Transcriptional regulation of phospholipid biosynthesis is linked to fatty acid metabolism by an acyl-CoA-binding-protein-dependent mechanism in *Saccharomyces cerevisiae*

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In the present study, we have used DNA microarray and quantitative real-time PCR analysis to examine the transcriptional changes that occur in response to cellular depletion of the yeast acyl-CoA-binding protein, Acb1p. Depletion of Acb1p resulted in the differential expression of genes encoding proteins involved in fatty acid and phospholipid synthesis (e.g. *FAS1*, *FAS2*, *ACC1*, *OLE1*, *INO1* and *OPI3*), glycolysis and glycerol metabolism (e.g. *GPD1* and *TDH1*), ion transport and uptake (e.g. *ITR1* and *HNMI*) and stress response (e.g. *HSP12*, *DDR2* and *CTT1*). In the present study, we show that transcription of the *INO1* gene, which encodes inositol-3-phosphate synthase, cannot be fully repressed by inositol and choline, and UAS_{INO1} (inositol-sensitive upstream activating sequence)-driven transcription is enhanced in Acb1p-depleted cells. In addition, the reduction in inositol-mediated repression of *INO1* transcription observed after depletion of Acb1p appeared to be

independent of the transcriptional repressor, Opi1p. We also demonstrated that *INO1* and *OPI3* expression can be normalized in Acb1p-depleted cells by the addition of high concentrations of exogenous fatty acids, or by the overexpression of *FAS1* or *ACC1*. Together, these findings revealed an Acb1p-dependent connection between fatty acid metabolism and transcriptional regulation of phospholipid biosynthesis in yeast. Finally, expression of an Acb1p mutant which is unable to bind acyl-CoA esters could not normalize the transcriptional changes caused by Acb1p depletion. This strongly implied that gene expression is modulated either by the Acb1p–acyl-CoA ester complex directly or by its ability to donate acyl-CoA esters to utilizing systems.

Key words: acyl-CoA-binding protein (ACBP), fatty acid, inositol, phospholipid, transcriptional regulation.

INTRODUCTION

In *Saccharomyces cerevisiae*, fatty acids and fatty acid derivatives have been suggested to function as regulators of gene expression. This suggestion is supported by the fact that two oleate-activated transcription factors, Oaf1p and Pip2p, have been shown to be essential for fatty-acid-mediated induction of genes encoding peroxisomal enzymes and proteins [1]. Furthermore, transcriptional regulation of the yeast Δ^9 desaturase *OLE1* by unsaturated fatty acids involves Spt23p and Mga2p [2].

It has been suggested that the regulatory properties of fatty acids are mediated through their activation to acyl-CoA esters, since fatty acids are unable to repress the expression of *ACC1* [3] and *OLE1* [4] and induce expression of the peroxisomal enzymes, acyl-CoA oxidase (*POX1*) and acyl-CoA synthetase (*FAA2*) [5] in *S. cerevisiae* depleted of the two acyl-CoA synthetases, Faa1p and Faa4p. However, disruption of both *FAA1* and *FAA4* affects fatty acid import, and does not prove that fatty acid activation is necessarily required for fatty-acid-mediated regulation of gene expression.

In *S. cerevisiae*, LCACoA (long-chain acyl-CoA) esters are either synthesized endogenously by the FAS (fatty acid synthase) complex or produced from exogenous fatty acids imported by the fatty acid import machinery. Inside the cell, acyl-CoA esters are bound to proteins in order to increase their solubility and to protect

them from hydrolysis by thioesterases. A number of intracellular proteins have been reported to bind LCACoA esters, including L-FABP (liver fatty-acid-binding protein), SCP (sterol carrier protein)-2 and ACBP (acyl-CoA-binding protein). L-FABP is predicted to have a role in LCACoA ester metabolism, and this is highlighted by the fact that L-FABP-knockout mice display an altered cytosolic LCACoA ester-binding capacity and ester acyl-chain distribution [6]. Similarly, SCP-2/SCP-X-knockout mice demonstrate impaired catabolism of methyl-branched fatty acyl-CoAs, inefficient transport of phytanoyl-CoA into peroxisomes and defective thiolytic cleavage of 3-oxopristanoyl-CoA [7]. However, *S. cerevisiae* does not express fatty-acid-binding protein [8] or a SCP-2 homologue [9]. Thus *S. cerevisiae* is suited for use as a model to study the function of ACBP, since potential compensatory effects caused by L-FABP and/or SCP-2 can be avoided.

Yeast ACBP, Acb1p, belongs to a large multi-gene family encoding a protein approx. 10 kDa in size. Acb1p binds LCACoA esters in a non-covalent and reversible manner with very high affinity (K_d of 1–10 nM) and specificity [10]. ACBP has been very highly conserved throughout evolution, and it has been detected in all eukaryotic organisms examined so far [11]. The high degree of sequence and structural conservation of ACBP among eukaryotic species, and the fact that human ACBP can complement ACBP function in yeast (S. Feddersen, J. Knudsen and N. J. Færgeman,

Abbreviations used: ACBP, acyl-CoA-binding protein; COP, cytidyl diphosphate; Cy3, indocarbocyanine; Cy5, indocarbocyanine; ER, endoplasmic reticulum; ESI, electrospray ionization; FAS, fatty acid synthase; HA, haemagglutinin; HNF-4 α , hepatocyte nuclear factor-4 α ; HRP, horseradish peroxidase; LCACoA, long-chain acyl-CoA; L-FABP, liver FABP; LC, liquid chromatography; Opi[−], overproduction of inositol; PA, phosphatidic acid; PC, phosphatidylcholine; PE, phosphatidylethanolamine; PG, phosphatidylglycerol; PI, phosphatidylinositol; PS, phosphatidylserine; Q-RT-PCR, quantitative real-time PCR; SC, synthetic complete; SCP, sterol carrier protein; UAS_{INO1}, inositol-sensitive upstream activating sequence; UPR, unfolded protein response; YNB, yeast nitrogen base; YNBD, YNB and dextrose; YPD, yeast extract, peptone and dextrose; YPgal, yeast extract, peptone and galactose.

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Table 1 Strains used in the present study

Strain	Genotype	Source or reference
Y700	<i>MATa, ade2-1 trp1 can1-100 leu2-112 his3-11 his3-15 ura3</i>	[16]
Y700 <i>pGAL-ACB1</i>	<i>MATa, ade2-1 trp1 can1-100 leu2-112 his3-11 his3-15 ura3, pGAL-ACB1::KanMX</i>	[16]
Y700 <i>INO1-HA₃</i>	<i>MATa, ade2-1 trp1 can1-100 leu2-112 his3-11 his3-15 ura3, INO1-HA₃::His3MX</i>	Present study
Y700 <i>pGAL-ACB1 INO1-HA₃</i>	<i>MATa, ade2-1 trp1 can1-100 leu2-112 his3-11 his3-15 ura3, pGAL-ACB1::KanMX, INO1-HA₃::His3MX</i>	Present study
Y700 <i>scs2Δ</i>	<i>MATa, ade2-1 trp1 can1-100 leu2-112 his3-11 his3-15 ura3, SCS2::TRP1</i>	Present study
Y700 <i>pGAL-ACB1 scs2Δ</i>	<i>MATa, ade2-1 trp1 can1-100 leu2-112 his3-11 his3-15 ura3, pGAL-ACB1::KanMX, SCS2::TRP1</i>	Present study
AID	<i>MATa/MATα, ade1/ade1 ino1/ino1</i>	S. Henry*
Y700 <i>pld1Δ</i>	<i>MATa, ade2-1 trp1 can1-100 leu2-112 his3-11 his3-15 ura3, PLD1::TRP1</i>	Present study
Y700 <i>pGAL-ACB1 pld1Δ</i>	<i>MATa, ade2-1 trp1 can1-100 leu2-112 his3-11 his3-15 ura3, pGAL-ACB1::KanMX, PLD1::TRP1</i>	Present study
Y700 <i>hac1Δ</i>	<i>MATa, ade2-1 trp1 can1-100 leu2-112 his3-11 his3-15 ura3, HAC1::URA3</i>	Present study
Y700 <i>pGAL-ACB1 hac1Δ</i>	<i>MATa, ade2-1 trp1 can1-100 leu2-112 his3-11 his3-15 ura3, pGAL-ACB1::KanMX, HAC1::URA3</i>	Present study

* AID was a gift from Professor Susan Henry, Department of Molecular Biology and Genetics, Cornell University, Ithaca, NY, U.S.A. [22]

unpublished work), suggest that its function is associated with one or more basal cellular function(s) common to all cells [11].

In vitro experiments have shown that ACBP can attenuate the inhibitory effect of LCACoA esters on acetyl-CoA carboxylase and the mitochondrial adenine nucleotide translocase, stimulate the mitochondrial long-chain acyl-CoA synthetase, and extract membrane-imbedded acyl-CoA esters and donate them to utilizing systems such as glycerolipid synthesis and β -oxidation [12,13]. Furthermore, it has been demonstrated that overexpression of Acb1p and bovine ACBP in *S. cerevisiae* increased the acyl-CoA ester pool size [14,15]. Thus ACBP may act as an acyl-CoA ester pool former and transporter *in vivo*. Despite this, the precise *in vivo* function of ACBP is currently unknown. Yeast cells which lack Acb1p activity display a variety of defects, including fragmented vacuoles, a multi-layered plasma membrane, accumulation of vesicles of variable sizes, a 3–5.5-fold elevation in the *OLE1* mRNA level, and reduced levels of very-long-chain fatty acids, ceramides and sphingolipids [4,16,17]. Owing to the high affinity of ACBP for acyl-CoA esters and the presence of cytosolic acyl-CoA hydrolases, the intracellular free acyl-CoA ester concentration is predicted to be very low [10]. It can therefore be expected that the Acb1p–acyl-CoA ester complex can modulate processes which are regulated by acyl-CoA esters and processes which consume acyl-CoA esters, either directly or indirectly through the donation of acyl-CoA esters.

In the present study, we used whole genome cDNA microarrays and Q-RT-PCR (quantitative real-time PCR) in order to identify transcriptional changes in response to the cellular depletion of Acb1p. Current knowledge suggested that depletion of Acb1p changes the expression of genes encoding proteins involved in fatty acid and phospholipid synthesis, glycolysis, glycerol metabolism, ion transport/uptake and the stress response. Moreover, the present study has identified an Acb1p-dependent connection between fatty acid metabolism and transcriptional regulation of phospholipid biosynthesis in yeast. It also demonstrated that the regulatory effects of Acb1p depended on its ability to bind acyl-CoA esters.

EXPERIMENTAL

Chemicals

Bacto peptone, yeast extract and YNB (yeast nitrogen base) were purchased from Difco (BD Diagnostic Systems). HRP (horseradish peroxidase)-conjugated goat anti-(mouse IgG) and goat anti-(rabbit IgG) secondary antibodies were obtained from Promega. Restriction enzymes and DNA/RNA-modifying

enzymes were either from New England Biolabs or Invitrogen. All chemicals were of analytical grade.

Yeast strains, media and growth

The *S. cerevisiae* strains used in the present study are listed in Table 1. Y700 *pGAL-ACB1* is a conditional knockout strain constructed by insertion of the *GAL1* promoter in front of the *ACB1* gene, as described by Gaigg et al. [16]. In the wild-type strain, Y700, and the Y700 *pGAL-ACB1* strain, single disruptions in *SCS2* and *PLD1* (*SPO14*), in which the target ORF (open reading frame) was disrupted by the *TRP1* marker, were constructed by PCR-mediated gene replacement using pFA6a-*TRP1* as a template (a gift from Professor John R. Pringle, Department of Biology, The University of North Carolina, Chapel Hill, NC, U.S.A.) as described previously [18]. Disruption of the *HAC1* gene was achieved by homologous recombination of the BamHI-linearized pHAO1 plasmid (a gift from Professor Peter Walter, Howard Hughes Medical Institute, University of California, San Francisco, San Francisco, CA, U.S.A.) into the wild-type strain, Y700, and the Y700 *pGAL-ACB1* strain. Correct integration of *TRP1* PCR products and pHAO1 in Y700 and Y700 *pGAL-ACB1* cells was confirmed by PCR. Media used to grow yeast included YPD (yeast extract, peptone and dextrose), YPgal (yeast extract, peptone and galactose) and YNBD (YNB and dextrose), made as described previously [17], and glucose-supplemented SC (synthetic complete) medium, made as described by Sherman [19]. *Myo*-inositol was either omitted from SC medium or added to the concentration indicated as required. Choline was added to a final concentration of 1 mM, and palmitic acid or palmitoleic acid was added to a final concentration of 500 μ M where indicated. Yeast strains were established on YPgal or YNBgal (YNB and galactose) medium and cultivated at 30 °C. Prior to experimentation, glucose-supplemented SC, YNBD or YPD medium was inoculated with yeast cells to a D_{600} (attenuance) of 0.005–0.01 and grown for 24 h. These cultures were diluted into fresh medium to a D_{600} of 0.05–0.1 and grown to the required D_{600} . Depletion of Acb1p was confirmed by Western blotting after each experiment.

Protein extraction, electrophoresis and Western blotting

Protein extraction, electrophoresis and Western blotting was performed as described in [17], with the exception that PVDF membranes were used instead of nitrocellulose membranes. Mouse influenza HA (haemagglutinin) antiserum was obtained from Roche Diagnostics and used at a 1:5000 dilution. HRP-conjugated goat anti-(mouse IgG) and goat anti-(rabbit IgG) secondary antibodies were both used at a concentration of 1:8000.

DNA microarray

Yeast DNA microarrays were purchased from MWG. A total of 6250 gene-specific oligonucleotide probes (40-mers) representing the complete *S. cerevisiae* genome were spotted on to one single epoxy-coated slide (1.8 cm × 3.6 cm). In addition, each MWG Yeast Array contained 86 replicated spots and 32 *Arabidopsis* control spots.

Yeast culture and total RNA isolation

Wild-type (Y700) and conditional *ACB1*-knockout (Y700 *pGAL-ACB1*) cells were grown to a D_{600} of 0.9–1.3 in YPD, YNBD or glucose-supplemented SC medium as described above. Following growth, cells were collected by centrifugation at 4500 *g* for 10 min at 4 °C in 50 ml Falcon tubes half-filled with crushed ice. The supernatant was removed, and cells were transferred to 1.5 ml microcentrifuge tubes and pelleted by centrifugation at 12 800 *g* for 3 min at 4 °C. Harvested cells were frozen in liquid nitrogen and stored at –80 °C. Total RNA was extracted using the hot phenol method as described previously [20]. DNase I digestion of the extracted RNA was performed following the manufacturer's instructions (MWG).

cDNA synthesis and post-labelling

RNA was isolated from Y700 and Y700 *pGAL-ACB1* cells grown in YPD medium. For cDNA synthesis and post-labelling, the CyScribe cDNA post-labelling kit (Amersham Biosciences) was used. cDNA synthesis and post-labelling was performed as described by the manufacturer (Amersham Biosciences), except that amino allyl-modified cDNA was purified using the Qiagen QIAquick PCR purification kit, in which the supplied wash and elution buffers were replaced with phosphate wash buffer [5 mM potassium phosphate buffer, pH 8.0, and 80% (v/v) ethanol] and phosphate elution buffer (4 mM potassium phosphate buffer, pH 8.5) respectively to avoid production of free amines.

Hybridization, washing and scanning

For hybridization, a formamide-based hybridization buffer (MWG) was used. Cy3 (indocarbocyanine)- and Cy5 (indocarbocyanine)-labelled cDNA was pooled, dried, resuspended in 30 µl of pre-heated (42 °C for 10 min) formamide-based buffer (MWG) and then incubated at 95 °C for 3 min. This hybridization mixture was centrifuged briefly at 10 000 *g* for 15 s at 22 °C and applied to the yeast DNA microarray under a clean coverslip. Slides were then placed into a wet hybridization chamber (GeneMachines®) and incubated for 18–22 h in a 42 °C water bath. After incubation, the microarrays were washed as described by the manufacturer (MWG). The arrays were then dried by centrifugation at 500 *g* for 2 min at room temperature (22 °C) and scanned at 532 nm (Cy3) and 635 nm (Cy5) in a GMS-418 scanner (Genetic Microsystems). Overlay images were constructed using the microarray spot-finding and quantification software, ImaGene version 4.2 (BioDiscovery).

Experimental setup and data analysis

RNA isolated from five different cultures of either Y700 or Y700 *pGAL-ACB1* cells grown in YPD medium was pooled in two different tubes. cDNA synthesized from Y700 and Y700 *pGAL-ACB1* RNA was labelled with Cy3 and Cy5 respectively. The experiment was repeated on independent cultures with opposite Cy3/Cy5 labelling. Overlay images were processed using ImaGene version 4.2 software (BioDiscovery). Fluorescent spots were located, and extracted intensity values were corrected for local background by subtracting the intensity level around

each spot from the spot's signal intensity. Intensities corrected for background were processed and analysed further using Acuity version 4.0 software (Axon Instruments). Data from each microarray were normalized using print-tip locally weighted linear regression (lowess) normalization, and average $\log_2 (R/G)$ values (M values) were calculated. In dual-colour experiments, R and G represent the processed and normalized intensity values measured using the red and green channel respectively. Subsequently, data were filtered in order to remove unreliable data from the dataset. Quality measurements were collected using ImaGene 4.2 and used to identify low-quality spots. Two quality parameters were measured for each spot: the number of pixels that touch the signal area which are neither signal nor background and all of the ignored and background pixels that fall within the circle defined by the spot-finding method. Spots defined by this quality control method to be 'bad quality' and data with intensity levels below the highest intensity level of the 32 *Arabidopsis* control spots at 532 nm (Cy3) and 635 nm (Cy5) were removed. Furthermore, data that were not present in both microarrays were also excluded. M values were then transferred to Microsoft Excel and the S.D. was calculated. Data with S.D. > 0.6 were removed. However, to avoid removing data which represented genes which were highly up- or down-regulated in both arrays but had S.D. > 0.6, the list of removed data was checked for such important data. Finally, genes considered to be differentially expressed were identified using a fold-change filter, in which the cut-off was set to an absolute M value of 0.70 (which corresponded to a 1:1.6 ratio of expression between wild-type and *Acb1p*-depleted cells).

Q-RT-PCR, RNA isolation and cDNA synthesis

Total RNA was isolated from cells cultured and harvested as described above. RNA was extracted using the hot phenol method as described previously [20] and treated with DNase I, following the manufacturer's instructions (MWG). cDNA was synthesized using Superscript II reverse transcriptase (Invitrogen) and random hexamers (deoxy-NTP₆) (Amersham Biosciences). Q-RT-PCR was performed on an ABI PRISM 7700 RealTime PCR-machine (Applied Biosystems) using 2×SYBR Green JumpStart™ Taq ReadyMix™ and Sigma Reference Dye (Sigma–Aldrich), following the manufacturer's instructions. Reactions were incubated at 95 °C for 2 min, followed by 40 cycles of 95 °C for 15 s, 60 °C for 45 s and 72 °C for 45 s. Fluorescence signals were measured during the elongation step, and all reactions were performed in duplicate. The level of the target mRNAs was examined and normalized to the level of the *ACT1* gene (actin) using the $2^{-\Delta\Delta CT}$ method [21]. Primers for Q-RT-PCR were designed using Primer Express version 2.0 (Applied Biosystems), and the efficacy and specificity of the primers was tested by dilution experiments and melting curves respectively. Used primer sets are listed in Supplementary Table S1 (<http://www.BiochemJ.org/bj/407/bj4070219add.htm>). Primer pairs were designed to amplify a fragment of approx. 50 bp, except for the *HAC1* primer pairs which amplified a fragment of 91 bp.

β-Galactosidase assay in yeast

Y700 and Y700 *pGAL-ACB1* cells transformed with plasmid pJH359 carrying an *INO1-CYC1-lacZ* fusion (a gift from Professor Susan Henry, Department of Molecular Biology and Genetics, Cornell University, Ithaca, NY, U.S.A. [22]) were grown overnight in an appropriate medium, diluted to a D_{600} of 0.05–0.1 and grown to a D_{600} of 0.6–0.8. β-Galactosidase activity in yeast was determined as described previously [23]. β-Galactosidase

activity (in units) was calculated according to the following equation:

$$\text{Activity} = A_{420}/(D_{600} \text{ of assayed culture} \times \text{volume assayed} \times \text{time})$$

where A_{420} is the absorbance of ONP (*o*-nitrophenol), D_{600} is the attenuation of the culture at the time of assay, volume is the amount of the culture used in the assay (in ml) and time is measured in min.

Complementation of *INO1* expression by increased fatty acid synthesis

Increased fatty acid synthesis was achieved by transforming Y700 (wild-type) and the Y700 *pGAL-ACB1* strains with high-copy plasmids expressing either *ACC1* (pG1M-ACC1, a gift from Keiji Furukawa, Kiku-Masamune Sake Brewing Co., Kobe, Japan [24]) or *FAS1* (pJS229, a gift from Professor Hans-Joachim Schüller, Universität Greifswald, Greifswald, Germany [25]). The corresponding empty vectors were transformed to be used as controls. Total RNA was extracted from exponentially growing cells cultured in selective SC medium supplemented with 0.2 mM inositol and used for Q-RT-PCR analysis.

Assay for the Opi⁻ (overproduction of inositol) phenotype

To test for the Opi⁻ phenotype, Acb1p-depleted cells and wild-type cells were dotted on to inositol-free SC plates and grown at 30 °C. Plates were then sprayed with a suspension of a diploid tester strain, AID (provided by Professor Susan Henry), which is homozygous for *ino1* and *adel1*, and incubated at 30 °C as previously described [26].

LC (liquid chromatography)–ESI (electrospray ionization) MS of lipids

The number of Acb1p-depleted cells at a given D_{600} , is reduced compared with wild-type cells, implying that Acb1p-depleted cells are bigger than wild-type cells (S. Feddersen, J. Knudsen and N. J. Færgeman, unpublished work). Thus whenever quantifying the relative phospholipid content of Acb1p-depleted cells compared with wild-type cells, we corrected for differences in cell numbers. Lipids were extracted from 25 D_{600} units of exponentially growing cells (D_{600} of ~1) using procedure IIIA as described by Hanson and Lester [27]. Aliquots corresponding to 10 D_{600} units of cells were dissolved in 65 °C solvent A (chloroform/methanol/water, 16:16:5, by vol.) and mixed with internal standard lipids extracted from 10 D_{600} units of cells grown on ^{13}C -glucose as the sole carbon source. The solvent was evaporated under a stream of nitrogen, and the lipids were redissolved in 10 μl of solvent A (heated to 65 °C) by stirring using a rotating pestle at 30 000 rev./min for 5 min. Immediately after mixing, 1 μl of the lipid solution was injected on to a PVA SIL HPLC column (1 mm $\times$ 150 mm, YMC Europe) and eluted at 45 $\mu\text{l}/\text{min}$. Solvents B (hexane/propan-2-ol, 98:2, v/v), C (chloroform/propan-2-ol, 65:35, v/v) and D (methanol) were changed linearly over time to give the proportions (by vol.) of B/C/D at the following time points: 0 min, 100:0:0; 22 min, 12:88:0; 30 min 10:74:16; 45 min, 8:61:31; 50–65 min, 0:0:100; 70–72.5 min, 0:100:0, and 75–82.5 min, 100:0:0. Ions in the effluent were ionized by ESI with an electrode potential of 3500 V, and the masses were detected using a Bruker Esquire-LC ion-trap mass spectrometer. The $^{12}\text{C}/^{13}\text{C}$ ratio of ion counts for each individual ^{12}C species and one selected ^{13}C species in each lipid class were calculated for both strains. Differences in the relative amount of individual lipid species between wild-type and Acb1p-depleted cells were calculated by dividing the wild-type $^{12}\text{C}/^{13}\text{C}$

ratio with the corresponding $^{12}\text{C}/^{13}\text{C}$ ratio of Acb1p-depleted cells. Species percentage composition, based on the sum of ion counts for all species within a certain lipid class, was calculated for all lipid classes. The content of a given phospholipid class in Acb1p-depleted cells was normalized to wild-type levels (wild-type = 100 %) and calculated using the following equation:

$$\text{Content} = \sum \{ \% \text{ individual phospholipid species in Wt} / [(\text{Wt}^{12}\text{C}/^{13}\text{C}) / (\Delta^{12}\text{C}/^{13}\text{C})] \}$$

where Δ is Y700 *pGAL-ACB1* and wild-type (Wt) is Y700.

RESULTS

Genome-wide changes in gene expression of *S. cerevisiae* in response to Acb1p depletion

DNA microarray profiling revealed that the effect of Acb1p depletion on gene expression is not restricted to *OLE1* [4,16]. A total of 134 genes were identified as being differentially expressed more than 1.6-fold, of which 44 genes were down-regulated and 90 genes were up-regulated in Acb1p-depleted cells compared with wild-type cells.

Confirmation of microarray data by Q-RT-PCR

Expression of 22 selected genes, of which ten had been shown to be differently expressed by microarray analysis, was examined by Q-RT-PCR (Table 2). The ten genes identified by microarray analysis were all shown to be differentially expressed to the same degree by Q-RT-PCR analysis (Table 2). We propose that the identified changes in gene expression measured by microarray analysis may reflect actual changes in mRNA levels, in response to Acb1p depletion. However, in the present study, we primarily focus on microarray data which have been confirmed by Q-RT-PCR.

Genes associated with lipid metabolism

The present microarray analysis and Q-RT-PCR analysis show that genes encoding proteins involved in fatty acid biosynthesis (*ACC1*, *FAS1*, *FAS2* and *OLE1*) all are up-regulated in Acb1p-depleted cells compared with wild-type cells (Table 2 and see Supplementary Table S2 at <http://www.BiochemJ.org/bj/407/bj4070219add.htm>). Consistent with previous findings, the gene encoding the yeast Δ^9 desaturase *OLE1*, was found to be up-regulated approx. 4-fold in Acb1p-depleted cells cultured in YNBD medium (results not shown) and approx. 2-fold in cells cultured in YPD medium (Table 2) [4,16].

The microarray and Q-RT-PCR data also revealed that genes encoding enzymes involved in phospholipid biosynthesis (*INO1*, *CHO1*, *CHO2* and *OPI3*) are up-regulated in Acb1p-depleted cells (Table 2). Moreover, the expression of both *PSD1* and *PSD2*, which also encode phospholipid biosynthesis enzymes, was increased 1.3-fold in Acb1p-depleted cells compared with wild-type cells (Table 2), as observed by Q-RT-PCR. This suggests that synthesis of glycerophospholipids may be affected in cells depleted of Acb1p. In particular, the *INO1* gene encoding the inositol-3-phosphate synthase (Ino1p) that catalyses synthesis of inositol-3-phosphate, the rate-limiting and committed step in the synthesis of PI (phosphatidylinositol), was strongly affected by Acb1p depletion. Although below the detection level of the microarrays, *INO1* mRNA levels were detected by Q-RT-PCR and found to be up-regulated approx. 150-fold after Acb1p depletion (Table 2). *INO1* mRNA and Ino1p levels were investigated further under different growth conditions.

Table 2 Comparison of fold change (Acb1p depleted cells compared with wild-type cells) obtained by microarray and Q-RT-PCR analysis

Gene ID	Gene	Function	Process	Fold change*	
				Microarray analysis	Q-RT-PCR†
Stress response					
ygr088w	<i>CTT1</i>	Catalase T, cytosolic	Response to stress	No data‡	3.1 ± 0.14
ybr072w	<i>HSP26</i>	Heat-shock protein	Response to stress	No data‡	5.9 ± 1.4
yol052c-a	<i>DDR2</i>	Heat-shock protein DDRA2	Response to stress	6.9 ± 0.71	2.7 ± 0.66
yil026w	<i>HSP104</i>	Heat-shock protein	Response to stress	No data‡	3.1 ± 0.41
yfi014w	<i>HSP12</i>	Heat-shock protein	Response to stress	No data‡	6.8 ± 1.6
C-compound and carbohydrate metabolism					
ydl022w	<i>GPD1</i>	Glycerol-3-phosphate dehydrogenase (NAD ⁺), cytoplasmic	Intracellular accumulation of glycerol	6.4 ± 1.8	5.5 ± 0.89
yjl052w	<i>TDH1</i>	Glyceraldehyde-3-phosphate dehydrogenase 1	Glycolysis; gluconeogenesis	4.8 ± 0.16	3.3 ± 0.91
Lipid metabolism					
ygl055w	<i>OLE1</i>	Stearyl-CoA desaturase	Fatty acid desaturation	1.9 ± 0.75	1.9 ± 0.45
yjl196c	<i>ELO1</i>	Fatty acid elongation protein	Fatty acid elongation	No data‡	1.2 ± 0.11
ykl182w	<i>FAS1</i>	Fatty-acyl-CoA synthase, β chain	Fatty acid biosynthesis	1.6 ± 0.25	1.6 ± 0.41
ypl231w	<i>FAS2</i>	Fatty-acyl-CoA synthase, α chain	Fatty acid biosynthesis	1.8 ± 0.17	1.6 ± 0.26
yhr007c	<i>ERG11</i>	Cytochrome P450 lanosterol 14 α -demethylase	Ergosterol biosynthesis	No data‡	-1.9 ± 0.10
yjl153c	<i>INO1</i>	Myo-inositol-3-phosphate synthase	PI biosynthesis	No data‡	151 ± 10.7
ygr170w	<i>PSD2</i>	Phosphatidylserine decarboxylase 2	PS biosynthesis	No data‡	1.3 ± 0.11
ygr157w	<i>CHO2</i>	Phosphatidylethanolamine <i>N</i> -methyltransferase	PC biosynthesis	1.8 ± 0.51	1.6 ± 0.02
yjr073c	<i>OPI3</i>	Methylene-fatty-acyl-phospholipid synthase	PC biosynthesis	2.9 ± 0.68	6.5 ± 0.55
ynl169c	<i>PSD1</i>	Phosphatidylserine decarboxylase 1	PS biosynthesis	No data‡	1.3 ± 0.17
ypr113w	<i>PIS1</i>	CDP-diacylglycerol-inositol 3-phosphatidyltransferase	PI biosynthesis	No data‡	1.1 ± 0.07
yer026c	<i>CHO1</i>	CDP-diacylglycerol serine <i>O</i> -phosphatidyltransferase	PS biosynthesis	1.8 ± 0.24	1.7 ± 0.14
Others					
ypl154c	<i>PEP4</i>	Aspartyl protease	Vacuolar protein, catabolism	1.8 ± 0.14	1.7 ± 0.32
yor061w	<i>CKA2</i>	CK2 α' chain	G ₂ /M and G ₁ /S transition etc.	No data‡	-1.2 ± 0.12
yfi031w	<i>HAC1</i>	Transcription factor	Unfolded protein response	No data‡	5.6 ± 0.39
ylr350w	<i>ORM2</i>	Strong similarity to ORM1	Response to unfolded protein	2.8 ± 0.43	Not determined
ycl043c	<i>PDI1</i>	Protein disulfide-isomerase precursor	Protein folding	2.2 ± 0.17	Not determined
yjl034w	<i>KAR2</i>	Nuclear fusion protein	Response to unfolded protein	1.9 ± 0.98	Not determined

* Positive values represent fold up-regulation in Acb1p-depleted cells compared with wild-type cells, whereas negative values represent fold down-regulation in Acb1p-depleted cells compared with wild-type cells. Data are means ± S.D. Q-RT-PCR, $n \geq 3$; microarray analysis, $n = 2$.

† For each Q-RT-PCR analysis, RNA from three or four different cell cultures were used for cDNA synthesis and mean values were calculated. *ACT1* was used as endogenous reference.

‡ Data are not given because value for intensities were below that of *Arabidopsis* controls or because data were not present in both (dye-reversal) arrays.

Figure 1 *INO1* mRNA and protein levels during different growth conditions in *Acb1p*-depleted cells

Q-RT-PCR analysis of *INO1* mRNA levels in wild-type (Y700, black bars) and *pGAL-ACB1* (*Acb1p*-depleted Y700, white bars) cells grown in minimal medium (SC) supplemented with choline and/or inositol as indicated (**A**) or in rich medium (YPD) (**C**). Immunoblot analysis of protein extracts from *S. cerevisiae* containing Ino1p-HA. Wild-type (Y700 Ino1p-HA) and *pGAL-ACB1* (*Acb1p*-depleted Y700 Ino1p-HA) cells were grown in SC medium supplemented with choline and/or inositol as indicated (**B**) or in YPD medium (**D**). The blots in (**B**) and (**D**) were re-probed with rabbit anti-Vac8p antibody (a gift from Professor Lois S. Weisman, Life Sciences Institute and Department of Cellular and Developmental Biology, University of Michigan, Ann Arbor, MI, U.S.A) (1:2000 dilution) to serve as loading control. Data in (**A**) and (**C**) are means $\pm$ S.D. ($n=3$) and Western blots are representative of three independent experiments.

Regulation of *INO1* expression in *Acb1p*-depleted cells

Most of the genes involved in phospholipid synthesis, e.g. *INO1*, *OPI3*, *PIS1*, *CHO1* and *CHO2*, are characterized by the presence of a *cis*-acting UAS_{INO1} (inositol-sensitive upstream activating sequence), also called an ICRE (inositol/choline-responsive element), in their promoters (for a review see [28]). Transcription of UAS_{INO1}-containing genes is positively regulated by Ino2p and Ino4p, which bind to UAS_{INO1} elements and activate transcription when cells are grown in the absence of inositol and choline. Repression occurs in the presence of high concentrations of inositol and choline and is mediated by Opi1p, which acts as a negative regulator (for a review see [29]). In the absence of inositol and choline, Opi1p is sequestered in the ER (endoplasmic reticulum) membrane by binding to PA (phosphatidic acid) and Scs2p [30]. On addition of inositol and choline, PA is quickly consumed for the synthesis of PI and PC (phosphatidylcholine), followed by translocation of Opi1p to the nucleus [30] where it represses UAS_{INO1}-dependent expression indirectly, presumably by interacting with the Ino2p transcriptional activator.

INO1 mRNA and the corresponding Ino1p protein level were both found to be significantly up-regulated in *Acb1p*-depleted cells in comparison with wild-type cells, when grown in both YPD and SC medium supplemented with 0.2 mM inositol (Figure 1). This suggests that full repression of *INO1* mRNA and Ino1p by inositol requires the presence of *Acb1p*. Consistent with this, we did not observe a significant difference in UAS_{INO1}-driven β -galactosidase activity between wild-type and *Acb1p*-depleted cells when cultured in inositol-free medium (results not shown). In contrast, when grown in YNBD medium supplemented with

Figure 2 UAS_{INO1}-driven gene expression is increased in *Acb1p*-depleted cells

Wild-type (Y700, black bars) and *pGAL-ACB1* (*Acb1p*-depleted Y700, white bars) cells were grown in YNBD medium (without inositol) or supplemented with inositol as indicated. Cells were transformed with the *INO1-CYC1-lacZ* plasmid and β -galactosidase activity was measured to assess the ability of UAS_{INO1} elements to drive *lacZ* expression in wild-type and *Acb1p*-depleted cells. β -Galactosidase activity was defined as described in the Experimental section. Data are means $\pm$ S.D. ($n=3$).

either 0.02 mM inositol or 0.2 mM inositol, UAS_{INO1}-driven β -galactosidase expression was approx. 3-fold higher in *Acb1p*-depleted cells compared with wild-type cells (Figure 2). Up-regulation of *INO1* transcription in *Acb1p*-depleted cells may be caused by increased UAS_{INO1}-driven transcription as a result of reduced inositol-mediated repression. This notion is supported by the fact that other genes containing UAS_{INO1} elements in

their promoter, including *OPI3*, *CHO1* and *CHO2*, are also up-regulated in Acb1p-depleted cells (Table 2). In a recent microarray study, a total of 36 Ino2p and Ino4p target genes were identified [31]. On Acb1p depletion, 20 of these genes were found to be up-regulated (at least 1.5-fold), three were down-regulated (at least 1.5-fold), and 13 genes were not shown to be differently expressed when compared with wild-type cells. This indicates that inositol-mediated repression of Ino2p and Ino4p target genes at steady-state depends on Acb1p. Since Ino2p, Ino4p and Opi1p regulate transcription of *UAS_{INO1}*-containing genes, mRNA levels of these transcription factors were measured by Q-RT-PCR. We found that the mRNA level of *INO2* and *INO4* were increased approx. 2-fold in Acb1p-depleted cells on comparison with wild-type cells grown in rich medium, whereas no significant changes was observed for cells grown in minimal medium (results not shown). The mRNA level of *OPI1* was increased approx. 1.3-fold in Acb1p-depleted cells compared with wild-type cells when grown in rich medium, whereas the mRNA level was decreased approx. 1.3-fold when cells were grown in minimal medium (results not shown).

Acb1p depletion reduces inositol-mediated repression of *INO1* by an Opi1p-independent regulatory mechanism

Since *INO1*, *CHO1*, *CHO2* and *OPI3* were identified to be up-regulated in Acb1p-depleted cells, we speculate that the transcriptional repressor Opi1p is sequestered in the ER membrane of these cells, as a result of higher levels of PA and/or Scs2p. Disruption of *SCS2* reduced *INO1* expression in both wild-type and Acb1p-depleted cells, but *INO1* mRNA levels were still increased approx. 60-fold (Figure 3A) and approx. 5.5-fold (Figure 3B) in Acb1p-depleted cells grown in YPD medium or SC medium containing 0.2 mM inositol respectively. A similar expression pattern was observed for *OPI3* on disruption of *SCS2* (results not shown). Moreover, *SCS2* mRNA levels were unaffected by Acb1p depletion (results not shown), and relative PA levels were unchanged or lower in Acb1p-depleted cells compared with wild-type cells (Table 3). Finally, overexpression of *OPI1* did not reduce *INO1* mRNA levels of Acb1p-depleted cells to that of wild-type cells (results not shown). Taken together, these observations suggest that Acb1p depletion reduces inositol-mediated repression of *INO1* and other Ino2p and Ino4p target genes by an Opi1p-independent regulatory mechanism.

Acb1p-depleted *S. cerevisiae* exhibit an Opi⁻ phenotype, but do not accumulate PA

The Opi⁻ phenotype has previously been associated with deregulation of the *INO1* gene [32]. Therefore we tested Acb1p-depleted cells for an Opi⁻ phenotype [32]. Growth of an inositol-dependent strain around Acb1p-depleted cells grown on synthetic inositol-free plates, shows that Acb1p depletion leads to inositol secretion (Figure 4A). Moreover, it has been shown that strains carrying deletions in *CHO1*, *CHO2* or *OPI3* have defects in PC synthesis, are unable to repress *INO1* in the presence of inositol and exhibit a conditional Opi⁻ phenotype, owing to the accumulation of PA [32,33]. Expression of *INO1* and the Opi⁻ phenotype can be normalized in these strains by the addition of exogenous choline, which alleviates PA accumulation by increasing PC synthesis via the CDP (cytidyldiphosphate)-choline pathway [28,34]. Thus if accumulation of PA causes increased *INO1* expression in Acb1p-depleted cells, supplementation of both choline and inositol would be expected to repress *INO1* mRNA and Ino1p levels further in both wild-type and Acb1p-depleted cells, compared with addition of inositol alone. However, addition of choline to

Figure 3 *INO1* mRNA levels in wild-type and Acb1p-depleted cells with *SCS2*, *PLD1* or *HAC1* disrupted

Q-RT-PCR analysis of *INO1* mRNA levels in wild-type (Y700) and pGAL-ACB1 (Acb1p-depleted Y700) cells disrupted in *SCS2*, *PLD1* or *HAC1* genes, following growth in rich (YPD) medium (A) or minimal (SC) medium supplemented with 0.2 mM inositol (B). Data are means $\pm$ S.D. ($n = 3$).

inositol-containing medium resulted in only a minor decrease in *INO1* mRNA and Ino1p levels in Acb1p-depleted cells compared with wild-type cells (Figures 1A and 1B). Hence increased *INO1* expression in Acb1p-depleted cells may not be caused by the accumulation of PA as a result of reduced PC synthesis by the PE (phosphatidylethanolamine) methylation pathway. Increased phospholipase D₁ (Pld1p)-mediated turnover of PC has also been reported to result in derepression of the *INO1* gene [26] probably by accumulation of PA. However, in a pld1-null background, *INO1* levels were still increased approx. 90-fold (Figure 3A) and approx. 4.5-fold (Figure 3B) in Acb1p-depleted cells grown in YPD medium and SC medium containing 0.2 mM inositol respectively. Increased PC turnover does not seem to cause derepression of *INO1* in Acb1p-depleted cells. This is in agreement with our previous data, which showed that PC turnover is only slightly reduced in Acb1p-depleted cells, compared with wild-type cells [16]. No increase was found in the relative levels of PC and PA in Acb1p-depleted cells when compared with wild-type cells (Table 3). Taken together, these results indicate that changes in PC metabolism or accumulation of PA are not responsible for increasing *INO1* mRNA and Ino1p levels in cells depleted of Acb1p.

The mechanism causing induction of *INO1* in Acb1p-depleted cells is independent of *HAC1* expression

Eukaryotic cells respond to the accumulation of unfolded proteins in the ER by activating a stress-response pathway known as

Table 3 The relative phospholipid content of Y700 *pGAL-ACB1* cells compared with Y700 cells

The content of a given phospholipid class in *Acblp*-depleted cells (Y700 *pGAL-ACB1*) was normalized to wild-type (Y700) levels (wild-type = 100%) and calculated using the following equation: Content = $\sum \{ \% \text{ individual phospholipid species in Wt} / [(Wt^{12C}/^{13C}) / (\Delta^{12C}/^{13C})] \}$, where Δ is Y700 *pGAL-ACB1* and wild-type (Wt) is Y700. Data are means $\pm$ S.D. ($n = 2$).

Medium	Phospholipid content (%)					
	PA	PI	PG	PS	PE	PC
YPD	95 $\pm$ 7.8	128 $\pm$ 15.2	89 $\pm$ 18.9	78 $\pm$ 2.5	129 $\pm$ 7.8	98 $\pm$ 15.5
SC medium + 0.2 mM inositol	81 $\pm$ 6.8	80 $\pm$ 4.4	46 $\pm$ 7.6	72 $\pm$ 20.8	80 $\pm$ 19.1	96 $\pm$ 18.1
SC medium without inositol	120 $\pm$ 22.5	223 $\pm$ 0.9	66 $\pm$ 8.4	125 $\pm$ 18.9	139 $\pm$ 23.5	100 $\pm$ 15.9

Figure 4 *Acblp*-depleted cells exhibit an *Opi⁻* phenotype

(A) Wild-type (Y700) and *pGAL-ACB1* (*Acblp*-depleted Y700) cells were tested for their ability to secrete inositol to the surrounding medium when supplemented with or without 0.5 mM palmitic acid. (B) Q-RT-PCR analysis of *INO1* mRNA levels in wild-type (Y700, black bars) and *pGAL-ACB1* (*Acblp*-depleted Y700, white bars) cells grown in YNB medium (without inositol) supplemented with or without 0.05% (v/v) Brij 58 and 0.5 mM palmitic acid as indicated. (C) Immunoblot analysis of protein extracts from *S. cerevisiae* expressing *Ino1p-HA*. Wild-type (Y700 *Ino1p-HA*) and *pGAL-ACB1* (*Acblp*-depleted Y700 *Ino1p-HA*) cells were grown in YNB medium (without inositol) supplemented with or without 0.05% (v/v) Brij 58 detergent and 0.5 mM palmitic acid as indicated. The blot was re-probed with an antibody against *Vac8p* to serve as a loading control. Data in (B) are means $\pm$ S.D. ($n = 3$) and the Western blot is representative of three independent experiments.

the UPR (unfolded protein response) pathway (for a review see [35]). In *S. cerevisiae*, the UPR pathway may also be activated in response to inositol starvation and it is coupled to inositol biosynthesis, as cells defective in UPR activation require exogenous inositol for growth and express low levels of *INO1* [36]. In yeast, it has been proposed that *Hac1p* negatively regulates the activity of *Opi1p* [37]. Interestingly, *HAC1*, *PDII*, *KAR2* and *ORM2* were identified in the present study as being up-regulated in cells depleted of *Acblp* (Table 2), which is indicative of

induction of the UPR pathway. Disruption of *HAC1* diminished *INO1* mRNA levels in wild-type cells and cells depleted of *Acblp* (Figure 3). However, *INO1* mRNA levels were still increased approx. 40-fold (Figure 3A) and approx. 5-fold (Figure 3B) in *Acblp*-depleted cells compared with wild-type cells when grown in YPD and SC medium containing 0.2 mM inositol respectively. These results clearly demonstrate that the observed reduction in inositol-mediated repression of *INO1* in cells lacking *Acblp* is not the result of induction of the UPR pathway. This suggestion is supported by a recent microarray study which showed that *Opi1p*-mediated repression of *INO1* in response to inositol cannot be prevented by constitutive activation of the UPR pathway [31].

Phospholipid analysis by LC-ESI MS/MS

To examine whether the observed induction of genes encoding enzymes involved in phospholipid biosynthesis in *Acblp*-depleted cells is caused by changes in the overall phospholipid content, we measured the relative levels of PC, PI, PE, PG (phosphatidylglycerol), PS (phosphatidylserine) and PA using LC-ESI-MS/MS, with ^{13}C -labelled yeast lipids used as internal standards. *Acblp*-depleted cells grown in YPD medium, exhibited an approx. 30% increase in the relative level of the major glycerolphospholipids, PI and PE, and unchanged levels of PC, PG and PA compared with wild-type cells (Table 3). However, the relative level of PS was decreased by approx. 20% in the *Acblp*-depleted cells. For *Acblp*-depleted cells grown in SC medium supplemented with 0.2 mM inositol, it was found that the relative levels of PA, PI, PS and PE was reduced by approx. 20%, whereas the PC level was unchanged compared with wild-type cells. In contrast, the relative PG level was decreased by approx. 50% (Table 3). These results indicate that a reduction in the relative phospholipid content is not solely responsible for the induction of phospholipid biosynthesis genes in *Acblp*-depleted cells, since only the relative PS level was consistently reduced under all conditions. Therefore *Acblp* may act downstream or independently of the signal produced through phospholipid metabolism to affect *INO1* expression. However, it should be noted that the relative phospholipid species composition is altered in *Acblp*-depleted cells compared with wild-type cells (see Supplementary Tables S3–S8 at <http://www.BiochemJ.org/bj/407/bj4070219add.htm>). *Acblp* depletion causes a shift from $C_{34:1}$ species towards $C_{34:2}$ and $C_{32:1}$ or $C_{32:2}$ species in the PI, PG, PS and PA classes, independently of whether cells are grown in YPD medium or SC medium supplemented with 0.2 mM inositol (see Supplementary Tables S3–S8). These results confirm that *Acblp* depletion leads to changes in the relative species composition of phospholipids, as reported previously [17]. Thus it cannot be excluded that changes in the relative species composition of phospholipids in *Acblp*-depleted cells results in the observed alteration in expression of genes encoding enzymes involved in phospholipid biosynthesis.

Figure 5 Fatty acid supplementation normalizes *INO1* and *OPI3* expression on *Acb1p* depletion

Q-RT-PCR analysis of *INO1* (A) and *OPI3* (B) mRNA levels in wild-type (Y700, black bars) and *pGAL-ACB1* (*Acb1p*-depleted Y700, white bars) cells is shown. Cells were grown in rich medium (YPD) containing 0.05% (v/v) Brij 58 detergent in the presence or absence of 0.5 mM palmitic acid ($C_{16:0}$) or 0.5 mM palmitoleic acid ($C_{16:1}$) as indicated. Data are means $\pm$ S.D. ($n=3$).

Supplementation with fatty acids or overexpression of *FAS1* or *ACC1* can normalize inositol-mediated repression of *INO1* in *Acb1p*-depleted cells

A link between fatty acid metabolism and *INO1* expression has previously been reported by Shirra et al. [38], who found that inhibition of acetyl-CoA carboxylase (*Acc1p*) activity restored expression of *INO1* in *SNF1* disrupted cells. Moreover, it was shown that supplementation with palmitoleic acid ($C_{16:1}$) relieved inositol auxotrophy and permitted *INO1* expression in cells lacking *SNF1* [38]. In addition, Gaspar et al. [39] suggested that the observed burst in PI synthesis following the addition of inositol depends on *de novo* fatty acid synthesis, further indicating a link between fatty acid biosynthesis and regulation of phospholipid biosynthesis. Addition of palmitic acid ($C_{16:0}$) to inositol-containing medium normalized *INO1* and *OPI3* expression in *Acb1p*-depleted cells to that of wild-type cells (Figure 5). In contrast, palmitoleic acid only partly normalized *INO1* and *OPI3* expression in *Acb1p*-depleted cells to levels observed in wild-type cells (Figure 5). Notably, palmitoleic acid increased *INO1* and *OPI3* expression in wild-type cells (Figure 5). This shows that palmitic acid and palmitoleic acid affect inositol-mediated repression of *Ino2p* and *Ino4p* target genes differently in both wild-type and *Acb1p*-depleted cells. However, a signal generated from fatty acid metabolism may be involved in the regulation of inositol-mediated repression of *Ino2p* and *Ino4p* target genes. Wenz et al. [25] showed previously that *FAS1*, but not *FAS2*, increased FAS activity 2.8-fold when expressed from a high-copy plasmid. We therefore hypothesized that ectopic expression from a high-copy plasmid of *FAS1* and *ACC1*, the rate-limiting enzyme in fatty acid biosynthesis, would increase fatty acid/acyl-

Figure 6 Ectopic overexpression of *ACC1* and *FAS1* decreases *INO1* expression in *Acb1p*-depleted cells

(A) Q-RT-PCR analysis of *INO1* mRNA levels in wild-type (Y700, black bars) and *pGAL-ACB1* (*Acb1p*-depleted Y700, white bars) cells transformed in pairs with combinations of empty vectors or construct to be expressed as indicated. Transformants were grown in selective medium (SC – uracil – tryptophan + 0.2 mM inositol). (B) Q-RT-PCR analysis of *INO1* mRNA levels in wild-type (Y700, black bars) and *pGAL-ACB1* (*Acb1p*-depleted Y700, white bars) cells transformed with either empty vector or construct to be expressed as indicated. Transformants were grown in selective medium (SC – tryptophan + 0.2 mM inositol). Data are means $\pm$ S.D. ($n=3$).

CoA ester availability and thereby reduce *INO1* expression in *Acb1p*-depleted cells. Overexpression of *FAS1* reduced the ratio of *INO1* mRNA levels in wild-type cells and *Acb1p*-depleted cells from approx. 1:6.5 to approx. 1:2.6 (Figure 6A). No further effect on *INO1* expression was observed when *FAS1* and *ACC1* were overexpressed at the same time (Figure 6A). Notably, when *ACC1* was overexpressed in cells transformed with the empty *FAS1* vector, the ratio of *INO1* mRNA levels in wild-type cells and *Acb1p*-depleted cells was only reduced from approx. 1:6.5 to approx. 1:4.7 (Figure 6A). However, ectopic expression of *ACC1* from a high-copy plasmid in cells only transformed with this plasmid reduced the ratio of *INO1* mRNA levels in wild-type cells and *Acb1p*-depleted cells from approx. 1:7.4 to approx. 1:2.2 (Figure 6B). A possible explanation for this observation is that transformation with two plasmids might perturb the copy number of each one, and therefore the level of expression of individual proteins may be affected. *INO1* mRNA levels could not be reduced in *Acb1p*-depleted cells when *FAS2* was overexpressed, further supporting the hypothesis that the observed effects are caused by an increased availability of fatty acids/acyl-CoA esters (results not shown). Together, these results show that increased availability of fatty acids/acyl-CoA esters in *Acb1p*-depleted cells normalizes the expression of genes involved in phospholipid biosynthesis. Thus *Acb1p* might facilitate removal of *de novo* synthesized acyl-CoA esters from the yeast FAS under normal physiological conditions, thereby making them available for further metabolism and regulatory processes.

Figure 7 Acb1p affects gene expression in an acyl-CoA ester-dependent manner

Q-RT-PCR analysis of *INO1* mRNA levels in wild-type (Y700), pGAL-ACB1 (Acb1p-depleted Y700), pGAL-ACB1 + ACB1 (Acb1p-depleted Y700 complemented with ACB1) and pGAL-ACB1 + ACB1^{Y28F/K32A} (Acb1p-depleted Y700 complemented with mutant *acb1*) cells cultured in SC medium supplemented with 0.2 mM inositol. Data are means $\pm$ S.D. ($n=3$).

Interestingly, the addition of 500 μ M palmitic acid to inositol-free plates repressed the Opi⁻ phenotype (Figure 4A), in accordance with the observed effect of palmitic acid on inositol uptake [16] and membrane abnormalities (S. Feddersen, J. Knudsen and N. J. Færgeman, unpublished work) in Acb1p-depleted cells. However, palmitic acid was unable to reduce the high *INO1* mRNA and Ino1p levels of both wild-type and Acb1p-depleted cells cultured in inositol-free medium (Figures 4B and 4C). This implies that high *INO1* expression is not necessarily associated with the Opi⁻ phenotype.

The *in vivo* function of Acb1p depends on ligand binding

Depletion of Acb1p in yeast results in slow growth and aberrant organelle morphology, including fragmented vacuoles, multi-layered plasma membranes and the accumulation of vesicles of variable sizes [16,17]. Ectopic expression of Acb1p in Acb1p-depleted cells normalizes vacuole morphology (S. Feddersen, J. Knudsen and N. J. Færgeman, unpublished work) and *INO1* mRNA levels (Figure 7), whereas expression of Acb1p^{Y28F/K32A}, which exhibited a 1000-fold reduced affinity for acyl-CoA esters compared with wild-type Acb1p [40], was unable to restore *INO1* mRNA levels (Figure 7) and vacuole morphology (S. Feddersen, J. Knudsen and N. J. Færgeman, unpublished work). The minor effect of Acb1p^{Y28F/K32A} on *INO1* expression in Acb1p-depleted cells is likely to be a result of the protein being overexpressed and binding acyl-CoA esters with very low affinity. Ectopic expression of rat L-FABP, which binds acyl-CoA esters with a K_d of 14–1000 nM [41,42] in Acb1p-depleted cells was also unable to normalize the *INO1* mRNA level (results not shown). This demonstrates that the function of Acb1p is closely linked to its ability to bind acyl-CoA esters with high affinity and supports the hypothesis of Acb1p playing a role in acyl-CoA trafficking.

DISCUSSION

In the present study, we have identified transcriptional changes in response to Acb1p depletion in genes encoding proteins which are involved in fatty acid and phospholipid synthesis, glycolysis, stress response, glycerol metabolism and ion transport and uptake (Table 2 and Supplementary Table S2). Depletion of Acb1p was shown to increase transcription of *INO1*, *CHO1*, *CHO2* and *OPI3*, all of which encode proteins required for phospholipid

biosynthesis. A common feature of all of these genes is the presence of UAS_{INO1} elements in their promoter regions (for a review see [28]). Regulation via UAS_{INO1} elements involves the transcriptional activators Ino2p and Ino4p, and the transcriptional repressor Opi1p [30]. Nuclear localization of Opi1p is regulated by PA and Scs2p, which bind and detain Opi1p to the ER membrane. We show that the observed reduction in inositol-mediated repression of UAS_{INO1}-driven transcription in Acb1p-depleted cells is not caused by altered *OPI1* levels or changes in PA and *SCS2* levels. Moreover, *INO2* and *INO4* mRNA levels were not significantly altered in Acb1p-depleted cells grown in minimal medium compared with wild-type cells (results not shown). The approx. 3-fold increase observed in *INO1* mRNA levels of Acb1p-depleted cells grown in minimal medium (Figure 3B) therefore may not be caused by increased levels of these transcriptional activators. Furthermore, Ino2p and Ino4p bind constitutively to the *INO1* promoter, and it has been suggested that occupancy of the *INO1* promoter by Ino2p and/or Ino4p is insufficient for its activation [37]. Taken together, these results strongly suggest that the reduction in inositol-mediated repression of Ino2p and Ino4p target genes in Acb1p-depleted cells occurs by an Ino2p, Ino4p and Opi1p-independent mechanism. However, it cannot be excluded that the Acb1p–acyl-CoA ester complex regulates the activity of Ino2p, Ino4p and Opi1p, either directly or indirectly. A possible regulatory mechanism is that Acb1p depletion alters Opi1p phosphorylation, which has been shown to affect its regulator activity [43].

Activation of the UPR pathway has previously been linked to increased phospholipid biosynthesis in yeast [37]. However, we did not find any connection between induction of the UPR pathway and increased expression of Ino2p and Ino4p target genes in Acb1p-depleted cells. Moreover, *HAC1* disruption reduced the expression of *OPI3* approx. 2-fold in Acb1p-depleted cells grown in YPD medium, whereas *OPI3* expression in wild-type cells was unaffected by *HAC1* disruption (results not shown). The absence of an effect of *HAC1* disruption on *OPI3* expression in wild-type cells confirms that the UPR pathway does not seem to be directly involved in transcriptional regulation of Ino2p and Ino4p target genes, as was recently suggested [31].

A large body of evidence suggests that Acb1p functions as an acyl-CoA ester transporter and pool former. We investigated the possibility that the altered expression of genes encoding enzymes involved in phospholipid synthesis, including *INO1*, in Acb1p-depleted yeast is the result of an absence of fatty acid metabolites. Supplementation of exogenous palmitic acid to Acb1p-depleted cells normalized *INO1* and *OPI3* mRNA levels to those of wild-type cells (Figure 5), in contrast with palmitoleic acid, which was found to partially normalize *INO1* and *OPI3* levels in Acb1p-depleted cells compared with wild-type cells (Figure 5). The difference in the effect of palmitic acid and palmitoleic acid may be a result of palmitic acid having additional effects on Acb1p-depleted cells.

Supplementation of palmitic acid to Acb1p-depleted cells partly restores inositol uptake [16] and reduced inositol secretion (Figure 4A). This may suggest that the reduction in inositol-mediated repression of Ino2p and Ino4p target genes, including *INO1*, in Acb1p-depleted cells, is simply a cellular response to low intracellular inositol levels. However, this is not likely to be the case since Acb1p-depleted cells, although unable to completely normalize *INO1* levels on addition of inositol, do respond to exogenous inositol by repressing *INO1* transcription (Figure 1A). Disruption of *SCS2*, which has previously been shown to reduce *INO1* expression in response to inositol depletion [44], did not normalize *INO1* expression in Acb1p-depleted cells (Figure 3). This further supports the hypothesis that *INO1* deregulation in

Acb1p-depleted cells is caused by means other than a lack of inositol. The steady-state expression of *GPD1* and *TDH1*, which is only slightly affected by inositol [45], is dramatically increased in Acb1p-depleted cells (Table 2), suggesting that at least part of the transcriptional response accompanying Acb1p depletion is inositol-independent.

Overall, the above observations present the possibility that reduced inositol-mediated repression of Ino2p and Ino4p target genes in Acb1p-depleted cells is the result of decreased availability of fatty acids and/or acyl-CoA esters. This is strongly supported by the result that ectopic overexpression of *FAS1* and *ACC1* reduced ratios of *INO1* mRNA levels in wild-type cells and Acb1p-depleted cells from approx. 1:6.5 and 1:7.4 to approx. 1:2.6 and 1:2.2 respectively (Figure 6). Notably, overexpression in Acb1p-depleted cells of *FAS2*, which is unable to enhance FAS activity [25], did not reduce *INO1* mRNA levels (results not shown). This suggests that the Acb1p-mediated removal of *de novo* synthesized acyl-CoA esters from the yeast FAS [16] can be overcome by increasing the availability of fatty acids and/or acyl-CoA esters through enhancement of *de novo* fatty acid biosynthesis or supplementation with fatty acids (Figures 5 and 6). These results strongly support the hypothesis that the altered availability of fatty acids and/or fatty acid derivatives may be involved in the mechanism by which Acb1p depletion alters inositol-mediated repression of Ino2p and Ino4p target genes.

In addition to affecting the expression of phospholipid biosynthesis genes, Acb1p depletion also increased the expression of fatty acid synthetic genes, including *ACC1*, *FAS1*, *FAS2* and *OLE1* (Table 2 and Supplementary Table S2). This indicates that the expression of these genes is also regulated by the availability of specific acyl-CoA esters, made available by Acb1p-acyl-CoA ester complexes. We have shown previously that depletion of Acb1p in the temperature-sensitive mutant of *ACC1*, *mtr7-1*, results in lethality at the permissive temperature [16], supporting the notion that Acb1p affects the availability of acyl-CoA esters.

The acyl species composition of phospholipids is altered on Acb1p depletion (Supplementary Tables S3–S8). Thus it cannot be ruled out that these alterations can affect the expression of Ino2p and Ino4p target genes. One possible model, consistent with our results, is that Acb1p recruits LCACoA esters to phospholipid-remodelling enzymes, thereby affecting the acyl species composition of phospholipids, which then affects the expression of Ino2p and Ino4p target genes.

Several studies have shown that fatty acids and fatty acid derivatives affect gene expression directly in both prokaryotes and eukaryotes [46,47]. A number of these studies suggest that the active species is acyl-CoA ester. In *Escherichia coli*, fatty acid synthesis and degradation is transcriptionally co-ordinated by the acyl-CoA ester-regulated transcription factor, FadR (for a review see [46]). Eukaryotic PPARs (peroxisome-proliferator-activated receptors) are activated by fatty acids and induce transcription of multiple genes involved in lipid metabolism [47], whereas LCACoA esters have been reported to antagonize this effect [48]. Immunogold EM (electron microscopy) has shown that ACBP/Acb1p is found in the nucleus of a number of mammalian [49,50] and yeast cells (S. Feddersen, J. Knudsen and N. J. Faergeman, unpublished work). The Acb1p-acyl-CoA complex may therefore modulate gene expression by donating acyl-CoA esters to gene regulatory components in the nucleus. Such a transport function is compatible with the recent suggestion implying that ACBP and HNF-4 α (hepatocyte nuclear factor 4 α) interact, and ACBP stimulates HNF-4 α -mediated transactivation of HNF-4 α target genes in hepatocytes [50].

In conclusion, we suggest that fatty acid derivatives are involved in the regulation of Ino2p and Ino4p target genes and that increased

expression of these target genes in Acb1p-depleted cells is caused by a decrease in the availability of acyl-CoA esters, as a result of the absence of Acb1p.

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