Dynamic simulation of cardiolipin remodeling: greasing the wheels for an interpretative approach to lipidomics

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Abstract Cardiolipin is a class of mitochondrial specific phospholipid, which is intricately involved in mitochondrial functionality. Differences in cardiolipin species exist in a variety of tissues and diseases. It has been demonstrated that the cardiolipin profile is a key modulator of the functions of many mitochondrial proteins. However, the chemical mechanism(s) leading to normal and/or pathological distribution of cardiolipin species remain elusive. Herein, we describe a novel approach for investigating the molecular mechanism of cardiolipin remodeling through a dynamic simulation. This approach applied data from shotgun lipidomic analyses of the heart, liver, brain, and lung mitochondrial lipidomes to model cardiolipin remodeling, including relative content, regiospecificity, and isomeric composition of cardiolipin species. Generated cardiolipin profiles were nearly identical to those determined by shotgun lipidomics. Importantly, the simulated isomeric compositions of cardiolipin species were further substantiated through product ion analysis. Finally, unique enzymatic activities involved in cardiolipin remodeling were assessed from the parameters used in the dynamic simulation of cardiolipin profiles. Collectively, we described, verified, and demonstrated a novel approach by integrating both lipidomic analysis and dynamic simulation to study cardiolipin biology. We believe this study provides a foundation to investigate cardiolipin metabolism and bioenergetic homeostasis in normal and disease states.—Kiebish, M. A., R. Bell, K. Yang, T. Phan, Z. Zhao, W. Ames, T. N. Seyfried, R. W. Gross, J. H. Chuang, and X. Han. Dynamic simulation of cardiolipin remodeling: greasing the wheels for an interpretative approach to lipidomics. J. Lipid Res. 2010. 51: 2153–2170.

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Lipids are critical for the regulation and control of cellular function, membrane dynamics, and signal transduction. The systematic identification and analysis of all cellular lipids in a biological sample is embodied by the rapidly developing field of lipidomics, which has been enabled by recent advances in mass spectrometry (1–5). A multidimensional MS-based shotgun lipidomics (MDMS-SL) approach for analysis of cellular lipidomes has been demonstrated as one of the most systematic and quantitative high-throughput platforms for global lipidomics capable of analyzing thousands of lipid molecular species directly from the lipid extracts of biological tissues (4, 5). Analysis of lipidomic data requires extensive use of bioinformatics and computational algorithms for the accurate identification and quantification of lipid molecular species and classes (5–9). However, most advances in bioinformatics in lipidomics have largely focused on lipid identification rather than interpretation of alterations in biological function resulting in adaptive or pathologic changes in lipid metabolism (10). Thus, development of bioinformatic and systems biology approaches for the interpretation of lipidomic networks would significantly advance the understanding of the roles of lipids in biological systems (3).

Cardiolipin is a complex polyglycerophospholipid found almost exclusively in mitochondrial membranes of eukaryotic organisms (11). Cardiophilin is intricately involved in the efficient production of ATP through utilizing the proton gradient as well as maintaining mitochondrial functionality (12–14). The diversity of cardiolipin molecular species varies markedly between tissues as well as among

Abbreviations: CDP-DAG, cytidine diphosphate-diacylglycerol; MDMS-SL, multidimensional MS-based shotgun lipidomics; PC, phosphatidylcholine; PE, phosphatidylethanolamine; PG, phosphatidylglycerol; TOF, time of flight.

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This diversity is regulated by differences in cardiolipin biosynthesis and catabolism, remodeling acyl selectivities, and transacylase as well as acyltransferase activities (20, 21). Cardiolipin is de novo synthesized in the mitochondrial membrane by the condensation of phosphatidylglycerol (PG) and cytidine diphosphate-diacylglycerol (CDP-DAG) (Fig. 1). The newly formed polyglycerophospholipid is referred to as immature cardiolipin, which is then deacylated and extensively remodeled using acyltransferase(s) and/or transacylase(s) (21, 22). Cardiolipin remodeling utilizes acyl CoA and acyl chains from the sn-2 position of choline glycerophospholipids (PC) and ethanolamine glycerophospholipids (PE) accordingly as acyl donors (20, 23, 24). The balance of acyltransferase and transacylase activities as well as acyl chain selectivities is reflected in the acyl chain pool, which sculpts nascent cardiolipins into mature cardiolipin molecular species.

Based on known biological pathways and MDMS-SL-derived data, we developed a dynamic simulation approach that recapitulates the temporal reconstruction of immature cardiolipins into mature cardiolipins by remodeling from available donor acyl chains. The sequential remodeling events can be visualized graphically for intuitive comparison of the dynamic aspects of cardiolipin remodeling. The biological parameters in these simulations can be modified to compare cardiolipin profiles obtained from MDMS-SL. Moreover, the algorithmic approach utilized in this simulation not only demonstrates the presence and concentration of isomeric cardiolipin species, which have been identified by MDMS-SL, but also predicts the existence of a variety of very low abundance cardiolipin molecular species, many of which cannot be accurately determined using currently available technologies. Furthermore, experimentally measured profiles of cardiolipin molecular species can be recapitulated with this simulation utilizing specific distributions of acyl chains reflecting the coordinated activities of phospholipases, acyltransferase(s), and/or transacylase(s). Collectively, we demonstrated that the cardiolipin dynamic simulation combined with MDMS-SL analyses of PG, phosphatidylcholine (PC), phosphatidylethanolamine (PE), and acyl CoA is a powerful new approach for determining the presence of isomeric cardiolipin molecular species and assessing the enzymatic activities that mediate the extent of cardiolipin remodeling. We contend that this dynamic simulation will provide novel insight into the mechanisms regulating the complex tissue-specific cardiolipin molecular species distributions and that new insight into the molecular mechanisms underlying alterations in acyl chain selectivity in health and disease can be accrued.

**MATERIALS AND METHODS**

**Mitochondrial isolation**

C57BL/6J wild-type male mice (4 months of age) were purchased from the National Institute on Aging (Bethesda, MD). All animal procedures were performed in accordance with the Guide for the Care and Use of Laboratory Animals and were approved by the Animal Studies Committee at Washington University. Mice were euthanized by asphyxiation with carbon dioxide. Mouse organs were perfused with phosphate-buffered saline plus heparin to remove possible plasma contamination before dissection. Brain cortex, heart, liver, and lung of mice were harvested immediately after perfusion.

Nonsynaptic brain mitochondria were isolated as previously described in detail using discontinuous Ficoll and sucrose gradients (25, 26). Myocardial, hepatic, and lung mitochondria were isolated as previously described (27). All experiments were performed on ice in a 4°C cold room. Briefly, the heart, liver, or lung was diced using a razor blade and homogenized in homogenization buffer [HM; 0.21 M mannitol, 70 mM sucrose, 0.1 mM EDTA-K, 0.5% BSA, 10 mM Tris-HCl, 1× protease inhibitor cocktail (complete®, Roche, Germany) pH 7.4] in a ratiometric volume of HM buffer to tissue of 8:1. The homogenate was then centrifuged at 700 g for 5 min. The supernatant was collected and the pellet was homogenized and recentrifuged. The supernatant was collected, and the pooled supernatants were centrifuged at 900 g for 5 min. The pellet was discarded, and the supernatant was collected and centrifuged at 10,000 g for 10 min. The pellet was resuspended in HM buffer and layered on a 1.6, 1.2, 1.0, and 0.8 M discontinuous sucrose gradient and spun for 50,000 g for 2 h in a SW 28 rotor. The highly enriched mitochondrial fraction was collected at the 1.6/1.2 M sucrose interface and recentrifuged at 18,000 g for 15 min. The pellet was resuspended in HM buffer without BSA and recentrifuged at 8,000 g for 10 min. The pellet was resuspended in HM buffer without BSA, and the protein concentration of the resuspended mitochondrial fractions was determined using the BCA protein assay (Thermo Scientific, Rockford, IL).
Instrumentation and MS

A triple-quadrupole mass spectrometer (Thermo Scientific TSQ Quantum Ultra Plus, San Jose, CA) equipped with a Nanomate device (Advin Biosciences Ltd, Ithaca, NY) and Xcalibur system software was utilized for MDMS-SL analysis as previously described (28, 29). Lipid molecular species were identified, quantified, and acyl chain designation determined using a MDMS-SL approach as previously described (1, 33, 34). Product-ion analyses of cardiolipin molecular species by using a high mass accuracy/high resolution mass spectrometer [i.e., Orbitrap-LTQ mass spectrometer (Thermo Scientific San Jose, CA)] were also performed as previously described (32) to verify the presence of very low abundance cardiolipin isomers.

MDMS-SL analysis of mitochondrial lipids

The murine mitochondrial lipidome from brain, liver, lung, and heart was determined using MDMS-SL, as previously described (26). Briefly, a 1 mg aliquot of mitochondrial protein was transferred to a disposable borosilicate glass tube (16 × 100 mm). Internal standards were added based on protein concentration (in nmol/mg protein) and included 16:1-16:1 PE [38 (liver or heart), 19 (lung), and 50 (brain)], 14:1-14:1 PC [30 (liver or heart), 15 (lung), and 22:5 (brain)], 15:0-15:0 PG [4 (liver or heart), 2 (lung), and 6 (brain)], and 1T14:0 cardiolipin [8 (liver or heart), 9 (lung), and 50 (brain)]. A modified Bligh and Dyer procedure was used to extract lipids from each mitochondrial preparation as previously described (33). Each lipid extract was reconstituted with a volume of 200 μl/mg mitochondrial protein in chloroform/methanol (1:1, v/v). The lipid extracts were flushed with nitrogen, capped, and stored at −20 °C for ESI/MS analysis. Each lipid solution was diluted 25- to 100-fold depending on experimental conditions.

Acyl CoA extraction and analysis

Extraction and analysis of acyl CoA was performed as previously described (34) with modification. Briefly, 4-month-old C57BL/6j male mice were sacrificed, and tissue samples were immediately harvested and clamp frozen in liquid nitrogen. The rapid rate of tissue harvest (~15–30 s) is critical for accurate assessment of acyl CoA levels and distribution. Tissue samples were then pulverized in liquid nitrogen, and 50–80 mg of tissue powder was weighed and homogenized in chloroform/methanol (1:2, v/v) in an ice/ethyl alcohol bath to immediately denature enzymes. An internal standard was added (17.0 μg) at a concentration of 50, 90, 50, and 60 pmol/mg protein for the liver, heart, lung, and cortex, respectively. This internal standard was selected, because the endogenous mass of this species was minimal, as demonstrated by MALDI-time of flight (TOF)/MS lipid analysis (supplementary Fig. 1). Chloroform and deionized water were added, yielding a ratio of chloroform/methanol/deionized water of 2:2:1 (v/v/v) and a volume of at least 10 μl/mg of tissue powder. The samples were then homogenized in an ice/ethyl alcohol bath for 2 min and then incubated on the ice/ethyl alcohol bath for 2 min, vortexing occasionally. The homogenate was then centrifuged at 23,000 g for 10 min in a 4 °C cold room. The methanol and deionized water layer (upper phase) was collected.

Modeling the rate, catabolism, and sn-1 and sn-2 selective remodeling of cardiolipin molecular species. To quantitatively describe the cardiolipin remodeling process, we developed software that simulates the dynamics of transferring acyl chains into cardiolipin. The simulation is based on incorporation of determined fatty acids from sn-2 positions of PC and PE or acyl CoA at designated concentrations into cardiolipin molecular species under mass action kinetics, with all four cardiolipin acyl chain positions treated equivalently. We first describe the rate of change of a given cardiolipin species with the chains α,α,α,α, at the sn-1, sn-2, sn-1’, and sn-2’ positions, respectively. The process of chain substitution from PC, PE, and acyl CoA into cardiolipin corresponds to a dynamic equation:

\[ \frac{d[a_1a_2a_3a_4]}{dt} = k_{pc}^{(1)} \left[ a_1 \right]_{pc} \sum_{[\beta]} \left[ a_2 \beta a_3 a_4 \right]_{cl} + k_{pc}^{(2)} \left[ a_1 \right]_{pc} \sum_{[\beta]} \left[ a_2 \beta a_3 a_4 \right]_{cl} + k_{pc}^{(1')} \left[ a_1 \right]_{pc} \sum_{[\beta]} \left[ a_2 a_3 \beta a_4 \right]_{cl} + k_{pc}^{(2')} \left[ a_1 \right]_{pc} \sum_{[\beta]} \left[ a_2 a_3 a_4 \beta \right]_{cl} - \left[ a_1 a_2 a_3 a_4 \right] \left[ \beta \right]_{pc} + k_{pc}^{(1')} \left[ a_1 \right]_{pc} \sum_{[\beta]} \left[ a_2 \beta a_3 a_4 \right]_{cl} + k_{pc}^{(2')} \left[ a_1 \right]_{pc} \sum_{[\beta]} \left[ a_2 a_3 \beta a_4 \right]_{cl} + k_{pc}^{(1)} \left[ a_1 \right]_{pc} \sum_{[\beta]} \left[ a_2 a_3 a_4 \beta \right]_{cl} \right] + (\text{analogous PE terms}) + (\text{analogous acyl CoA terms}) \]

Here \( k_{pc}^{(1)} \) is the reaction rate constant between PC and one of the sn-1 positions of cardiolipin, and \( k_{pc}^{(2)} \) is the reaction rate constant between PC and one of the sn-2 positions. Thus, the model can accommodate differential kinetic flux present at the sn-1 and sn-1’ positions versus sn-2 and sn-2’ positions in cardiolipin that mimic the phospholipase activities that contribute to the generation of monohyocardiolipin and cardiolipin remodeling. For the analogous PE terms, the PE-specific values \( k_{pe}^{(1)} \) and \( k_{pe}^{(2)} \) replace the corresponding PC rate parameters and likewise for the acyl CoA terms. For simplicity, our model assumes that the relative reactivity of the sn-1 and sn-2 positions is a property of the cardiolipin molecule, i.e., \( k_{pc}^{(1)} / k_{pc}^{(2)} = k_{pc}^{(1')} / k_{pc}^{(2')} \) and likewise for the acyl CoA parameters.
To account for the steady synthesis of immature cardiolipin from PG, which will replace the proportion of cardiolipin that is catabolized, we added two additional terms to the simulation. One term corresponds to the constant catabolism of a small fraction of the cardiolipin, described by an exponential decay parameter $\gamma$. This catabolism is assumed to be nonspecific, i.e., all cardiolipin species are catabolized at the same rate. The second term corresponds to a constant injection of immature cardiolipin that keeps the total quantity of cardiolipin in the system constant. Inclusion of these terms allows for the simulation of the relative importance of rates of cardiolipin turnover (catabolism and de novo synthesis) versus rates of acyl chain substitution. Overall, the equation describing the kinetics is given by:

$$\frac{d[a_{\alpha_1}\alpha_2\alpha_3\alpha_4]_{\text{CL}}}{dt} = \text{chain swapping} - \gamma[a_{\alpha_1}\alpha_2\alpha_3\alpha_4]_{\text{CL}} + \text{incorporation rate of immature cardiolipin},$$

where the chain exchange terms are described in the equation above.

This equation is integrated numerically with time step $\Delta t$, a parameter that determines the amount of cardiolipin that will be remodeled in a single iteration using designated acyl chains from PC and PE sn-2 positions, as well as acyl CoA. In practice, the numerical integration is performed using dimensionless variables, with a timestep small enough that only a small fraction of the cardiolipin is remodeled in any given iteration to maintain realistic behavior. In these dimensionless variables, this condition is typically satisfied as long as $\Delta t \ll 1$ (with some dependence on the distribution of species concentrations of each of the acyl chain sources), and as a conservative criterion in most simulations we use a timestep of $\Delta t \approx 0.05-0.15$. Whenever practical, a larger value of $\Delta t$ is used to reduce the simulation computation time.

The dynamic simulation can determine the cardiolipin concentrations at each time point in three output formats: 1) the concentrations of all molecular species as specified by the acyl chains in each of the four acyl positions (full chain output), 2) the aggregate concentrations of all molecular species at each designated number of carbons and double bonds, a format that can be easily compared with experimental mass spectrometry measurements (mass spectrometry output), or 3) a hybrid format in which the cardiolipin species are sorted by the numbers of carbons and double bonds, which allows for simplified analysis of the isomeric species that correspond to each ion of cardiolipin species determined at a certain $m/z$ by MS.

**Acyl chain distributions used for cardiolipin remodeling.** The acyl chains utilized for the formation of mature cardiolipin molecular species could originate from a transacylation reaction from PC or PE or via an acyltransferase reaction from acyl CoA (38–40). To model these selective aspects of the remodeling machinery, we devised an approach that allows for the input of PC and PE sn-2 positional acyl chain composition as well as acyl CoA chain data utilizing designated acyl chain selectivity (Fig. 2 (3)). Alternatively, it is possible to manually select the acyl chains from PC and PE as well as from acyl CoA to be contributed to cardiolipin remodeling. This aspect of the remodeling program allows for the capability of defining the possible ratios of acyltransferase versus transacylase activities based on the fatty acids that are present in discrete cardiolipin molecular species, in addition to the interpretation of acyl selectivity.
Visualization and dynamic simulation output for cardiolipin remodeling. To view the sequential remodeling of cardiolipin molecular species, a graphical viewer was written in JAVA for enhanced portability across operating systems. This viewer displays the percentage distribution of cardiolipin species as organized by the carbon:double bond composition (an output of mass spectrometric format). Sequential remodeling of cardiolipin species can be viewed on a frame-by-frame basis or as an animated movie. All software is available at http://bioinformatics.bc.edu/~jchuang/dynamic.html.

RESULTS

Analysis of cardiolipin molecular species and their acyl chain regioisomers in myocardial, hepatic, lung, and brain mitochondria revealed a diverse distribution that provided insight into the biochemical processes that mediated the distinct distributions of cardiolipins in these tissues (supplementary Tables I and VII). Based on the premise that the diversity of cardiolipin molecular species in various organs was dependent on the availability of different acyl chains in PC and PE as well as the acyl CoA composition and the acyl selectivities of the involved acyltransferases and transacylases, a dynamic model was constructed to identify the critical reactions necessary for the generation of mature cardiolipins.

Using the MDMS-SL measured concentrations of individual cardiolipin species as well as those of the donor acyl chains from PC, PE, and acyl CoA, we computed the relative contributions from each of the donor substrates to the observed cardiolipin distribution. A distribution of acyl chains was generated from PC, PE, and acyl CoA that resembled the acyl composition of myocardial, hepatic, lung, or brain cardiolipin, and this distribution was used for the chain substitution process (see “Materials and Methods”). Based on the analysis of predicted immature cardiolipin from PG molecular species, the proportional amount of newly synthesized cardiolipin was inferred to contribute to the observed cardiolipin distribution in each tissue. Generally, the amount as well as ratio of saturated acyl chains found in PG molecular species served as reliable indicators of the relative content of immature cardiolipin molecular species being represented in the mature pattern of remodeled cardiolipin. This assumption seems reasonably justified, because the content of saturated acyl chains in mature cardiolipins is quite low and these are likely derived from either de novo cardiolipin biosynthesis or from saturated acyl chains derived from acyl CoA during remodeling. Because the sn-2 positions of PC and PE contain almost solely unsaturated acyl chain species, it is unlikely that the saturated acyl chains in mature cardiolipin transacylated from either the PC or PE pools. Using this approach, we examined the dynamic time course of altered cardiolipin acyl chain distributions of four bioenergetically diverse organs: the heart, liver, and brain (see next sections), as well as lung (see supplementary materials).

Dynamic simulation of myocardial cardiolipin molecular species

Examination of the predicted immature cardiolipin molecular species generated by the condensation of myocardial mitochondrial PG (supplementary Table II) revealed that there is only a minimal contribution of cardiolipin molecular species originating from de novo biosynthesis in the mature remodeled myocardial cardiolipin distributions identified by MDMS-SL (Fig. 3A). Thus, the amount of decay and replacement estimated in the dynamic simulation for myocardial cardiolipin was minimal due to the lack of immature cardiolipin species in the mature cardiolipin distribution. We determined by MDMS-SL as well as experimentation a distribution of acyl chains as 16:0 (0.27%), 16:1 (0.6%), 16:2 (0.03%), 18:0 (0.13%), 18:1 (9.92%), 18:2 (67.79%), 18:3 (0.19%), 20:1 (0.09%), 20:2 (1.98%), 20:3 (2.1%), 20:4 (0.84%), 22:5 (1.43%), and 22:6 (14.36%) and used this distribution to simulate myocardial cardiolipin remodeling that generated a cardiolipin profile best representing the profile obtained by MDMS-SL. A Δ value (see “Materials and Methods”) of 0.1 was used in the simulation. Figure 3 demonstrates the sequential remodeling of cardiolipin from the predicted immature cardiolipin distribution after 0, 50, 100, and 250 iterations (Fig. 3A–D, respectively). Simulation data can be found in supplementary materials and viewed by downloading the customized cardiolipin remodeling viewer.

During the progression of remodeling steps, the stepwise iterations demonstrate the structural remodeling of cardiolipin molecular species at different stages of the remodeling process. The process of cardiolipin remodeling from predicted immature state to a mature state involves catabolism of the majority of species in immature cardiolipin (Fig. 3A), which then progress into a broad distribution of cardiolipin molecular species (Fig. 3B). For example, a similar broad distribution of cardiolipin molecular species was observed between day 5 and 15 for normal mouse cardiac development (data not shown) as well as in mature mouse brain (16, 26). It is interesting that this broad distribution of cardiolipin molecular species is also observed in various disease states, culture conditions, and transgenic models of enzymes that catalyze cardiolipin remodeling (23, 41–45). This suggests a common remodeling event or role of various acyltransferases, transacylases, or phospholipases involved in this stage of the remodeling process. After 100 iterations of in silico remodeling, the mature myocardial cardiolipin distribution becomes more evident (Fig. 3C) and reaches an equilibrium state after 250 iterations (Fig. 3D). This molecular species distribution corresponds closely to the distribution of cardiolipin molecular species obtained from MDMS-SL in mouse myocardial mitochondria (Fig. 3E, Table 1). Similarities in designated molecular species can also be viewed in the hybrid output of the dynamic simulation for the heart (supplementary materials), which demonstrates the accuracy in identification of the molecular species obtained by MDMS-SL and the dynamic simulation approach. In all, the dynamic simulation output demonstrates that the level of detection of cardiolipin molecular species by current technologies is around the threshold of 0.1% of total cardiolipin from myocardial mitochondrial extracts. The dynamic simulation suggests other extremely low abundance species that likely exist in the cardiolipin distribution but are below the detection limit of MDMS-SL under the experimental conditions (Table 1).
Fig. 3. Dynamic simulation of myocardial cardiolipin molecular species. A: Predicted myocardial immature cardiolipin molecular species were generated representing time point 0. Generated distributions of cardiolipin molecular species were represented at iteration 50 (B), 100 (C), and 250 (D). E: A representative mouse myocardial cardiolipin distribution as determined by MDMS-SL. The asterisks indicate the individual ion peaks of mouse myocardial cardiolipin (15). The mouse myocardial cardiolipin molecular species detected and quantified by MDMS-SL as well as determined by the dynamic simulation were tabulated in Table 1.
### TABLE 1. Mouse myocardial cardiolipin molecular species distribution analyzed by MDMS-SSL and dynamic simulation

| Carbon: Double Bond | Major Species | Mean Mass Content | Mean Distribution of Species (%) | Distribution Generated by Dynamic Simulation (%) |
|--------------------|--------------|------------------|----------------------------------|-----------------------------------------------|
| 68:6               |              |                  |                                  |                                               |
| 68:5               |              |                  |                                  |                                               |
| 68:4               |              |                  |                                  |                                               |
| 70:8               |              |                  |                                  |                                               |
| 70:7               | 18:2:18:2:18:2:16:1 | 710.97 | 0.40 | 0.76 | 0.79 |
| 70:6               | 18:2:18:2:18:1:16:1 | 711.98 | 0.35 | 0.66 | 0.75 |
| 70:5               | 18:2:18:1:18:1:16:0 | 712.99 | 0.20 | 0.39 | 0.24 |
| 70:4               | 18:2:18:1:18:1:16:0 | 713.99 | 0.11 | 0.21 | 0.03 |
| 72:9               | 18:2:18:18:2:8:5 | 722.97 | 0.37 | 0.70 | 0.28 |
| 72:8               | 18:2:18:18:2:8:2 | 723.98 | 11.18 | 21.29 | 21.42 |
| 72:7               | 18:2:18:2:18:1 | 724.99 | 7.11 | 13.54 | 12.65 |
| 72:6               | 18:2:18:1:18:1:18:1 | 725.99 | 2.31 | 4.41 | 3.01 |
| 72:5               | 18:2:18:1:18:1:18:1 | 727.00 | 0.16 | 0.31 | 0.37 |
| 72:4               |              |                  |                                  |                                               |
| 74:12              |              |                  |                                  |                                               |
| 74:11              | 18:2:18:2:16:1:22:6 | 734.97 | 0.25 | 0.49 | 0.51 |
| 74:10              | 18:2:18:2:16:1:22:6 | 735.98 | 0.74 | 1.40 | 1.53 |
| 74:9               | 18:2:18:2:16:1:22:6 | 736.99 | 2.14 | 4.08 | 3.26 |
| 74:8               | 18:1:18:2:18:2:20:3 | 737.99 | 2.20 | 4.19 | 3.75 |
| 74:7               | 18:1:18:2:18:2:20:2 | 739.99 | 1.09 | 2.08 | 1.41 |
| 74:6               | 18:1:18:1:18:2:20:2 | 740.01 | 0.25 | 0.48 | 0.24 |
| 74:5               |              |                  |                                  |                                               |
| 76:14              |              |                  |                                  |                                               |
| 76:12              | 18:2:18:2:16:1:22:6 | 747.98 | 9.69 | 18.45 | 18.10 |
| 76:11              | 18:2:18:1:18:1:22:6 | 748.99 | 4.85 | 9.23 | 9.91 |
| 76:10              | 18:1:18:2:18:2:20:4 | 749.99 | 0.85 | 1.63 | 2.35 |
| 76:9               | 18:1:18:2:18:2:20:4 | 751.00 | 0.14 | 0.27 | 0.53 |
| 76:8               | 18:1:18:1:18:2:20:2 | 752.01 | 0.09 | 0.16 | 0.21 |
| 76:7               |              |                  |                                  |                                               |
| 78:15              | 18:2:18:3:20:4:22:6 | 758.97 | 0.06 | 0.12 | 0.11 |
| 78:14              | 18:2:18:2:18:2:22:6 | 759.98 | 0.27 | 0.51 | 0.77 |
| 78:13              | 18:2:18:2:20:3:22:6 | 760.99 | 1.02 | 1.95 | 1.97 |
| 78:12              | 18:2:18:2:20:3:22:6 | 761.99 | 1.32 | 2.54 | 2.29 |
| 78:11              | 18:2:18:2:20:1:22:6 | 763.00 | 0.35 | 0.66 | 0.80 |
| 78:10              |              |                  |                                  |                                               |
| 78:9               |              |                  |                                  |                                               |
| 80:17              |              |                  |                                  |                                               |
| 80:16              | 18:2:18:2:22:6:22:6 | 771.98 | 3.33 | 6.34 | 5.74 |
| 80:15              | 18:2:18:1:22:6:22:6 | 772.99 | 0.93 | 1.78 | 2.87 |
| 80:14              | 18:2:18:0:22:6:22:6 | 773.99 | 0.30 | 0.58 | 0.65 |
| 80:13              |              |                  |                                  |                                               |
| 80:12              |              |                  |                                  |                                               |
| 80:11              |              |                  |                                  |                                               |
| 82:18              |              |                  |                                  |                                               |
| 82:17              | 18:2:20:3:22:6:22:6 | 784.99 | 0.05 | 0.10 | 0.41 |
| 82:16              | 18:2:20:2:22:6:22:6 | 785.99 | 0.07 | 0.14 | 0.46 |
| 82:15              |              |                  |                                  |                                               |
| 82:14              |              |                  |                                  |                                               |
| 84:20              | 18:2:22:6:22:6:22:6 | 795.98 | 0.28 | 0.54 | 0.81 |
| 84:19              | 18:2:22:6:22:6:22:5 | 796.99 | 0.04 | 0.07 | 0.36 |
| 84:18              |              |                  |                                  |                                               |
| 84:17              |              |                  |                                  |                                               |
| 86:22              |              |                  |                                  |                                               |
| 86:21              |              |                  |                                  |                                               |
| 86:20              |              |                  |                                  |                                               |
| 88:24              |              |                  |                                  |                                               |
| 88:23              |              |                  |                                  |                                               |

Mouse myocardial mitochondrial lipids were extracted by a modified Bligh and Dyer procedure as described in the section of "Materials and Methods." The cardiolipin molecular species in the lipid extracts were identified and quantified as previously described (15). The mass content and distribution of cardiolipin species represent the mean of four separately prepared mitochondria from different animals. The distribution of cardiolipin molecular species determined by the dynamic simulation represents the one frame output 0.1Ex 250 iterations for heart cardiolipin. This data was obtained using the MS output function. Values < 0.01 percent are not displayed but can be found in supplementary tables. CL, cardiolipin.
The equilibrium reached in the simulation is dynamic, in which remodeling maintains the cardiolipin distribution, and acyl chains are proportionally represented in both the mature cardiolipin profile and the available pool of acyl chains. This is likely because the organs generally maintain their cardiolipin pattern after development, although cardiolipin remodeling continues during aging with only subtle changes to the acyl chain composition or distribution (16, 46–48).

Dynamic simulation of hepatic cardiolipin molecular species

Visualization of the distribution of predicted immature cardiolipin molecular species generated by the distribution of hepatic mitochondrial PG molecular species revealed a more diversified profile of species compared with those of myocardial immature cardiolipin species (Fig. 4A). Additionally, because molecular species from immature cardiolipin are minimally represented in the mature cardiolipin, only a minimal decay rate was used (between 0.00001 and 0.00005) for the dynamic simulation representing a fraction of de novo cardiolipin biosynthesis. The selection of these decay rates, however, do not address specific kinetic flux of the enzymes involved in synthesis and are only designed to reflect the relative representation of de novo cardiolipin synthesis in the cardiolipin distribution. The acyl chain distribution of hepatic cardiolipin contained more linoleic acyl chains compared with that of myocardial cardiolipin. The seeded acyl chain distribution used for hepatic cardiolipin dynamic simulation consisted of: 16:0 (0.35%), 16:1 (2.56%), 18:0 (0.12%), 18:1 (10.65%), 18:2 (81.32%), 18:3 (0.05%), 20:1 (0.06%), 20:2 (1.78%), 20:3 (2.07%), 20:4 (0.29%), 22:5 (0.08%), and 22:6 (0.56%), and a \( \Delta t \) value of 0.1 was selected for the rate of remodeling. This acyl chain distribution generated the best-fit equilibrium cardiolipin profile compared with that determined by MDMS-SL. As demonstrated by Fig. 4B, a broad distribution of cardiolipin species is rapidly formed from these determined acyl chains. Due to the abundance of linoleic and to a much lesser extent oleic acyl chains relative to the abundance of the rest of the acyl chains, the peaks at carbon:double bond 72:8 and 72:7 become predominant at 100 iterations, majorly representing the species 18:2-18:2-18:2-18:2 (72:8) and 18:1-18:2-18:2-18:2 (72:7) (Fig. 4C). The simulation achieved an equilibrium around 250 iterations (Fig. 4D). This equilibrium state resembles the exact distribution for hepatic cardiolipin obtained from MDMS-SL analysis of the hepatic mitochondrial lipidome (Fig. 4E). The major and minor molecular species identified by MDMS-SL also match the species generated by the dynamic simulation (Table 2 and supplementary materials). Moreover, the level of detection of hepatic cardiolipin molecular species demonstrated by the dynamic simulation is similar to that of the heart (i.e., between 0.1% and 0.2%). Thus, the simulation strategically facilitates detection of low to very low abundance isomers of cardiolipin species, the levels of which are below the detection limitation of MDMS-SL analysis at its current stage.

Dynamic simulation of brain cardiolipin molecular species

The analysis of the distribution of brain cardiolipin molecular species is more complicated than that of myocardial or hepatic cardiolipin. This is due to the broad distribution, the complex acyl chain composition, and the presence of numerous isomeric molecular species compared with the cardiolipin species present in the other organs. Additionally, immature cardiolipin molecular species formed from PG could account for a portion, small but significant, relative to the mature cardiolipin distribution (Fig. 5). Thus, the residual portion of cardiolipin biosynthesis being represented in the overall pattern would complicate the analysis of predicted species being generated in the overall distribution, depending on the rate and stage of remodeling.

To incorporate the proportion of immature cardiolipin species from de novo cardiolipin biosynthesis into the cardiolipin remodeling simulation, a decay rate of 0.01 was used with an initial acyl chain distribution of 16:0 (3.76%), 16:1 (5.08%), 16:2 (1.2%), 18:0 (0.59%), 18:1 (46.64%), 18:2 (7.25%), 18:3 (2.47%), 20:1 (1.12%), 20:2 (1.71%), 20:3 (2.21%), 20:4 (13.18%), 22:2 (0.23%), 22:3 (0.12%), 22:4 (0.6%), 22:5 (0.3%), and 22:6 (13.46%). This distribution generated the best-fit cardiolipin distribution compared with that obtained by MDMS-SL. A \( \Delta t \) of 0.1 was used in the simulation. The sequential remodeling of brain cardiolipin using these designated acyl chains led to a greater abundance of species that contain longer chains and greater unsaturation (Fig. 5B, C) compared with the distribution of heart and liver cardiolipin. Experimental attempts to simulate brain cardiolipin molecular species with a lower decay rate than 0.01 resulted in less diversity of molecular species and an increase in the percentage distribution of the major molecular species (data not shown) that did not accordingly fit the actual mature cardiolipin distribution determined by MDMS-SL (Fig. 5D, E). Again, the profile generated by the dynamic simulation resembled the percentage distribution determined by MDMS-SL (Table 3) and maintained an approximate threshold of detection around 0.15% of total brain cardiolipin.

Validation of isomeric cardiolipin species determined by the dynamic simulation

To substantiate the accuracy of our dynamic simulation approach, we performed product ion analysis of ions containing very low abundance cardiolipin isomers by using a high mass accuracy/resolution mass spectrometer, which could not be detected by MDMS-SL at its current stage but were determined by our simulation. Owing to the low background noise, high sensitivity, and high mass accuracy/resolution, the presence of very low abundance isomers of an ion can be identified through high mass accuracy/resolution mass spectrometry as demonstrated previously (32).

For example, selection and collision-induced dissociation of the cardiolipin species at \( m/z \) 723.48 present in mouse myocardial cardiolipins demonstrated an abundant fragment at \( m/z \) 279.23 in the fatty acyl fragment.
Dynamic simulation of hepatic cardiolipin molecular species. A: Predicted hepatic immature cardiolipin molecular species were generated representing time point 0. Generated distributions of cardiolipin molecular species were represented at iteration 50 (B), 100 (C), and 250 (D). E: A representative mouse hepatic cardiolipin distribution as determined by MDMS-SL. The asterisks indicate the individual ion peaks of mouse hepatic cardiolipin. The mouse hepatic cardiolipin molecular species detected and quantified by MDMS-SL as well as determined by the dynamic simulation were tabulated in Table 2.

region corresponding to 18:2 carboxylate, along with a fragment at m/z 415.22 corresponding to the deprotonated 18:2 lysophosphatidic acid, and a product ion at m/z 592.36 corresponding to the loss of a 18:2 ketene (Fig. 6A). These fragment ions identified the ion as a tetra-18:2 cardiolipin. In addition, we also observed several very low abundance fragments at m/z 253.21, 255.23, 277.21, 281.25, 303.23, and 305.25, corresponding to 16:1, 16:0, 18:3, 18:1, 20:4, and 20:3 carboxylates, respectively (Fig. 6A, inset a). This observation indicates the
presence of other very low abundance cardiolipin species corresponding to the ion at m/z 723.48 including 16:1-18:2-18:2-20:3, 16:0-18:2-18:2-20:4, and 18:1-18:2-18:2-18:3 cardiolipin isomers. This identification using product ion analysis is consistent with the results from the dynamic simulation (Fig. 6A, inset b).

By using this approach, we also validated the presence of very low abundance cardiolipin isomers as predicted by dynamic simulation residing at much lower abundance ions compared with the dominant ion at m/z 723.48. For example, product ion analysis of the cardiolipin ion at m/z 736.51 demonstrated abundant fatty acyl fragments at m/z 279.23 and 305.25 corresponding to 18:2 and 20:3 carboxylates in an approximate peak intensity ratio of 3:1 (Fig. 6B), identifying the presence of 18:2-18:2-18:2-20:3 cardiolipin. Expansion of this acyl fragment region shows the presence of multiple very low abundance fatty acyl carboxylates, including 16:0 (m/z 255.23), 18:1 (m/z 281.25), 20:4 (m/z 303.23), 22:6 (m/z 327.23), and 22:5 (m/z 329.25) (Fig. 6B, inset c). The presence of these acyl chains validated the presence of 18:1-18:2-18:1-18:6, and 16:0-18:1-18:1-18:6 cardiolipin isomers at m/z 736.51 as determined by the dynamic simulation results that indicate the value of this approach for the predictive analysis of very low abundance cardiolipin isomeric species that may not be readily detectable by the current stage of MS.

| Carbon: Double Bond | Major Species                   | [CL-2H]^− | Mean Mass Content | Mean Distribution of Species (%) | Distribution Generated by Dynamic Simulation (%) |
|---------------------|--------------------------------|-----------|------------------|----------------------------------|-----------------------------------------------|
| 68:6                |                                |           |                  |                                  |                                               |
| 68:5                |                                |           |                  |                                  |                                               |
| 68:4                |                                |           |                  |                                  |                                               |
| 70:8                |                                |           |                  |                                  |                                               |
| 70:7                | 18:2-18:2-18:2-16:1            | 710.97    | 1.49             | 5.99                            | 5.52                                          |
| 70:6                | 18:2-18:2-18:1-16:1            | 711.98    | 0.65             | 2.60                            | 3.04                                          |
| 70:5                | 18:2-18:1-18:1-16:1            | 712.99    | 0.15             | 0.62                            | 0.66                                          |
| 70:4                | 18:2-18:1-18:1-16:0            | 713.99    | 0.19             | 0.78                            | 0.07                                          |
| 72:9                |                                |           |                  |                                  |                                               |
| 72:8                | 18:2-18:2-18:2-18:2            | 723.98    | 11.26            | 45.30                           | 44.16                                         |
| 72:7                | 18:2-18:2-18:2-18:1            | 724.99    | 5.57             | 22.39                           | 23.48                                         |
| 72:6                | 18:2-18:2-18:1-18:1            | 725.99    | 1.41             | 5.68                            | 4.99                                          |
| 72:5                | 18:2-18:2-18:1-18:0            | 727.00    | 0.06             | 0.26                            | 0.54                                          |
| 72:4                |                                |           |                  |                                  |                                               |
| 74:11               | 18:2-18:2-16:1-18:2-20:4       | 734.97    | 0.03             | 0.10                            | 0.12                                          |
| 74:10               | 18:2-18:2-18:2-20:4            | 735.98    | 0.22             | 0.88                            | 0.71                                          |
| 74:9                | 18:2-18:2-18:2-20:3            | 736.99    | 1.34             | 5.39                            | 4.74                                          |
| 74:8                | 18:2-18:2-18:2-20:2            | 737.99    | 1.42             | 5.70                            | 5.66                                          |
| 74:7                | 18:1-18:2-18:2-20:2            | 739.00    | 0.47             | 1.87                            | 1.91                                          |
| 74:6                | 18:1-18:2-18:2-20:1            | 740.01    | 0.11             | 0.43                            | 0.29                                          |
| 74:5                |                                |           |                  |                                  |                                               |
| 76:12               | 18:2-18:2-18:2-22:6            | 747.98    | 0.32             | 1.29                            | 1.25                                          |
| 76:11               | 18:2-18:2-18:2-22:5            | 748.99    | 0.17             | 0.70                            | 0.73                                          |
| 76:10               |                                |           |                  |                                  |                                               |
| 76:9                |                                |           |                  |                                  |                                               |
| 76:8                |                                |           |                  |                                  |                                               |
| 76:7                |                                |           |                  |                                  |                                               |
| 78:14               |                                |           |                  |                                  |                                               |
| 78:13               |                                |           |                  |                                  |                                               |
| 78:12               |                                |           |                  |                                  |                                               |
| 78:11               |                                |           |                  |                                  |                                               |
| 78:10               |                                |           |                  |                                  |                                               |
| 80:16               |                                |           |                  |                                  |                                               |

Mouse hepatic mitochondrial lipids were extracted by a modified Bligh and Dyer procedure as described in the section of “Materials and Methods.” The cardiolipin molecular species in the lipid extracts were identified and quantified as previously described (15). The mass content and distribution of cardiolipin species represent the mean of four separately prepared mitochondria from different animals. The distribution of cardiolipin molecular species determined by the dynamic simulation represents the one frame output 0.1a 250 iterations for liver cardiolipin. This data was obtained using the MS output function. Values < 0.01 percent are not displayed but can be found in supplementary tables. CL, cardiolipin.
Dynamic simulation of cardiolipin remodeling is commonly believed that acyl CoA as well as PC and PE contribute to the pool of acyl chains used for cardiolipin remodeling in mitochondria (20, 21, 23, 24, 38, 43). In addition, Lands cycle remodeling utilizes the sn-2 positions of glycerophospholipids for remodeling (49, 50).

**Prediction of acyltransferase and transacylase activities as well as acyl selectivities in cardiolipin remodeling from different organs**

Analysis of the mitochondrial lipidome from the heart, liver, and brain allowed for the opportunity to characterize possible substrates used for cardiolipin remodeling. It is commonly believed that acyl CoA as well as PC and PE contribute to the pool of acyl chains used for cardiolipin remodeling in mitochondria (20, 21, 23, 24, 38, 43). In addition, Lands cycle remodeling utilizes the sn-2 positions of glycerophospholipids for remodeling (49, 50). MDMS-SL enables us to identify and quantify acyl chains.
| Carbon: Double Bond | Major Species | [CL-2H]^2 | Mean Mass Content | Mean Distribution of Species (%) | Distribution Generated by Dynamic Simulation (%) |
|-------------------|--------------|-----------|------------------|---------------------------------|-----------------------------------------------|
| 66:6              | 0.01         |           |                  |                                 |                                               |
| 66:5              | 0.03         |           |                  |                                 |                                               |
| 66:4              | 0.07         |           |                  |                                 |                                               |
| 66:3              | 0.10         |           |                  |                                 |                                               |
| 66:2              | 0.08         |           |                  |                                 |                                               |
| 66:1              | 0.03         |           |                  |                                 |                                               |
| 68:8              | 0.01         |           |                  |                                 |                                               |
| 68:7              | 0.05         |           |                  |                                 |                                               |
| 68:6              | 0.16         |           |                  |                                 |                                               |
| 68:5              | 0.41         |           |                  |                                 |                                               |
| 68:4              | 0.41         |           |                  |                                 |                                               |
| 68:3              | 0.41         |           |                  |                                 |                                               |
| 68:2              | 0.41         |           |                  |                                 |                                               |
| 70:10             | 0.01         |           |                  |                                 |                                               |
| 70:9              | 0.06         |           |                  |                                 |                                               |
| 70:8              | 0.23         |           |                  |                                 |                                               |
| 70:7              | 0.63         |           |                  |                                 |                                               |
| 70:6              | 1.28         |           |                  |                                 |                                               |
| 70:5              | 2.20         |           |                  |                                 |                                               |
| 70:4              | 3.10         |           |                  |                                 |                                               |
| 70:3              | 2.10         |           |                  |                                 |                                               |
| 70:2              | 0.07         |           |                  |                                 |                                               |
| 70:1              | 0.06         |           |                  |                                 |                                               |
| 72:10             | 0.29         |           |                  |                                 |                                               |
| 72:9              | 0.85         |           |                  |                                 |                                               |
| 72:8              | 1.88         |           |                  |                                 |                                               |
| 72:7              | 3.24         |           |                  |                                 |                                               |
| 72:6              | 3.85         |           |                  |                                 |                                               |
| 72:5              | 3.55         |           |                  |                                 |                                               |
| 72:4              | 5.17         |           |                  |                                 |                                               |
| 72:3              | 0.18         |           |                  |                                 |                                               |
| 74:13             | 0.04         |           |                  |                                 |                                               |
| 74:12             | 0.23         |           |                  |                                 |                                               |
| 74:11             | 0.80         |           |                  |                                 |                                               |
| 74:10             | 2.17         |           |                  |                                 |                                               |
| 74:9              | 4.33         |           |                  |                                 |                                               |
| 74:8              | 4.68         |           |                  |                                 |                                               |
| 74:7              | 6.34         |           |                  |                                 |                                               |
| 74:6              | 1.55         |           |                  |                                 |                                               |
| 74:5              | 0.98         |           |                  |                                 |                                               |
| 74:4              | 0.49         |           |                  |                                 |                                               |
| 74:3              | 0.01         |           |                  |                                 |                                               |
| 76:15             | 0.03         |           |                  |                                 |                                               |
| 76:14             | 0.16         |           |                  |                                 |                                               |
| 76:13             | 0.65         |           |                  |                                 |                                               |
| 76:12             | 1.93         |           |                  |                                 |                                               |
| 76:11             | 3.20         |           |                  |                                 |                                               |
| 76:10             | 5.50         |           |                  |                                 |                                               |
| 76:9              | 6.81         |           |                  |                                 |                                               |
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Mouse brain mitochondrial lipids were extracted by a modified Bligh and Dyer procedure as described in the section of "Materials and Methods." The cardiolipin molecular species in the lipid extracts were identified and quantified as previously described (15). The mass content and distribution of cardiolipin species represent the mean of four separately prepared mitochondria from different animals. The distribution of cardiolipin molecular species determined by the dynamic simulation represents the one frame output 0.1 Ha to 250 iterations for brain cardiolipin. This data was obtained using the MS output function. Values < 0.01 percent are not displayed but can be found in supplementary tables. CL, cardiolipin.

| Carbon: Double Bond | Major Species | [CL-2H]^2 | Mean Content | Mean Distribution of Species (%) | Distribution Generated by Dynamic Simulation (%) |
|---------------------|---------------|------------|--------------|-------------------------------|---------------------------------------------|
| 76:8                | 22:6:18:1:18:0 | 752.01     | 0.68         | 0.98                          | 1.20                                       |
|                     | 20:4:20:2:18:1 | 753.02     | 0.25         | 0.37                          | 0.83                                       |
|                     | 20:4:20:1:18:0 | 754.03     | 0.33         | 0.47                          | 0.22                                       |
| 76:7                | 20:4:20:1:18:1 | 755.03     | 0.29         | 0.43                          | 0.15                                       |
| 76:6                | 20:4:20:1:18:0 | 755.03     | 0.29         | 0.43                          | 0.15                                       |
| 76:5                | 20:4:20:1:18:1 | 755.03     | 0.29         | 0.43                          | 0.15                                       |
| 76:4                | 20:4:20:1:18:0 | 755.03     | 0.29         | 0.43                          | 0.15                                       |
| 78:16               | 22:6:20:4:20:4:18:1 | 770.97 | 0.41         | 0.60                          | 0.26                                       |
| 78:15               | 22:6:20:4:20:4:18:1 | 771.98 | 0.64         | 0.93                          | 0.70                                       |
| 78:14               | 22:6:20:4:20:4:18:1 | 772.99 | 2.02         | 2.92                          | 2.16                                       |
| 78:13               | 22:6:20:4:20:4:18:1 | 773.99 | 2.83         | 4.10                          | 2.96                                       |
| 78:12               | 18:0:20:5:22:5:22:5 | 775.00 | 0.60         | 0.87                          | 0.70                                       |
| 78:11               | 18:0:20:5:22:5:22:5 | 776.01 | 0.35         | 0.51                          | 0.58                                       |
| 78:10               | 18:0:20:5:22:5:22:5 | 777.02 | 0.22         | 0.32                          | 0.19                                       |
| 78:9                | 18:0:20:5:22:5:22:5 | 778.02 | 0.41         | 0.59                          | 0.60                                       |
| 78:8                | 18:0:20:5:22:5:22:5 | 779.02 | 0.60         | 0.87                          | 0.70                                       |
| 80:17               | 22:6:20:4:20:4:18:1 | 770.97 | 0.41         | 0.60                          | 0.26                                       |
| 80:16               | 22:6:20:4:20:4:18:1 | 771.98 | 0.64         | 0.93                          | 0.70                                       |
| 80:15               | 22:6:20:4:20:4:18:1 | 772.99 | 2.02         | 2.92                          | 2.16                                       |
| 80:14               | 22:6:20:4:20:4:18:1 | 773.99 | 2.83         | 4.10                          | 2.96                                       |
| 80:13               | 18:0:20:5:22:5:22:5 | 775.00 | 0.60         | 0.87                          | 0.70                                       |
| 80:12               | 18:0:20:5:22:5:22:5 | 776.01 | 0.35         | 0.51                          | 0.58                                       |
| 80:11               | 18:0:20:5:22:5:22:5 | 777.02 | 0.22         | 0.32                          | 0.19                                       |
| 80:10               | 18:0:20:5:22:5:22:5 | 778.02 | 0.41         | 0.59                          | 0.60                                       |

TABLE 3. Continued.
To rebuild the acyl chain distribution of cardiolipin, the overall distribution of acyl chains from the PC and PE sn-2 positions was calculated assuming no acyl selectivity originating from a transacylation reaction. This approach was followed determining the contribution of acyl CoA fatty acids with designated acyl selectivity for particular fatty acids. There are multiple reasons for taking this approach. First, we were attempting to determine if there was common acyl selectivity for acyl CoA between organs with different molecular species at the sn-1 and sn-2 positions, allowing us to infer Lands cycle remodeling based on acyl chain identification. Determination of the original sources (i.e., acyl CoA, PC sn-2, and PE sn-2 acyl chains, and PG molecular species) of the acyl pool of cardiolipin would facilitate the assessment of the enzymatic activities involved in cardiolipin biosynthesis and remodeling. In addition, modeling the molecular mechanism in which acyl chains are transferred into the sn-1, sn-1’ or sn-2, sn-2’ positions of cardiolipin either equally or with designated selection identifies the potential components reorganizing the remodeling machinery. Furthermore, through rebuilding the acyl chain distribution that serves as the acyl pool in the remodeling process by proper selection of acyl chains from different sources, the dynamic simulation allows us to assess these activities. Specifically, the decay rate of the synthesized immature cardiolipin species calculates the de novo synthesis activity, the amounts of acyl chains contributed from PC and PE sn-2 fatty acyl pools indicate the transacylase activities, the amounts of acyl CoA used in the remodeling determine the acyltransferase activities, and the differential utilization of the different acyl chains from the PC, PE, or acyl CoA pool represents the selectivities of these enzymatic activities to the individual pools. It should be emphasized that there exist multiple acyltransferases and transacylases that have demonstrated cardiolipin remodeling activity, and indeed a variety of isoforms have been detected of these enzymes (20, 38, 51–53). The current dynamic simulation approach is unable to differentiate these isoform activities yet proposes the relative selectivity of the combined action of various acyltransferases or transacylases. Additionally, we do not infer any selection toward remodeling specific de novo synthesized molecular species; instead, we infer that all immature cardiolipin species are remodeled equally and that partially and mature remodeled cardiolipin molecular species are treated with equal affinity.

Fig. 6. Validation of very low abundance cardiolipin isomeric species by product ion analysis using an Orbitrap-LTQ mass spectrometer. Product ion mass spectra of representative myocardial cardiolipin species were acquired using a high mass accuracy/resolution instrument as described under “Materials and Methods.” A: A product ion mass spectrum of cardiolipin species at m/z 723.48 demonstrates the presence of several very low abundance fragments (inset a) corresponding to fatty acyl carboxylates in addition to the abundant 18:2 carboxylate (m/z 279.25). B: A product ion mass spectrum of cardiolipin species at m/z 736.51 demonstrates the presence of several very low abundance fragments (inset c) corresponding to fatty acyl carboxylates in addition to the abundant 18:2 and 20:3 carboxylates (m/z 279.23 and 305.25, respectively). Insets b and d list the cardiolipin isomeric species and their composition determined by the dynamic simulation.
different acyl CoA compositions as well as cardiolipin distributions. If similarities existed in acyl selectivity, then it would suggest a common enzyme or isoform of an enzyme likely involved in the remodeling process among tissues. Indeed, it is not unlikely that numerous acyltransferases exist and that the relative selectivity reflects the coordinated activities of these acyltransferases; however, inferring common selectivity between tissues will allow for the identification of the function of acyltransferase(s) in various tissues. Second, PC and PE are mainly comprised of 20:4 and 22:6 acyl chains at the sn-2 positions as demonstrated by the abundant species of 16:0-20:4, 16:0-22:6, and 18:0-22:6 PC as well as 16:0-20:4, 16:0-22:6, 18:0-20:4, and 18:0-22:6 PE (supplementary Tables III and IV).

Fig. 7. The contribution and selectivity of acyl chains for cardiolipin remodeling from different fatty acyl donor pools determined by dynamic simulation. Based on the acyl chains used to generate mature myocardial, hepatic, and brain cardiolipin, inference was used to determine the possible contribution from acyltransferase, transacylase, or cardiolipin biosynthesis activities. The possible proportion of acyl chains from PG, PC sn-2 positions, PE sn-2 positions, and acyl CoA as determined by MDMS-SL are represented in the chart that would match the acyl chain distribution used in the dynamic simulation. Selectivity of specific acyl chains for acyltransferase dependence (acyl CoA) (A) or transacylase dependence (PC or PE sn-2 acyl chains) (B) was inferred in the heart, liver, and brain. This inference approach allows the comparison of different acyl selectivities for both acyltransferase and transacylase activities.
Cardiolipin is intricately involved in maintaining bioenergetic efficiency and homeostasis of cellular metabolism (11, 13). The complex biophysical properties as well as the complex heterogeneity of the cellular lipidome.

Recent advances in lipidomic analysis now allow for the accurate quantification of hundreds to thousands of lipid molecular species. However, development of bioinformatics and algorithmic approaches to integrate lipidomic data into a systems biology interpretation will be required to transition lipidomics from a descriptive to a mechanistic paradigm. In this study, we utilized the MDMS-SL platform to structurally identify and quantify the lipid molecular species. However, development of bioinformatic analysis as well as extend the range and detection of lipidomic analysis. This composite experimental and bioinformatic approach represents a major advance in interpreting the complex heterogeneity of the cellular lipidome.

**DISCUSSION**

Alternatively, we attempted to fit the acyl chain pool used for cardiolipin remodeling with a proportion of acyl chains from sn-2 positions of PC and PE as well as acyl CoA inferring some acyl chain selectivity. This approach allowed us to increase the percentage contribution of selected acyl chains as well as cardiolipin species from predicted immature cardiolipin if these species were visualized in the mature cardiolipin profile. The distribution of acyl chains used for the myocardial cardiolipin simulation could be formed by contributing 4% and 3% each from PC and PE sn-2 positions, respectively, without acyl selectivity. The remaining 93% of the acyl chains would be donated from acyl CoA with the designated acyl specificity (Fig. 7A). This specificity was most evident in 18:2, 20:3, and 22:6 acyl chains and less selective for 18:1, 20:2, and 22:5 acyl chains.

Due to the lower amount of 20:4 and 22:6 containing acyl chains in hepatic cardiolipin, the level of transacylation from PC and PE appeared to be lower in the hepatic than in the myocardial cardiolipin. This was represented by 1% each for both glycerophospholipid classes without acyl selectivity. The remaining 98% of the acyl chains were contributed by acyl CoA with selectivity assigned to 18:2 and 20:3. Less selectivity was designated to 16:1, 18:1, 20:2, and 20:3 (Fig. 7A).

The composition of acyl chains in brain cardiolipin was more complex than that found in myocardial and hepatic cardiolipin. The most abundant acyl chains in brain cardiolipin are 18:1, 20:4, and 22:6, with lesser amounts of 18:2. These aforementioned acyl chains are abundant in the sn-2 position of PC and PE. This suggests that transacylation activity is greater for brain cardiolipin than for myocardial or hepatic cardiolipin. In addition, the proportion of saturated acyl chains was greater in brain cardiolipin than hepatic and myocardial cardiolipin, suggesting that immature cardiolipin contributes significantly to the species composition of mature cardiolipin in brain than it does for hepatic and myocardial cardiolipin. Based on these criteria, it was determined that 10% of the acyl chains of brain cardiolipin molecular species likely originated from predicted immature cardiolipin, 22% and 14% of the acyl chains nonselectively originated from the sn-2 position of PC and PE, respectively, and the remaining 64% of the acyl chains originated from acyl CoA with acyl selectivity. The acyl chains that displayed the greatest selectivity were 18:2, 22:6, 16:1, and to a lesser extent 22:5 and 18:1 (Fig. 7A).

Analysis of the designated acyl selectivity in diverse acyl CoA distributions revealed an unexpected finding when comparing all three organs (Fig. 7). Aligning the determined acyl selectivities for acyl CoA revealed the possibility that there are consistent selectivities among tissues as well as some differences. The relative proportional selectivity for 18:1 CoA and 18:2 CoA appeared to be consistent among all tissues. This was even more surprising for brain cardiolipin, which contains more 18:1 CoA and less 18:2 CoA content than that in other tissues. The proportional selectivity for acyl CoA in 22:5 and 22:6 acyl chains was evident in myocardial and brain cardiolipin, but not hepatic cardiolipin. In contrast, the proportional selectivity for 20:2 and 20:3 was only found in myocardial and hepatic cardiolipin and not brain cardiolipin. Additionally, there appeared to be selectivity for 16:1 acyl chains in acyl CoA for hepatic cardiolipin and to even a greater extent for brain cardiolipin; however, no selectivity for 16:1 could be concluded for myocardial cardiolipin. The similarities and differences in acyl selectivity between tissues may result from tissue-specific splicing variants or post-translation modifications of various known acyltransferases (20, 38, 41, 45, 52, 54). Recently, the acyltransferase MLCL AT demonstrated a relative selectivity of 18:2 > 18:1 > 16:1 (40), which is consistent with the acyl CoA selectivity in the examined heart and liver tissue and also in the examined brain tissue except for 16:1 CoA selectivity.

We also modeled the acyl selectivity of transacylase activity (Fig. 7B). This approach demonstrated variability in the selectivity for 16:1, 18:1, 18:2, 20:2, and 20:3 acyl chains between different organs with no distinct pattern. Because it is likely that cardiolipin remodeling might be relegated by the coordinated activities of both acyltransferases and transacylases, the acyl selectivity from both activities might generate the acyl chain pool used for cardiolipin remodeling. In order to not infer any intrinsic bias in this analysis, we attempted to highlight the relative selectivities of acyl chains in solely acyltransferase or transacylase dependence in the current study. In all likelihood, the combination of the two mechanisms occur in synchrony to facilitate the highly specific acyl chain distributions present in different tissues. Additional experimentation might also reveal the relative proportion of acyltransferase versus transacylase activities contributing to the various cardiolipin distributions.
diverse molecular species of cardiolipin have simulated the investigation of numerous analytical and mathematical approaches to determine the importance of cardiolipin in biological function. Coarse-grained molecular dynamics, thermodynamics and thermal response, atomic force microscopy, and atomic-scale simulations have all been used to characterize structural orientation and interaction of cardiolipin in membrane dynamics (55–60). More recently, mathematical models using linear algebraic representation have been used to explain the acyl selectivity of cardiolipin remodeling based on free energies (61). Although that approach demonstrated novel insight into potential mechanisms governing acyl selectivity, it did not directly address the importance of kinetics in the generation of cardiolipin acyl chain distributions. In addition, that approach only addressed distributions that are relegated by tetralinoleic abundance of cardiolipin; however, brain cardiolipin does not conform to that model (16, 26). Moreover, alterations in free energies are complex functions of enthalpy and entropy driven by altering molecular membrane symmetry and the order/disorder at the membrane interface.

A critical point addressed by the cardiolipin dynamic simulation in the current study was imposing the option for sn-1 or sn-2 selectivity in cardiolipin acyl chain remodeling. This option allowed for the inference of the rate as well as pattern distribution of cardiolipin molecular species if acyl chain selectivity was employed or if all four chains were remodeled equally to reach the equilibrium distribution. Our findings indicate that treating all four chains equally for remodeling generated a cardiolipin distribution most representative of that obtained by MDMS-SL. This assumption supports the findings of Malhotra et al. (62), who demonstrated that sn-1 and sn-2 positions of cardiolipin are remodeled equally using purified recombinant Drosophila tafazzin.

The use of a dynamic simulation approach for identifying and modeling the biological parameters regulating cardiolipin synthesis and remodeling represents an important advance in integrating lipidomic data with the biochemical mechanisms of cardiolipin remodeling. This approach exploits differences in cardiolipin composition and metabolism between tissues (63) to gain mechanistic insight into cardiolipin remodeling pathways and kinetics. Similar approaches using algorithmic inference and in silico kinetic modeling to describe the enzymatic parameters governing the molecular species distribution found in other lipid classes are also possible through this approach. The integration of a flux-based approach for lipidomics has been utilized in describing eicosanoid metabolism; however, the number of parameters used to ascertain the flux of biological pathways can result in overfitting the experimental data, depending on the model (64).

The dynamic simulation provides a formidable tool for analyzing the complex organization of cardiolipin molecular species as controlled by biosynthesis and remodeling. The strategy that we presented allows for the inference of acyltransferase and transacylase activities as well as acyl chain selectivities that would be needed to generate the acyl chains used during the remodeling process. Furthermore, it provides precise predictions as to the relative composition of isomeric molecular species, which can clarify the identification of low abundance molecular species of cardiolipin, which may not be detectable by current analytical technologies. This approach offers a systematic method of analyzing similarities between cardiolipin remodeling among different tissues, a key step toward dissecting the functions of the enzymes involved in lipid regulation.

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