A role for Phospholipase D in Drosophila embryonic cellularization

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Abstract

Background: Cellularization of the Drosophila embryo is an unusually synchronous form of cytokinesis in which polarized membrane extension proceeds in part through incorporation of new membrane via fusion of apically-translocated Golgi-derived vesicles.

Results: We describe here involvement of the signaling enzyme Phospholipase D (Pld) in regulation of this developmental step. Functional analysis using gene targeting revealed that cellularization is hindered by the loss of Pld, resulting frequently in early embryonic developmental arrest. Mechanistically, chronic Pld deficiency causes abnormal Golgi structure and secretory vesicle trafficking.

Conclusion: Our results suggest that Pld functions to promote trafficking of Golgi-derived fusion-competent vesicles during cellularization.

Background

Embryogenesis in Drosophila melanogaster commences with 13 nuclear divisions in the absence of cytokinesis, generating a syncytium of ~6,000 nuclei located immediately beneath the plasma membrane. Cellularization then ensues, resulting in the synchronous formation of lateral and then basal membranes around each nucleus through rearrangement of the actin cytoskeleton and extensive formation of de novo membrane. It is estimated that a 25-fold increase in plasma membrane is necessary to complete the process. The de novo membrane comes from the resorption of microvilli on the outer surface of the blastoderm [1,2] and the incorporation of ER- and/or Golgi-derived secretory vesicles [3-7].

Genetic screens have identified numerous proteins that regulate cytoskeletal reorganization during cellularization, several proteins of unknown function including SLAM [7] that are required at the leading edge of the extending lateral membrane (the furrow canal), and a plasma-membrane associated component of the exocytic membrane trafficking machinery, the SNARE protein Syntaxin1 (Sx1A, reviewed in Mazumdar and Mazumdar, 2002). Both Sx1A and SLAM enable the fusion of...
incoming membrane vesicles into the expanding membranes. In the absence of SLAM, membrane vesicles accumulate in the apical cytoplasm. However, the accumulating vesicles contain Rab11, a marker for recycling endosomes, suggesting that the incoming Golgi-derived secretory vesicles either first fuse into the apical plasma membrane, and then endocytosis and traffic through the recycling endosomal compartment prior to fusing into the expanding lateral membranes, or traffic directly to the lateral membranes via the Rab11 endosomal pathway [8].

Little is known about the nature of the incoming Golgi-derived secretory vesicles or how their apical trafficking is regulated, aside from the findings that Lava lamp (Lva), a microfilament/microtubule-associated protein that potentially couples the Golgi to the microtubule network [6], and the regulated small GTPase Rho [9,10] which is known for its role in cytoskeletal reorganization and membrane vesicle trafficking, are required for the progression of cellularization. Disruption or brefeldin A-mediated inhibition of exchange factors for the small GTPase ARF1 also inhibits cellularization [6,11].

A protein that facilitates Golgi-derived vesicle production, trafficking, and fusion into target membranes in yeast and mammalian cells is the signaling enzyme Phospholipase D (PLD), which hydrolyzes the membrane phospholipid phosphatidylcholine to yield the lipid second messenger phosphatidic acid [reviewed in [12]]. PA is a pleiotropic lipid that functions in membrane vesicle trafficking by promoting membrane budding from the Golgi complex [13,14], by facilitating exocytic trafficking [15,16], and by promoting vesicle fusion into target membranes [17-19]. In addition, through regulation of phosphatidylinositol 4,5-bisphosphate (PI4,5P2) production, PLD/PA regulates actin cytoskeleton reorganization [20]. Isolation of PLDs from yeast and mammals led to the identification of an eukaryotic PLD family: a Phox homology (PX) domain is noteworthy for a non-conserved "acidic patch" (residues 283–298), and a conserved catalytic domains (CR I-IV), a loop sequence, a hinge region between the two catalytic half-domains, as well as a Pleckstrin homology (PH) domain, conserved catalytic domains (CR I-IV), a loop sequence, a P[4,5]P2-interacting motif, and a conserved carboxy terminus [reviewed in [12]]. The PLD amino-terminus is encoded a 1278 amino acid protein that exhibits ~35% sequence similarity to human PLD2 (Fig. 1B, C), and slightly less to human PLD1. The protein contains the conserved regions characteristic of other members of the eukaryotic PLD family: a Phox homology (PX) domain followed by a Pleckstrin homology (PH) domain, conserved catalytic domains (CR I-IV), a loop sequence, a P[4,5]P2-interacting motif, and a conserved carboxy terminus [reviewed in [12]]. The Pld amino-terminus is noteworthy for a non-conserved "acidic patch" (residues 101–112), whereas the PX domain, which in the mammalian isoforms regulates subcellular localization [16], contains an unusual "basic patch" (residues 283–298). PX domain functioning in Pld may be regulated uniquely by interactions between the acidic and basic patches. Although the central loop region in Pld is in the same position as the loop found in mammalian PLD1, it lacks similarity to it, suggesting that this region may serve as a hinge region between the two catalytic half-domains, as has been shown for the mammalian PLD enzymes [25], rather than to convey structural or functional information. Taken together, the presence of the characteristic PLD domains suggested that Pld should be a bona fide PLD homologue.

Pld encodes a 1278 amino acid protein that exhibits ~35% sequence similarity to human PLD2 (Fig. 1B, C), and slightly less to human PLD1. The protein contains the conserved regions characteristic of other members of the eukaryotic PLD family: a Phox homology (PX) domain followed by a Pleckstrin homology (PH) domain, conserved catalytic domains (CR I-IV), a loop sequence, a P[4,5]P2-interacting motif, and a conserved carboxy terminus [reviewed in [12]]. The Pld amino-terminus is noteworthy for a non-conserved "acidic patch" (residues 101–112), whereas the PX domain, which in the mammalian isoforms regulates subcellular localization [16], contains an unusual "basic patch" (residues 283–298). PX domain functioning in Pld may be regulated uniquely by interactions between the acidic and basic patches. Although the central loop region in Pld is in the same position as the loop found in mammalian PLD1, it lacks similarity to it, suggesting that this region may serve as a hinge region between the two catalytic half-domains, as has been shown for the mammalian PLD enzymes [25], rather than to convey structural or functional information. Taken together, the presence of the characteristic PLD domains suggested that Pld should be a bona fide PLD homologue.

Pld is activated by components of signaling pathways that are important in cellularization

Although yeast PLD is constitutively active (Rose et al., 1995), mammalian PLD is regulated by ARF and Rho GTPases. Accordingly, the report that PLD activity in Dro-
Pld gene and protein structure. A. The Pld genomic locus is shown (42A15), as well as the closely located bicoid-interacting protein 3 (bin3) region (black). The Pld transcript is represented as boxes with exon numbers inside. The coding region is indicated in gray. ESTs representative of different 5' UTR splice isoforms found in BDGP are shown (GH: adult head; RE: embryos 0–22 h; GM: ovaries; LP: larvae and early pupae). B. The Pld domain organization is shown in comparison with mammalian PLD1 and 2. C. Identical and highly similar amino acids in human PLD2 are highlighted in red. (*) indicates the position of the mutant amino acid in the Pld-H1095N allele.
sophila could be stimulated by GTPγS [26] suggested that Drosophila Pld might be regulated similarly. To investigate this, Pld, and a putatively inactive allele of Pld containing a point mutation (H1095N) in the second HKD domain that renders all other known PLD proteins inactive [25], were expressed in mammalian cells.

Mammalian PLD1 exhibits little activity in the absence of cellular stimulation [22]. PMA, a protein kinase C agonist that activates PLD1 through both direct and indirect pathways, potently increases PLD activity. In contrast, only a small increase in activity, reflecting activation of endogenous PLD, is observed when inactive PLD1 (K898R) is overexpressed and assayed [25]. Similarly, Drosophila Pld activity, but not Pld-H1095N activity, significantly increased following PMA stimulation [see supplemental figure in 24].

To determine whether ARF and Rho stimulate Drosophila Pld, we used a cell-based system in which increased levels of the GTPases exaggerate mammalian PLD responses to G-protein receptor-coupled signaling. HEK293 cells expressing the m3 muscarinic acetylcholine receptor [27] were co-transfected with pCAG:Pld and mammalian ARF1, ARF6, or RhoA expression plasmids (Fig. 2A). A small amount of activity was observed in cells overexpressing the inactive Pld-H1095N protein, which derived from the endogenous mammalian PLD1s present in the HEK293 cells. Similar levels of activity are observed after transfection of inactive human PLD1 (hPLD1-K898R) or empty vectors (supplemental figure in citation 24 and data not shown). Wild-type Drosophila Pld was activated by receptor signaling through carbachol stimulation and this was enhanced by overexpression of ARF1, as has been observed for mammalian PLD2 [19]. In contrast, hPLD1 activity was enhanced more readily by overexpression of RhoA. Taken together, these findings demonstrate that Pld is a bona fide PLD and can be regulated by ARF GTPases, which mediate vesicle trafficking and actin cytoskeleton reorganization in many settings.

Pld is maternally and zygotically expressed and peaks prior to cellularization

Many of the proteins required for cellularization are synthesized from maternal transcripts, although a few are transcribed zygotically and peak just prior to or at the onset of cellularization [28]. To determine the temporal sequence of Pld mRNA expression in early embryogenesis, we used quantitative RT-PCR (Fig. 2B). Pld is expressed maternally, since it was observed in the non-fertilized (NF) 0–1 hr after egg deposition (AED) collection. It is also expressed zygotically, based on the dramatic increase observed when zygotic transcription initiates, i.e. between the fertilized 0–1 hr and the 2 hr AED collection. Pld mRNA levels plummeted between 2 and 4 hours AED and increased again modestly around 9 hours AED. Thus, the temporal pattern of Pld expression is consistent with that of genes that have roles in cellularization.

Pld localizes to small cytoplasmic vesicles during cellularization

In some mammalian cell types, PLD1 functions at sites of exocytosis on the plasma membrane to promote the fusion of secretory granules [18,19]. In others, though, PLD1 localizes to the Golgi and/or peri-nuclear vesicles and facilitates budding from the trans-Golgi to generate Golgi-derived secretory vesicles [e.g. [29]], inter-endosomal or ER-to-Golgi trafficking [30], or trafficking of secretory vesicles to the plasma membrane [15]. Determining where Pld localizes would therefore suggest potential roles for its function during cellularization.

In cellularizing embryos, Pld localized to the cytoplasm and/or to cytosolic vesicles of uniform small size evenly distributed between apical and basal regions of the blastoderm (Fig. 3A–D). Pld was not observed to localize specifically to the plasma membrane, membrane furrows, or extending lateral membranes where membrane vesicle fusion takes place. Detection of Pld overexpressed by the NGT40 promoter (Fig. 3C) confirmed that the antisera recognizes Pld protein. When overexpressed, Pld exhibits a preferential localization for the apical region of the cell; the reason for this is not known but is suggestive with respect to the overexpression phenotype discussed below. Pld also did not co-localize with Rab11 as a marker for recycling endosomes (not shown) or the 120-kD Golgi integral membrane protein, p120 (Fig. 3E, F). During early cellularization, Pld partially co-localized with actin at the sites where cytoskeletal reorganization and membrane formation are taking place (Fig. 3G, H). This overlap ceased by the end of cellularization, as indicated by the clearing of Pld within the established actin tracks (Fig. 3I, J). Taken together, this pattern of localization would be consistent with a role for Pld in translocation of fusion-competent Golgi-derived vesicles to the embryonic cortex or regulation of the actin cytoskeleton, although roles in budding or fusion are not ruled out.

To compare the localization of Drosophila Pld to its mammalian homologs, Drosophila Pld was expressed in (mammalian) COS7 cells. Pld localized to cytoplasmic vesicles of varying size and to the plasma membrane (Fig. 4). The vesicular staining did not co-localize with the ER-marker calreticulin or the recycling endosome proteins Rab11 and the transferrin receptor (data not shown). The pattern of subcellular localization of dPld was intermediate between that of mammalian PLD1 and PLD2: PLD1 localized to peri-nuclear vesicles, whereas PLD2 localized to the plasma membrane. The unique aspects of the Drosophila PLD pattern of localization could ensue from the
Figure 2
Pld exhibits classical Phospholipase D activity regulated by signaling pathways and peaks in expression prior to cellularization. A. HEK-293 cells stably expressing the m3 muscarinic acetylcholine receptor were co-transfected with pCGN:Pld and with ARF1, ARF6 or RhoA. After 24 hours, the cells were stimulated with carbachol, an activator of G-protein coupled muscarinic receptors, and assayed in vivo for PLD activity. The assay is based on the ability of PLD to use primary alcohols, such as butanol, in place of water, in the PC hydrolysis reaction. This leads to the production of phosphatidyl-alcohol (-butanol, PtdBut). Assaying PtdBut is preferable to attempting to quantitate PA, since PA is highly labile and turns over quickly, whereas PtdBut is relatively inert, accumulates over the course of the 30' assay, and thus provides an estimate of the total amount of PLD activity that took place during the assay period. See citation 41 for assay details. The data represent means of triplicate measurements. Standard deviations averaged 3–5%. The experiment is representative of three independent experiments. *, significant difference, p < 0.05.

B. Temporal expression of endogenous Pld as analyzed by quantitative PCR. Embryonic stages are given in hours after egg deposition (AED). 0–1 NF, non-fertilized eggs between 0 and 1 hour AED. Cellularization occurs between 2 and 3 h AED during the 14th mitotic cycle. Pld cDNA concentration is expressed as a ratio of Pld to rp49, a ribosomal RNA expressed at constant levels.
Figure 3
Pld localizes to cytoplasmic vesicles during cellularization. A. Pld localization in heat-fixed y^1 w^67c^23 control embryos using an affinity-purified anti-Pld antisera (sagittal image generated at or near the embryonic equator). a, apical; n, nuclei; b, basal. B. Affinity-purified pre-immune serum does not generate a visible signal. C. Pld localization in transgenic lines overexpressing Pld as a maternal transcript using NGT-GAL4. D. Pld localization in transgenic lines expressing Pld RNAi. E, F. Pld does not co-localize with p120-marked Golgi vesicles (red) in wild-type (sagittal section) or Pld; NGT40+A embryos (surface view). G-J. Cellularizing embryos immunostained with anti-Pld antisera (green), monoclonal anti-actin (red), and Propidium iodide (blue). Pld in early cellularizing embryos (G, H) co-localizes with actin at sites where adjacent cells are forming junctions (inset). Later in cellularization (I-J), Pld does not localize to sites at which actin tracks have been assembled (inset).
non-conserved regions within its amino-terminus and/or loop region.

**Pld overexpression during early embryogenesis reduces viability**

The effect of increasing PLD activity during cellularization was examined using the P[Gal4-nos.NGT] transgenic binary expression system [31] to deliver wild-type or catalytically-inactive (H1095N) Pld at the start of embryogenesis. Immunostaining indicated that both isoforms were expressed at elevated levels, particularly in the apical region of the newly forming cells (Fig. 3C). Pld overexpression did not affect Golgi localization or morphology (Fig. 3E, F and data not shown). Embryonic lethality averaging 30.4% (in three experiments) was observed for all P[UAS-Pld] transgenic lines driven by the strong GAL4 driver P[Gal4-nos.NGT]40; P[Gal4-nos.NGT]A. Embryonic lethality was dependent on expression level since no lethality was observed with the weaker P[Gal4-nos.NGT]11 driver. In addition, lethality was not observed for the strong Gal4-driven overexpression of the catalytically-inactive P[UAS-Pld-H1095N] transgene, indicating that the phenotype ensues from increased PLD activity and thus PA production, rather than from a non-enzymatic consequence of PLD overexpression.

**Genetic ablation of Pld reduces viability in early embryogenesis**

The insertional, or ends-in, homologous recombination method [32] was used to mutate the Pld gene. For gene targeting (Fig. 5), a 4.8-kb genomic fragment of Pld beginning in exon 2 at amino acid 149 of the ORF and terminating 18 amino acids before the stop codon in exon 9 was engineered to carry a recognition site for the I-SceI endonuclease and two stop codons within exon 5. Seven independent Pld targeting events were recovered after screening 60,000 flies. Of the 7 events, only 3 had the expected genomic locus organization (Fig. 5) and the presence of the two introduced stop codons. The remaining 4 repaired the inserted DNA sufficiently distal to the insertion site that one or both stop codons were converted back to the wild-type sequence. The recombination event generated a tandem duplication of the Pld locus resulting in two truncated (C-terminally truncated and N-terminally truncated, respectively) Pld genes. Analysis of Pld mRNA from Pldnull mutant flies by RT-PCR indicated that only the first gene (which contained a functional promoter) was expressed, and trans-splicing did not occur (not shown). The corresponding truncated Pld protein generated would be 1001 amino acids long and contain only 1 HKD domain, and thus would be inactive, since as demonstrated previously [25], two functional HKD domains are necessary for PLD activity. Expression of this truncated protein should not result in phenotypic consequences, since, as described above, overexpression of a presumptively inactive Pld allele (mutated in the second HKD motif (Pld-H1095N)) did not increase PLD activity in tissue culture systems and did not cause embryonic lethality.

Pldnull flies could be recovered as homozygotes, but displayed a 56% ± 6% reduction in expected viability in comparison to heterozygous flies as determined by mating heterozygous null flies (Pldnull/CyO × Pldnull/CyO; CyO is

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**Figure 4**

*Drosophila Pld recombinant protein localizes to cytoplasmic vesicles and the plasma membrane in COS7 cells.*

COS7 cells were transiently transfected with either pCGN:dPld, pCGN:mPLD1, or pCGN:mPLD2. The cells were immunostained with a rat monoclonal anti-HA antibody (3F10). mPLD1 localized to peri-nuclear vesicles, while mPLD2 was predominantly localized to the plasma membrane. *Drosophila* Pld staining marked intracellular vesicles and the plasma membrane.
the curly wing balancer). This result indicates that zygotic transcription of \textit{Pld} is important, since the maternal \textit{Pld} mRNA was present. A larger decrease in viability (90\% ± 3\% reduction) was observed when homozygous null females, collected from the heterozygous stock, were crossed with heterozygous males (\textit{Pldnull/Pldnull} × \textit{Pldnull/CyO}), indicating that maternal \textit{Pld} is also important. It should be noted, however, that the effects on viability progressively decreased after maintaining the line for several generations as a homozygous \textit{Pldnull} stock (data not shown), suggesting that compensatory mechanisms that offset the loss of \textit{Pld} become activated or selected for.

To determine the embryonic developmental stage at which the \textit{Pldnull} flies were arresting, we monitored the progressive development of living \textit{Pldnull} embryos. Of 53 \textit{y¹\textit{w67c23}} control embryos, almost all progressed normally through cellularization and gastrulation (the 3 that did
not were either unfertilized or arrested early in mitosis. Of 78 \textit{Pld}null embryos, almost half (37) exhibited visible morphological defects prior to or during cellularization, and 15 displayed abnormal gastrulation (which could have ensued from abnormalities initiated during cellularization). However, it should be noted that of the 15 that exhibited abnormal gastrulation (as characterized by a thin and shortened germ band), 3 were still able to progress to the larval stage and hatch. Taken together, these results indicate that \textit{Pld} facilitates the normal progression through early embryogenesis.

### \textbf{Pld deficiency results in Golgi and/or Golgi-derived membrane vesicles increased in size}

To examine potential mechanisms responsible for the abnormal progression of \textit{Pld}null embryos during embryogenesis, we assessed the subcellular localization of Lava Lamp (Lva), a Golgi-associated protein essential for cellularization that facilitates interaction between Golgi secretory vesicles and the cytoskeleton [6]. In wild-type cellularizing embryos, Lva localized to Golgi vesicles, particularly in basal regions of the forming cells (Fig. 6A, B). In contrast, \textit{Pld}null embryos exhibited increased fluorescence intensity (1.7±0.2-fold per similar vesicle area) upon staining with anti-Lva antibody, and the Lva-stained Golgi vesicles were increased in size (8.3 ± 1.9-fold). This observation of abnormal Golgi structure suggests a role for \textit{Pld} in facilitating the Golgi fissioning required to generate the vesicles destined to be transported via Lva to the embryonic cortex.

### \textbf{Pld deficiency results in accumulation of Neurotactin-containing vesicles at or near the plasma membrane}

The transmembrane protein Neurotactin (Nrt), which is synthesized de novo during cellularization, traffics through the secretory pathway and translocates to the apical plasma membrane via Golgi-derived vesicles, following which it undergoes endocytosis and re-fusion into lateral membranes via the recycling endosomal pathway. Vesicles bound for lateral membranes may also travel there via the Rab11 endosomal pathway without first trafficking to the plasma membrane [8]. In \textit{Pld}-deficient embryos, Nrt-containing vesicles were observed to accumulate throughout the apical cytoplasm and in a large subapical/apicolateral compartment, in large part juxtaposed to the plasma membrane (compare Fig. 6D with 6C). A similar appearing pattern of mis-localization is observed in \textit{shibire} mutant embryos [8]; in this setting, the aggregating vesicles co-localize with Rab11, suggesting that they are composed of recycling endosomes unable to fuse into lateral membranes. \textit{shibire} is a dynamin isoform required for budding from recycling endosomes. However, the aggregating vesicles in \textit{Pld}-deficient embryos did not co-localize with Rab11 (data not shown), nor were the Rab11-containing vesicles observed to be altered significantly in localization or morphology (Fig. 6E, F). Moreover, the Nrt vesicle aggregates did not co-localize with a marker for Golgi (data not shown). Thus, Nrt appears to exit the Golgi into secretory vesicles and translocate apically, but then is not inserted into the plasma membrane, thereby suggesting that, in addition to its role in Golgi vesicle fission, \textit{Pld} mediates vesicle fusion at the plasma membrane during cellularization. As another pos-

![Figure 6](http://www.biomedcentral.com/1471-213X/6/60)
sibility, Pld may mediate fusion of post-Rab11 vesicles into the lateral membrane.

**Cellularization is delayed in Pldnull embryos**

Since the Pldnull mutants displayed cellularization defects consisting of altered Golgi structure and post-Golgi trafficking, we examined whether these defects were detrimental for cellularization. Single embryos from an outcrossed Pldnull stock were analyzed using video microscopy starting at pole cell formation through the end of cellularization. Three of the six Pldnull embryos exhibited a substantial delay during cellularization (Table 1), taking an average of 28% longer to complete the process (87 minutes versus 68 minutes), and a fourth was significantly delayed. Two of the six Pldnull embryos were wild-type in their progression through cellularization, suggesting incomplete penetrance of the phenotype. Nonetheless, the delay in cellularization in the majority of Pld-deficient embryos indicates that the presence of Pld facilitates the progression of cellularization in a timely manner.

**Discussion and Conclusions**

The role of lipid signaling enzymes in the regulation of membrane trafficking during cellularization is largely unknown. We report here that Phospholipase D plays an important role in Drosophila embryonic cellularization, using the experimental approaches of overexpression and gene targeting. The severity of the gene targeting phenotype is potentially understated since Pldnull flies appear to activate compensatory mechanisms: heterozygous null flies, when first crossed, generate few homozygous offspring (10% of the expected frequency), but continued breeding of the surviving null flies yields a stock with wild-type levels of viability. Such compensation has been observed for lipid signaling [e.g. [33,34]] and in general in other Drosophila systems [35,36]. Supporting the possibility of compensatory mechanisms, preliminary experiments using PLD RNAi on wild-type embryos, a setting in which compensatory mechanisms would not easily be activated, elicited in a nearly complete (82%) inhibition of cellularization (data not shown). Although there is only one PLD isof orm in Drosophila, which rules out redundant coverage via a closely-related family member, PA levels could be supported through at least three other mechanisms – increasing activity of DAG Kinase (which generates PA through phosphorylation of DAG), decreasing activity of phosphatidic phosphohydrolases (which mediate the reverse reaction), or increasing activity of LysoPhosphatidic Acetyl Transf erases (which convert Lyso-Phosphatidic acid to PA). A question for the future will be to determine if any of the multiple members of these large gene families are altered in expression in Pld-deficient flies.

Only recently has the importance of targeted membrane secretion in cellularization been recognized. Lecuit and Wieschaus (2000) showed that membrane growth during cellularization occurs in large part through the regulated mobilization of ER- and/or Golgi-derived membrane populations that insert into precise locations along the plasma membrane. We report here that on a gross level, loss of Pld impairs progression through cellularization. Presumably underlying this observation, two consequences of Pld deficiency on vesicle trafficking were observed: altered morphology of the Golgi apparatus, from which secretory vesicles are derived, and diminished insertion of the secretory vesicles into the apical and apicolateral plasma membrane.

| Embryo   | Total time from cellularization start to pole cell migration (min:sec) | increase in time required for cellularization |
|----------|---------------------------------------------------------------------|---------------------------------------------|
| yw #1    | 65:49                                                               |                                             |
| yw #2    | 68:09                                                               |                                             |
| yw #3    | 71:15                                                               |                                             |
| Pldnull #1| 69:42                                                               | 1.9%                                        |
| Pldnull #2| 86:26                                                               | 26.4%                                       |
| Pldnull #3| 69:30                                                               | 1.6%                                        |
| Pldnull #4| 88:33                                                               | 29.4%                                       |
| Pldnull #5| 75:12                                                               | 9.9%                                        |
| Pldnull #6| 86:21                                                               | 26.2%                                       |

The embryonic development of single embryos from the time of pole cell formation to the start of gastrulation was recorded using a stereomicroscope with an attached video recorder. The videos were analyzed from the start of cellularization (marked by initial membrane invagination and pole cell formation) to the beginning of pole cell migration and gastrulation. Pldnull embryos were collected from cages set up with homozygous Pldnull males and females from an outcrossed heterozygous Pldnull stock.
PLD has been reported to play roles in vesicle trafficking in other organisms. In yeast, PLD is required to form the prospore membrane during meiosis via fusion of secretory pathway-derived vesicles [17] and for the generation of secretory vesicles in mutant strains lacking Sec14p [33]. In mammalian cells, PLD1 facilitates secretory vesicle generation [14] and fusion into the plasma membrane [18], whereas PLD2 has been proposed to facilitate endocytosis [37]. The results we describe here would be consistent with the roles reported for PLD1 and for yeast PLD, and establish the first animal model that can be used to explore the function of PLD in vivo. We have described one other use of this mode, in the exploration of the role of PLD in phototransduction [24].

Of note, Pld does not exhibit steady-state localization in the Golgi. However, studies on mammalian PLD have suggested that both PLD1 and PLD2 cycle through a succession of subcellular membrane locations including the plasma membrane, recycling and outward bound endosomes, and the Golgi, and that they undertake functional roles at multiple steps during the transit [16,38]. The steady-state pattern of localization reflects just the slowest step in the cycle, which differs in different cell types or as a consequence of the initiation of signaling events. Our observation that Pld exhibits both vesicular and plasma membrane localization in mammalian cells (Fig. 4) suggests that it too is likely to function at multiple subcellular sites under appropriate circumstances.

Future questions include how Pld is activated during cellularization and how alterations in Pld activity change the morphological structure of the Golgi. Numerous questions remain as to how fusion-competent vesicles are generated from the Golgi and undergo translocation to sites of incorporation into the extending membranes. Further exploration of the role of Pld in this process should lead to new insights.

Methods

General reagents

Phospholipids were purchased or prepared as described [27]. All other reagents were obtained from standard sources.

Plasmid construction

The pCGN:Pld and pCGN:Pld-H1095N mammalian expression plasmids were constructed using the coding region of cDNA GH07346. From the above plasmids, the resulting 3.9-kb Pld coding region was excised using Xba I and BamH I and ligated into the EcoRI/Bgl II sites of pUAST-T (Brand and Perrimon, 1993) to generate pUAS:Pld and pUAS-Pld-HN, respectively. Prior to ligation, the Xba I and EcoRI I sites were filled in.

Homologous recombination targeting construct P{w<sup>hs</sup>,Pld*}. See Fig. 5 for schematic. A BamH I fragment of BAC clone RCP198.08AII that contained Pld was inserted into pCaSpeR4. A 4.8-kb BamH I-Sac I fragment containing Pld exons 2–9 was cloned into pBS(Not) (gift of K. Golic, U. Utah). Two mutated pBS(Not): Pld plasmids, each containing a different stop codon in exon 5 (at Pld amino acids 616 and 1002, respectively) were generated using QuickChange. Pld fragments were generated from both plasmids by PCR. Each fragment encoded a different stop codon, terminated at the exon 5 end in a phosphorylated half I-SceI recognition site (ttacctca or cagggtta, respectively), and contained either Not I or Acc65 I at the opposite end. The fragments were digested with either Not I or Acc65 I and inserted as a 3-way ligation into a similarly restricted pIV2 vector (gift of K. Golic). The blunt ligation of the half-I-SceI sites generated the full 18-bp site.

Fly strains and crosses

Drosophila melanogaster strains were maintained on standard cornmeal/yeast/sugar and agar media. Transgenic lines carrying P{UAS-Pld}, P{UAS-Pld-H1095N}, or P{w<sup>hs</sup>,Pld*} were generated by embryo transformation of y<sup>1</sup>wo<sup>72</sup>23 flies [39]. Flies carrying the P[GALA-nos.NGT]<sup>11</sup> [31] and the P[GALA-nos.NGT]<sup>40</sup>; P[GALA-nos.NGT]<sup>A</sup> [40] drivers were used for viability assays as described by Wheeler et al. (2002). Three different P{UAS-Pld} and P{UAS-Pld-H1095N} insertion lines were used. All counts were performed in duplicate at 25°C. For homologous recombination experiments, targeting was performed using a single P{w<sup>hs</sup>,Pld*} donor on the third chromosome. P{w<sup>hs</sup>,Pld*} virgin females were crossed to w (v); P{<sup>ry</sup>77, 70FLP} P{<sup>v</sup>77, 70I-SceI}/TM3, Ser (gift of K. Golic), generating P{<sup>ry</sup>77, 70FLP} P{<sup>v</sup>77, 70I-SceI}/P{w<sup>hs</sup>,Pld*} progeny. Eggs were collected for 24 hours and heat shocked between 24 and 48 hours for 1 hour at 38°C. After hatching, the flies carrying one copy of P{w<sup>hs</sup>,Pld*} and 70FLP and 70I-SceI were crossed to a 70FLP line that has strong constitutive expression of FLP (gift of K. Golic). Flies were screened for w non-mosaic eyes, followed by translocation of the <sup>hs</sup> from the third to the second chromosome by crossing to Balancer stocks. Outcrosses against wild-type strains were performed prior to functional analyses.

Molecular analysis of the Pld<sup>null</sup>

Genomic DNA was extracted using standard protocols. Southern blot hybridization was performed using the probes shown in Fig. 5. Allele specific PCR and sequencing were performed to assay for the presence of the introduced stop codon mutations.
Quantitative PCR
Total RNA was extracted from Canton-S embryos at different embryonic stages using the StrataPrep Total RNA Miniprep Kit (Stratagene). cDNA was synthesized using the First Strand cDNA Synthesis Kit for RT-PCR (AMV) (Roche). Quantitative PCR measurements were performed using a LightCycler and the LightCycler Faststart DNA Master SYBR Green I kit (Roche). Primers used for the amplification of Pld cDNA were 5’-agaggccagagcatgtagag (exon 2) and 5’-cgtgtgtcagatgg (spans the exon 2-exon 3 boundary). Primers 5’-tagggcacaagctggaa and 5’-tctcttgctcgcctga were used for the amplification of rp49. Samples were run in triplicate and standardized using Pld and rp49 plasmid DNA.

PLD assays
PLD cDNAs were transiently overexpressed in COS7 or HEK-293 cells using the mammalian expression vector pCGN. In vivo assays were performed as described previously [41].

Immunological techniques and confocal microscopy
A 405-bp Pld cDNA fragment corresponding to the amino-terminal 135 amino acids of the Pld protein was cloned into the pET32a bacterial expression vector (Novagen). The resulting His-Trx-Pld fusion protein was purified on Ni⁺-resin and used to immunize rabbits (Covance). The antisera was antigen affinity-purified. An anti-Pld peptide antisera directed against the C-terminus (Covance). The antisera was antigen affinity-purified. An anti-Pld peptide antisera directed against the C-terminus (Covance). The antisera was antigen affinity-purified. An anti-Pld peptide antisera directed against the C-terminus (Covance). The antisera was antigen affinity-purified. An anti-Pld peptide antisera directed against the C-terminus (Covance). The antisera was antigen affinity-purified. An anti-Pld peptide antisera directed against the C-terminus (Covance). The antisera was antigen affinity-purified. An anti-Pld peptide antisera directed against the C-terminus (Covance). The antisera was antigen affinity-purified. An anti-Pld peptide antisera directed against the C-terminus (Covance). The antisera was antigen affinity-purified. An anti-Pld peptide antisera directed against the C-terminus (Covance). The antisera was antigen affinity-purified. An anti-Pld peptide antisera directed against the C-terminus (Covance). The antisera was antigen affinity-purified. An anti-Pld peptide antisera directed against the C-terminus (Covance). The antisera was antigen affinity-purified. An anti-Pld peptide antisera directed against the C-terminus (Covance).

Transfection and immunofluorescent staining of COS7 cells
COS7 cells were maintained, transfected, and immunostained using standard protocols. Primary antibodies included: rat anti-HA (1:300), rabbit anti-HA (1:100), rabbit anti-calreticulin (1:200, StressGen), and mouse anti-transferrin receptor (1:150). Fluorescent secondary antibodies were from Molecular Probes.

Embryonic staging for the Pldnull mutant
γΔμ67c23 and Pldnull mutant flies were allowed to lay eggs on apple juice plates for 1 hour. The plates were then aged for 1.5 hours, following which the eggs were transferred to new plates and covered with halocarbon oil 57. The embryos were staged, followed through the initial stages of gastrulation using a dissecting microscope, placed at 25°C for 24 hours, and analyzed for the presence of larva.

Video microscopy on homozygous Pldnull embryos from the outcrossed Pldnull stock
The Pldnull stock was outcrossed for five generations and then balanced over CyO. Homozygous Pldnull embryos were collected from a cage set up with homozygous (straight-winged) males and females collected from the heterozygous Pldnull stock. Embryos prior to pole cell formation were transferred to a drop of halocarbon oil57 on a Petriperm dish and cover slipped. Embryonic development from the time of pole cell formation to the start of gastrulation was recorded using a stereomicroscope with an attached video recorder.

Authors' contributions
ML generated the Pld antisera, performed the viability, videomicroscopy, western blot analysis, Pld immunolocalization and marker analyses, generated the transgenic rescue strain, and co-drafted the manuscript. HJ generated the Pldnull strain using homologous recombination, performed viability and videomicroscopy analyses, and co-drafted the manuscript. SJ performed the RNAi experiments and analysis. IC cloned the Pld gene. OR performed 5' RACE to obtain full length cDNA clones, performed transcriptional initiation analysis, and examined IRES activity of the 5' untranslated region. VO performed the RT-PCR analysis of embryonic expression. YMA and GD analyzed the biochemical activity of Pld in mammalian cells. JPG and MAF made substantial contributions to the conception and design of the project, the interpretation of the data, and the subsequent revision of the manuscript for publication. All of the authors reviewed the final version of the manuscript and gave approval for it.

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