Accumulation of long-chain bases in yeast promotes their conversion to a long-chain base vinyl ether

Fernando Martinez-Montanes  
*University of Fribourg, Switzerland*

Museer A. Lone  
*University of Fribourg, Switzerland*

Fong-Fu Hsu  
*Washington University School of Medicine in St. Louis*

Roger Schneiter  
*University of Fribourg, Switzerland*

Follow this and additional works at: [https://digitalcommons.wustl.edu/open_access_pubs](https://digitalcommons.wustl.edu/open_access_pubs)

Recommended Citation  
[https://digitalcommons.wustl.edu/open_access_pubs/5448](https://digitalcommons.wustl.edu/open_access_pubs/5448)

This Open Access Publication is brought to you for free and open access by Digital Commons@Becker. It has been accepted for inclusion in Open Access Publications by an authorized administrator of Digital Commons@Becker. For more information, please contact engeszer@wustl.edu.
Accumulation of long-chain bases in yeast promotes their conversion to a long-chain base vinyl ether

Fernando Martínez-Montañés,* Moseer A. Lone,** Fong-Fu Hsu,† and Roger Schneiter*‡

Department of Biology,* University of Fribourg, 1700 Fribourg, Switzerland; and Department of Internal Medicine,† Washington University School of Medicine, St. Louis, MO 63110

Abstract Long-chain bases (LCBs) are the precursors to ceramide and sphingolipids in eukaryotic cells. They are formed by the action of serine palmitoyl-CoA transferase (SPT), a complex of integral membrane proteins located in the endoplasmic reticulum. SPT activity is negatively regulated by Orm proteins to prevent the toxic overaccumulation of LCBs. Here we show that overaccumulation of LCBs in yeast results in their conversion to a hitherto undescribed LCB derivative, an LCB vinyl ether. The LCB vinyl ether is predominantly formed from phytosphingosine (PHS) as revealed by conversion of odd chain length tracers C17-dihydrosphingosine and C17-PHS into the corresponding LCB vinyl ether derivative. PHS vinyl ether formation depends on ongoing acetyl-CoA synthesis, and its levels are elevated when the LCB degradative pathway is blocked by deletion of the major LCB kinase, LCB4, or the LCB phosphate lyase, DPL1. PHS vinyl ether formation thus appears to constitute a shunt for the LCB phosphate- and lyase-dependent degradation of LCBs. Consistent with a role of PHS vinyl ether formation in LCB detoxification, the lipid is efficiently exported from the cells.—Martínez-Montañés, F., M. A. Lone, F-F. Hsu, and R. Schneiter. Accumulation of long-chain bases in yeast promotes their conversion to a long-chain base vinyl ether. J. Lipid Res. 2016. 57: 2040–2050.

Supplementary key words ceramide • sphingolipids • Saccharomyces cerevisiae • mass spectrometry

Sphingolipids are an essential class of lipids greatly enriched in the plasma membrane of eukaryotic cells. They have been implicated in the formation and maintenance of lateral membrane domains, important for protein sorting and signaling along the compartments of the secretory pathway. Apart from these structural roles, their biosynthetic precursor and intermediates, such as long-chain bases (LCBs) and ceramide, exert important signaling functions to coordinate complex processes, for example, cell cycle progression, apoptosis, and inflammation. Hence, the synthesis and turnover of these lipids must be precisely controlled (1–3).

Sphingolipid synthesis starts in the endoplasmic reticulum (ER), where serine palmitoyl-CoA transferase (SPT) catalyzes the first step in formation of LCBs (4). Variations in chain length of the condensing acyl-CoA and the incorporation of alternative amino acids can result in the synthesis of a chemically heterogeneous set of sphingoid bases (5, 6). The activity of SPT, the rate-limiting enzyme of the pathway, is negatively regulated by Orm proteins, conserved integral ER membrane proteins whose phosphorylation relieves inhibition of SPT activity (7, 8). Kinases that phosphorylate Orm proteins thus integrate multiple signals to maintain sphingolipid homeostasis, including heat and ER stress, and availability of nutrients (9–13).

The major LCBs in yeast are dihydrosphingosine (DHS) and phytosphingosine (PHS), which upon ceramide formation condense with a CoA-activated C26 very long-chain fatty acid (14, 15). This reaction is catalyzed by the ER-localized ceramide synthase (CerS). Upon transport to the Golgi apparatus, ceramides are converted to a set of complex sphingolipids: inositol phosphorylceramide, mannosyl-inositol phosphorylceramide, and mannosyl-diinositol phosphorylceramide (16–19).

In addition to these biosynthetic routes, complex sphingolipids, ceramide, and LCBs are also subject to degradation. Complex sphingolipids are cleaved by an inositol phosphosphingolipid phospholipase C, Isc1 (20). Ceramide, on the other hand, is degraded through alkaline ceramidases Ydc1 and Ypc1 (21, 22). Phosphorylated LCBs, finally, can be cleaved by a sphingosine-1-phosphate lyase, Dpl1, to ethanolamine phosphate and fatty aldehyde (23). The activity of components of this degradative branch, Isc1, Ydc1, and Ypc1, is controlled by the target of rapamycin complex 1

 Abbreviations: CerS, ceramide synthase; DHS, dihydrosphingosine; ER, endoplasmic reticulum; LCB, long-chain base; OD, optical density; PHS, phytosphingosine; SPT, serine palmitoyl-CoA transferase; YPD, yeast peptone dextrose.

*Present address of M. A. Lone: Institute for Clinical Chemistry, University Hospital Zurich, 8091 Zurich, Switzerland.
†To whom correspondence should be addressed.
‡To whom correspondence should be addressed.
§The online version of this article (available at http://www.jlr.org) contains a supplement.
(24). Importantly, the transient intermediates of the pathway, LCB, LCB phosphate, and ceramide, not only act as biosynthetic precursors but also have important signaling functions in stress response (see Fig. 1A for an overview of the pathway) (25, 26).

Here we describe a novel LCB derivative, identified as an LCB vinyl ether. This LCB vinyl ether is generated mainly from PHS in cells that accumulate high levels of LCBs either due to deregulated de novo synthesis, a block in the degradative pathway, or uptake of externally provided PHS. Conversion of PHS to the vinyl ether derivative appears to act as a shunt for the catabolic pathway because PHS vinyl ether levels are greatly elevated in mutants that cannot phosphorylate LCBs or in mutants lacking the sphingosine-1-phosphate lyase. Consistent with a potential role in PHS detoxification, the vinyl ether is excreted from cells.

MATERIALS AND METHODS

Yeast strains and growth conditions

Yeast strains and their genotypes are listed in supplemental Table S1. Strains were cultivated in yeast peptone dextrose (YPD)-rich medium (1% Bacto yeast extract, 2% Bacto peptone; US Biological, Swampscott, MA) or synthetic dextrose medium lacking uracil (SD-URA) synthetic medium (0.67% yeast nitrogen base without amino acids; US Biological, Salem, MA), 2% glucose, and the following amino acids: 20 mg/l of each adenine, arginine, histidine, methionine, and tryptophan; 60 mg/l leucine, 230 mg/l lysine, and 300 mg/l threonine. Double-mutant strains were generated by crossing of single mutants and by gene disruption, using PCR deletion cassettes and a marker rescue strategy (27). Myriocin (Sigma Aldrich, St. Louis, MO) was diluted in DMSO and used from a 1,000× stock, and Fumonisin B1 (Enzo Life Sciences, Farmingdale, NY) was diluted in water and used from a 5× stock. Sphingosine, 1-deoxysphinganine, C18- and 2-deoxyglucose were from Sigma. 1,2-Dihydroceramide, C17/C18-DHS, and C17/C18-PHS were obtained from Avanti (Avanti Polar Lipids, Alabaster, AL); 1,3-threo-DHS and 2-deoxyxyphosphine were from Sigma.

Lipid extraction and analysis by MS

For lipid analysis, overnight cultures of strains were diluted into fresh YPD (Figs. 1B, 2, 3, 4, 5A, 7A) or SD-URA (Fig. 5B, 6B, 7B) media, and cells were grown at 30°C to an optical density (OD)600nm of approximately 2. Temperature-sensitive strains were grown at 24°C in YPD (Fig. 7A). Lipids were extracted from 10 OD600nm units of cells with CHCl3 and methanol (2:1 by volume) or from the culture supernatant with 2 vol of diethyl ether. C17-DHS (1 nmol) was used as internal standard (28). LC/MS was conducted on a Bruker Esquire HCT ion trap mass spectrometer in the positive ion mode, ions at m/z 344.3159. In the negative-ion mode, ions at m/z 342.3013, were observed. These results indicate that the lipid at m/z 344.3 correlated with the abundance of PHS in wild-type and elo3 mutant cells (436 pmol/OD; Fig. 1B, C). Elo3 is a component of the ER-associated acyl chain elongase complex required for the synthesis of C26 very long-chain fatty acids (29, 30). elo3 mutant cells make C22 instead of the normal C26 fatty acids. Shorter acyl-CoAs, however, are a poor substrate for CerS, the enzyme that catalyzes the Nacylation of LCBs to form ceramide (31, 32). As a consequence, elo3 mutant cells display greatly elevated levels of PHS. While wild-type cells have about 19 pmol/OD of PHS, elo3 mutant cells have up to 451 pmol/OD of PHS. The fact that the relative abundance of the lipid at m/z 344.3 correlated with the abundance of PHS in wild-type and elo3 mutant cells suggested that it might be derived from PHS.

To test this hypothesis, we characterized the structure of this lipid by high-resolution MS. When subjected to ESI in the positive-ion mode, [M + H]+ ions were observed at m/z 344.3164, which corresponds to an elemental composition of C20H42O3N (calculated m/z = 344.3159). In the negative-ion mode, ions at m/z 342.3014, corresponding to an elemental composition of C19H36O5N (calculated m/z = 342.3013), were observed. These results indicate that the compound had an elemental composition of C20H42O3N, representing a 1-Oethenyl-2-amino-4-octadecene-1,3-diol, that is a PHS derivative containing a vinyl ether at the C1 hydroxyl group of PHS (Fig. 2).

Fragmentation (MS2) of the [M + H]+ ions at m/z 344.3164 gave rise to ions of 326 and 308, arising from consecutive losses of water (Fig. 2A, route a), along with ions of m/z 300 arising from loss of CH2=CHOH, and ions of m/z 282 (300 - H2O), arising from an additional loss of water (supplemental Fig. S1). Two pathways leading to the elimination of CH2=CHOH are proposed. The first pathway involves the participation of the hydrogen of the 3-OH group to

RESULTS AND DISCUSSION

Identification and characterization of a PHS vinyl ether

In the course of analyzing LCB levels by MS in lipid extracts from various yeast mutants, we noticed an uncharacterized peak at m/z 344.3. This lipid was of low abundance in wild-type cells (29 pmol/OD), but its concentration was greatly elevated in elo3 mutant cells (346 pmol/OD; Fig. 1B, C). Elo3 is a component of the ER-associated acyl chain elongase complex required for the synthesis of C26 very long-chain fatty acids (29, 30). elo3 mutant cells make C22 instead of the normal C26 fatty acids. Shorter acyl-CoAs, however, are a poor substrate for CerS, the enzyme that catalyzes the Nacylation of LCBs to form ceramide (31, 32). As a consequence, elo3 mutant cells display greatly elevated levels of PHS. While wild-type cells have about 19 pmol/OD of PHS, elo3 mutant cells have up to 451 pmol/OD of PHS. The fact that the relative abundance of the lipid at m/z 344.3 correlated with the abundance of PHS in wild-type and elo3 mutant cells suggested that it might be derived from PHS.

To test this hypothesis, we characterized the structure of this lipid by high-resolution MS. When subjected to ESI in the positive-ion mode, [M + H]+ ions were observed at m/z 344.3164, which corresponds to an elemental composition of C20H42O3N (calculated m/z = 344.3159). In the negative-ion mode, ions at m/z 342.3014, corresponding to an elemental composition of C19H36O5N (calculated m/z = 342.3013), were observed. These results indicate that the compound had an elemental composition of C20H42O3N, representing a 1-Oethenyl-2-amino-4-octadecene-1,3-diol, that is a PHS derivative containing a vinyl ether at the C1 hydroxyl group of PHS (Fig. 2).

Fragmentation (MS2) of the [M + H]+ ions at m/z 344.3164 gave rise to ions of 326 and 308, arising from consecutive losses of water (Fig. 2A, route a), along with ions of m/z 300 arising from loss of CH2=CHOH, and ions of m/z 282 (300 - H2O), arising from an additional loss of water (supplemental Fig. S1). Two pathways leading to the elimination of CH2=CHOH are proposed. The first pathway involves the participation of the hydrogen of the 3-OH group to
Fig. 1. Identification of a putative LCB derivative. A: Schematic overview of the yeast sphingolipid biosynthetic and degradative pathways. Key enzymes and lipid intermediates are shown. Mutants used in this study are indicated in bold, and drugs that were used are shown in red. The pathway leading to dihydroceramide is highlighted in orange, the maturation of phytoceramide to the complex sphingolipids is highlighted in green, and the degradative pathway is highlighted in blue. B, C: ESI/MS profile of LCBs present in wild-type (WT; B) and elo3Δ (C) mutant cells. Lipid extracts prepared from wild-type and elongase (elo3Δ) mutant cells were analyzed by ESI/MS in the positive ion mode using the odd chain-length C17-DHS (m/z 288.3) as internal standard (indicated in blue). The major PHS species present in both strains, C18-PHS (m/z 318.3) is indicated in green. The [M + H]+ ion at m/z 344.3, indicated in red, represents a putative novel LCB derivative.
form a long alkyl chain with a terminal oxetane species of m/z 300 (Fig. 2A, route b), which gave rise to ions of m/z 282 via further loss of water (Fig. 2A, route b1). The second pathway involves the hydrogen of the secondary amino group, leading to the formation of an alkyl chain with a terminal aziridine (Fig. 2A, route c), which undergoes further loss of water to yield ions of m/z 282 (Fig. 2A, route c1). These fragmentation pathways were further supported by MS³ spectrum of the ions of m/z 282 (344 → 282; supplemental Fig. S1B), which is identical to the MS³ spectrum of m/z 282 (344 → 326 → 282; data not shown). The spectrum (supplemental Fig. S1B) also contained ions of m/z 265 and 252 arising from losses of NH₃ (Fig. 2A, route b2, structures highlighted in blue) and HCHO (Fig. 2A, route b3), respectively. The ion of m/z 264 is a hallmark of the sphingosine LCB structure and arises from loss of water from the aziridine precursor ions (Fig. 2A, route c2, structures highlighted in orange) (33, 34). The above fragmentation processes were supported by high resolution MS, from which the deduced elemental composition of the fragment ions are consistent with the suggested structures (data not shown).
by ions of m/z 223 and 221 representing a terminally conjugated diene and triene, respectively, arising from consecutive losses of H₂. The spectrum also contained ions at m/z 197, 183, 169, 155, and so forth, and at m/z 111, 97, 83, arising from cleavages of the C-C bond of the LCB via charge-remote fragmentation (Fig. 2B). These results are consistent with the assignments of the suggested structure of 1-O-ethenyl-2-amino-4-octadecene-1,3-diol.

We note that the structure of the proposed PHS vinyl ether has the same elemental composition as C₂-dihydroceramide and is thus isobaric with C₂-dihydroceramide. The fragmentation pattern of the PHS vinyl ether in both positive and negative ion mode, however, is clearly distinct from that of C₂-dihydroceramide (supplemental Fig. S3).

Fragmentation of acetylated LCBs typically results in a characteristic loss of m/z 42, corresponding to the loss of a ketene (supplemental Fig. S3). This is not observed upon fragmentation of PHS vinyl ether, which instead loses a fragment of m/z 44, corresponding to a vinyl alcohol. Thus,

The structural assignment was further supported by high resolution LIT MS⁶ of the corresponding [M – H]⁻ ions (Fig. 2B). The MS² spectrum of the [M – H]⁻ ion at m/z 342 contained ions at m/z 324 and 306 arising from consecutive losses of water, and prominent ions of m/z 255, arising from cleavage of C₅(OH)-C₄(NH₂) bond of the LCB, together with ions of m/z 225 arising from cleavage of the C₄(OH)-C₃(OH) bond (35) (supplemental Fig. S2A). The cleavage of this latter bond is consistent with the formation of the ions of m/z 116, in which the anionic charge site is located at the oxygen atom attached to C₃ of the LCB (Fig. 2B). The presence of the ions of m/z 116 also supports the notion of the presence of the 1-O-ethenyl group in the molecule. Further dissociation of the ions of m/z 255 (342 → 255; supplemental Fig. S2B) gave rise to the terminally conjugated ions of m/z 253 via loss of H₂, the prominent ions of m/z 225 arising from loss of HCHO, and ions of m/z 237 by loss of water. The MS³ spectrum of the ions of m/z 225 (342 → 255 → 225; supplemental Fig. S2C) are dominated

Fig. 3. The LCB vinyl ether is predominantly produced from PHS. A: PHS but not DHS is efficiently converted to the vinyl ether derivative. Lipids were extracted from eloΔ and eloΔ surΔ double mutant cells and LCBs were analyzed by ESI/MS with C17-DHS as an internal standard. eloΔ mutant cells have about equimolar levels of both PHS and PHS vinyl ether. eloΔ surΔ double mutant cells accumulate high levels of DHS but show only low levels of the corresponding DHS vinyl ether. B: Externally added PHS but not DHS is efficiently converted to the PHS vinyl ether. Wild-type cells were grown to OD ~2 and supplemented with 10 µM of the indicated LCB for 30 min, lipids were extracted and analyzed by ESI/MS. The ratio of the externally added LCB to the corresponding vinyl ether derivative is plotted. Values represent means ± SD of three independent determinations. n.d.; not detectable.
Wild-type cells incubated with 10 µM PHS during 30 min displayed high levels of PHS vinyl ether. In these cells, levels of PHS vinyl ether were ~1.7-fold higher than free PHS levels (Fig. 3B). Cells incubated with DHS, however, displayed only low levels of the DHS vinyl ether, supporting the conclusion that PHS is the preferred substrate for formation of the LCB vinyl ether.

To distinguish between the conversion of internally synthesized LCBs and that of externally added LCBs to the vinyl ether, we challenged cells with a synthetic, odd chain length LCB tracer. We have previously shown that these C17-LCBs are efficiently taken up and incorporated into ceramide and complex sphingolipids (28). Wild-type cells converted C17-PHS efficiently to the C17-PHS vinyl ether, whereas C17-DHS was only inefficiently transformed to the C17-DHS vinyl ether (Fig. 3B). Other LCBs, such as sphingosine, or the stereoisomer of the natural DHS, l-threo-DHS, were also only very inefficiently converted to the corresponding vinyl ether derivatives. Deoxysphinganine, on the other hand, was not converted to the vinyl ether, which is consistent with the fact that the vinyl ether group is bound to the C1 hydroxyl group, which is missing in deoxysphinganine. Taken together, these results thus show the fragmentation pattern of the PHS vinyl ether is not compatible with that of either an N- or O-acetylated LCB, including C2-dihydroceramide.

**PHS is efficiently converted to PHS vinyl ether**

To confirm this structural assignment and to test whether DHS could also be converted to a DHS vinyl ether, we analyzed the formation of the LCB vinyl ether in cells that cannot form PHS due to a deletion of the Sur2 hydroxylase, which converts DHS into PHS (36). Compared with elo3Δ mutant cells, the elo3Δ sur2Δ double mutant had greatly elevated levels of DHS. Despite these elevated DHS levels, the elo3Δ sur2Δ double mutant produces only very low levels of the corresponding DHS vinyl ether. PHS, accumulating in the elo3Δ single mutant, however, is efficiently converted to the PHS vinyl ether as elo3Δ mutant cells display about equal levels of both PHS and PHS vinyl ether (Fig. 3A). We thus conclude that PHS rather than DHS is the preferred substrate for formation of the LCB vinyl ether.

To test whether conversion of PHS to the vinyl ether derivative is a general reaction of cells to high levels of PHS, we challenged wild-type cells with externally added LCBs. Wild-type cells incubated with 10 µM PHS during 30 min displayed high levels of PHS vinyl ether. In these cells, levels of PHS vinyl ether were ~1.7-fold higher than free PHS levels (Fig. 3B). Cells incubated with DHS, however, displayed only low levels of the DHS vinyl ether, supporting the conclusion that PHS is the preferred substrate for formation of the LCB vinyl ether.

To distinguish between the conversion of internally synthesized LCBs and that of externally added LCBs to the vinyl ether, we challenged cells with a synthetic, odd chain length LCB tracer. We have previously shown that these C17-LCBs are efficiently taken up and incorporated into ceramide and complex sphingolipids (28). Wild-type cells converted C17-PHS efficiently to the C17-PHS vinyl ether, whereas C17-DHS was only inefficiently transformed to the C17-DHS vinyl ether (Fig. 3B). Other LCBs, such as sphingosine, or the stereoisomer of the natural DHS, l-threo-DHS, were also only very inefficiently converted to the corresponding vinyl ether derivatives. Deoxysphinganine, on the other hand, was not converted to the vinyl ether, which is consistent with the fact that the vinyl ether group is bound to the C1 hydroxyl group, which is missing in deoxysphinganine. Taken together, these results thus show...
phosphatase that dephosphorylates exogenously imported LCB phosphates, and this activity is necessary for the incorporation of exogenous LCBs into sphingolipids (21, 37–39). Levels of free PHS and those of PHS vinyl ether were reduced to wild-type concentrations upon deletion of Lcb3 in the \( \text{orm1} \Delta \text{orm2} \Delta \) double mutant (Fig. 4A). Similarly, upon inhibition of SPT activity by myriocin, both PHS and PHS vinyl ether levels were significantly reduced in both\( \text{elο3} \Delta \text{orm1} \Delta \text{orm2} \Delta \) and \( \text{elo3} \Delta \text{elo3} \Delta \) mutant cells (Fig. 4A, B).

Consistent with the notion that levels of the vinyl ether parallel those of free PHS, wild-type cells treated with the CerS inhibitor fumonisin B1, which results in increased PHS levels, displayed a slight but significant increase in PHS vinyl ether levels (28) (Fig. 4C). Deletion of the catalytic components of the CerS, Lag1 and Lac1, on the other hand, resulted in very high levels of PHS vinyl ether, consistent with the fact that \( \text{lag1} \Delta \text{lac1} \Delta \) double mutant cells have high levels of free PHS (14) (Fig. 4C). Repression of the phosphatase that dephosphorylates exogenously imported LCB phosphates, and this activity is necessary for the incorporation of exogenous LCBs into sphingolipids (21, 37–39). Levels of free PHS and those of PHS vinyl ether were reduced to wild-type concentrations upon deletion of Lcb3 in the \( \text{orm1} \Delta \text{orm2} \Delta \) double mutant (Fig. 4A). Similarly, upon inhibition of SPT activity by myriocin, both PHS and PHS vinyl ether levels were significantly reduced in both \( \text{elο3} \Delta \text{orm1} \Delta \text{orm2} \Delta \) and \( \text{elo3} \Delta \text{elo3} \Delta \) mutant cells (Fig. 4A, B).

Consistent with the notion that levels of the vinyl ether parallel those of the free PHS, wild-type cells treated with the CerS inhibitor fumonisin B1, which results in increased PHS levels, displayed a slight but significant increase in PHS vinyl ether levels (28) (Fig. 4C). Deletion of the catalytic components of the CerS, Lag1 and Lac1, on the other hand, resulted in very high levels of PHS vinyl ether, consistent with the fact that \( \text{lag1} \Delta \text{lac1} \Delta \) double mutant cells have high levels of free PHS (14) (Fig. 4C). Repression of the that both internally as well as externally provided PHS are efficiently converted to the vinyl ether and that the levels of free PHS and those of PHS vinyl ether reach similar levels irrespective of whether PHS is endogenously synthesized or whether it is taken up by the cells. DHS, on the other hand, is not efficiently converted to the DHS vinyl ether, indicating that the converting enzyme(s) has high substrate specificity for PHS over DHS.

**PHS vinyl ether levels parallel those of free PHS**

To test whether mutants other than \( \text{elο3} \Delta \) known to accumulate high levels of internal PHS also show elevated levels of PHS vinyl ether and that levels of free PHS and those of PHS vinyl ether reach similar levels irrespective of whether PHS is endogenously synthesized or whether it is taken up by the cells. DHS, on the other hand, is not efficiently converted to the DHS vinyl ether, indicating that the converting enzyme(s) has high substrate specificity for PHS over DHS.

**PHS vinyl ether levels parallel those of free PHS**

To test whether mutants other than \( \text{elο3} \Delta \) known to accumulate high levels of internal PHS also show elevated levels of PHS vinyl ether and that levels of free PHS and those of PHS vinyl ether reach similar levels irrespective of whether PHS is endogenously synthesized or whether it is taken up by the cells. DHS, on the other hand, is not efficiently converted to the DHS vinyl ether, indicating that the converting enzyme(s) has high substrate specificity for PHS over DHS.

**PHS vinyl ether levels parallel those of free PHS**

To test whether mutants other than \( \text{elο3} \Delta \) known to accumulate high levels of internal PHS also show elevated levels of PHS vinyl ether and that levels of free PHS and those of PHS vinyl ether reach similar levels irrespective of whether PHS is endogenously synthesized or whether it is taken up by the cells. DHS, on the other hand, is not efficiently converted to the DHS vinyl ether, indicating that the converting enzyme(s) has high substrate specificity for PHS over DHS.

**PHS vinyl ether levels parallel those of free PHS**

To test whether mutants other than \( \text{elο3} \Delta \) known to accumulate high levels of internal PHS also show elevated levels of PHS vinyl ether and that levels of free PHS and those of PHS vinyl ether reach similar levels irrespective of whether PHS is endogenously synthesized or whether it is taken up by the cells. DHS, on the other hand, is not efficiently converted to the DHS vinyl ether, indicating that the converting enzyme(s) has high substrate specificity for PHS over DHS.

**PHS vinyl ether levels parallel those of free PHS**

To test whether mutants other than \( \text{elο3} \Delta \) known to accumulate high levels of internal PHS also show elevated levels of PHS vinyl ether and that levels of free PHS and those of PHS vinyl ether reach similar levels irrespective of whether PHS is endogenously synthesized or whether it is taken up by the cells. DHS, on the other hand, is not efficiently converted to the DHS vinyl ether, indicating that the converting enzyme(s) has high substrate specificity for PHS over DHS.

**PHS vinyl ether levels parallel those of free PHS**

To test whether mutants other than \( \text{elο3} \Delta \) known to accumulate high levels of internal PHS also show elevated levels of PHS vinyl ether and that levels of free PHS and those of PHS vinyl ether reach similar levels irrespective of whether PHS is endogenously synthesized or whether it is taken up by the cells. DHS, on the other hand, is not efficiently converted to the DHS vinyl ether, indicating that the converting enzyme(s) has high substrate specificity for PHS over DHS.

**PHS vinyl ether levels parallel those of free PHS**

To test whether mutants other than \( \text{elο3} \Delta \) known to accumulate high levels of internal PHS also show elevated levels of PHS vinyl ether and that levels of free PHS and those of PHS vinyl ether reach similar levels irrespective of whether PHS is endogenously synthesized or whether it is taken up by the cells. DHS, on the other hand, is not efficiently converted to the DHS vinyl ether, indicating that the converting enzyme(s) has high substrate specificity for PHS over DHS.

**PHS vinyl ether levels parallel those of free PHS**

To test whether mutants other than \( \text{elο3} \Delta \) known to accumulate high levels of internal PHS also show elevated levels of PHS vinyl ether and that levels of free PHS and those of PHS vinyl ether reach similar levels irrespective of whether PHS is endogenously synthesized or whether it is taken up by the cells. DHS, on the other hand, is not efficiently converted to the DHS vinyl ether, indicating that the converting enzyme(s) has high substrate specificity for PHS over DHS.

**PHS vinyl ether levels parallel those of free PHS**

To test whether mutants other than \( \text{elο3} \Delta \) known to accumulate high levels of internal PHS also show elevated levels of PHS vinyl ether and that levels of free PHS and those of PHS vinyl ether reach similar levels irrespective of whether PHS is endogenously synthesized or whether it is taken up by the cells. DHS, on the other hand, is not efficiently converted to the DHS vinyl ether, indicating that the converting enzyme(s) has high substrate specificity for PHS over DHS.
containing 2-deoxyglucose instead of glucose and NaN3/NaF (10 mM) for 15 min. Lipids were then extracted and quantified from the cell pellet and the culture medium. Released LCB levels are expressed as percentage of total LCBs. Asterisks denote statistical significance (* P < 0.05; ** P < 0.001; *** P < 0.0001). n.s.; not significant.

Taken together, these results indicate that the levels of the vinyl ether consistently parallel those of the free PHS under the various conditions tested here, indicating that the half-life of the two lipids are similar, arguing thus against the possibility that the vinyl ether acts as an inert storage form for the free LCB. In addition, the fact that LCBs of different chain length are converted to the respective vinyl ether indicates that the converting enzymes do not discriminate between these chain length variants of PHS.

**PHS vinyl ether formation acts in parallel to the degradative pathway**

LCBs enter the degradative pathway by first being phosphorylated by either Lcb4 or Lcb5, the two LCB kinases in yeast. Lcb4 is the major kinase located in the ER, whereas Lcb5 has only minor activity in exponentially growing cells (42, 43). The resulting LCB-phosphates (LCB-P) can then either be dephosphorylated by Lcb3/Ysr3 or they are cleaved by the LCB-P lyase, Dpl1, to ethanolamine phosphate and 1-hexadecanal (23). To test whether formation of the vinyl ether depends on prior phosphorylation of the LCB or whether it acts in parallel to this catabolic pathway, we analyzed LCB levels in double mutants of elo3Δ with either lcb4Δ or lcb5Δ. Deletion of these LCB kinases in the elo3Δ mutant background resulted in slightly elevated levels of DHS and PHS (Fig. 5A). Deletion of the main LCB kinase, Lcb4, in the elo3Δ mutant background, however, resulted in a dramatic accumulation of PHS vinyl ether, suggesting that a block in the catabolic pathway shunts PHS toward the formation of the vinyl ether derivative. Deletion of the minor kinase activity, Lcb5, on the other hand, did not significantly increase vinyl ether levels (Fig. 5A).

Given that blocking the degradative pathway through deletion of the major LCB kinase Lcb4 resulted in greatly elevated levels of PHS vinyl ether, we examined whether deletion of the lyase would lead to a similar increase in PHS vinyl ether levels. Deletion of Dpl1 in an elo3Δ mutant background again gave rise to slightly elevated levels of DHS and PHS; PHS vinyl ether levels, however, were not significantly increased. Upon overexpression of the LCB phosphate phosphatase, Lcb3, however, PHS vinyl ether levels became significantly elevated (Fig. 5B). Overexpression of Lcb3 is expected to reduce the efficiency of the degradative pathway and hence to further increase levels of free LCBs. The fact that this resulted in accumulation of PHS vinyl ether is thus consistent with a shunt function of the pathway, which diverges free PHS into formation of the vinyl ether.

Taken together, these data indicate that vinyl ether synthesis is independent of the degradative pathway and that it acts in parallel to the catabolic pathway, possibly as a shunt for the degradative pathway under conditions of great excess of free intracellular LCBs. In addition, this shunt is possibly important to prevent the detrimental accumulation of LCB-phosphate (44, 45). On the other hand, the presence of the vinyl group on the C1 hydroxyl shield this LCB derivative from phosphorylation and subsequent degradation through the lyase pathway.

**PHS vinyl ether formation depends on ongoing acetyl-CoA synthesis**

To examine how the PHS vinyl ether is synthesized, we hypothesized that PHS may first be acetylated and the ketone group may subsequently be reduced, first to the hydroxyl...
and then to the vinyl ether (Fig. 6A). This hypothesis would predict that formation of PHS vinyl ether is decreased in cells that have low acetyl-CoA levels. Acetyl-CoA can be produced by at least three major pathways in yeast: through the mitochondrial pyruvate dehydrogenase (PDH) complex, through peroxisomal β-oxidation, and through the two acetyl-CoA synthetases, Acs1 and Acs2 (46–48). Under aerobic conditions and on glucose-containing media, Acs2 is essential and Acs1 and the β-oxidation genes, e.g., \( \text{POT1} \), are repressed, and the \( \text{PDH} \) genes are not essential (46, 47, 49). To examine the requirement of acetyl-CoA for the conversion of PHS into PHS vinyl ether, we challenged wild-type and \( \text{acs} \) mutant cells with externally provided PHS and monitored the appearance of the vinyl ether. Wild-type and \( \text{acs}2\Delta \) mutant cells carrying a plasmid borne copy of \( \text{ACS2} \) (\( \text{acs}2\Delta + \text{pACS2} \)) displayed an essentially equimolar ratio between PHS and PHS vinyl ether (Fig. 6B). The \( \text{acs}2\Delta \) mutant rescued by a temperature-sensitive (ts) allele of \( \text{ACS2} \) (\( \text{pACS2-1ts} \)) and the \( \text{acs}1\Delta \text{acs}2\Delta \) double mutant carrying the same ts allele of \( \text{ACS2} \), however, displayed significantly decreased levels of PHS vinyl ether compared with free PHS. 

These data thus indicate that normal acetyl-CoA levels are required for the efficient conversion of free PHS into PHS vinyl ether and hence that vinyl ether formation may proceed through the formation of an \( \text{O} \)-acytlated LCB intermediate. It is interesting to note that acetylated LCBs are found in certain microorganisms, such as \( \text{Wickerhamomyces ciferrii} \) (50). However, deletion of the acetyltransferases implied in the formation of the \( \text{W. ciferrii} \) acetylated LCBs, \( \text{SLI1} \) and \( \text{ATF2} \), in \( \text{Saccharomyces cerevisiae} \) did not affect PHS vinyl ether synthesis (data not shown). In mammals, on the other hand, 3-\( \text{O} \)-acetyl-sphingosine is present in so called fast migrating forms of cerebrosides. They appear during myelinogenesis and may play critical functions in myelin structure and function (51). The possibility that PHS vinyl ether synthesis occurs through an acetylated intermediate renders its synthesis analogous to that of the mammalian ether containing glycerophospholipids, which occurs through the exchange of sn-1 bound acyl group on dihydroxyacetone phosphate by an alkyl group. The resulting alkyl ether can then be further reduced to a vinyl ether, as typically found in the plasmalogens (52).

### The PHS vinyl ether is excreted into the culture medium

Given that a PHS vinyl ether is considerably more hydrophobic than PHS itself, we wondered whether synthesis of the vinyl ether derivative might be a means to detoxify the cells from the detrimental effects of high PHS levels. We thus analyzed PHS and PHS vinyl ether levels in the cell pellet and the culture supernatant of elongase double mutant cells, \( \text{tsc13-1 elo3} \), which have high levels of free PHS and PHS vinyl ether (Fig. 7A). \( \text{TSC13} \) encodes for the enoyl reductase that catalyzes the last step in each very long-chain fatty acid elongation cycle (53). Interestingly, PHS and PHS vinyl ether levels present in the cell pellet were similar to the levels of these two lipids present in the culture supernatant, indicating that both of these LCBs can be exported by the cells. PHS vinyl ether levels, however, far exceeded PHS levels in both the cell pellet and the culture supernatant, consistent with a possible role of the PHS vinyl ether in PHS detoxification.

Because intra- and extracellular levels of both PHS and PHS vinyl ether were comparable, it is conceivable that export of these lipids occurs by passive, energy-independent transport pathways. PHS export, on the other hand, has previously been described to be ATP dependent and to rely on \( \text{Rsb1} \), a seven transmembrane protein, whose overexpression rescues the LCB sensitivity of \( \text{dpl1} \Delta \) mutant cells (54, 55). To examine the energy requirement of the export of the vinyl ether, we challenged \( \text{dpl1} \Delta \) mutant cells overexpressing \( \text{RSB1} \) with C17-PHS for 20 min, switched cells to medium containing 2-deoxyglucose and NaN\(_3\)/NaF to deplete ATP levels for 15 min, and then analyzed C17-PHS and C17-PHS vinyl ether levels in the cell pellet and the culture supernatant. Levels of both C17-PHS and that of the C17-PHS vinyl ether in the extracellular medium dropped significantly upon energy depletion of the cells, consistent with the notion that the export of both of these LCBs is dependent on an active, energy-requiring process (Fig. 7B).

Taken together, the data presented here indicate that free PHS is efficiently converted to a PHS vinyl ether derivative. This conversion is dependent on acetyl-CoA levels, and both free PHS and PHS vinyl ether are efficiently excreted by the cells. Based on these observations, we propose that the synthesis of the vinyl ether containing LCB may act to reduce levels of endogenous free LCBs and thus relieve cells of the growth inhibition of these LCBs. In this model, PHS vinyl ether synthesis and its export may thus act as a detoxification pathway to reduce levels of endogenous PHS (Fig. 8). If this were correct, one would predict that unlike free PHS, the PHS vinyl ether would not be taken up by the cells. A prediction that can be tested once a synthetic PHS vinyl ether will be available. In any case, conversion of free PHS into the vinyl ether derivative provides an additional means to regulate and fine tune the levels of free LCBs and thus to sustain cell proliferation under adverse conditions.

The authors thank J. D. Boeke and T. Dunn for mutant strains; S. Reddy Polu and S. Schürch for initial analysis of the compound; A. Conzelmann, T. Hornemann, and S. G. Gowda for helpful discussions; and S. Cottier for comments on the manuscript.
REFERENCES


