, a possibility that needs further assessment. On the other hand, our data clearly show that the epistatic interaction between PI3K-C2α-dependent PtdIns3P production and Rab11 activation at the PRE is required at earlier developmental stages and is critical for Rab8 and Smo translocation to the plasma membrane and, ultimately, for primary cilium signaling functions.

In line with this view, the phenotype of Pik3c2a-null embryos indicates that PI3K-C2α is a major player in cargo delivery to the primary cilium. This is particularly evident at the ventral node, where Smo did not reach the ciliary shaft. Our results with the PI3K-C2α CIII and Rab11Q70L mutants restoring the response to Shh stimulation demonstrate a critical role for Rab11/PRE-dependent traffic in Smo translocation to the cilium. Although part of the ciliary Smo can derive from the endocytic recycling compartment (Kim et al., 2010), another source is provided by lateral transport on the plasma membrane (Milenkovic et al., 2009). Given the Rab11 delocalization/deactivation induced by the loss of PI3K-C2α-derived PtdIns3P, it is possible that the trafficking dysfunction affects the pool of Smo at the plasma membrane as well. In line with this hypothesis, Shh signaling was fully restored in PI3K-C2α-deficient cells by either the PI3K-C2α CIII mutant or the constitutively active form of Rab11.

In agreement with the profound effect on Smo localization and Shh signaling, embryos lacking PI3K-C2α showed a series of developmental abnormalities typically detected in the loss of Hh signaling. Although a previous report suggested that the loss of PI3K-C2α causes lethality because of abnormal endothelial cell function (Yoshioka et al., 2012), several of these phenotypes appeared earlier than vasculogenesis. Phenotypes detected in Pik3c2a−/− embryos do not match those detected in conditions where the primary cilium is absent that show, for example, bilateral expression of Nodal and Lefty and consequent randomization of situs (Huangfu et al., 2003; Murcia et al., 2000; Nonaka et al., 1998) as well as defective cleavage of Gli3 (Huangfu and Anderson, 2005). Consistent with this observation, the loss of PI3K-C2α did not abolish cilia, which are yet shorter and swollen, likely as a consequence of defective trafficking of ciliary components. On the contrary, phenotypes of Pik3c2a−/− embryos largely overlapped with those of embryos lacking Smo or showing defective Hh signal transduction activation. These embryos, for example, show smaller size, no turning, no cardiac looping, complete loss of expression of Nodal and Lefty, and increased Gli3 cleavage (Huangfu and Anderson, 2005; Zhang et al., 2001). These observations suggest that loss of PI3K-C2α does not affect Gli3 cleavage but blocks PtdIns3P as well as Rab11-dependent ciliary targeting of Smo, thus impairing Hh signal activation.

These findings place PI3K-C2α in an epistatic interaction with the Hh signaling machinery and explain a large set of the developmental abnormalities found in Pik3c2a−/− embryos. Our data cannot rule out that the multifaceted functions of PI3K-C2α at the plasma membrane (Posor et al., 2013) and at adherens junctions (Yoshioka et al., 2012) contribute to the in vivo phenotype. Nonetheless, we provide genetic evidence that PI3K-C2α crucially regulates a PtdIns3P-dependent membrane traffic at the PRE that acts upstream of Rab11 localization/activation and promotes Smo ciliary targeting and Shh signaling.

**EXPERIMENTAL PROCEDURES**

**Animal Models**

A lacZ (bacterial β-galactosidase)-neoR cassette was inserted in-frame with the ATG start codon of Pik3c2a via bacterial recombination. Constructs were electroporated in embryonic stem cells and chimeras obtained by standard procedures. Mice were backcrossed for eight generations in the C57Bl/6; wild-type littermates from heterozygous crosses were used as controls. The experiments involving mutant and wild-type mice were carried out according to European Union (86/609/CEE, CE Off J n° L358, 18 December 1986) and institutional animal welfare guidelines and legislation and approved by the local ethics committee.

**MEF Derivation, Culture, and Treatments**

Primary cultures of MEFs were generated from individual E11.5 littermates by trypsinization of eviscerated embryonic bodies and expanded and maintained in Dulbecco’s modified Eagle’s medium supplemented with 10% fetal bovine serum. Early-passage cells (less than four passages) were used for analysis. Starvation and cillum assembly were achieved by 72 hr of serum deprivation. MEFs at 80% confluency were transfected on coverslips in 24-well plates with 0.5 μg of plasmid DNA using the Effectene. For Shh experiments, MEFs and NIH 3T3 cells were plated at near-confluent densities and serum starved for 48 hr prior to treatment to allow ciliation. Rm-Shh-N (R&D Systems) stimulation lasted 24 hr. Other procedures are reported in Supplemental Experimental Procedures.
Plasmids containing shRNA sequences (arrest GAPZ lentiviral shRNAmir) against Mm Pik3c2a were purchased from Thermo Scientific and used to generate lentiviral particles to stably infect NIH 3T3 and IMCD3 cells as described in Supplemental Experimental Procedures. The Sh1 (V2LMM_73461) target sequence was 5'-GGCAAGATATGTTAGCTT-3', and the Sh2 (V2LMM_66190) target sequence was 5'-CAAGTCTTCTTTAACCTCTTT-3'. Cells at early passages after infection (second to third) were used for experiments. Sh1-resistant wild-type, kinase-inactive (KD), and class III (cIII) PI3KC2a were generated by creating three silent mutations in the human pEGFP–PI3KC2a with the QuikChange site-directed mutagenesis kit (Stratagene) using the following primer: 5'-GTTTAAGGTTGGTGAAGATCTTCGCCAGGACATGTTAG-3' and transfected using the Effectene transfection reagent.

**Supplemental Experimental Procedures**

determinants of ciliogenesis. The Sh1 (V2LMM_73461) target sequence was 5'-GGCAAGATATGTTAGCTT-3', and the Sh2 (V2LMM_66190) target sequence was 5'-CAAGTCTTCTTTAACCTCTTT-3'. Cells at early passages after infection (second to third) were used for experiments. Sh1-resistant wild-type, kinase-inactive (KD), and class III (cIII) PI3KC2a were generated by creating three silent mutations in the human pEGFP–PI3KC2a with the QuikChange site-directed mutagenesis kit (Stratagene) using the following primer: 5'-GTTTAAGGTTGGTGAAGATCTTCGCCAGGACATGTTAG-3' and transfected using the Effectene transfection reagent.

**Generation of the Rab11 Binding Domain Probe**

The C-terminal portion of the Rab11 effector FIP3 (Rab11 binding domain) was cloned in pGEX 4T2 vector. Recombinant protein was produced in bacteria (4 hr induction at room temperature), purified (elution 10 mM glutathione, HCl 50 mM, 5 mM MgCl2, 100 mM NaCl) at 80°C. Cells were washed in ice-cold PBS and lysed in 1 ml of MBL buffer (25 mM HEPES [pH 7.5], 150 mM NaCl, 1% Igepal CA-630, 10% glycerol, 25 mM NAF, 10 mM MgCl2, 1 mM EDTA, 1mM sodium orthovanadate, and protease inhibitor cocktail). Supernatant was collected after 15 min centrifugation at 13,000 rpm. A total of 500 μg of protein extract was incubated with 30 μg of recombinant protein coupled with glutathione S-transferase agarose. The reaction mixture was gently rocked for 1 hr at 4°C. Beads were washed four times with lysis buffer. Samples were resuspended in Laemmli buffer for SDS-PAGE and immunoblot analysis. Endogenous content of total Rab11 in cell lysates was measured by loading 50 μg of total extracts in a different gel followed by immunoblot and used to normalize measurements of active Rab11. For quantification analysis, pictures were taken ensuring that intensity was within the linear range and the Quantity One 1-D analysis software (Bio-Rad) was used.

**Statistical Analyses**

Values are reported as the mean ± SEM. Statistical significance was calculated with one-way ANOVA and Bonferroni post hoc tests. One, two, and three asterisks in all the figures indicate significance with a p value <0.05, <0.01, and <0.001, respectively. All the analyses were performed with the software Prism5 (GraphPad Software).

All other procedures are reported in Supplemental Experimental Procedures.

**SUPPLEMENTAL INFORMATION**

Supplemental Information includes Supplemental Experimental Procedures, seven figures, and one table and can be found available at this website online.

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