Transcript profiling and lipidomic analysis of ceramide subspecies in mouse embryonic stem cells and embryoid bodies

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Abstract  Ceramides (Cers) are important in embryogenesis, but no comprehensive analysis of gene expression for Cer metabolism nor the Cer amounts and subspecies has been conducted with an often used model: mouse embryonic stem cells (mESCs) versus embryoid bodies (EBs). Measuring the mRNA levels by quantitative RT-PCR and the amounts of the respective metabolites by LC-ESI/MS/MS, notable differences between R1 mESCs and EBs were: EBs have higher mRNAs for CerS1 and CerS3, which synthesize C18- and C≥24-carbons dihydroceramides (DH)Cer, respectively; EBs have higher CerS2 (for C24:0- and C24:1-); and EBs have lower CerS5 + CerS6 (for C16-). In agreement with these findings, EBs have (DH)Cer with higher proportions of C18-, C24- and C26- and less C16-fatty acids, and longer (DH)Cer are also seen in monohexosylCers and sphingomyelins. EBs had higher mRNAs for fatty acyl-CoA elongases that produce C18-, C24-, and C26-fatty acyl-CoAs (Elovl3 and Elovl6), and higher amounts of these cosubstrates for CerS4. Thus, these studies have found generally good agreement between genomic and metabonomic data in defining that conversion of mESCs to EBs is accompanied by a large number of changes in gene expression and subspecies distributions for both sphingolipids and fatty acyl-CoAs.—Park, H., C. A. Haynes, A. V. Nair, M. Kulik, S. Dalton, K. Moremen, and A. H. Merrill, Jr.  Transcrip

Supplementary key words  embryonic stem cell • embryoid body • sphingolipid • differentiation • ceramide synthase • fatty acyl-CoA elongase

Complex sphingolipids (SLs), including glycosphingolipids (GSLs), are a highly diverse family of compounds (1) that have long been regarded to be important in biological processes such as embryogenesis (2–4). The ceramide (Cer) moieties of SLs and GSLs also change in amount during embryogenesis (2–6), and appear to participate in developmental signaling (4, 7–10).

Ceramides are biosynthesized de novo (1, 11) and by recycling (12, 13) pathways that have been well characterized with respect to the intermediates and genes, as depicted in a scheme (Fig. 1) recently described in this journal (14). De novo biosynthesis from serine and palmitoyl-CoA by serine palmitoyltransferase (SPT) (15) first forms 3-ketosphinganine (3-Keto-Sa), which is reduced to sphinganine (Sa, which is also abbreviated d18:0), then acylated to dihydroceramides (DHCer) by a family of ceramide synthases, CerS, that differ in fatty acyl-CoA selectivity as shown in the expanded pathway diagram, Fig. 1B (16–18). DHCer are incorporated into more complex dihydroSL [dihydrospingomyelins (DHSM), dihydroglucosylceramides (DHGlCer), etc.] or converted to Cer by addition of a 4-trans double bond to produce the sphingosine backbone (abbreviated d18:1) (19), then incorporated into more complex SLs.

From this diagram, one would predict that shifts in the expression of the various CerS, and possibly of the availability

Abbreviations: Cer, ceramide; CMH, ceramide monohexose; DH-Cer, dihydroceramide; DHCMH, dihydroceramide monohexoses; DHGalCer, dihydrogalactosylceramide; DHGlCer, dihydroglucosylceramide; (DH)SL, dihydrospingolipids; DHSM, dihydrospingomyelin; EB, embryoid bodies; GalCer, galactosylceramide; GlcCer, glucosylceramide; GSL, glucosylsphingolipid; IS, internal standard; 3-Keto-SA, 3-ketosphinganine; LIF, leukemia inhibitory factor; mESC, mouse embryonic stem cell; qRT-PCR, quantitative RT-PCR; Sa, sphinganine; Sa1P, sphinganine-1-phosphate; SL, sphingolipid; So, sphingosine; So1P, sphingosine-1-phosphate.

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2 The online version of this article (available at http://www.jlr.org) contains supplementary data in the form of two figures.

This work was funded by the National Institutes of Health grant RR018502 for the Intrgrated Technology Resource for Biomolecular Glycomics at the University of Georgia and GM069338 (Lipid Maps). Its contents are solely the responsibility of the authors and do not necessarily represent the official views of the National Institutes of Health.

*Author’s Choice—Final version full access.

Manuscript received 23 September 2009 and in revised form 24 September 2009.

Published, JLR Papers in Press, September 24, 2009
DOI 10.1194/jlr.M000984

This article is available online at http://www.jlr.org
of the corresponding fatty acyl-CoAs, can affect the N-acyl chain length subspecies of DHCer and Cer (and downstream more complex SLs and GSLs) in cells. To test this hypothesis, we chose mouse embryonic stem cells (mESCs) and embryoid bodies (EBs) because this is a widely studied developmental model (20). EBs are aggregates of cells derived from ESCs that, to some extent, recapitulate early stages of mammalian embryonic development, including

![Fig. 1. Depiction of the early steps of sphingolipid biosynthesis using a modified KEGG pathway scheme (Sphingo-GenMAPP) (14) to facilitate visualization of differences in gene expression in R1 mESCs and EBs. A: Beginning in the upper left of this scheme, serine and palmitoyl-CoA are condensed by serine palmitoyltransferase (represented by 3 SPT genes) to form 3-ketosphinganine (3-Keto-Sa), which is reduced to sphinganine (Sa) by 3-keto-Sa reductase (3KSR, the FVT1 gene) and either N-acylated by ceramide synthases (CerS) to dihydroceramides (DHCer) (shown by arrows leading to the right) or phosphorylated by sphingosine kinase (SphK1 or SphK2) to Sa 1-phosphate (Sa1P), which can be hydrolyzed back to Sa (by phosphatases Ppap2a or Sgpp2) or cleaved to hexadecanal (C16:0-al) and ethanolamine phosphate (EP) by the shown lyase (sphingosine 1-phosphate lyase). DHCer can be hydrolyzed to Sa by ceramidases (Asah1, 2 or 3); incorporated into more complex sphingolipids (DHCer 1-phosphate, DHCer1P, by Cer kinase, CERK), glycosylated to dihydrogalactosylCer (DHGalCer), dihydroglucosylCer (DHGlcCer), converted to dihydrosphingomyelin (DHSM by SM synthases SMS1 or SMS2); or converted to Cer by DHCer desaturase (DES1 or DES2) (or hydroxylated to phytoceramide by DES2, not shown) and either hydrolyzed by ceramidases or converted to downstream metabolites, etc. Also shown are the fatty acyl-CoA elongases (Elovl) because some of these are involved in production of the cosubstrates for CerS. B: Expansion of the pathway to depict the individual molecular species made by different CerS isoforms (formerly named Lass), with abbreviation of the sphingoid base as d18:0 for sphinganine (citing the number of hydroxyl groups, d for dihydroxyl), d18:1 for sphingosine, and the N-acyl chain length and number of double bonds as Cx:y, respectively. As described in the text, the “heat” scale depicts the relative amounts of transcripts for each of these genes for R1 EBs versus mESCs as measured by qRT-PCR (see Fig. 2).]
cell types derived from endoderm, mesoderm, and ectodermal lineages (20). Described in this report are the expression of the CerS genes (16, 21) and the amounts and types of Cer subspecies as part of a broader investigation of all of the known genes for glycoprotein and (glyco)sphingolipid biosynthesis (22). Also analyzed were the cosubstrate fatty acyl-CoAs and expression of transcripts for fatty acid elongases and desaturases as these may also influence the types of Cer subspecies. These studies found that there were shifts in the expression of many of the genes of this pathway as well as in the associated metabolites in EBs versus ESCs.

MATERIALS AND METHODS

Growth and characterization of mESCs and EBs

The R1 and D3 mESCs (23, 24) (ATCC, Manassas, VA) were cultured in the absence of feeder cells on tissue-culture-grade plastic ware precoated with 0.1% gelatin-PBS, as described previously (25). The mESC culture medium consisted of DMEM (Gibco BRL) supplemented with 10% fetal calf serum (GIBCO BRL), 1 mM L-glutamine, 0.1 mM 2-mercaptoethanol, 100 U/ml penicillin, 100 U/ml streptomycin, and 1000 U/ml recombinant human LIF (leukemia inhibitory factor; ESGRO, Chemicon International) at 37°C under 10% CO2.

Differentiation of mESCs into EBs was carried out as described previously (26). mESCs were harvested by trypsinization to convert suspensions of single ESCs into aggregates that were seeded into 10-cm bacteriological dishes at a density of 1 × 10⁵ cells/ml, in 10 ml mESC-medium lacking LIF. EBs were harvested daily as the medium was changed every 2 days and cultures were split one into two at day 4. Cell populations of mESCs and EBs were characterized by flow cytometry using the lineage specific marker CD9 and by transcript analysis using qRT-PCR. Changes in morphology and lineage-specific gene expression for mESC and EB cell populations produced by the Dalton laboratory were previously described (27).

Measurement of gene expression levels using qRT-PCR

Primer design and validation. Primer pairs for SL synthesis genes and control genes were designed within a single exon using conditions described in Nairn et al. (22). Primer sequences and accession numbers for genes in this study are presented in Nairn et al. (27). Individual quantitative (q)RT-PCR reactions were also checked for efficiency 100 ± 5% using reaction raw fluorescence data in the LinReg PCR program (28).

Total RNA isolation and cDNA synthesis. mESCs and EBs (day 6) cell pellets were harvested and flash-frozen in liquid nitrogen and stored at −80°C until use. Cell pellets were homogenized followed by RNA isolation and cDNA synthesis, which were performed as previously described (27).

qRT-PCR reactions and calculation of relative gene expression levels. Triplicate reactions (20 μl each) containing 5 μl of diluted cDNA or genomic DNA template (primer validation reactions), 5 μl of primer pair mix (125 μM final concentration), and 10 μl of IQ™ SYBR Green Supermix (BioRad, Hercules, CA) were assembled in 96-well microtiter plates. Amplification reactions and the calculation of relative gene expression levels were carried out as previously described (27). The normalization gene was selected by determining the expression level for several housekeeping genes across all samples to determine the most stable gene as described in (22). The control gene, Ribosomal Protein L4 (RPL4, NM_024212) was included on each plate to control for run variation and to normalize individual gene expression.

Integration of gene expression data with Sphingo-GenMAPP pathway

GenMAPP (Gene MicroArray Pathway Profiler) v2.1 (29) is a tool to visualize global gene-expression profiles in the context of cell signaling pathways or KEGG metabolic pathways (31). In the context of the present study, the transcript expression data for R1 mESCs and EBs from the qRT-PCR analysis was imported into a GenMAPP pathway for SL metabolism that had been modified (14) based on the current literature (32) (Fig. 1).

LC-ESI MS/MS

SLs. The SLs were extracted from cells using published methods (33–35). Briefly, ~1 to 10 × 10⁶ cells were scraped from culture dishes in a small volume of PBS, then centrifuged to remove excess PBS for a final volume of 0.05–0.1 ml. The cells were suspended in 0.5 ml of CH₂OH and 0.25 ml of CHCl₃, spiked with a single chain-length internal standard for each category of SL being monitored (which we have found to allow quantitative analysis of all of the compounds in this study, as reported in reference 35) (see details below), disrupted using a bath-type sonicator at room temperature, and incubated overnight in a heating block at 48°C. After cooling the tubes to room temperature, 75 μl of 1 M KOH in CH₂OH was added and the mixture was sonicated for 30 s, then incubated for 2 h at 37°C. The solution was neutralized with several μl of glacial acetic acid, 1 ml of CHCl₃ was added followed by 2 ml of water, and sonicated. After centrifugation, the upper phase was carefully removed and discarded. The lower phase was dried using a SpeedVac and redissolved in 0.3 ml of 98% of 97:2:1 CH₃CN/CH₂OH/CH₃COOH (v/v/v) (mobile-phase A) and 2% of 64:15:20 CH₃OH/CH₃CN/CH₂OH/CH₃COOH (v/v/v) (mobile-phase B); both contained 5 mM ammonium acetate.

The methods used for LC-ESI MS/MS analysis of SL metabolites have been reported in detail (34, 35). SL data was collected using a Perkin Elmer Series 200 MicroPump system (PerkinElmer, Norwalk, CT) coupled to a PE Sciex API 3000 triple quadrupole mass spectrometer equipped with a turbo ion-spray source (Applied Biosystems, Foster City, CA). Complex SLs such as Cers, CMHs (ceramide monohexoses, the sum of glucosylceramide and galactosylceramide), and SMs were separated by normal-phase LC using 2.1 mm × 5 cm Supelco NH₂ column (Supelco, Bellefonte, PA) at a flow rate of 1.5 ml/min. The elution protocol was: 0.5 min of pre-equilibration of the column with 98:2 A/B (v/v), sample injection, elution with 98:2 A/B (v/v) for 1.1 min followed by a linear gradient to 82:18 A/B (v/v) in 0.2 min, elution at this composition for 0.4 min, then a linear gradient to 100% B over 0.8 min. Afterwards, the column was reequilibrated by a 0.5 min linear gradient to 98:2 A/B (v/v) and a wash of the column with 98:2 A/B (v/v) for 0.3 min. Quantitation was performed by comparison of each unknown with the spiked internal standard (IS) for that SL class (the IS for each class was an uncommon chain length homolog that has been validated for analysis of other chain lengths) (35): C12-ceramide, C12-sphingomyelin, and C12-glycosylceramide for CMH, all of which were obtained from Avanti Polar Lipids (Alabaster, AL). The ISs were added as 10 μl of a 50 μM of stock for a total of 500 pmol of each IS molecular species per sample (see above). Student’s t-test was used for statistical analyses.
Fatty Acyl-CoAs. The fatty acyl-CoAs were extracted from cells using a published method (36). The medium from cells cultured in 100 mm dishes was aspirated and cells were gently washed twice with 5 ml of 4°C PBS, ∼1 ml of ice-cold PBS was added, and the cells were scraped from the dishes using a cell scraper and transferred to chilled 15 × 100 mm screw-cap borosilicate tubes with Teflon-lined caps (Kimble Chase, Vineland, NJ). An additional ∼1 ml of PBS was used to recover the remaining cells by this same procedure and these were added to the first. After centrifugation (1500 rpm, 3 min), the PBS was removed by aspiration and 0.5 ml of CH₃OH containing 1 mM EDTA was added together with 10 μl of an internal standard mixture that contained 100 pmol each of C₁₇:0- and C₂₅:0-CoA, which were prepared in CH₃OH/CHCl₃ (2:1 v/v) containing 30 mM triethylammonium acetate. After sonication (three times at ∼0.5 min each), 0.25 ml of CHCl₃ was added followed by another brief sonication.

The single-phase extraction mixture was incubated in a 50°C heating block for 30 min. After cooling to room temperature, CHCl₃ and water (0.25 ml each) were added with mixing by vortexing after each addition. After brief centrifugation using a table-top centrifuge, the fatty acyl-CoAs were in the upper phase and interface and most of the other lipids (which would interfere with the subsequent reverse-phase LC) were in the lower phase. The upper phase was removed with a Pasteur pipette and transferred into a clean screw-cap glass tube. The upper layer was adjusted with 5% CH₃OH (v/v), 5% butanol (v/v), and 5% triethylamine, the sample was injected, and mobile phase C and 52% mobile phase D (CH₃CN containing 0.05% triethylamine), the sample was injected, and mobile phase C was used to maintain the column temperature at 40°C.

The flow rate was 0.2 ml/min in binary gradient mode with the following elution program: the column was equilibrated with mobile phase C [water/CH₃CN (85:15, v/v), containing 0.05% triethylamine], the sample was injected, and mobile phase C was continued for 5 min, followed by a 15 min gradient to 48% mobile phase C and 52% mobile phase D (CH₃CN containing 0.05% triethylamine) during which the long-chain- and very long-chain-fatty acyl-CoAs eluted. Afterward, the column was washed by a 1 min gradient to 100% D and a 5 min hold at 100% D, followed by reequilibration of the column by a 1 min gradient to 100% C and a 5 min hold at 100% C before injection of the next sample.

MS/MS of fatty acyl-CoAs in positive ion mode was performed on a 4000 QTrap triple quadrupole linear ion trap mass spectrometer (Applied Biosystems) as recently described (36).

RESULTS

Characterization of R1 mESCs and EBs

The R1 mESCs and EBs were examined for gross morphological changes and transcript abundances as markers for these cells. The mESCs expressed Zinc finger protein 42 (zfp42/Rex1), Gastrulation brain homeobox (GBX2), and Nanog homeobox (Nanog) as pluripotent markers, and the qRT-PCR transcript profiles for Goosecoid (Gsc), Zinc finger protein of the celebellum 1 (Zic1), SRY-box containing gene 17 (Sox17), Alpha fetoprotein (Afp), and GATA binding protein 6 (Gata6) were typical for the transition from mESCs to EBs (22).

Differences in the expression of SL metabolism genes in R1 mESCs and EBs using Sphingo-GenMAPP

The relative transcript expression levels of R1 EBs versus mESCs are depicted in the pathway diagram (Fig. 1) and with the fold differences shown by the “heat” scale defined on the right side of the figure. There were no differences in the transcript levels for serine palmitoyltransferases (SPT1 and SPT2), the first enzymes of the pathway, but EBs had higher expression of transcripts for 3-ketosphinganine reductase and some of the CerS isoenzymes (CerS1, 3, and 6), as is also shown by bar graphs in Fig. 2 (upper panel). The transcripts were also higher in EBs for enzymes that use Cer for biosynthesis of more complex SLs, i.e., both of the SM synthases and the glycosyltransferases that produce GlcCer and GalCer (Fig. 1). EBs had lower transcript abundances for the major desaturase that synthesizes Cer from DHCer (dihydroCer desaturase, DES1) and transcripts for the other desaturase (DES2), which makes phytoceramides (37) were barely detectable in both mESCs and EBs. Because the CerS enzymes use very long-chain fatty acyl-CoAs that are made by elongation of palmitoyl- and stearoyl-CoAs, the transcripts for the pertinent fatty acyl-CoA elongases were also analyzed by qRT-PCR. As shown in Fig. 1 and 2B, the transcript level for Elovl6, a fatty acyl-CoA elongase that catalyzes the elongation of saturated and monosaturated acyl-CoAs with 12-16 carbons (38, 39), was 4-fold higher for EBs than mESCs. A fatty acyl-CoA elongase that extends C₁₆₀-CoA to C₂₄₀-CoA, Elovl3 (38, 40, 41), showed 3-fold higher transcript levels for EBs than mESCs, although lower transcript levels were observed for Elovl1, which is also responsible for elongating very long-chain saturated fatty acyl-CoAs (38, 41). EBs also had ∼3.5-fold higher Elovl2 (37, 42) transcript levels, no evident change in Elovl4 (38, 44) transcripts, and slightly higher Elovl5 (38, 43) transcript abundance (which are all elongases for polysaturated fatty acids).

These differences in gene expression imply that the SL backbones of EBs might differ from mESCs by having: i) higher proportions of subpecies with longer acyl chains (i.e., >C₁₆) because EBs have a higher transcript abundance of CerSI, which is relatively specific for stearoyl-CoA (C₁₈:0) (45), and CerS₃, which is selective for very-long-chain fatty acyl-CoAs (>C₂₀) (46); ii) higher, lower, or unchanged C₁₆-Cer because an elevation in (DH)Cer as CerS₅ (47) increases in DHSL as DES1 is lower and DES2 unchanged; and iv) possibly lower amounts of (DH)Cer because the transcripts encoding most of the enzymes that use (DH)Cer were elevated (including the transcripts for one of the ceramidases, Asah2). The gene expression data also suggest that the EBs have the capacity to elongate fatty acyl-CoAs to produce the cosubstrates that are used to make these Cer
would predict that the relative amounts of the different sub-
species might correlate with the relative abundances of the
respective CerS transcripts, which have been regraphed in
Fig. 3B to show the differences between mESCs and EBs.
The order of the CerS genes has been arranged so they cor-
respond to the respective types of Cer that are made, i.e.,
C16-Cer (CerS6 and CerS5), C18-Cer (CerS1), C20 ± 2 Cer
(CerS4), C24-Cer (CerS2), and C26-Cer (CerS3) (c.f. Fig. 1).

Fig. 2. Relative transcript abundance of CerS (Lass) genes (A) and fatty acyl-CoA elongase (Elovl) genes (B) by qRT-PCR for R1 mouse embryonic stem cells and day 6 EBs. A: Upper panel, fold change in relative transcript abundance of CerS genes for EBs/mESCs. A value of zero represents no change in transcript abundance. Values greater than zero represent an increase in transcript abundance in EBs relative to mESCs and values less than zero represent a decrease in EBs relative to mESCs. Values that are off-scale are indicated by a hatched bar and the fold-change value is given above the bar. Lower panel, bars represent average Ct values from triplicate samples normalized to the housekeeping gene RPL4 (as described in the Materials and Methods). Relative transcript abundance for mESCs (white bars) and EBs (black bars) are plotted on a log scale for each gene assayed. Error bars represent 1 SD from the mean. Transcript values less than 1 x 10^-6 are below the threshold of detection. B: Relative transcript abundance of Elovl genes in R1 mESCs versus EBs determined by qRT-PCR. Upper panel, fold change in relative transcript abundance of fatty acyl-CoA elongase gene family members (Elovl1-6) for EBs/mESCs as described for A above. Lower panel, relative transcript abundance for mESCs (white bars) and EBs (black bars) are plotted as described for A above.

Characterization of the Cer backbone subspecies of R1 mESCs and EBs

Fig. 3A shows the relative proportions of the Cer subspecies in R1 mESCs (day 0) and EBs on days 4 and 6 as “do-
nut” charts. The rationale for the latter depiction is that
CerS have approximately the same Km for the sphingoid base substrate (48); therefore, as a first approximation, one

subspecies. To test these hypotheses, the Cer subspecies
(and fatty acyl-CoAs) were analyzed by LC-ESI MS/MS.

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2 Bar graphs with the amounts of each subspecies of (DH)Cer, (DH)CMH, and (DH)SM for R1 and D3 mESCs and EBs are available as Fig. I and II in the supplementary data.
Analysis of the fatty acyl-CoA species in R1 mESCs and EBs

The amounts of the fatty acyl-CoAs in R1 mESCs and EBs are shown in Fig. 5. During the time course for conversion of mESCs to EBs, C18:0- and C18:1-fatty acyl-CoAs increased by approximately 2-fold, as did many of the very long-chain (C22:0-, C24:0- and C24:1-) fatty acyl-CoAs. In contrast, palmitoyl-CoA derivatives (C16:0-) were approximately the same in mESCs and EBs. Thus, these results are consistent with the observed changes in the higher amounts of C18:1-, C22:0-, C24:0-, and C24:1-fatty acyl-CoAs in EBs versus mESCs. These results additionally indicate that the CerS that would use these fatty acyl-CoAs to produce the pertinent Cer subspecies are not hampered by a lack of cosubstrate availability during EB development.

Characterization of the DHCer and Cer backbone subspecies of (DH)SL in R1 mESCs and EBs

The DHCer of mESCs and day 6 EBs also exhibited shifts in fatty acyl-chain length (Fig. 4A) that correlated with the differences in CerS expression that were similar to the changes in Cer (c.f., Fig. 4A and Fig. 4D), namely, lower proportions of C16-DHCer and more C18-and C24/24:1 DHCer with the greatest increase in C24:1-Cer. Because DHCer is the first N-acylated intermediate of de novo sphingolipid biosynthesis (Fig. 1) versus Cer, which is formed next and during the turnover of more complex sphingolipids, perhaps this accounts for the closer resemblance between the chain length subspecies of DHCer and the transcript abundance of the CerS isoforms (c.f., Fig. 3A and 4).

The (DH)CMH (Fig. 4B and E) and SM (Fig. 4F) of EBs displayed higher proportions of the C24/C24:1 backbones, however, most of the other subspecies differences that were seen in Cer and DHCer were not evident in these downstream metabolites.

There were also differences in the proportion of DHCer versus Cer backbones in EBs versus mESCs (which are not evident in Fig. 3 or 4 because the data are shown only as proportions within each category). When summed across all chain lengths and categories, the total amount of these sphingolipids did not differ between mESCs and day 6 EBs cells (i.e., 780 vs. 740 pmol/mg protein, respectively), but the DHSL percentage increased from ~8% for mESCs to 19% for EBs. This finding was consistent with the lower DES1 expression in EBs versus mESCs (Fig. 1).

DISCUSSION

The goal of these studies was to explore whether mESCs undergo changes in the mRNA levels for the early steps of de novo SL biosynthesis upon conversion to EBs, then to compare these with the Cer subspecies that are present.
There were increases in the mRNAs for many of the early enzymes of SL biosynthesis (Fig. 1), including some of the CerS (1, 3, and 6) and downstream synthases for SMs and GSLs, and decreases in other CerS (4 and 5, with 2 unchanged), the sphingosine kinases, and two of three ceramidases. Although all of these are worthy of further investigation, this study focused on the lipid backbone subspecies because they can be readily analyzed by LC-ESI MS/MS and serve as a relatively easy way to determine how closely changes in gene expression match changes in these important compounds.

For all of the mESC and EB SLs investigated in this study, the prevalent N-acyl-linked fatty acid was C16:0 (palmitate), which is added by CerS5 and CerS6. Consistent with this, the transcript abundance of CerS5 plus CerS6 is higher than that for the other CerSs (1, 3, and 4) except CerS2 (Fig. 2). Considering the relative abundance of CerS2 mRNA, it is somewhat surprising that the C24-chain-length backbones are not more prevalent (supplementary Fig. II). We do not know if this is because CerS2 protein amount or enzymatic activity are not equivalently high, but it is worth noting that another factor might be a limited availability of the cosubstrate C24- and C24:1-fatty acyl-CoAs as these are <10% of the levels of the C16- and C18-fatty acyl-CoAs (Fig. 5). Consistent with this possibility, the higher proportions of C24-chain length Cer in EBs versus mESC occur without an increase in CerS2 mRNA whereas the amounts of the very long-chain fatty acyl-CoAs doubled.

The findings for the fatty acyl-CoAs are, by themselves, interesting because there have been no previous studies of whether mESC and EB differ in the mRNA levels for this pathway. As shown in Fig. 1, the mRNA for Elovl6 was higher for EBs than mESCs, which matched the ~2-fold higher stearoyl- and oleoyl-CoAs (Fig. 5). Furthermore, the higher C24-chain-length fatty acyl-CoAs correlated with only one of the two elongases (Elovl3, which increased, vs. Elovl1, which decreased), therefore, it is likely that Elovl3 is responsible for the observed increases in very long-chain fatty acyl-CoAs.

Decreases in the proportion of C16- and increases in very long-chain DHCer and Cer were two of the most consistent shifts in backbone composition not only for these compounds but also the downstream (DH)SM and (DH)CMH (Figs. 3 and 4) and were even more evident in the D3 cell line (Fig. 6). The functional significance of these subspecies shifts for embryogenesis, if any, is not known; however, synthesis of C16-Cer is elevated during the apoptosis (49–51), and differences in the time course of changes in C16- versus C24-Cer have been associated with different phases of apoptosis (52, 532). Changes in C18-Cer might also be important because this subspecies is associated with cell growth control (53).

EBs have been widely used as a model system for studies of the formation of the inner cell mass of the mouse embryo shortly before implantation, which has been proposed to involve apoptosis (54) and, thus, might involve Cer. Another important function for Cer-induced apoptosis...
might be to eliminate residual, pluripotent EBs in later stages of differentiation (9).

It is also interesting that the ratio of DHCer to Cer was higher in most of the SLs of EBs than mESCs (c.f. left vs. right columns of supplementary Figs. I and II), which correlated with the lower DHCer desaturase (DES1) mRNA in EBs versus mESCs (Fig. 1). Dihydroceramides are generally thought not to play an important role in apoptosis, but can participate in other biological processes such as autophagy (55), which has also been proposed to be involved in the formation of the inner cell mass of the mouse embryo (56).

In addition to shifts in the backbone subspecies, the total amounts of (DH)Cers were higher in R1 and D3 EBs than mESCs (supplementary Figs. I and II), which might participate in the regulation of primitive ectoderm cell polarity and/or Cer-induced apoptosis in embryonic development (8, 57). In contrast, the CMH of R1 and D3 EBs were much lower than for mESCs, despite increases in the transcripts for the synthases for GlcCer and GalCer (Fig. 1), which may suggest that these simple GSLs are being used for the biosynthesis of more complex GSLs that are important during development (3, 58, 59). This would be an interesting direction for future studies.

Since the development of tools for analysis of mRNAs using microarrays and high-throughput qRT-PCR for transcript profiling, there have been many studies of changes in gene expression during development (60–62); however, prior to this study, the CerS gene family has not been included because it was only recently characterized (16). There have also been no previous studies of this pathway that have combined analysis of the mRNAs for these genes with the quantitation of the related metabolites. Thus, it is encouraging that, for the most part, there was reasonable agreement between the predictions from the genomic analysis and what was found by this focused lipidomic survey. This type of approach might uncover new ways in which SLs are involved in stem cell biology and embryonic development, including where disruption of SL metabolism results in birth defects (63, 64).

The authors thank Dr. Michael Pierce at the University of Georgia for useful comments throughout the course of this work, and Dr. Akemi Suzuki (Tokai University) for helpful discussions concerning annotation of sphingolipid metabolic genes.

Fig. 6. Subspecies distributions of (dihydro)ceramides, (dihydro)ceramide monohexoses and (dihydro)sphingomyelins of D3 mouse embryonic stem cells and day 6 embryoid bodies. The sphingolipids were analyzed by LC ESI-MS/MS as described in reference 34–36 and are shown with the ESCs as the upper member of each pair and EBs as the lower dataset. A: Dihydroceramides (DH-Cer); B: dihydroceramide monohexoses (DHCMH); C: dihydroceramidesphingomyelins (DHSM); D: ceramides (Cer); E: ceramide monohexoses (CMH); and F: sphingomyelins (SM).

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