Redundant roles of the phosphatidate phosphatase family in triacylglycerol synthesis in human adipocytes

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Abstract

Aims/hypothesis In mammals, the evolutionary conserved family of Mg2+-dependent phosphatidate phosphatases (PAP1), involved in phospholipid and triacylglycerol synthesis, consists of lipin-1, lipin-2 and lipin-3. While mutations in the murine \textit{Lpin1} gene cause lipodystrophy and its knockdown in mouse 3T3-L1 cells impairs adipogenesis, deleterious mutations of human \textit{LPIN1} do not affect adipose tissue distribution. However, reduced \textit{LPIN1} and PAP1 activity has been described in participants with type 2 diabetes. We aimed to characterise the roles of all lipin family members in human adipose tissue and adipogenesis.

Methods The expression of the lipin family was analysed in adipose tissue in a cross-sectional study. Moreover, the effects of lipin small interfering RNA (siRNA)-mediated depletion on in vitro human adipogenesis were assessed.

Results Adipose tissue gene expression of the lipin family is altered in type 2 diabetes. Depletion of every lipin family member in a human Simpson–Golabi–Behmel syndrome (SGBS) pre-adipocyte cell line, alters expression levels of adipogenic transcription factors and lipid biosynthesis genes.

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in early stages of differentiation. Lipin-1 knockdown alone causes a 95% depletion of PAP1 activity. Despite the reduced PAP1 activity and alterations in early adipogenesis, lipin-silenced cells differentiate and accumulate neutral lipids. Even combinatorial knockdown of lipins shows mild effects on triacylglycerol accumulation in mature adipocytes. 

Conclusions/interpretation Overall, our data support the hypothesis of alternative pathways for triacylglycerol synthesis in human adipocytes under conditions of repressed lipin expression. We propose that induction of alternative lipid phosphate phosphatases, along with the inhibition of lipid hydrolysis, contributes to the maintenance of triacylglycerol content to near normal levels.

Keywords Basic science · Cell lines · Human · Lipid metabolism

Abbreviations
DAG Diacylglycerol
DNL De novo lipogenesis
ER Endoplasmic reticulum
FAO Fatty acid oxidation
LPP Lipid phosphate phosphatases
ORO Oil Red O staining
PAP Phosphatidate phosphatase
PPAR Peroxisome proliferator activated receptor
SGBS Simpson–Golabi–Behmel syndrome
siRNA Small interfering RNA
SAT Subcutaneous adipose tissue
TAG Triacylglycerol
VAT Visceral adipose tissue

Introduction
Triacylglycerols (TAGs) are neutral lipids that act as the major energy storage molecules, repository for fatty acids, and phospholipid precursors [1]. Adipocytes are the specialised cells for neutral lipid storage and one of their important physiological functions is to buffer the toxicity caused by NEFAs. Excessive calorie intake or genetic disorders can lead to lipid deposition in ectopic tissues, impair their function and lead to dyslipidaemia, insulin resistance and type 2 diabetes [2, 3].

Lipins are Mg$^{2+}$-dependent phosphatidate phosphatases (PAP1) with a central role in lipid metabolism, and catalyse the dephosphorylation of phosphatidate to diacylglycerol (DAG), which can be (1) acylated to form TAG, or (2) used in phospholipids synthesis [4, 5]. A second type of PAP activity is mediated by Mg$^{2+}$-independent transmembrane lipid phosphate phosphatases (LPPs, also known as PAP2), which are thought to regulate signalling properties of phosphatidate and DAG [6].

Fungi, nematodes and insects express one lipin, whereas mammals express three paralogues called lipin-1, -2 and -3 that exhibit distinct but overlapping expression in many mouse and human tissues [5]. Consistent with their key metabolic role, loss of lipin function disrupts TAG production, membrane organisation and phospholipid synthesis in several model organisms [7]. Interestingly, besides their enzymatic functions, lipins also regulate transcription [8–13]. For instance, lipin-1 transcriptional co-regulation of the peroxisome proliferator activated receptor (PPAR)A/PPAR coactivator 1α axis modulates fatty acid oxidation (FAO) in liver [9].

Lpin1 gene was originally identified as the deficient gene causing lipodystrophy, insulin resistance, peripheral neuropathy and neonatal fatty liver in the fld (also known as Lpin1) mouse model [14]. Loss of lipin-1 in mice blocks adipogenesis at an early stage preceding TAG accumulation, suggesting a distinct role of lipin-1 in differentiation [15, 16]. Consistently, small interfering RNA (siRNA)-mediated silencing of lipin-1 in mouse 3T3-L1 cells potently inhibits adipogenesis [10, 17]. In contrast, deleterious mutations in the LPIN1 gene in humans, which lead to recurrent rhabdomyolysis in childhood, do not compromise adipose tissue [18]. The basis of this difference between mice and humans is unknown; it has been hypothesised that it is due to compensation by the other two lipins [18]. Nevertheless, genetic variation in LPIN1 and LPIN2, and reduced LPIN1 expression levels and PAP1 activity in human adipose tissue have been associated with type 2 diabetes [19–25], which suggests a loss of their protective role against lipotoxicity.

While most of the studies have been performed on mice, very little is known on the roles of lipin-1 in human adipocyte physiology, while there is virtually no information on lipin-2 and -3. This prompted us to investigate the functions of the three lipins in type 2 diabetes and in human adipocytes by studying the effects of loss-of-function.

Methods
Reagents Unless otherwise stated, all reagents were supplied by Sigma-Aldrich Corporation (St. Louis, MO, USA). Human insulin was purchased from NovoNordisk (Bagsværd, Denmark); rosiglitazone from Cayman Chemical (Ann Arbor, MI, USA); and cell culture media from Gibco (Thermo Fisher Scientific, Waltham, MA, USA).

Human adipose tissue biopsy collection The cross-sectional study has been previously described [26, 27]. We used a cohort of 71 participants for the gene expression analysis, and of 28 males for protein analysis. They were grouped as:
(1) normoweight (BMI between 18.50 and 24.99), (2) obesity (BMI ≥25) and (3) obesity with type 2 diabetes (hereafter type 2 diabetes). Group characteristics are described in electronic supplementary material (ESM) Table 1A and 1B, respectively. Samples were obtained at the Joan XXIII University Hospital (Tarragona, Spain). VAT and SAT samples (gene expression) and SAT samples (protein expression) were obtained during abdominal elective surgical procedures for benign pathologies (cholecystectomy or surgery for abdominal hernia). The Ethics Committee approved the study and informed consent was obtained from all participants. Participants had no systemic disease other than obesity [28] or type 2 diabetes [29]. See ESM Methods for further details.

**Cell culture and differentiation** Simpson–Golabi–Behmel syndrome (SGBS) cells, a well-established system for studies of human adipocyte biology [30, 31], were differentiated as described [30], except that 1 nmol/l of insulin, 0.1 μmol/l cortisol, 2 μmol/l rosiglitazone and 25 mmol/l dexamethasone were used. See ESM Methods for detailed information.

Adipose-derived stem cells were isolated from adipose tissue (n = 3 female donors; age (years) 37.4 ± 6.4, BMI (kg/m²) 25.9 ± 3.0) from patients undergoing elective liposuction surgery. See ESM Methods for the isolation, proliferation and adipogenic procedure details.

**Quantification of expression levels** Protein content in cell lysates (lysis buffer: 50 mmol/l HEPES pH 7.4, 150 mmol/l NaCl, 4 mmol/l MgCl₂, 1% Triton X-100 and protease inhibitors) was quantified by using the bicinchoninic acid assay (Pierce, Rockford, IL, USA). See ESM for western blot procedures. Antibodies against lipin family members were previously described [17, 32]. Quantitative PCR was performed as previously described [27], and expressed relative to cyclophilin A and to control. See ESM for detailed information and ESM Table 2A and 2B for commercial reagents.

**Cell Fractionation** For cell fractionation, SGBS cells were grown and differentiated until day 10. Subcellular fractions were obtained by using hypotonic lysis followed by high salt extraction of nuclei. See ESM Methods for further details.

**Gene silencing** Transfections of SGBS cells with siRNA oligonucleotides were carried out by using Lipofectamine RNAiMAX transfection reagent (Thermo Fisher Scientific). Pre-adipocyte knockdowns were performed as a two-shots transfection of a mix of two siRNA duplexes per gene (12.5 nmol/l of each duplex) (see ESM Table 2C): reverse transfection at the start of the experiment and forward transfection the day before confluence. The non-targeting control concentration depended on total siRNA amount of the single and multiple knockdowns.

**Neutral lipid accumulation and metabolism** Data were obtained from differentiated SGBS cells and normalised by protein content. For TAG analysis, cells were processed as previously described [32], except that the supernatant fraction was analysed with the Serum Triglyceride Determination kit (Sigma). Glycerol release to cell culture media was quantified by using the Free Glycerol Determination Kit (Sigma). Fatty acid and glucose incorporation into TAGs, and FAO analysis were performed as previously described [33], with slight modifications (see ESM Methods for details).

**Enzyme assay** Cells lysates (lysis buffer: 50 mmol/l Tris-HCl pH 7.5, 0.25 mol/l sucrose, 10 mmol/l 2-mercaptoethanol, protease inhibitors) were subjected to centrifugation at 1000 g for 10 min at 4°C to remove cell debris. Protein concentration was determined by the method of Bradford [34] using BSA as a reference protein. Preparation of the substrate and measure of PAP activity was as previously described [35–37]. See ESM Methods for details. The Mg²⁺-independent LPP activity was measured in the same reaction mixture except that 2 mmol/l EDTA was substituted for 0.5 mmol/l MgCl₂. The Mg²⁺-dependent PAP activity was determined by subtraction of LPPs activity from PAP activity. A unit of PAP activity (expressed as units/mg protein) was defined as the amount of enzyme catalysing the formation of 1 nmol of product/min.

**Metabolomic analysis** SGBS cells were grown, transfected with siRNA as explained above, and differentiated to day 4. In brief, lipids were extracted from lyophilised samples by using dichloromethane/methanol and water. The organic phase (lipidic) was collected, dried under a stream of nitrogen, and resuspended in acetonitrile/isopropanol/water for untargeted LC-MS analysis. Differentially regulated lipids (p value <0.05 and fold >2) were retained for compound identification by MS/MS analyses. See ESM Methods for detailed procedures.

**Statistical analysis** Statistical analysis was performed by using the SPSS software version 15 (Chicago, IL, USA). ANOVA, Kruskal–Wallis, Pearson χ², Spearman correlation and Linear Stepwise Regression tests were performed for the human cohort analysis, and the General Linear Model Univariate test for in vitro experiments. Statistical power in the cohort analysis was ≥80%. The level of significance was set at α = 0.05.

**Results** Adipose tissue expression levels of the lipin family is altered in type 2 diabetes To investigate the roles of lipin paralogues in adipose tissue, we started by examining their gene expression in paired abdominal subcutaneous (SAT) and visceral (VAT) adipose tissue biopsies. As shown in
and sex as confusing and interacting variables, and showed that (1) SAT LPIN1 expression depends negatively on HOMA-IR ($R = 0.466$, $p = 0.005$, excluded variables: BMI, TAG, age and sex; $LPIN1_{SAT} = 0.37 \times \log_{10}(\text{HOMA} + 0.984)$), and (2) SAT LPIN3 expression depends positively on plasma glucose levels ($R = 0.414$, $p = 0.002$, excluded variables: age and sex; $LPIN3_{SAT} = 1 \times 10^{0.202 \times \log_{10}(\text{Glucose} - 0.961)}$).

The observed changes of the lipin family expression in type 2 diabetes participants may account for the altered PAP1 activity in adipocytes from these patients. Moreover, LPIN3 expression is associated with fasting glucose levels. Alterations in lipin-3 protein levels in SAT of participants with type 2 diabetes may be masked by its presence also in the stromal vascular fraction (data not shown).

### The three lipin family members have a role in early human adipogenesis

Next, we examined the expression of lipins during adipogenesis. Lipin-1 was induced, and lipin-2 and lipin-3 levels were present along SGBS adipogenesis with slight variation (Fig. 2a, b). Thus, SGBS adipocytes, similar to differentiated adipose-derived stem cells (ESM Fig. 1d), express the three lipin paralogues. Finally, under baseline conditions, the three lipins partitioned between the cytosolic, intranuclear and membrane-bound forms in SGBS adipocytes (ESM Fig. 1e).

To address the effects of decreased PAP1 activity on adipogenesis, we depleted each lipin member prior to the induction of adipogenesis in SGBS pre-adipocytes (by using siRNA; see Methods and Fig. 2c). Cells were analysed at day 4 after differentiation to assess adipogenic early events. Protein expression analysis was used to confirm knockdowns and evaluate possible compensatory mechanisms among lipins. Lipin-1 depleted cells responded by compensatory upregulation of lipin-2 protein (Fig. 2d and ESM Fig. 1f), but not LPIN2 transcript levels (data not shown). Conversely, single lipin-2 and lipin-3 knockdowns led to lower protein levels of the other family members (Fig. 2d).

### Table 1 Correlation analysis in abdominal SAT and VAT adipose tissue biopsies from 71 participants

| Variable     | SAT         | VAT         |
|--------------|-------------|-------------|
|              | LPIN1 | LPIN2 | LPIN3 | LPIN1 | LPIN2 | LPIN3 |
| BMI          | $-0.349^{**}$ | $-0.021$ | $0.220$ | $-0.283^{†}$ | $-0.091$ | $0.034$ |
| HOMA-IR      | $-0.471^{**}$ | $-0.104$ | $-0.006$ | $-0.329^{*}$ | $-0.139$ | $-0.006$ |
| Glucose      | $-0.349^{**}$ | $0.253^{†}$ | $0.357^{**}$ | $-0.119$ | $0.189$ | $0.298^{†}$ |
| Insulin      | $-0.522^{***}$ | $-0.169$ | $-0.039$ | $-0.340^{**}$ | $-0.179$ | $0.017$ |
| Triglycerol  | $-0.319^{**}$ | $-0.018$ | $0.072$ | $-0.202$ | $0.131$ | $0.238$ |
| NEFA         | $-0.095$ | $0.129$ | $0.128$ | $-0.029$ | $0.125$ | $0.336^{**}$ |
| Glycerol     | $-0.188$ | $-0.105$ | $0.144$ | $-0.113$ | $0.097$ | $0.274^{†}$ |

Spearman coefficient of the correlation analysis is shown. $^* p < 0.05$, $^{**} p < 0.01$, $^{***} p < 0.001$

Where statistical power is lower than 80%; $^{†} p < 0.05$
Analysis of PAP1 activity showed that lipin-1 accounted for almost all PAP1 activity (lipin-1-depleted cells showed 5% of PAP1 activity in the control), with lipin-2 and lipin-3 single knockdowns reducing it to 49% and 61%, respectively (Fig. 2e).

Next, we analysed incorporation of fatty acids and glucose into TAGs. While esterification of fatty acids was downregulated in lipin-1- and lipin-2-depleted cells, glucose incorporation into TAGs decreased in all three lipin-depleted cells (Fig. 2f).

Finally, we analysed the expression levels of transcription factors that promote early adipogenesis. CEBPA was downregulated in cells depleted of any lipin family member, and CEBPB was decreased upon lipin-2 deficiency (Fig. 2g). Under these conditions, gene expression of two key transcription factors regulated by CCAAT/enhancer binding proteins, PPARG and SREBP1 were downregulated (Fig. 2g), and also at the protein level (ESM Fig. 2a, b). CEBPD showed a significant upregulation in cells depleted of any lipin family member (Fig. 2g), probably due to the triggering of a compensation mechanism.

Given the role of lipins in neutral lipid biosynthesis, we next explored expression of lipogenic genes. The expression of these genes was significantly downregulated in cells depleted of any lipin family member (Fig. 2h). In contrast, glycerol-3-phosphate acyltransferase was significantly
upregulated in the lipin-1 and lipin-2 knockdowns (Fig. 2h).
Lipid quantification by mass spectrometry showed that,
among the lipid species that were altered, the levels of most
DAG (ESM Fig. 2c) and TAG (ESM Fig. 2d) species were
downregulated in cells depleted of any lipin compared with
controls. Interestingly, phosphatidate levels did not change
(data not shown).

Overall, despite the crosstalk between lipin family
members, lipin-1 silencing leads to a 95% depletion of PAP1
activity. However, our results point to a role of all three
members on early stages of human adipogenesis.

Single lipin silencing does not block TAG accumulation
in fully differentiated adipocytes Next, we analysed the
effects of lipin silencing in SGBS pre-adipocytes on late
adipogenesis. Similarly to the above experiments, we depleted
each lipin member prior to induction of adipogenesis in SGBS
pre-adipocytes, and cells were analysed at day 10 after
differentiation (Fig. 3a), where protein downregulation of
lipins still persisted (Fig. 3b and ESM Fig. 3a).

At day 10, lipin-1 depleted cells still responded by
compensatory upregulation of lipin-2 protein (Fig. 3b
and ESM Fig. 3a). Additionally, both single lipin-2 and lipin-3
knockdowns led to lower protein levels of the other lipins.
PAP1 activity analysis showed that lipin-1 accounted for
almost all PAP1 activity (lipin-1-depleted cells showed 3% of
the activity in the control). Lipin-2 and lipin-3 single
knockdowns reduced PAP1 activity only slightly (84% and
74%, respectively) (Fig. 3c).

Next, we assessed neutral lipid accumulation. Depletion of
any lipin triggered a slight reduction (15% to 34%) in TAG
accumulation (Fig. 3d). Expression of key transcription
factors for lipogenesis (ESM Fig. 3a, b) and genes of lipid
biosynthesis (ESM Fig. 3c) were altered to a lesser extent
compared with the effects observed at day 4.

Finally, fatty acid esterification into TAGs and α-glycerophosphate synthesis/de novo lipogenesis (DNL)
were downregulated in cells depleted of any lipin (Fig. 3e).
Compared with day 4, lipin-3 depletion also showed reduced
fatty acid esterification into TAGs at day 10.

Overall, the effects of lipin knockdown were mild on fully
differentiated adipocytes suggesting that cells overcame the
initial impairment of lipid gene expression. This may be due
to the existence of enough remnant protein amounts of lipins,
although PAP1 activity at day 10 was still almost absent in the
lipin-1 knockdowns.

Combinatorial silencing of lipins does not further alter the
phenotype of fully differentiated adipocytes We then asked
whether compensation by other lipin family members might
help cells to recover the initial alterations shown at day 4 after
adipogenesis. We used a combinatorial knockdown to deplete
combinations of two of the three lipins in SGBS pre-
adipocytes (see Methods), and analysed cells at day 10 of
differentiation (as in Fig. 3a).

Double lipin-1 and lipin-3 knockdowns led to an
upregulation of lipin-2 protein levels (Fig. 4a and ESM
Fig. 4). This compensatory pattern hindered the efficiency of
lipin-2 knockdown in the double (lipin-1 and -2) and triple
knockdown cells. Nevertheless, PAP1 activity in the triple
knockdown was residual compared with the control
(0.18 ± 0.31 vs 39.29 ± 3.42 nmol min\(^{-1}\) mg\(^{-1}\))
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Despite the almost complete lack of PAP1 activity,
near neutral lipid accumulation was not totally compromised in
the triple knockdown (Fig. 4b). This is in agreement with
the fact that deleterious mutations in LPIN1 do not affect
adipose tissue development in humans, and it suggests that
it is unlikely to be due to a compensatory upregulation of
lipin-2 (Fig. 2d), since the double lipin-1 and -2
knockdown shows a similar reduction in the content of
neutral lipids (Figs 3d, 4b).
Moreover, depletion of the three lipins did not significantly affect fatty acid esterification into TAGs and only slightly downregulated α-glycerophosphate synthesis/DNL (Fig. 4c), pointing to an improvement compared with day 4 (day 4 incorporation of fatty acids: 0.52 ± 0.35, p = 0.006, and of glucose: 0.32 ± 0.15, p < 0.001, fold over the control level).

**Consumption of TAGs is reduced in lipin-depleted SGBS adipocytes** Lipid droplets accumulate neutral lipids in a dynamic manner, with a balance among lipid biosynthesis, lipid hydrolysis and FAO in mitochondria. To test whether cells protect intracellular lipids by reducing its consumption, we analysed the rates of palmitate oxidation. The results show similar FAO rates in the control, single (Fig. 5a) and triple knockdown (Fig. 5b). In contrast, levels of glycerol release were downregulated compared with controls in the single knockdown (Fig. 5a) and in the triple knockdown (Fig. 5b). This was confirmed by using other commercial sources of lipin-1 siRNA (ESM Fig. 5b) and analysing LPP expression at day four of differentiation (ESM Fig. 5c).

Propranolol, a nonspecific β-blocker, is an effective PAP1 activity inhibitor and modestly effective as LPP inhibitor, with the LPP3 isoform being more sensitive to inhibition [36]. Propranolol treatment during adipogenesis (Fig. 6e) altered lipogenic gene expression (Fig. 6f) and blocked lipid droplet formation in SGBS adipocytes (ESM Fig. 5d).

**Discussion**

Herein we have addressed the contribution of the lipin paralogues to human adipogenesis. Decreased lipin expression levels and PAP1 activity in adipose tissue have been linked with insulin resistance [23–25]. In our cohort, adipose tissue expression confirmed that lipin-1 is altered in type 2 diabetes and that LPIN1 is negatively associated with insulin resistance. Moreover, we show that LPIN3 transcript levels are positively related with fasting glucose levels. This prompted us to analyse separately the effects of gene silencing of each lipin parologue in adipogenesis.

Previous studies have established that lipin-1 plays a major role in fat metabolism in rodents, with lipin-1 deficiency causing lipodystrophy features in mice and rats [14, 38]. The role of lipins in human adipogenesis is still undefined: mutations in the LPIN1 gene have not yet been detected in human lipodystrophy [39, 40]. Moreover, deleterious LPIN1 mutations cause paediatric rhabdomyolysis while fat distribution, average weight and plasma biochemical variables are normal [18]. Our data suggest that the lack of an essential role for PAP1 activity in human adipogenesis is not due to...
compensation by lipin-2 or lipin-3. We find that the triple lipin silencing in SGBS pre-adipocytes maintains the ability to incorporate fatty acids into TAG and accumulate neutral lipids despite loss of nearly all PAP1 activity.

SGBS pre-adipocytes are differentiated in the absence of serum and, therefore, neutral lipids should be obtained from DNL. Besides, although glyceroneogenesis is the major pathway for glycerol synthesis in the mature adipocyte [41], during adipogenesis glycolysis may account for an important source of α-glycerophosphate. Thus, glucose incorporation into TAGs may account for DNL, but also for α-glycerophosphate synthesis via glyceroneogenesis and glycolysis. Downregulated α-glycerophosphate synthesis/DNL points to broader effects of lipin silencing other than TAG synthesis. Moreover, the putative intranuclear roles of lipin-1, -2 and -3 in SGBS adipocytes add some complexity to the above.

The lipin family members cooperate for optimal PAP activity in mouse adipose tissue [42], liver [43, 44], brain [43] and human primary myoblasts [45]. Our data show that human lipin-1 accounts for most PAP1 activity in SGBS adipocytes and that upregulation of lipin-2 in lipin-1-depleted cells does not compensate for PAP1 activity. By contrast, downregulation of PAP1 activity in lipin-2 and lipin-3 single knockdowns cannot be fully attributed to their own depletion since they also downregulate lipin-1.
Activation of non-lipin compensatory pathways may also mask the lipin phenotypes. Nevertheless, (1) we found no evidence, at the mRNA level, of upregulation of the monoacylglycerol O-acyltransferase pathway that can generate DAG from monoacylglycerol [46, 47] (data not shown); and (2) cholesteryl esters levels are likely not upregulated since the effect of lipin knockdown on neutral lipid content looks similar by Oil Red O staining (ORO) and TAG content (ORO: 0.76 ± 0.06, and TAG: 0.85 ± 0.13 in the lipin-1 knockdown, and ORO: 0.74 ± 0.12, and TAG: 0.54 ± 0.33 in the triple lipin knockdown, compared with the control set as 1). In contrast, LPP activity was induced in the lipin knockdowns, possibly due to an earlier upregulation of their transcript levels (observed by day 4). Moreover, inhibition of both lipins and LPPs with propranolol completely blocked lipid droplet formation, although this fact must be taken with caution because of the non-specificity of this compound. LPPs hydrolyse phosphatidate as well as different substrates, and act on the outer surface of plasma membrane and in the luminal surface of endoplasmic reticulum (ER) and Golgi membranes [6] and thus, it is not clear if they can have access to the phosphatidate formed from glycerol phosphate and acyl-CoA. However, in yeast, TAG synthesised both in the cytosolic and luminal leaflets of the ER membranes are efficiently packed into lipid droplets [48].

Another strategy to protect the lipid storage when glycerolipid synthesis is compromised might be downregulation of lipid hydrolysis and FAO. Lipin-1 regulates basal lipolysis [49], and modulates FAO transcript expression levels [9]. We show that all combinatorial knockdowns of lipins downregulate basal lipolysis to a greater extent than reduction of neutral lipid content.

In summary, we confirm that adipose tissue expression of the lipin family is altered in type 2 diabetes. Furthermore, loss of nearly all PAP1 activity, due to combined lipin-1, -2 and -3 knockdown, has only mild effects on final adipogenesis and lipid accumulation in SGBS cells. Conversely, lipin-2 and lipin-3 may contribute little to total PAP activity, but still play a role in early adipogenesis. Our results suggest a compensation strategy to accumulate near normal neutral lipid levels, activating other pathways (such as LPPs) and inhibiting TAG hydrolysis. More work is clearly required to decipher whether LPPs or other unknown pathways may compensate lack of PAP1 activity.

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Author contributions SS, AZ and MM were the main contributors in the conception, design and interpretation of the data, and in writing the manuscript. AT, HS, G-SH, DS, JC, CM and MM performed the experiments and data analysis, and revised the manuscript. JG, MW, CR, OY and GMC were responsible for acquisition of data and analysis, and revised the manuscript. All authors had final approval of the submitted and published versions. MM is the guarantor of this work and, as such, had full access to all the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

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