RESEARCH ARTICLE

Phosphatidylserine synthase regulates cellular homeostasis through distinct metabolic mechanisms

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Abstract

Phosphatidylserine (PS), synthesized in the endoplasmic reticulum (ER) by phosphatidylserine synthase (PSS), is transported to the plasma membrane (PM) and mitochondria through distinct routes. The in vivo functions of PS at different subcellular locations and the coordination between different PS transport routes are not fully understood. Here, we report that Drosophila PSS regulates cell growth, lipid storage and mitochondrial function. In pss RNAi, reduced PS depletes plasma membrane Akt, contributing to cell growth defects; the metabolic shift from phospholipid synthesis to neutral lipid synthesis results in ectopic lipid accumulation; and the reduction of mitochondrial PS impairs mitochondrial protein import and mitochondrial integrity. Importantly, reducing PS transport from the ER to PM by loss of PI4KIIIα partially rescues the mitochondrial defects of pss RNAi. Together, our results uncover a balance between different PS transport routes and reveal that PSS regulates cellular homeostasis through distinct metabolic mechanisms.

Author summary

Phosphatidylserine (PS), a membrane phospholipid synthesized in the endoplasmic reticulum (ER) by the enzyme phosphatidylserine synthase (PSS), is transported to the plasma membrane (PM) and mitochondria through different paths. The cellular functions of PS at different places in the cell and the mechanisms that coordinate the different PS transport paths are not fully understood. Here, we identified that PSS regulates cell growth, lipid storage and mitochondrial function in the fruit fly larval salivary gland. We showed that loss of pss function has three effects: (1) reduced levels of PS lead to reduced levels of plasma membrane Akt, a key component in the insulin pathway, which is important for cell growth; (2) it causes a shift from phospholipid synthesis to neutral lipid synthesis, which results in excess lipid accumulation; and (3) it reduces the level of mitochondrial PS, which impairs mitochondrial protein import and mitochondrial morphology. We also found that reducing the transport of PS from the ER to PM partially rescues the mitochondrial defects caused by loss of pss function. Together, our results reveal...
Introduction

Phospholipids make up the membranes that separate cells from extracellular environments and enclose subcellular compartments. Besides their structural role in membranes, phospholipids and their modification products also have specific intracellular and/or intercellular roles in many cellular processes [1]. The synthesis and the subcellular distribution of phospholipids are important for their function.

Phosphatidylserine (PS) is synthesized in regions of the endoplasmic reticulum (ER) called MAMs (mitochondria-associated membranes) [2, 3], and is then imported into mitochondria for phosphatidylethanolamine (PE) synthesis by mitochondrial-localized phosphatidylserine decarboxylase [4], or transported to the plasma membrane (PM) [5, 6]. In the PM, PS mainly resides in the inner leaflet, and loss of this asymmetry acts as an “eat me” signal to trigger apoptotic cell death [7].

In mammals, PSS1 and PSS2 utilize phosphatidylcholine (PC) and PE, respectively, as substrates to synthesize PS [8–12], while in yeast, CHO1 uses CDP-diacylglycerol (CDP-DAG) as the precursor [13, 14]. In mammals, ex vivo alteration of the expression of PS metabolic enzymes is the major approach to revealing the functions of PS and the enzymes related to its metabolism. However, disturbing one enzymatic reaction may cause differential changes in the levels of substrate, product and product-derived metabolites. It is hard to tease apart the specific contribution of individual metabolite changes in vivo. Moreover, there is strong redundancy and compensation of PS metabolic pathways in mammals. For instance, in mice, deficiency of PSS1 or PSS2 is viable, while double deficiency of PSS1 and PSS2 is embryonic lethal [9, 10]. Therefore, although extensive ex vivo studies have revealed many functions of PS at different subcellular locations [15–17], the detailed underlying mechanisms and in vivo functions of PS remain to be fully understood.

Besides the metabolic enzymes, the intracellular lipid trafficking routes are also important for the function of phospholipids [18–22]. From the ER, PS is transported to the PM by oxysterol-binding protein (OSBP) family proteins utilizing the phosphatidylinositol 4-phosphate (PI4P) gradient between the ER and PM, which is generated by PI4KIIIα, also known as PI4Kα or Sst4 [23, 24]. PS can also be imported into the mitochondria through ER-mitochondrion connections and the Ups2-Mdm35 complex [25–29]. How the cell coordinates different PS transport routes remains to be investigated.

In this study, we identified the sole Drosophila phosphatidylserine synthase, PSS, from an in vivo RNAi screen for genes affecting lipid storage. Besides an ectopic lipid storage phenotype, pss RNAi caused defects in cell growth and mitochondrial integrity, including mitochondrial protein import. We reveal distinct metabolic causes of these phenotypes and, more importantly, we show that there is a balance between PS transport from the ER to the PM and from the ER to mitochondria.

Results

Loss of CG4825 reduces cell size and causes ectopic lipid storage in Drosophila salivary gland

We previously performed an RNAi screen in Drosophila 3rd instar larval salivary gland and fat body for aberrant lipid storage by using pumpless-Gal4 (ppl-GAL4) as a driver to achieve
specific RNAi expression in salivary gland and fat body[30]. In the screen, dissected 3\textsuperscript{rd} instar larval salivary glands were stained with the neutral lipid dye BODIPY or Nile red. We found that ppl-GAL4-driven RNAi knockdown of CG4825 (ppl>CG4825 KK105709 RNAi) causes ectopic lipid accumulation in salivary gland and reduces salivary gland size compared with the ppl-GAL4 control or control RNAi group (ppl>control RNAi) (Fig 1A and 1B and S1 Fig). The reduced organ size could be the result of reduced cell number and/or decreased cell size. We found that the salivary gland cell number is not changed in CG4825 RNAi (S1 Fig), while the cell size is reduced significantly (S1 Fig). This suggests a defect in cell growth but not cell proliferation. Compared with the control, RNAi dramatically reduces the mRNA level of CG4825 in 3\textsuperscript{rd} instar larval salivary gland (Fig 1C), confirming the knockdown effect.

Three other independent ppl-GAL4-driven CG4825 RNAi lines (CG4825\textsuperscript{G02753}, CG4825\textsuperscript{NIG4825R-1} and CG4825\textsuperscript{NIG4825R-3}) resulted in similar phenotypes, validating the specific effects of CG4825 knockdown (S1 Fig). Because ppl-GAL4 is highly expressed in both fat body and salivary gland, we also used the salivary gland-specific driver AB1-GAL4 to knock down the expression of CG4825 in the salivary gland but not the fat body. Similar to ppl-GAL4-mediated RNAi knockdown, AB1>CG4825\textsuperscript{KK105709} RNAi increases salivary gland lipid storage and reduces cell size (Fig 1D).
CG4825 RNAi driven by ubiquitously expressed tub-GAL4 leads to lethality at the 1st instar larval stage, which indicates that CG4825 is essential for viability. Besides these RNAi lines, we also examined CG4825 mutants. CG4825\textsuperscript{M101234} is a loss-of-function mutant of CG4825 (Fig 1E), and the homozygous CG4825\textsuperscript{M101234} mutation is lethal during the 1st instar larval stage, which precludes us from directly examining the ectopic lipid phenotype in the salivary gland. CG4825\textsuperscript{KG06018} is a hypomorphic allele of CG4825 with a transposon element inserted into the transcription start site (Fig 1E). The CG4825 transcription level in CG4825\textsuperscript{KG06018} salivary gland is reduced to about 20% of that of wild type (Fig 1F). Importantly, similar to CG4825 RNAi, there are ectopic lipid droplets in CG4825\textsuperscript{KG06018} 3rd instar larval salivary gland cells (Fig 1G). Together, these results demonstrate the tissue-autonomous function of CG4825 in regulating cell growth and lipid storage.

CG4825 encodes the sole Drosophila phosphatidylserine synthase (PSS)

CG4825 contains a phosphatidylserine synthase (PSS) domain and is conserved from yeast to mammals based on protein sequence alignment (Fig 1E and Fig 2A). From bacteria to mammals, there are different PSS proteins for the synthesis of PS from different substrates (Fig 2B). CG4825 is the only PSS domain-containing protein in Drosophila, so we refer to it as PSS hereafter. Drosophila PSS is similar to mammalian PSS1 in the phylogenetic tree (Fig 2A). It has been reported that PSS activity is required for embryonic viability in mice [9, 10]. This is consistent with the lethal phenotype in pss\textsuperscript{M101234} mutants and the global knockdown of pss, and suggests that Drosophila PS synthesis is mainly through pss.

To determine which substrate is used by Drosophila PSS, we pulse-traced NBD-labeled PA (NBD-PA), PC (NBD-PC) and PE (NBD-PE) in Drosophila S2 cells and used thin layer chromatography (TLC) to examine their conversion. The NBD-labeled PA was converted to PE, PC and PS (Fig 2C). Labeled PS appeared later than PE and PC, which suggests that it may be derived from PE or PC. Treating cells with NBD-PC did not yield any labeled PS (Fig 2C), which indicates that S2 cells may not be able to convert PC to PS. However, NBD-PS did appear after the cells were pulse-labeled with NBD-PE (Fig 2C). Together, these results suggest that PA can be converted to PE and subsequently to PS in Drosophila S2 cells.

We further analyzed whether RNAi of pss affects the level of PS in vivo. We dissected wild-type and pss RNAi salivary glands and measured the levels of PS and other lipids through lipidomic profiling. As expected, the level of PS is dramatically reduced in pss RNAi to around 20% of wild type (Fig 2D). The level of PE is increased, while the PC level is not significantly changed (Fig 2E). Moreover, along with the marked reduction of PS, the levels of PA and PI are significantly increased, while the levels of phosphatidylglycerol (PG) and cardiolipin (CL) are reduced (Fig 2E). The lipidomic data also show that the levels of triacylglycerol (TAG) and diacylglycerol (DAG) are increased in pss RNAi, consistent with the BODIPY staining result (Fig 2E). Put together, these results indicate that as the sole PS synthase in Drosophila, PSS, likely uses PE as the substrate for PS synthesis.

pss knockdown depletes plasma membrane Akt and reduces cell growth, at least in part, via the insulin pathway

We next explored the mechanisms underlying pss-mediated cell growth and neutral lipid homeostasis. Previous reports show that the insulin pathway regulates Drosophila salivary gland cell growth [30, 31]. We examined the activity of the insulin pathway. tGPH is a GFP reporter that reflects insulin pathway activity through the PM:cytoplasm ratio of GFP signal [31]. We found that the plasma membrane tGPH signal is decreased and the PM:cytoplasm
Fig 2. CG4825 is the sole Drosophila phosphatidylserine synthase (PSS). (A) The evolutionary phylogenetic tree of PSS and its homologs in different species. The scale bar indicates amino acid divergence between sequences. Bootstrap number is 1,000 for resampling. (B) Schematic of the phospholipid and glycerolipid synthesis pathways. The black arrows indicate the common pathways in different species, the green arrow shows the pathway in yeast, the red arrows indicate the pathways in mammals, and the blue arrows indicate the pathways that occur in other species. The black arrows indicate the common pathways in different species, the green arrows indicate the pathways in mammals, and the blue arrows indicate the pathways that occur in other species. The scale bar indicates amino acid divergence between sequences. Bootstrap number is 1,000 for resampling. (C) S2 cells were cultured with medium containing 10 μM NBD-PA, NBD-PC, or NBD-PE for pulse-chase analysis. The S2 cells were harvested before the pulse-chase treatment (P) and 0, 1, 2, and 4 hours after pulse-chase. The extracted total lipids from S2 cells were separated on TLC plates for 1~2 hour. NBD-PE is converted to NBD-PS while NBD-PC is not. M, marker. Noted that a contaminating arrow shows the pathway in yeast, the red arrows indicate the pathways in mammals, and the blue arrows indicate the pathways that occur in other species. The scale bar indicates amino acid divergence between sequences. Bootstrap number is 1,000 for resampling. (D-E) Lipid profiling of control and pss RNAi 3rd instar larval salivary glands. The levels of the lipid species were normalized by calculating the mole fraction of the total polar lipids. In pss RNAi, the levels of PS, PG and CL are reduced, the levels of PE, PA, PI, DAG and TAG are increased, and the level of PC is unchanged. Data are shown as mean ± SEM. Data were compared with the unpaired Welch Two Sample t-test. * p < 0.05, ** p < 0.01, *** p < 0.001.

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Overexpressing phosphatidylserine decarboxylase enhances the cell growth defect and suppresses the ectopic lipid storage phenotype of pss knockdown

If a reduced level of plasma membrane PS is the cause of impaired insulin pathway activity in pss RNAi, elevating the level of plasma membrane PS might rescue the small cell size phenotype, and further reduction of the PS level should exacerbate the reduced cell size phenotype in salivary glands. Phosphatidylserine decarboxylase, which is encoded by the Pisd gene in mice and by PSD in yeast (Fig 2B), is localized in mitochondria under normal condition and...
Fig 3. The cell growth defect of *pss* RNAi is caused by impaired insulin pathway activity. (A) Images of the tGPH reporter in control and *pss* RNAi 3rd instar larval salivary glands. The same laser power and exposure time were used during the imaging. Scale bar represents 100 μm. (B) Quantification of the PM:cytoplasm PIP3 ratio in control and *pss* RNAi salivary glands (n = 5). Data are shown as mean ± SEM and were compared with Welch Two Sample t-test. *** p < 0.001. (C) Western blot of Akt and pAkt in *ppl* > control RNAi, and *pss* RNAi 3rd instar larval salivary glands. Three replicates were performed and a representative result from one replicate is shown here. A total of 10 μg protein was loaded. (D)
Immunofluorescent staining of Akt in 3rd instar larval salivary gland cells. Scale bar represents 25 μm. (E) Overexpression of AktCA rescues the reduced salivary gland size, but not the ectopic lipid accumulation, of pss RNAi. Scale bar represents 100 μm. (F) Quantification of the size of salivary gland cells with RNAi of ps and over-expression of Akt (n = 5 for each group). Data are shown as mean ± SEM. Data were compared with One-way ANOVA. Multiple comparisons of means were conducted with Tukey Contrasts. * p < 0.05, ** p < 0.001, ns: not statistically significant. (G) Overexpression (OE) of Pisd (CG5991) in pss RNAi suppresses the lipid accumulation and enhances the cell size reduction. In E and F, BODIPY (green) stains lipid droplets and DAPI (blue) stains the nuclei. Scale bar represents 100 μm. (H) Quantification of the size of salivary gland cells in different genetic backgrounds (n = 5 for each group). Data are shown as mean ± SEM. Data were compared with One-way ANOVA. Multiple comparisons of means were conducted with Tukey Contrasts. * p < 0.05, ** p < 0.001.

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converts PS to PE. We examined the phenotype of pss RNAi with knockdown or overexpression of Drosophila Pisd (CG5991). We hypothesized that in ps RNAi salivary glands, RNAi of Pisd may increase the PS level in the ER and subsequently restore the plasma membrane PS level, while overexpression of Pisd may further decrease the plasma membrane PS level. We found that RNAi of Pisd did not suppress the cell size phenotype caused by pss RNAi (S2 Fig), although the RNAi reduced the expression level of Pisd (S2 Fig). However, overexpressing Pisd further reduced the size of 3rd instar larval salivary gland cells in pss RNAi (Fig 3G and 3H). Together, these results provide additional evidence that a reduced level of plasma membrane PS in pss RNAi impairs insulin pathway activity and affects cell growth.

Is the ectopic lipid storage phenotype in pss RNAi also caused by reduced PS and impaired insulin pathway activity? Interestingly, Pisd overexpression completely suppresses the ectopic lipid accumulation phenotype of pss RNAi (Fig 3G), which demonstrates that these two phenotypes are caused by different mechanisms. Accordingly, although elevating insulin pathway activity by overexpression of either AktCA or PI3KCA suppresses the cell growth defect, it only marginally reduces the ectopic lipid storage in ps RNAi (Figs 3E and S2B).

**Loss of pss affects mitochondrial protein import and mitochondrial integrity**

The above results prompted us to further investigate the underlying mechanism of pss RNAi-induced ectopic lipid accumulation. Another important trafficking route of PS, besides moving from the ER to the PM, is its import from the ER into mitochondria. Previous reports showed that mitochondrial PS is mainly used for the production of mitochondrial PE, which is important for mitochondrial morphology and function [26, 33]. In the lipidomic analysis of pss RNAi, our attention was also drawn to mitochondria by the marked reduction of CL (Fig 2E), a special phospholipid that is highly enriched in mitochondria. Mitochondrial fatty acid oxidation is important for the catabolism of lipids and therefore disturbing the function of mitochondria may lead to lipid accumulation [34].

Does knockdown of *Drosophila pss* affect mitochondria? We used a mitoEYFP reporter to label the mitochondria in *in vivo* [35]. In control 3rd instar larval salivary glands, mitoEYFP appears as fluorescent puncta (Fig 4A). Surprisingly, the fluorescent signal is almost completely absent in pss RNAi (Fig 4A). MitoTimer, a GFP-based mitochondrial marker [36], shows the same phenotype in pss RNAi (S3 Fig). We found that pss RNAi does not significantly affect the transcription of mitoEYFP in salivary glands (S3 Fig). In addition, pss RNAi does not prevent the expression of other GFP or myr-mRFP reporters tested (S3 Fig). To explore whether the lack of the mitoEYFP fluorescent signal is due to the absence of mitochondria in pss RNAi, we immunostained mitochondria with an antibody against ATP5A, the alpha subunit of mitochondrial ATP synthase. The ATP5A signal is present in both control and pss RNAi salivary gland cells (Fig 4B), indicating that mitochondria are present in pss RNAi cells. The ATP5A signal appears as punctate structures in wild type, while in pss RNAi, the signal is more condensed, which could be due to packed mitochondria in small cells (Fig 4B). Both
The ectopic lipid storage phenotype in pss RNAi is likely due to the metabolic shift from phospholipid synthesis to neutral lipid synthesis

What is the relationship between mitochondrial abnormality and ectopic lipid storage? The increased levels of PE, DAG and PA, along with the reduced conversion of PE to PS (Fig 2E), may provide an explanation for the ectopic lipid storage in pss RNAi. In our previous study [39], a similar ectopic lipid storage phenotype was found in CdsA RNAi and bbc RNAi, which shifted PA-(CDP-DAG) synthesis to PA-DAG synthesis and DAG-PE synthesis to DAG-TAG synthesis, respectively. Therefore, it is possible that the ectopic lipid storage phenotype in pss RNAi is caused by the metabolic shift from phospholipid synthesis, namely DAG-PE-PS, to neutral lipid synthesis (Fig 2B). To test this hypothesis, we performed genetic analysis. RNAi of Lipin, the gene encoding phosphatidic acid phosphatase which generates DAG from PA,
fully suppresses the lipid storage phenotype of pss RNAi (Fig 5A). Similarly, CdsA overexpression completely suppresses the ectopic lipid phenotype of pss RNAi (Fig 5B). Together, these results indicate that the metabolic shift from phospholipid synthesis to neutral lipid synthesis in pss RNAi likely results in ectopic lipid storage.

We also analyzed the lipid profile in pss RNAi with Pisd overexpression. The level of TAG is marginally reduced, although not statistically significant, in cells with pss RNAi and Pisd overexpression (Fig 5C), which is consistent with our observation that lipid droplets are not found in cells of pss RNAi with Pisd overexpression. Pisd overexpression further increases the level of PE in pss RNAi salivary gland, indicating that Pisd overexpression increases overall lipid flow from PS to PE (Fig 5C). Intriguingly, the PS level in salivary gland cells with RNAi of pss and overexpression of Pisd is marginally higher compared to Pisd overexpression alone or pss single RNAi (Fig 5C). The lipidomic data also show a further reduction of the CL level in the pss RNAi salivary gland with overexpression of Pisd (Fig 5C).

**Decreasing lipid storage or promoting cell growth cannot rescue the mitochondrial dysfunction caused by pss RNAi**

Since Pisd overexpression completely suppresses the ectopic lipid accumulation phenotype of pss RNAi, we next examined whether Pisd overexpression rescues the mitochondrial defect of pss RNAi. Overexpression of Pisd did not restore the mitoEYFP signal in pss RNAi (Fig 6A). In addition, compared to wild type or pss RNAi alone, some large ATP5A-positive puncta appeared in pss RNAi with Pisd overexpression (Fig 6B). We also conducted ultra-structural analysis by EM (Fig 6C). Compared with pss RNAi, the structural abnormalities of mitochondria seem much worse in pss RNAi with Pisd overexpression. The regular tubular cristae structures have almost completely disappeared. Instead, filamentous mesh resembling matrix condensation or crista fragments were found in the mitochondrial matrix (Fig 6C). In addition, autophagosomes was frequently found (Fig 6C). These results indicate that although Pisd overexpression rescued the ectopic lipid storage phenotype of pss RNAi, it did not rescue the mitochondrial defects. Similarly, overexpression of CdsA does not rescue the mitoEYFP import defect in pss RNAi, even though it completely rescues the ectopic lipid storage phenotype (Fig 6D). Along the same line, we found that overexpression of AktCA does not rescue the loss of the mitoEYFP signal (Fig 6E), but it does rescue the cell growth defect of pss RNAi (Fig 3E). Together, these results indicate that PSS regulates mitochondrial function and cell growth or lipid storage through distinct mechanisms.

**Reducing PS transport from the ER to PM by loss of PI4KIIIα partially rescues the mitochondrial defects of pss RNAi**

None of the above manipulations (overexpression of CdsA, AktCA or Pisd) rescues the mitochondrial defects of pss RNAi. This observation is consistent with the idea that reduced PS import into mitochondria is probably the underlying cause of the mitochondrial defects, because overexpression of CdsA, AktCA or Pisd is unlikely to increase the level of mitochondrial PS. After its synthesis in the ER, PS is transported to other cellular membranes such as PM, mitochondria and Golgi. It is unknown whether these different PS transport routes are coordinated. In particular, it is not known whether there is a balance between different PS transport routes and, if there is, whether disrupting one PS transport route may promote other PS transport routes. If that is the case, blocking PS transport from the ER to PM in pss RNAi may divert more PS to mitochondria and therefore suppress the mitochondrial defect of pss RNAi. PS transport from the ER to the PM relies on the PI4P gradient between the PM and ER [5, 6]. At the PM, PI4KIIIα phosphorylates PI to PI4P (Fig 6F). RNAi of PI4KIIIα may
Fig 5. The metabolic shift from phospholipid synthesis to TAG synthesis contributes to ectopic lipid storage in salivary glands with pss RNAi. (A) Staining of lipid droplets with BODIPY in 3rd instar larval salivary gland with
reduce the level of PI4P in the PM, leading to loss of the PM-ER PI4P gradient required for PS transport from the ER to PM. We knocked down the expression of PI4KIIIα in pss RNAi salivary gland cells. Interestingly, the loss of the mitoEYFP signals in pss RNAi salivary gland cells is partially reversed by RNAi of PI4KIIIα, suggesting that reducing PS transport from the ER to PM may induce a compensatory increase in PS transport from the ER to mitochondria and rescue the mitochondrial defects of pss RNAi (Fig 6G). Furthermore, this result indicates that there is a balance between PS transport from the ER to PM and from the ER to mitochondria.

Discussion

There are numerous difficulties in revealing the in vivo cellular and physiological roles of phospholipids, including redundancy of metabolic genes, interconnected metabolic pathways, and different contributions of substrate, product and further metabolites derived from the product. In this study, through genetics and lipidomic analysis, we reveal that distinct mechanisms underlie the pleiotropic cellular defects caused by knocking down Drosophila PS synthase, PSS. Our detailed phenotypic and mechanistic analyses of pss knockdown provide a clear example of how altering lipid homeostasis contributes to different cellular phenotypes.

PS can be synthesized from PC or PE in mammals. In Drosophila, we did not detect the conversion of NBD-PC to NBD-PS in S2 cells and it is likely that Drosophila PSS utilizes PE, the major phospholipid in Drosophila, as the substrate for PS synthesis based on the lipidomic results from 3rd instar larval salivary glands (Fig 2D and 2E). We found that pss knockdown leads to cell growth defects, ectopic lipid accumulation and loss of mitochondrial integrity. We further showed that these three defects are likely due to different metabolic impacts of pss knockdown (Fig 7A–7D). We propose that PSS regulates cell growth, at least in part, via the insulin pathway, by affecting the level of plasma membrane PS and subsequently the membrane recruitment of Akt (Fig 7B). It is also possible that similar to PI4KIIIα mutants [40], pss RNAi leads to a defect in PM integrity which may cause the reduced Akt recruitment in PM. This notion is further supported by the aberrant myr-mRFP localization in the ppl> pss RNAi cells (S3 Fig). Interestingly, a previous study in Drosophila also found that both Lipin and GPAT are important for insulin pathway activity [41]. Both GPAT and Lipin act upstream of PSS in PS synthesis (Fig 2B). Therefore, it is possible that the level of PS contributes to the GPAT/Lipin-mediated regulation of insulin pathway activity. Considering the additive effect of pss RNAi and elevating insulin pathway activity on cell growth, we cannot rule out the possibility that other cellular defects, such as the mitochondrial dysfunction, may also contribute to the abnormal cell growth in pss RNAi.

The ectopic lipid storage phenotype of pss knockdown is mainly attributed to the metabolic shift of the glycerolipid synthetic program. Glycerophospholipids and TAG share similar synthetic pathways (Fig 2B). Phospholipid homeostasis and lipid storage are highly interconnected [39, 42]. Besides the overall compensatory DAG-TAG metabolic shift, changes in the levels of specific phospholipids, such as PE, PA and PI, may also contribute to ectopic lipid storage in pss RNAi. For example, PE acts as a feedback regulator of SREBP-mediated lipogenesis [43]. The increased level of PA may increase the size of lipid droplets, where TAG is stored [44].

The loss of mitochondrial integrity is the most dramatic consequence of pss knockdown. Mitochondria exchange lipids with the ER and other organelles. PS is imported into
A  

B  

C  

D  

E  

F  

G
mitochondria for the synthesis of mitochondrial PE, which is known to be important for mitochondrial morphology and function. In flies, *pss* RNAi affects the mitochondrial import of *mitoEYFP* and the morphology of mitochondria. In mammalian cells, although the mitochondrial import of *mitoEYFP* is not affected by double knockdown of *PTDSS1* and *PTDSS2*, the morphology of mitochondria is aberrant. A recent study reported that the reduction of mitochondria for the synthesis of mitochondrial PE, which is known to be important for mitochondrial morphology and function. In flies, *pss* RNAi affects the mitochondrial import of *mitoEYFP* and the morphology of mitochondria. In mammalian cells, although the mitochondrial import of *mitoEYFP* is not affected by double knockdown of *PTDSS1* and *PTDSS2*, the morphology of mitochondria is aberrant. A recent study reported that the reduction of

**Fig 6. The mitochondrial defect of *pss* RNAi is not rescued by *Pisd* overexpression.** (A) Fluorescence images of *mitoEYFP* in 3rd instar larval salivary glands of different genetic backgrounds. The absence of *mitoEYFP* signal in *pss* RNAi salivary glands is not rescued by *Pisd* (CG5991) overexpression. *mitoEYFP* (green) marks mitochondria and DAPI (blue) stains the nuclei. Scale bar represents 20 μm. (B) Immunostaining of mitochondria with anti-ATP5A antibody in 3rd instar larval salivary glands of different genetic backgrounds. Anti-ATP5A antibody (green) marks mitochondria and DAPI (blue) stains the nuclei. Scale bar represents 20 μm. (C) The EM structure of mitochondria in 3rd instar larval salivary gland cells of control, *pss* RNAi, and *pss* RNAi with *Pisd* overexpression (*PisdOE*). Scale bar represents 1 μm. Red arrows in *ppl-GAL4/+* and *ppl > pss* RNAi salivary gland cells mark the cristae junction sites, and the red arrow in the salivary gland cell with *pss* RNAi and *PisdOE* marks the filamentous meshes or cristae fragments. (D) Images of *mitoEYFP* in 3rd instar larval salivary glands of different genetic backgrounds. Overexpression of *CdSα* does not restore the *mitoEYFP* signal in salivary glands of *pss* RNAi. Scale bar represents 20 μm. (E) Images of *mitoEYFP* in 3rd instar larval salivary glands of different genetic backgrounds. The loss of *mitoEYFP* signal in *pss* RNAi is not rescued by the overexpression of AktCA. Scale bar represents 20 μm. (F) Illustration of the cellular PS transport pathways. (G) Images of *mitoEYFP* in 3rd instar larval salivary glands of different genetic backgrounds. The loss of *mitoEYFP* signal in *pss* RNAi is partially rescued by *PI4KIIIα* RNAi.

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**Fig 7. Proposed models of the mechanisms underlying the different *pss* RNAi phenotypes.** (A-B) Loss of *pss* function leads to reduced Akt, increased DAG and TAG, and impaired mitochondrial structure and function. (C) The overexpression of *Pisd* in *pss* RNAi may lead to increased engagement of phospholipids in the PS-PE-PS cycle locally, probably at the ER or mitochondria. Therefore, *Pisd* overexpression reduces the level of PS at the PM and the amount of DAG available for TAG synthesis, leading to smaller cells and decreased lipid storage compared to *pss* RNAi alone (B). (D) RNAi of *PI4KIIIα* rescues the defective mitochondrial import of *mitoEYFP* in *pss* RNAi by increasing the transport of PS to the mitochondria.

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mitochondrial PE promotes the proteolysis of mitochondrial proteins and this may explain the aberrant mitochondria in pss RNAi salivary gland cells [45]. Previous reports showed that mitochondrial import of preproteins is impaired in PE-depleted [46] or CL-deficient mitochondria [47, 48]. The difference in mitoEYFP mitochondrial import between pss RNAi in Drosophila and PTDSS1/PTDSS2 knockout in mammalian cells may reflect a difference in mitochondrial protein import between flies and mammals or insufficient knockdown of PTDSS1/PTDSS2 in mammalian cells.

At first glance, it is puzzling that overexpression of Pisd rescues the ectopic lipid accumulation phenotype of pss RNAi, but not the mitochondrial defects. The lipid metabolic changes in pss RNAi with Pisd overexpression is intriguing. Pisd overexpression should increase the level of mitochondrial PE. Salivary glands are small, and therefore we were unable to isolate mitochondria from them to measure the mitochondrial PE level. However, the total PE level is indeed increased in pss RNAi with Pisd overexpression. If PS in the mitochondria is used for the synthesis of mitochondrial PE, why then did overexpression of Pisd fail to rescue the mitochondrial defect of pss RNAi? It is possible that there are unidentified functions of mitochondrial PS in maintaining mitochondrial integrity and protein import. Alternatively, it is possible that the impairment of mitochondrial protein import in pss RNAi prevents a sufficient amount of Pisd entering into mitochondria, and Pisd instead stays in the ER (or other locations), leading to increased PE in the ER (Fig 7C). In support of this hypothesis, a recent report showed that PSD (yeast Pisd) localizes and functions in both mitochondria and ER [49]. However, the elevation of PE in the ER seems contradictory to the metabolic shift in pss RNAi and the suppression of the ectopic lipid accumulation phenotype in pss RNAi by Pisd overexpression (Fig 7B and 7C). The slightly increased level of PS in pss RNAi with Pisd overexpression compared to pss RNAi alone further complicates the analysis. It appears that the elevated PE level caused by Pisd overexpression also increases the flow from PE to PS, resulting in the slightly increased PS level in pss RNAi with Pisd overexpression. It is possible that when Pisd is overexpressed, more phospholipids are engaged in the PS-PE-PS cycle locally, probably at the ER or mitochondria. To achieve a full understanding of the underlying metabolic flow changes, organelle-specific lipid profiling combined with metabolic flux analysis may be required.

Our findings may further explain the early lethality of PSS1/2 deficiency in mouse [9, 10]. Defective mitochondrial function or impaired insulin pathway activity could both lead to embryonic lethality. The lethality of the Drosophila pss mutant and the pleiotropic phenotypes caused by RNAi of pss suggest that dietary/maternally derived PS is either insufficient or cannot be delivered to internal cells/tissues to ensure normal physiological function.

After synthesis, PS can be transported from the ER to the PM or to mitochondria. Little is known about other PS transport routes or how cells coordinate or prioritize the different PS trafficking routes [50]. The genetic suppression of pss RNAi by PI4KIIIα RNAi indicates that there is a balance between PS transport from the ER to PM and from the ER to mitochondria. In support of this transport balance, the enhancement of the cell growth defect in pss RNAi by Pisd overexpression can also be explained by diversion of PS away from the ER to PM transport path. Furthermore, the suppression of the mitoEYFP defects of pss RNAi by PI4KIIIα RNAi opens up a new possibility to screen for more suppressors. The identification and analysis of suppressors will be important for revealing other PS transport routes and their regulation.

Materials and methods

Drosophila husbandry and stocks

All the flies used in this study (S1 Table) were maintained on standard cornmeal food. The w1118, ppl-GAL4/+ or ppl>white RNAi was chosen as the wild type or control RNAi group. For
overexpression, UAS-GMA (GFP tagged actin-binding domain of Moe) or UAS-myr-mRFP (membrane RFP) was used as the UAS control. The CG4825KK105709 line was used in pss RNAi experiments if not specified. The fly stocks were obtained from the Bloomington Drosophila Stock Center (BDSC), the Vienna Drosophila Resource Center (VDRC), NIG Stock Center (NIG) and Tsinghua Fly Center (THFC).

**Tissue staining, microscopy and image analysis**

The lipid droplets of wandering 3rd instar larval salivary gland were stained by BODIPY, Nile red, or LipidTOX following the procedure described before [30, 39, 51]. After fixation with 4% paraformaldehyde (PFA) for 0.5 hour, the samples were stained with DAPI (2 ng/μl). For immunofluorescent staining of ATP5A or Akt, the dissected 3rd instar larval salivary glands were fixed in 4% PFA followed by treatment with 0.3% PBST (PBS + 0.3% TritonX-100), and blocked with 5% BSA for 1 hour. The samples were incubated with anti-ATP5A (dilated at 1:200; Abcam, ab14748) or anti-Akt (diluted at 1:200; Cell Signaling) overnight at 4˚C. Alexa Fluor 488-conjugated goat anti-mouse (1:1000; Invitrogen) or Alexa Fluor 488-conjugated goat anti-rabbit (1:1000; Invitrogen) was chosen as the secondary antibody, respectively. The stained samples were mounted in 80% glycerol after washing with 1×PBS three times. All the images were acquired by confocal microscopy (Leica SP8) using 20× and 63× objectives, with NA 0.75 and 1.4, respectively. The wavelengths of the laser are 405 nm, 488 nm, 553 nm and 638 nm for DAPI, BODIPY, Nile Red and LipidTOX, respectively. The quantifications of the salivary gland size and cell size were performed by measurement of the cross-sectional area with Image J software (1.51j8).

**Molecular biology and qRT-PCR**

For RT-PCR, total mRNAs were isolated from wandering 3rd instar larval salivary glands using Trizol reagent (Invitrogen) and the cDNA was generated using the Superscript II reverse transcriptase kit (Invitrogen). qRT-PCR was performed with the Stratagene MX300P system (Agilent) using Trans Start Green qPCR SuperMix (Transgene Biotech). The expression level of each gene was normalized to rp49. The primers used in this study are listed in S2 Table. To make the UAS-flag-pss transgene, the coding region of pss was cloned from w1118 and inserted into pUAST-attB-flag through the NotI and XbaI sites.

**Phylogenetic analysis**

The Drosophila PSS sequence was analyzed with Pfam [52]. PSS family members were aligned with the global sequence alignment software ClustalW, and the Neighbor-Joining phylogenetic tree of the PSS family was constructed with MEGA (6.06) [53]. The phylogenetic analysis was conducted with the bootstrap method using 1,000 bootstrap replications.

**NBD-labeled phospholipid chase and TLC**

The procedure for the pulse-chase analysis of the S2 cells was modified from Miyata [27] and Tamura [54]. The S2 cells were incubated with NBD-PA, NBD-PC or NBD-PE (Avanti Polar Lipids, Inc.) for 20 min. The S2 cells were washed, then incubated for different periods of time. After harvesting, the total lipids in the cells were extracted and resuspended in chloroform/methanol (1:2, vol/vol). The lipid samples were separated by TLC on silica gel 60 F254 plates (Merck,1.05729.0001) using a solvent system of chloroform/methanol/25% ammonia, 65:35:5 [26]. The TLC plates were detected with a Typhoon 9500 imager and the images were analyzed with Image J (1.51j8).
Cell culture and RNAi in cultured cells

The siRNAs for human PTDSS1 and PTDSS2 (S3 Table) were designed and synthesized by GenePharma Co., Ltd (www.genepharma.com). HeLa cells were cultured in high-glucose DMEM medium (HyClone) supplemented with 10% fetal bovine serum, 100 U/ml penicillin and 100 μg/ml streptomycin (HyClone) at 37°C. The HeLa cells were transfected with 100 pmole of PTDSS1 or/and PTDSS2 siRNA and 0.7 pmole mitoEYFP expression plasmid using Lipofectamine 2000 (Invitrogen) for 48 hours.

Lipidomic analysis

Lipids were extracted form salivary glands of wandering 3rd instar larvae and analyzed as previously described [55]. The samples for each genotype contained 25 pairs of salivary glands. The mole fraction of each lipid was normalized to the mole fraction of total polar lipids.

Western blot and quantification

The salivary glands were dissected from 60 wandering 3rd instar larvae. The samples were lysed in 240 μl of ice-cold 1% SDS lysis buffer. 10 μg of the sample protein were loaded and detected with the following rabbit antibodies: anti-Akt (Cell Signaling, diluted at 1:1000), anti-phospho-Akt (Ser473) (Cell Signaling, diluted at 1:1000), and rabbit anti-α-tubulin (Abcam, diluted at 1:4000). Quantification of the band intensities was conducted using Image J software (1.51j8) and the protein levels were normalized to tubulin.

High-pressure freezing (HPF) electron microscopy imaging

The 3rd instar larval salivary gland samples were loaded into carriers and cryofixed on a Leica Microsystems HPM 100 (EM ICE) at ~2,100 bar and automatically cooled into liquid nitrogen. After HPF, the samples were transferred under liquid nitrogen to a Leica Microsystems AFS-2 unit and incubated at -90°C for 72 h in freeze substitution solution: acetone with 2% (wt/vol) osmium tetroxide and 2% (vol/vol) water. The temperature of the samples was gradually increased according to the following timeline: increase by 8°C/h for 4 h; hold at -60°C for 12 h; increase by 5°C/h for 6 h; hold at -30°C for 10 h; increase at 4°C/h for 10 h; hold at 10°C for 10 h. Samples were washed four times in acetone, stained in 1% uranyl acetate for 1 h, and rinsed 3 times in pure acetone. Samples were infiltrated stepwise with increasing concentrations of Embed 812 resin: 2:1 (Embed 812:acetone) for 3 h, 1:1 for 5 h, then twice in 100% fresh resin for 8 h. The samples were then transferred to an embedding mold containing fresh resin and polymerized in a 60°C oven for 3 days. Ultrathin sections (60 nm) were produced with a diamond knife (Diatome) on an ultramicrotome (Ultracut UCT; Leica Microsystems). The sections were all collected on slot copper grids (EMS), then visualized with a JEM 1400 TEM (Hitachi 7700) operating at 80 kV. Pictures were recorded with a Gatan 832 4kX2.7k CCD camera.

Statistical analysis

All the data are shown as mean ± SEM. All the statistical analyses were conducted with R language (3.5.1) and R packages (Rcmdr). The graphs were drawn by GraphPad (version 7.00).

Supporting information

S1 Fig. pss RNAi reduces cell size. (A) Quantification of the cell number in ppl-GAL4/+ , ppl>control RNAi and pss RNAi 3rd instar larval salivary glands (n = 7). The salivary gland cell number is not changed when pss is knocked down.
(B) Quantification of the cell size in ppl-GAL4/+, ppl>control RNAi and pss RNAi 3rd instar larval salivary glands (n = 5). The salivary gland cell size is reduced in pss RNAi compared to ppl-GAL4/+.

(C) Lipid droplet staining in 3rd instar larval salivary glands from four CG4825 RNAi lines. Scale bar represents 100 μm. BODIPY (green) labels lipid droplets and DAPI (blue) labels nuclei.

(A and B) Data are shown as mean ± SEM. Data were compared with One-way ANOVA. *** p < 0.001.

(TIF)

S2 Fig. Genetic analysis of pss and Pisd. (A) The transcription level of Akt in 3rd instar larval salivary glands with pss RNAi (n = 3, each repeat contains RNA from 25 larvae). Data are shown as mean ± SEM. Data were compared with the unpaired Welch Two Sample t-test.

(B) Over-expression of PI3KCA in pss RNAi suppresses the reduced salivary gland size and but not ectopic lipid accumulation phenotypes. BODIPY (green) labels lipid droplets and DAPI (blue) labels nuclei. Scale bar represents 100 μm.

(C) RNAi of Pisd in pss RNAi does not suppress the reduced salivary gland size and ectopic lipid accumulation phenotypes. Nile Red (red) labels lipid droplets and DAPI (blue) labels nuclei. Scale bar represents 100 μm.

(D) The RNAi knockdown efficiency of Pisd (n = 3, each repeat contains RNA from 25 larvae). Data are shown as mean ± SEM. Data were compared with the unpaired Welch Two Sample t-test. * p < 0.05.

(TIF)

S3 Fig. pss RNAi impairs mitochondrial protein import. (A) Images of MitoTimer-labeled mitochondria in ppl-Gal4/+, ppl>control RNAi and pss RNAi 3rd instar larval salivary gland. MitoTimer is detected in two forms: GFP and dsRed. Scale bar represents 20 μm.

(B) The transcription level of mitoEYFP in 3rd instar larval salivary glands with pss RNAi (n = 3, each repeat contains RNA from 25 larvae). Data are shown as mean ± SEM. Data were compared with the unpaired Welch Two Sample t-test.

(C) pss RNAi does not affect the expression of other GFP/myr-mRFP reporters. Scale bar represents 50 μm.

(D) The cristae junction width in pss RNAi cell mitochondria is increased. Scale bar represents 0.2 μm.

(TIF)

S1 Table. The fly strains used in this study.

(DOCX)

S2 Table. The primers used in this study.

(DOCX)

S3 Table. The siRNAs used in this study.

(DOCX)

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