Inhibitory effect of dietary lipids on chaperone-mediated autophagy

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Cytosolic proteins can be selectively delivered to lysosomes for degradation through a type of autophagy known as chaperone-mediated autophagy (CMA). CMA contributes to intracellular quality control and to the cellular response to stress. Compromised CMA has been described in aging and in different age-related disorders. CMA substrates cross the lysosomal membrane through a translocation complex; consequently, changes in the properties of the lysosomal membrane should have a marked impact on CMA activity. In this work, we have analyzed the impact that dietary intake of lipids has on CMA activity. We have found that chronic exposure to a high-fat diet or acute exposure to a cholesterol-enriched diet both have an inhibitory effect on CMA. Lysosomes from livers of lipid-challenged mice had a marked decrease in the levels of the CMA receptor, the lysosome-associated membrane protein type 2A, because of loss of its stability at the lysosomal membrane. This accelerated degradation of lysosome-associated membrane protein type 2A, also described as the mechanism that determines the enhanced instability of LAMP-2A at the lysosomal membrane of the lipid-challenged animals that resemble those observed with age. Our findings identify a previously unknown negative impact of high dietary lipid intake on CMA and underscore the importance of diet composition on CMA malfunction in aging.

cathepsins | lipid load | lyso-bis phosphatidic acid | membrane microdomains | membrane proteins

Autophagy is the process that mediates degradation of intracellular components in lysosomes (1). Different autophagic pathways have been described to coexist in most mammalian cells, but they differ in the mechanisms involved in the delivery of cargo to the lysosomal compartment (2, 3). This study focuses on chaperone-mediated autophagy (CMA), a type of lysosomal degradation for a selective pool of cytosolic proteins all bearing a targeting motif (4, 5). Once this motif is recognized by the cytosolic chaperone heat shock cognate protein 70 (hsc70), it is delivered to the surface of the lysosomal membrane (6, 7), where it binds to the lysosome-associated membrane protein type 2A (LAMP-2A) (8). Binding of the substrate to the cytosolic tail of LAMP-2A promotes the assembly of this protein into a multimeric protein complex that mediates the translocation of the substrate protein across the lysosomal membrane (9). Translocation is attained on substrate unfolding, and it requires the assistance of a variant of hsc70 (lys-hsc70) that resides in the lysosomal lumen (10).

Basal levels of CMA activity are detectable in almost all cells (11) and contribute to the maintenance of cellular homeostasis, as well as to specialized functions depending on the cell type and substrate degraded. For example, CMA has been shown to participate in antigen presentation (12), regulation of cellular growth (13), modulation of neuronal survival (11), and control of specific transcriptional programs in response to nutritional challenges (14). CMA is maximally activated as part of the cellular response to different stressors, such as prolonged starvation (15), oxidative stress (16) and exposure to agents that lead to protein damage (17). In fact, compromised CMA in cultured cells renders cells more susceptible to stressors, such as oxidants, prooxidants, and UV light (18). CMA activity declines with age, and it is compromised in different age-related disorders, such as neurodegenerative diseases, metabolic disorders, and nephropathies, as well as in some lysosomal storage disorders (19).

Levels of LAMP-2A at the lysosomal membrane have been shown to be limiting for CMA activity because they directly determine rates of substrate binding and translocation through this pathway (20). Multiple nonexclusive mechanisms regulate LAMP-2A levels at the lysosomal membrane, including de novo synthesis of the protein, mobilization of a luminal pool of LAMP-2A to the lysosomal membrane, and its regulated degradation in this compartment (20–22). Mobilization of LAMP-2A to specific membrane microdomains of particular lipid composition (enriched in cholesterol and glycosphingolipids) initiates its sequential cleavage by a yet to be identified metalloprotease and cathepsin A, a luminal protease that dynamically associates with these microdomains (23). CMA regulation may depend on the dynamic interaction of LAMP-2A with the lysosomal membrane microdomains. Under conditions of low CMA activity, part of the membrane pool of LAMP-2A is mobilized to these regions for degradation, whereas when maximal activation of CMA occurs, LAMP-2A is excluded from these regions because its assembly into the multimeric translocation complex only occurs outside the microdomains (22). Alterations in the recruitment of LAMP-2A to these regions of selective cleavage have been identified as the main reason for the pronounced decrease in LAMP-2A levels in the lysosomal membrane of old organisms and the subsequent decline in the activity of this pathway (24). However, the exact mechanism that determines the enhanced instability of LAMP-2A at the lysosomal membrane with age remains unknown.

We have recently identified an inhibitory effect of different lipid challenges on another autophagic pathway, namely, macroautophagy, and we have narrowed the defect to changes in the lipid composition, particularly to the cholesterol content of the
membranes of the vesicular compartments involved in that process (25). Given that lateral mobility of proteins at the lysosomal membrane is essential for CMA, we hypothesized that modifications of the lysosomal membrane as a result of changes in the cellular availability of lipids could have a marked impact on CMA activity. In this work, we have analyzed the effect of different lipid challenges on CMA, both in cultured cells and in vivo by subjecting mice to defined lipid content diets. Our results have revealed reduced CMA activity in cells exposed to different lipid challenges because of accelerated degradation of the CMA receptor under these conditions in lysosomes. Qualitative and quantitative changes in the lipid composition of the membranes of lysosomes from animals exposed to dietary lipid challenges resemble those observed in old animals and favor higher mobilization of LAMP-2A toward the lipid membrane microdomains, where its degradation occurs. This negative effect of dietary lipids on the stability of LAMP-2A at the lysosomal membrane could be one of the main reasons for reduced CMA activity in aging.

Results

Different Lipid Challenges Exert an Inhibitory Effect on CMA. We first analyzed the effect on CMA of increasing concentrations of oleate in mouse fibroblasts in culture using a recently developed photoactivatable CMA reporter (KFERQ-PA-mCherry) (11). Photactivation converts the reporter protein present at that time in the cell to fluoresce in red, but any reporter synthesized de novo after phot activation will not fluoresce. This strategy allows tracking changes with time in the fluorescence pattern and degradation of the reporter protein independent of possible changes in synthesis. As CMA is activated, the reporter is delivered to lysosomes, resulting in a gradual change from a cytosolic diffuse pattern to a lysosomal punctate pattern. Comparison of the number of fluorescent puncta per cell provides a good estimate of the amount of substrate bound to the lysosomal membrane at any given time, which we have found to correlate well with CMA activity in cultured cells (11). As shown in Fig. 1A, exposure to increasing concentrations of oleate sufficient to induce intracellular accumulation of lipids in the form of lipid droplets [visualized by staining with BODIPY 493/503; Molecular Probes (Invitrogen)] had an inhibitory effect on the levels of CMA reporter bound to lysosomes in the treated cells. We confirmed that treatment with oleic acid also resulted in reduced lysosomal uptake of the reporter by blocking lysosomal proteolysis with leupeptin and using an antibody against the reporter (this is necessary because once internalized, the CMA reporter is no longer fluorescent as a result of its unfolding) (Fig. S1A). A similar inhibitory effect on CMA was observed when cells were exposed to palmitic acid, a saturated fatty acid (Fig. S1B), although in this case, only lower concentrations could be tested because of the higher toxicity of this lipid. Treatment with the inhibitor of desmosterol Δ24-reductase,
U18666A, which has been shown to alter cholesterol trafficking by preventing its exit from late endosomes and lysosomes (Fig. S2), had a similar inhibitory effect on basal and even inducible CMA (activated by prolonged serum removal) (Fig. 1B). In contrast, we did not observe any significant effect on CMA activity on treatment with the inhibitor of glycophospholipid synthesis, N-butyl-deoxygalactonojirimycin (Fig. 1B), which also leads to accumulation of intracellular lipids, mainly in the form of lipid droplets (Fig. S2B), without a significant effect on lysosomal/endosomal lipid content (Fig. S2C). These results support that intracellular buildup of lipids may not be enough to interfere with CMA activity but that direct changes in the cholesterol content at the lysosomal compartment may instead be responsible for the observed compromise in CMA.

To investigate the effect of lipid challenges on CMA further and to address the physiological relevance of this effect, we moved to an in vivo model and analyzed the consequences of changes in dietary lipid intake in mice on the rates of hepatic CMA activity. To this purpose, we analyzed the effect of acute (3-wk) exposure to a diet enriched in cholesterol (2% CHOL) and of chronic (16-wk) exposure to a lipid challenge in animals fed a high-fat diet (HFD; 60% calories from fat and ~5% calories from cholesterol), and compared these animals with control groups maintained on a regular diet (RD). We first analyzed the ability of intact lysosomes isolated from livers of these two groups of animals to degrade a radiolabeled pool of cytosolic proteins enriched in CMA substrates. We validated that purity of the fractions was comparable in the three groups of animals (determined as hexosaminidase enrichments of 28.1 ± 1.5, 21.4 ± 4.5, and 23.7 ± 1.9 in the RD, CHOL, and HFD, respectively) and that diets did not reduce the stability of the lysosomal membrane (determined as the percentage of hexosaminidase released into the media of 8.1 ± 1.2, 7.8 ± 1.4, and 4.1 ± 1.1 in the RD, CHOL, and HFD, respectively). Incubation of intact lysosomes with radiolabeled proteins recapitulates the three main steps of CMA: binding, uptake, and degradation once in the lysosomal lumen (26). Degradation of the cytosolic proteins was significantly lower in lysosomes isolated from animals exposed to either the CHOL or HFD (Fig. 1C). These observed changes in CMA seem to be primary at the level of binding/uptake rather than degradation, because we did not find differences in the efficiencies of intracellular degradation when experiments were performed with lysosomes in which the membranes have been disrupted to allow direct access of the lysosomal proteases to the substrates (Fig. S3). Interestingly, and confirming that the effect of the lipid diets was primarily on CMA, this treatment did not have an effect on the degradation of proteins by a subpopulation of lysosomes unable to perform CMA because they lack the luminal chaperone required for substrate translocation (Fig. S3B).

In agreement with reduced CMA activity in the lysosomes from the treated groups, levels of endogenous cytosolic proteins previously identified as CMA substrates, such as GAPDH (27), were higher in the cytosolic fraction (Fig. S4A) and markedly lower inside lysosomes (Fig. S4B) from CHOL- and HFD-treated animals compared with those maintained on the RD. Despite the higher cytosolic content of GAPDH, the specific activity of this enzyme was lower in the CHOL and HFD groups (Fig. S4C), supporting a possible gradual loss of function associated with the lower turnover of the enzyme under these conditions. To address the effect of the diets on substrate binding and uptake via CMA directly, we used a second well-established in vitro assay with isolated lysosomes incubated with specific CMA substrate proteins that allows us to analyze these two steps separately independent of proteolysis (26). When substrates are incubated with isolated intact lysosomes, the substrate translocated into the lumen is rapidly degraded and only that bound to the lysosomal membrane is detected. However, if lysosomes have been previously treated with protease inhibitors, the protein translocated into the lumen remains intact and the total amount of substrate recovered with lysosomes corresponds to that bound to the membrane and that present in the lumen. Substrate uptake can be calculated as the difference in levels of substrate in lysosomes treated or not treated with protease inhibitors. Comparison of binding and uptake of GAPDH in lysosomes isolated from the different groups of mice revealed no differences in the rate of lysosomal binding of the protein (Fig. 1 D and E). In contrast, uptake was significantly reduced in lysosomes isolated from the HFD and CHOL groups (Fig. 1 D and E). A similar decrease in lysosomal uptake was also observed for RNase A, another well-characterized CMA substrate (Fig. S4D). These results support that the reduced rates of CMA observed in the treated animals are mainly attributable to a reduced ability of these lysosomes to translocate cytosolic substrates into their lumen.

Effect of Different Lipid Challenges on CMA Components in Lysosomes. The essential components involved in substrate translocation via CMA are LAMP-2A, which organizes into the multimeric translocation complex (9), and lys-hsc70, which assists the substrate from the luminal side of the membrane (10) and also mediates the active dissociation of LAMP-2A from the multimeric complex (9). Analysis of the levels of these two proteins in lysosomes isolated from the oleate-treated cells in which CMA activity was gradually compromised revealed a dose-dependent decrease in the levels of these two proteins in lysosomes (Fig. 2A). Levels of other lysosome membrane components, such as LAMP-1, remained unchanged.

Similar reductions in lysosomal levels of LAMP-2A and hsc70 were observed in the subgroup of CMA-active lysosomes isolated from animals maintained on the CHOL or HFD (Fig. 2B and C). Isolation of lysosomal membranes using hypertonic shock and high-speed centrifugation confirmed that changes in the CMA-related proteins were more pronounced at the lysosomal membrane (Fig. 2D and E). The reduction in levels of LAMP-1, the most abundant protein at the lysosomal membrane, was less pronounced (25% compared with the 50% observed for LAMP-2A) (Fig. 2D and E).

Dietary Lipids Reduce the Stability of LAMP-2A at the Lysosomal Membrane. Because LAMP-2A is the limiting component for CMA at the lysosomal membrane, and changes in the levels of CMA chaperones in these membranes can be secondary to the reduced levels of this receptor, we investigated the mechanism behind the reduced levels of LAMP-2A observed in the treated animal groups. We did not find significant differences in the levels of LAMP-2A or hsc70 mRNA between controls and any of the treated groups of mice, supporting the lack of differences in de novo synthesis of these proteins (Fig. 3A). In contrast, analysis of the stability of LAMP-2A at the lysosomal membrane at different times of incubation in an isotonic buffer revealed a faster reduction in the level of this protein in the membrane of lysosomes isolated from animals maintained on either the CHOL (Fig. 3B) or HFD (Fig. 3C). The differences with control mice were completely abolished when lysosomes were incubated in the presence of protease inhibitors, supporting that the marked reduction in protein levels was mainly attributable to accelerated degradation in these compartments (Fig. 3B and C).

Interestingly, reduced stability of LAMP-2A because of its accelerated degradation in lysosomes has been described as the main cause for the functional decline of CMA in aging (24). To determine the possible contribution of dietary lipids to LAMP-2A instability with age, we compared the kinetics of degradation of LAMP-2A in lysosomes isolated from 22-mo-old mice subjected or not subjected to the CHOL for 3 wk (Fig. 3B) or to the HFD for 4 mo (initiated at 18 mo of age) (Fig. 3C). Changes in LAMP-1 were only noticeable in the older group when these animals were subjected to the diet and were rather discrete.
compared with changes in LAMP-2A. As expected, CMA substrate uptake was reduced (Fig. S5) and degradation of LAMP-2A was markedly accelerated (Fig. 3B and C) in the 22-mo-old group compared with 4-mo-old mice [about 65% decrease in LAMP-2A stability in the old group, comparable to the 60% decrease previously reported in old rat livers (24)]. However, subjecting these animals to the CHOL or HFD did not have an additive effect on the rates of substrate uptake (Fig. S5) or of LAMP-2A degradation (Fig. 3B and C), supporting that both aging and lipid load likely share common mechanisms for their effect on LAMP-2A stability at the lysosomal membrane.

To investigate the mechanisms behind the lipid-mediated accelerated degradation of LAMP-2A at the lysosomal membrane further, we analyzed the different components previously described to participate in the regulated degradation of this protein. Discrete cleavage of LAMP-2A by cathepsin A at the lysosomal membrane is the trigger that initiates the degradation of this protein in the lysosomal compartment (23). Measurement of the specific activity of cathepsin A in the isolated lysosomal membranes (Fig. 3D) and immunoblot analysis for cathepsin A revealed a marked increase in the levels of the mature and precursor forms of this enzyme among the different groups of animals. However, this rapid maturation does not seem to be a generalized feature of all hydrolases in these lysosomes but, instead, something specific for cathepsin A, because analysis of other lysosomal cathepsins (cathepsin D shown in Fig. 3F) did not reveal a significant increase in the lysosomal levels of the mature and precursor forms of this enzyme among the different groups of animals.

Cathepsin A-mediated cleavage of LAMP-2A in lysosomes occurs in discrete lipid microdomain regions at the lysosomal membrane (22). Molecules of LAMP-2A destined for degradation are retrieved to these regions, where cathepsin A preferentially binds to the lysosomal membrane. Using previously established detergent extraction and flotation in sucrose density gradient procedures (22), we isolated the lipid microdomains where LAMP-2A degradation occurs from lysosomal membranes of RD-, CHOL-, or HFD-maintained mice and analyzed LAMP-2A distribution in these fractions. The resistance of the microdomains containing LAMP-2A to detergent extraction allows for their recovery in the regions of lower density of the sucrose gradient. As shown in Fig. 4, we found a consistent increase in the percentage of LAMP-2A present in these detergent-resistant regions at any given time in animals on both the CHOL (Fig. 4A) and HFD (Fig. 4B). The increased association of LAMP-2A with these regions was even evident under normal feeding conditions, when LAMP-2A is usually degraded faster (Fig. S6), suggesting that continuous enhanced degradation in these regions may be the main reason for the low levels of lysosomal LAMP-2A during the lipid challenges. In support of higher degradation, levels of cathepsin A were also considerably higher in the lipid microdomains isolated from CHOL- and HFD-maintained animals (Fig. 4A and B). Analysis of the content of ganglioside GM1,
previously shown to locate preferentially in these regions, demonstrated not only higher absolute GM1 levels in the large lipid microdomains from CHOL- and HFD-treated animals but a broader distribution of this ganglioside into other smaller detergent-resistant regions (lower flotation ability) (Fig. 4A and B).

Association of LAMP-2A with lipid microdomains has been shown to determine both the degradation rate of this receptor at the lysosomal membrane and its ability to organize into the multimeric complex required for substrate translocation, which only occurs outside these regions (9, 22). Analysis of the multimeric status of LAMP-2A at the lysosomal membrane using blue-native electrophoresis revealed a marked reduction in the amount of LAMP-2A organized into the 700-kDa multimeric complex required for substrate translocation, which only occurs outside these regions (9, 22). Analysis of the multimeric status of LAMP-2A at the lysosomal membrane using blue-native electrophoresis revealed a marked reduction in the amount of LAMP-2A organized into the 700-kDa multimeric complex required for substrate translocation, which only occurs outside these regions (9, 22). Analysis of the multimeric status of LAMP-2A at the lysosomal membrane using blue-native electrophoresis revealed a marked reduction in the amount of LAMP-2A organized into the 700-kDa multimeric complex required for substrate translocation, which only occurs outside these regions (9, 22). Analysis of the multimeric status of LAMP-2A at the lysosomal membrane using blue-native electrophoresis revealed a marked reduction in the amount of LAMP-2A organized into the 700-kDa multimeric complex required for substrate translocation, which only occurs outside these regions (9, 22). Analysis of the multimeric status of LAMP-2A at the lysosomal membrane using blue-native electrophoresis revealed a marked reduction in the amount of LAMP-2A organized into the 700-kDa multimeric complex required for substrate translocation, which only occurs outside these regions (9, 22). Analysis of the multimeric status of LAMP-2A at the lysosomal membrane using blue-native electrophoresis revealed a marked reduction in the amount of LAMP-2A organized into the 700-kDa multimeric complex required for substrate translocation, which only occurs outside these regions (9, 22). Analysis of the multimeric status of LAMP-2A at the lysosomal membrane using blue-native electrophoresis revealed a marked reduction in the amount of LAMP-2A organized into the 700-kDa multimeric complex required for substrate translocation, which only occurs outside these regions (9, 22). Analysis of the multimeric status of LAMP-2A at the lysosomal membrane using blue-native electrophoresis revealed a marked reduction in the amount of LAMP-2A organized into the 700-kDa multimeric complex required for substrate translocation, which only occurs outside these regions (9, 22). Analysis of the multimeric status of LAMP-2A at the lysosomal membrane using blue-native electrophoresis revealed a marked reduction in the amount of LAMP-2A organized into the 700-kDa multimeric complex required for substrate translocation, which only occurs outside these regions (9, 22). Analysis of the multimeric status of LAMP-2A at the lysosomal membrane using blue-native electrophoresis revealed a marked reduction in the amount of LAMP-2A organized into the 700-kDa multimeric complex required for substrate translocation, which only occurs outside these regions (9, 22). Analysis of the multimeric status of LAMP-2A at the lysosomal membrane using blue-native electrophoresis revealed a marked reduction in the amount of LAMP-2A organized into the 700-kDa multimeric complex required for substrate translocation, which only occurs outside these regions (9, 22). Analysis of the multimeric status of LAMP-2A at the lysosomal membrane using blue-native electrophoresis revealed a marked reduction in the amount of LAMP-2A organized into the 700-kDa multimeric complex required for substrate translocation, which only occurs outside these regions (9, 22). Analysis of the multimeric status of LAMP-2A at the lysosomal membrane using blue-native electrophoresis revealed a marked reduction in the amount of LAMP-2A organized into the 700-kDa multimeric complex required for substrate translocation, which only occurs outside these regions (9, 22).

Effect of Dietary Lipids on the Lipid Composition of Lysosomal Membranes. To analyze the impact that dietary lipids had on the lipid composition of the lysosomal membrane directly and to compare these changes with those associated with aging, we isolated lysosomes from animals on the CHOL and HFD and from 22-mo-old animals maintained on the RD. To be able to analyze changes in the lipid composition at the lysosomal membrane, independent of the lysosomal content, we subjected lysosomes to hypotonic shock and separated the membrane fraction by high-speed centrifugation. In contrast to lysosomes involved in macrophtagocytosis, the subgroup of lysosomes active for CMA does not participate in degradation of organelles under normal conditions (28); consequently, contamination by membranes or organelles sequestered for degradation is unlikely. In fact, the lumen of these lysosomes contains mainly amorphous material, luminal membranes are not often observed (Fig. S7A), and immunoblot analysis reveals that only minimal traces of structural membrane proteins of different organelles are detected in this subgroup of lysosomes (Fig. S7B).

We performed comparative lipidomic analysis of lysosomal membranes from these four groups (RD, CHOL, HFD, and 22-mo-old mice) to gain a better understanding of the qualitative and quantitative changes in lipid composition imposed by dietary lipids and by aging. Analysis of the lysosomal membranes from the CHOL-treated animals revealed a statistically significant increase (~20%) in the percentage of free cholesterol in these membranes along with a statistically significant ~25% decrease in the percentage of phosphatidylethanolamine (PE) as trends for a reduction in major lipid groups at the lysosomal membrane, such as phosphatidylcholine (PC), phosphatidylinositol (PI),...
phosphatidylserine (PS), and sphingomyelin (SM) (Fig. 5A). Interestingly, we found a marked increase (~75%) in the percentage of lysobisphosphatidic acid (LBPA) (Fig. 5A, Inset), a phospholipid known for its ability to enhance transfer of cholesterol from and to membranes and to mediate the formation of intraluminal vesicles in the endolysosomal system (29). We then analyzed possible changes in lipid species composition (Fig. 5D shows the heat map of the comparative lipid profiles) and found that CHOL-treated animals have a significant decrease in saturated SM and ceramide (CER) chains and a significant increase in the forms with unsaturated chains, both in SM and in CER (Fig. 5D and Fig. S8 A–D). In contrast, we observed a significant increase in the percentage of long-chain saturated PC, along with a decrease in the PC chains, with a higher number of unsaturations (Fig. 5D and Fig. S9 A–D).

Comparative lipidomics in the lysosomal membranes from HFD-treated mice revealed that some of the changes in their lipid composition were shared with the CHOL group, whereas others were specific for this treatment. For example, we did not find changes in the percentage of free cholesterol in HFD lysosomal membranes but noticed that CER levels were significantly increased instead (Fig. 5B). These changes may have a similar effect on membrane organization, because CER has been shown to promote the order of lipid membranes and to segregate laterally into rigid, gel-like domains to a higher extent than cholesterol (30, 31). Interestingly, as in the CHOL group, the membranes from HFD-treated mice contained significantly higher levels of LBPA (Fig. 5B). This increase in LBPA cannot be attributed to a mere higher content of contaminating late endosomal compartments in the lysosomal fraction isolated from CHOL- or HFD-treated mice, because analysis of mannose-6-phosphate receptor, a marker of the recycling endocytic compartment, was comparable in lysosomes from mice maintained on the RD and high-lipid content diets (Fig. S7C). Also shared with the CHOL group was the decrease in PE (Fig. 5B), and some of the changes in SM and CER, such as those pertinent to long-saturated lipid species (heat map in Fig. 5D and Fig. S8 E–H). However, in clear contrast to the CHOL group, we observed a significant decrease of long-chain saturated and monounsaturated PC, PE, and PI, along with an increase in the PC, PE and PI chains, with a high number of unsaturations (heat map in Fig. 5D and Fig. S9 E–H). Changes in monounsaturated SM and CER species and in some of the forms with both chains unsaturated were also the reverse of those observed in CHOL-treated mice (Fig. 5D and Fig. S8 E–H). These results support that both diets favor formation of organized lipid microdomains at the lysosomal membrane, but of different lipid composition. This may explain why although both diets exert an inhibitory effect on CMA, the effect of the CHOL was consistently more pronounced (compare substrate uptake, LAMP-2A levels, and LAMP-2A stability in Fig. 1 D and E, Fig. 2 D and E, and Fig. 3 B and C).

Finally, we performed a similar lipidomic analysis in lysosomal membranes from 22-mo-old mice maintained on the RD. As in the case of the lipid diets, the lipid composition of the membranes from the older age group had some distinctive features unique for this group, but it also reproduced many of the changes observed in the lipid-challenged animals, overall being closer to the changes in the CHOL group. Thus, we observed a small (~5%) but significant increase in free cholesterol levels, as in the CHOL group, and increases in CER (~15%), glucosyl ceramide (~35%), and LysoPC (~45%) comparable to those observed in the HFD group (Fig. 5C). Strikingly, the increase in LBPA observed with both diets was also noticeable in the aged group (Fig. 5C). Overall, aging promoted changes in the composition of the acyl chains that resemble those of animals on the CHOL: an increase in SM, CHOL, and glucoceramide unsaturated in both chains (Fig. S8 I–L), an increase in the percentage of saturated PC, and a decrease
in the percentage of polyunsaturated PC (Fig. S9 I–L). The three treatments resulted in comparable changes in long saturated SM species (Fig. S8 A, E, and I), and the aging group shared with the HFD-treated animals an increase in the levels of short saturated species of SM (Fig. S8 I–L). This increase in short saturated species of PC and SM with age, along with the increase in free cholesterol, may promote the formation of cholesterol-enriched, detergent-resistant microdomains.

Overall, our findings are consistent with lipid challenges through the diet having a direct impact on the organization of the lysosomal membrane through qualitative and quantitative changes in its lipid composition. The lipidomic analysis also reveals high similarity between the diet-induced changes at the lysosomal membrane and those occurring with age. Among the numerous consequences that these changes can have on lysosomal function, in this work, we have characterized the negative effect that the high enrichment in cholesterol and perturbation of the lysosomal membrane lipid composition has on CMA, a pathway heavily dependent on membrane lateral mobility.

Discussion
In this work, we have identified a negative effect of dietary lipid challenges on CMA activity that is mediated, for the most part, by changes in the lipid composition of the lysosomal membrane as a result of the lipid exposure. The decreased stability of lysosomal membrane proteins observed under these conditions, particularly the higher susceptibility of LAMP-2A to these changes, unveils a unique mechanism for CMA compromise under these conditions, which could also be behind the functional loss in this pathway under different pathological conditions and in aging.

Our analysis also emphasizes the modulatory role of lipids on autophagy as part of the recently described interplay between this catabolic process and lipid metabolism. Thus, we have recently reported that macroautophagy can contribute to the mobilization of intracellular lipid deposits; in fact, this process, known as macrolipophagy, is up-regulated in response to moderate lipid challenges (32). However, chronic lipid challenges or acute exposure to abnormally high lipid concentrations exerts an inhibitory effect on macroautophagy (28). The main defect identified as responsible for macroautophagy failure under these conditions is the reduced fusion ability between the vesicular compartments involved in that process: autophagosomes and lysosomes. Interestingly, the compromise in vesicular fusion can be attributed to changes in the lipid composition of the membrane of these vesicles and, in fact, can be reproduced by artificially mimicking these lipid changes in isolated autophagosomes (i.e., using

Fig. 5. Changes in the lipid composition of the lysosomal membrane in response to dietary challenges and aging. Lysosomal membranes isolated from livers of mice maintained on the RD and CHOL (A), on the RD and HFD (B), or from 3- and 22-mo-old mice (C) were subjected to lipidomic analysis. Graphs show the mean values of the molar percentage of every lipid class with respect to the total amount of lipids (after excluding triglycerides and cholesterol esters). Values are the mean ± SEM of five different experiments. (D) Heat map showing the comparative lipid profile of lysosomal membranes isolated from livers of mice maintained on the CHOL vs. RD, on the HFD vs. RD, or in 22-mo-old vs. 3-mo-old mice (OLD/YOUNG). The three columns represent the normalized values of the individual lipid species. The color bar represents the log2 value of the ratio of each lipid species. Only changes that are statistically significant in the relative lipid amount are highlighted in the heat map. Relative increases and decreases are shown in red and green, respectively (n = 5). *P < 0.05 compared with the RD group. CE, cholesterol ester; CHOL FREE, free cholesterol; DAG, diacylglyceride; LPC, lysophosphatidyl choline; PA, phosphatidic acid; PG, phosphatidylglycerol; TAG, triglyceride.
chemical extractors of cholesterol) (28). Here, we show that high concentrations of cholesterol at the lysosomal membrane also exert a marked inhibitory effect on CMA, mainly by enhancing LAMP-2A mobilization to the specific microdomains where this protein is normally degraded. This dual compromise of macroautophagy and CMA in response to an abnormally high content of dietary lipids may underlie the basis of part of the cellular toxicity observed under these conditions. These autophagic pathways, along with the ubiquitin/proteasome system, are mainly responsible for the maintenance of cellular quality control. Thus, compromise of the two autophagic pathways may render cells particularly susceptible to stressors, such as oxidative stress, because of the inability to handle the damage associated with these stressors. In this respect, and in agreement with previous reports (33, 34), we found a moderate increase in the levels and some of the proteolytic activities of the proteasome in the liver of animals exposed to the diets with a high lipid content (Fig. S10 A and B). Up-regulation of the proteasome may be a cellular attempt to compensate for the loss of activity of the autophagic systems. However, the fact that levels of polyubiquitinated proteins increase during the high-lipid diets (Fig. S10C) already points toward an overall deficient cellular quality control.

In addition to the expected problems in cellular quality control, it is possible that the inhibitory effect of lipid challenges on CMA described in this work may have important implications in cellular metabolism. Thus, in studies with cancer cells, we have recently identified a role for CMA in the regulation of glycolysis and the need for functional CMA to maintain proper β-oxidation in these cells (35). Although the mechanisms behind this effect of CMA in β-oxidation remain unknown, it is interesting that CMA was consistently up-regulated in response to exposure to low concentrations of oleic acid (Fig. 1A). This up-regulation of CMA could be linked to the increase in β-oxidation necessary to accommodate the higher affluence of lipids under these conditions. In this respect, the inhibitory effect of dietary lipids on CMA may further contribute to intracellular lipid accumulation by reducing the mitochondrial catabolism.

Future studies are required to elucidate the reason for the higher susceptibility of LAMP-2A to lysosomal membrane lipid challenges, compared with other lysosomal membrane proteins. We have previously shown differences between the membrane dynamics of LAMP-2A and the other variants of this protein and the phospholipids of LAMP-1 (9, 21, 22). Although these other proteins can also be associated with lipid microdomains, they do not seem to coincide in the same ones in which LAMP-2A undergoes degradation (22). Likewise, LAMP-2B and LAMP-1 can also be detected in oligomeric complexes at the lysosomal membrane, but they are usually of smaller size than the 700-kDa complex enriched in LAMP-2A that is required for substrate translocation (9). It is possible that the diet-induced changes in the lipid composition of the lysosomal membrane affect specific microdomain regions where LAMPS locate differently.

We have found a marked change in the amount of cholesterol at the lysosomal membrane of animals maintained on the CHOL. When used in model membranes, similar concentrations of cholesterol have been shown to form mainly a liquid-ordered phase, equivalent to that in the LAMP-2A-enriched lysosomal microdomains (36). However, as revealed by the lipidomic analysis, changes in other lipids at the lysosomal membrane also contribute to decreased CMA. In this respect, we have found that not only quantitative but qualitative changes in different lipid species occur at the lysosomal membrane during the dietary cholesterol lipid challenge. Some of the noted changes seem to favor more compact cholesterol packing inside microdomains, as is the case with the relative increase in saturated PC, which occupies less space than saturated SM (which is actually decreased in CHOL lysosomes). In contrast, other changes may help to maintain the fluidity of the membrane outside the microdomain regions despite the high increase in cholesterol. For example, the observed switch toward unsaturated forms of SM and CER would allow for such high concentrations of cholesterol while maintaining fluidity, because unsaturated lipids with a double bond in the middle of the acyl chain remain fluid even in the presence of cholesterol (37).

The comparative analyses of lipid changes in both diets support that different lipid combinations may have a similar impact on the final properties of the lysosomal membrane. Thus, in the HFD group, we did not observe an increase in cholesterol levels; instead, the elevated levels of short saturated CER and SM species detected in this group may promote the formation of even tighter domains (38, 39). It is well established that CER and phospholipids with polyunsaturated fatty acid chains, which are also elevated in the lysosomal membrane of HFD mice, do not mix well with cholesterol and form domains with different lipid packing than those of cholesterol, which may promote the concentration of specific proteins in these domains (40). Therefore, although HFD- and CHOL-induced changes in the lipid composition of the lysosomal membrane were not identical, both diets favor the formation of detergent-resistant microdomains in which LAMP-2A is degraded. The qualitative differences in the composition of the microdomains promoted by each diet may explain why the inhibitory effect of the HFD on CMA is less prominent compared with the one observed after the CHOL.

The age-related changes in the lipid composition of the lysosomal membrane revealed features also observed in the other two groups (i.e., increase in LBPA or changes in the long saturated SM chains); however, overall, the changes in the aged group more closely resemble those induced by the CHOL, which are also the ones with a higher negative impact on CMA activity. As for both diets, age-dependent changes in lysosomal membrane lipids also favor the formation of microdomains. In this case, along with high CHOL, the increases in glycosylceramide and other glycosphingolipids and the enrichment in short and saturated forms of SM and PC also promote the formation of detergent-resistant microdomains. In previous studies, we have reported that in addition to the alteration in the regulated degradation of LAMP-2A that occurs in the microdomains, part of the membrane-resistant LAMP-2A is abnormally internalized into the lumen, where it is rapidly degraded (24). It is plausible that changes in the lipid composition of the lysosomal membrane with age observed in this study also contribute to that abnormal internalization. In this respect, the marked increase in LBPA observed in the three interventions is of great interest because this unconventional phospholipid can induce formation of small vesicles and invaginations in other membranes (29).

The different properties of the lipid microdomains could have a direct impact on the ability of different LAMPS to associate with these regions. Although all LAMPS have transmembrane regions of comparable length and highly homologous luminal regions that predict similar structural features (41), differences in posttranslational modifications among them, which are likely changes in the glycosylation pattern, could modulate their interaction with the lipids in different microdomains. Another possibility is that yet unidentified membrane proteins actively mobilize LAMPS in and out of these microdomain regions and that changes in the packing density of lipids or length of their lateral chains determine the affinity of these LAMP-targeting proteins for the lipid microdomain regions. In this respect, membrane-associated hsc70 has been previously shown to be necessary for the active insertion of LAMP-2A at the lysosomal membrane, because blocking antibodies against this chaperone are enough to inhibit this process (20). Finally, it is also possible that the increase in the content of known lysosomal lipidic cargos, such as cholesterol esters, contributes to the abnormal dynamics of LAMP-2A in the lysosomes of animals exposed to the
CHOL. In fact, a fraction of lysosomal LAMP-2A that resides in the lumen associated with lipids can be retrieved back to the lysosomal membrane under conditions requiring maximal CMA activation (20). It is plausible that changes in the lysosomal luminal lipids affect the efficiency of retrieval of this subtraction of lysosomal LAMP-2A toward the lysosomal membrane.

In this study, we have also identified a marked decrease in levels of hsc70 in the membranes of lysosomes from both CHOL- and HFD-maintained mice. It is possible that this decrease is just a consequence of the reduced levels of LAMP-2A at the lysosomal membrane in these animals, because hsc70 interacts with this receptor for substrate translocation. However, in light of the recently described direct association of hsc70 with lipids described at the membrane of late endosomes (42), we cannot discard the possibility that the reduced levels of hsc70 at the lysosomal membrane after the lipid challenges reflect reduced direct binding to membrane lipids. In the case of late endosomes, hsc70 binds directly to PS in the outer leaflet of the membrane. Although, we have not found significant differences in the PS content at the lysosomal membrane after the lipid challenges, we cannot discard the possibility that changes in other lipids could affect the organization of PS at the lysosomal membrane and, in this way, interfere with chaperone binding. In addition, the observed increase in LBPA could affect the association of hsc70 with the lysosomal membranes by competition with hsp70, which has been demonstrated to bind LBPA directly in membranes (29).

The pathophysiological implications of our findings are multiple, because these could help in elucidating the basis for CMA malfunction in different conditions. From the physiological point of view, this modulatory effect of intracellular lipids on CMA may directly or indirectly contribute to the previously described bidirectional cross-talk between macroautophagy and CMA (18, 43). Most cells respond to blockage in macroautophagy by up-regulating CMA (43), which has been proven beneficial because it helps in preserving cellular resistance to particular stressors (44). Based on the recently described contribution of macroautophagy to mobilization of intracellular lipid stores (32), it is anticipated that a blockage in macroautophagy will reduce the availability of intracellular cholesterol and reduce its levels in organelle membranes, including lysosomes. This reduction in lysosomal membrane cholesterol may contribute to the up-regulation of CMA observed when macroautophagy is compromised. Aging is associated with a decline in CMA activity, which seems, for the most part, to be attributable to reduced stability of LAMP-2A at the lysosomal membrane (24). Our early studies support that the rapid degradation of LAMP-2A in lysosomes from old animals is mainly attributable to changes in the lysosomal membrane with age rather than to direct changes in LAMP-2A, because the increased instability can be reproduced when the recombinant LAMP-2A is incorporated in resealed membranes from old mice lysosomes but not if the membranes originated from young mice (24). Our current study supports that changes in the lipid composition of the membrane with age are responsible for this destabilizing effect on LAMP-2A. In fact, the accelerated degradation of LAMP-2A in lysosomes isolated from animals maintained on the CHOL or HFD is comparable to the one observed in lysosomes from old mice (24). The fact that maintaining old animals on the HFD for 4 mo or on the CHOL diet for 3 wk did not have an additive effect on LAMP-2A instability further supports that changes in lipid composition of the lysosomal membrane with age are behind the observed decline in CMA activity in aging. Reduced CMA activity has also been described in different neurodegenerative diseases, such as Parkinson disease and some tauopathies (45–47), and in metabolic disorders, such as diabetes (13). Because aging has been shown to be an aggravating factor in all these diseases, it has been proposed that the primary defect in CMA could be further aggravated by the age-dependent decrease in the activity of this pathway, and thus contributes to accelerate the pathological changes. In this respect, modulation of dietary lipid intake may provide a way to slow down the decline in CMA with age and, consequently, to delay disease onset. Interestingly, alterations in intracellular lipids and in lipid metabolism in general have also been described in several of these conditions, leaving open the possibility of a perpetuating negative feedback between intracellular lipids and CMA in these pathological conditions and further reinforcing the possible beneficial effect of interventions aimed at modulating dietary lipid intake.

Methods

A detailed description of all methods is provided in SI Methods.

Animals, Cells, and Reagents. Male C57BL/6 mice (6–8 wk old) from the Jackson Laboratory were maintained on the RD (2018, Global 18% Protein Rodent Diet; Teklad), the HFD (D12492, 60% kcal in fat; Research Diets) for 16 wk once they reached 8 wk of age, or the 2% CHOL (2018 + 2% cholesterol TD.01383; Tekland) for 3 wk. For the aging studies, 3- and 22-mo-old mice from the National Institute on Aging age-controlled colony were used. For lysosomal isolation, the livers of two animals were pooled per condition in each experiment. Mouse fibroblasts (National Institutes of Health NIH 3T3) were from the American Type Culture Collection. Sources of chemicals are as described previously (8, 18, 20, 25) and as detailed in SI Methods.

Lysosome Isolation and CMA Measurements. Lysosomes were isolated from mouse liver and cultured cells by centrifugation in a discontinuous metrizamide density gradient as described (48). Uptake assays were performed by incubation of isolated lysosomes with radiolabeled cytosolic proteins and analysis of protein breakdown (49) or with single purified proteins and analysis of protein association by immunoblot as described (26). The stability of LAMP-2A was measured in intact lysosomes on incubation in isotonic medium for increasing periods of time (20). Detergent-resistant microdomains were isolated from lysosomal membranes by detergent extraction and flotation in sucrose gradients as described (22). Blue-native electrophoresis of solubilized lysosomal membranes was used to visualize the CMA translocation complex utilizing 3–12% (w/vol) NativePAGE Novex bis-Tris precast gels (Invitrogen).

Measure of CMA Activity in Intact Cells. Direct fluorescence microscopy was used to determine CMA activity in fibroblasts expressing the CMA reporter (KFERQ-photonactivable mCherry1) after photoactivation (11). Images were acquired with an Axiovert 200 fluorescence microscope (Carl Zeiss Ltd.), subjected to deconvolution with the manufacturer’s software, and prepared using Photoshop 6.0 software (Adobe Systems, Inc.).

Lipid Extraction and Analysis. A modified Bligh/Dyer extraction procedure was used for lipid extraction from organelle fractions before analysis by liquid chromatography-mass spectrometry utilizing multiple reaction monitoring (50, 51). Separated lipid classes were quantified via multiple reaction monitoring (MRM) mode on a triple-quadrupole instrument (API 3200; Applied Biosystems) using previously reported MRM transition pairs and instrument settings (50).

General Methods. BODIPY 493/503 was used to visualize lipid droplets in cultured cells (32). RT–PCR was used for mRNA quantification in total RNA prepared with the SuperScript II RNase H Reverse Transcriptase (Invitrogen) and oligo-(dT)18 primers. Details of amplification primers are provided in SI Methods.

Statistical Analysis. Two-way ANOVA, followed by the Bonferroni post hoc and Student t tests for unpaired data, was used for statistical analysis.

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Supporting Information

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SI Methods

Animals and Cells. Male C57BL/6 mice (6–8 wk old) were obtained from the Jackson Laboratory. Animals were maintained on the RD (2018, Global 18% Protein Rodent Diet; Teklad), the HFD (D12492, 60% kcal in fat; Research Diets) for 16 wk once they reached 8 wk of age, or the 2% CHOL (2018 + 2% cholesterol TD.01383; Teckland) for 3 wk. At the end of the treatments, some of the animals were starved for 24 h before isolation of lysosomes but they had free access to water. All studies were approved by the Animal Care and Use Committee of the Albert Einstein College of Medicine and followed the National Institutes of Health guidelines on the care and use of animals. Mouse fibroblasts [National Institute of Health (NIH) 3T3] were from the American Type Culture Collection. Cells were maintained in DMEM (Sigma) supplemented with 10% (vol/vol) newborn calf serum. Oleic acid and palmitic acid were conjugated to albumin as described (1), and cells were treated for 24 h.

Chemicals. Sources of chemicals were as described previously (2–5). The antibodies against the cytosolic tail of rat and mouse LAMP-2A and LAMP-2B were prepared in our laboratory (2). The antibodies against mouse LAMP-1 and M6PR were from the Developmental Studies Hybridoma Bank (Iowa University), the antibody against cathepsin D was from Santa Cruz Biotechnology, and the antibodies against hsc70 and hsp90 were from Stressgen. The antibodies against RNase A and cathepsin A were from Rockland Immunochemicals, and the antibodies against GAPDH and ABCB9 were from Abcam. The antibodies against the 19S and 20S proteasome subunits were from BioMol International, the antibody against ubiquitin was from Dako, and the antibody against GFP was from Roche. U18666A was from Sigma, and the antibody against the 19S and 20S proteasome subunits were from BioMol International. Lysosomal membranes were solubilized by octyl glucoside buffer (except where indicated), and incubating the supernatant can originate only from lysosomal disruption because secretion of this enzyme or permeation through the membrane is not possible. Consequently, the percentage of total lysosomal hydrolase value (lipid diet)/value (RD) [ad-particle centrifugation as described (6)].

Isolation of Subcellular Fractions. Mouse liver lysosomes were isolated from a light mitochondrial-lysosomal fraction in a discontinuous metrizamide density gradient, and lysosomal fractions with different activities for CMA were further separated by differential centrifugation as described (6). Lysosomes from cultured cells were isolated as described (7). A crude fraction containing lysosomes and mitochondria was prepared after cell lysis by differential centrifugation (2,500 × g for 15 min and 17,000 × g for 10 min). Lysosomal matrices and membranes were isolated after hypotonic shock (8). Right after isolation, lysosomes were subjected to centrifugation and the activity of the lysosomal hydrolyase β-hexosaminidase was measured in both the pellet (lysosomes) and the supernatant. The enzyme present in the supernatant can originate only from lysosomal disruption because secretion of this enzyme or permeation through the membrane is not possible. Consequently, the percentage of total β-hexosaminidase activity (pellet and supernatant) present outside lysosomes (supernatant) was used to monitor lysosomal breakage. We did not find differences in purity of the fractions or stability of the lysosomal membrane between fed and starved animals or those maintained on the RD, CHOL, or HFD [additional purity information is provided by Koga et al. (9)].

Uptake and Degradation of Substrate Proteins by Isolated Lysosomes. Mouse fibroblast cytosolic proteins were metabolically radio-labeled by incubation with [3H]leucine (2 μCi/mL) at 37 °C for 2 d (6). This pool of radiolabeled cytosolic proteins was incubated in 10 mM 3-(N-morpholino) propanesulfonic acid (MOPS) buffer (pH 7.3), 0.3 M sucrose, and 1 mM DTT with intact lysosomes at 37 °C for 30 min (10). Degradation of the radiolabeled substrates was measured after acid precipitation, and protein degradation was expressed as the percentage of the initial acid-insoluble radioactivity (protein) transformed into acid-soluble radioactivity (amino acids and small peptides) at the end of the incubation. Where indicated, lysosomes were disrupted by a hypotonic shock before incubation (8). GAPDH or RNase A was incubated in the MOPS buffer with untreated or protease inhibitor-treated lysosomes as described (10). After incubation for 20 min at 37 °C, lysosomes were collected by centrifugation and samples were subjected to SDS/PAGE and immunoblotted with an antibody against GAPDH or RNase A. Transport was measured, and uptake was calculated as the difference between the amount of substrate associated with lysosomes (protease inhibitor-treated lysosomes) and the amount of substrate bound to their membrane (untreated lysosomes). The inhibitory effect of the diets on the different steps of CMA activity was calculated by the formula [value (RD – value lipid diet)/value (RD)]-100 and expressed as a percentage.

Measure of CMA Activity in Intact Cells. Stable clones of mouse fibroblasts expressing either of the two CMA reporters (KFERQ-photoactivatable mCherry1 or KFERQ-photoactivatable CFP2 (11), treated with oleate or lipid-modifying drugs, as indicated, were photoactivated by light-emitting diode (Norlux) for 10 min with an intensity of 3.5 mA (constant current). After 16 h, cells were fixed with 4% (wt/vol) paraformaldehyde and mounted, and images were acquired with an Axiovert 200 fluorescence microscope (Carl Zeiss Ltd.), subjected to deconvolution with the manufacturer’s software, and prepared using Photoshop 6.0 software (Adobe Systems, Inc.). The number of fluorescent puncta per cell was quantified using ImageJ software (National Institutes of Health) in individual frames after thresholding.

Measurement of LAMP-2A Degradation. Rates of degradation of LAMP-2A in the isolated lysosomal membranes were determined by immunoblotting using a specific antibody against the cytosolic tail of LAMP-2A as previously described (3). Briefly, isolated lysosomal membranes were incubated in MOPS buffer at 37 °C, and aliquots were removed and subjected to SDS/PAGE and immunoblotting for LAMP-2A at different times.

Isolation of Detergent-Resistant Lysosomal Membrane Microdomains. Lysosomal membranes (150 μg of protein) from mouse liver were incubated with 1% Triton X-114 in 150 mM NaCl, 50 mM Tris-HCl, and 5 mM EDTA (pH 7.4) (incubation buffer) on ice for 30 min; they were then subjected to centrifugation in a stepwise discontinuous sucrose gradient (5–35%) and centrifuged at 200,000 × g for 19 h in an SW-60Ti rotor (Beckman) (12). Four samples of 500 μL collected starting from the detergent-resistant band were subjected to acid precipitation with 10% (vol/vol) trichloroacetic acid and BSA. Precipitates were washed with acetone, resuspended in electrophoresis sample buffer, and subjected to SDS/PAGE and immunoblotting.

Identification of LAMP-2A Multimeric Complexes by Blue-Native Electrophoresis. Lysosomal membranes were solubilized by resuspending them in 20 mM MOPS, 150 mM NaCl, and 0.5% octyl glucoside buffer (except where indicated), and incubating...
Protein concentration was determined by the 225. at 4 °C. The organic phase was recovered, 5910. Aldrich) to measure 272:18% (wt/vol) NativePAGE Novex bis-Tris precast gels (In- vitrogen), transferred to PVDF membranes, and immunoblotted for LAMP-2A and ABCB9.

Direct Fluorescence and Indirect Immunofluorescence Microscopy. To visualize lipid droplets, cells grown on coverslips were incubated with BODIPY 493/503 for 15 min (14). After extensive washing, cells were fixed with 4% (wt/vol) paraformaldehyde and mounted, and images were acquired with an Axiovert 200 fluorescence microscope, subjected to deconvolution with the manufacturer’s software, and prepared using Photoshop 6.0 software. The number of fluorescent puncta per cell was quantified using ImageJ software in individual frames after thresholding. For immunofluorescence, fixed cells were permeabilized and incubated with primary antibody, followed by fluorophore-conjugated secondary antibody, as described previously (11). The percentages of colocalization and colocalized pixels were measured using JACoP (just another colocalization plug-in; Susanne Bolte and Fabrice P. Cordelières) and colocalization plug-ins of ImageJ, respectively.

mRNA Quantification. Total RNA was extracted from mouse livers using the RNeasy Protect Mini Kit (Qiagen) following the manufacturer’s instructions and stored at −80 °C until use. The first-strand cDNA was synthesized from 0.5 µg of the total RNA with the SuperScript II RNase H Reverse Transcriptase (In-vitrogen) and oligo-(dT)12-18 primers. Actin and a region of exon 8 of LAMP-2A were amplified with specific primers (LAMP-2A, 5′-GCAGTGCAGATGAAGACAAC-3′; LAMP-2A, 5′-GTATGATGGC- GCTTGAGAC-3′; actin, 5′-AAGGACTCTCTATAATGTTG- AGCA-3′; 5′-ATCTTCCTCATGTCGTCCAGTTG-3′; hsc70, 5′-TCTCGGCAACACTCTCC-3′; 5′-CCGACTAGCGT- TTGC-3′) using the SYBR green PCR kit (PE Biosystems). Amplification of the LAMP-2A, hsc70, and actin DNA products (120, 196, and 108 bp, respectively) was measured in real-time in a SmartCycler (Cepheid). The expression levels of LAMP-2A in different samples were normalized with respect to those of actin in the same samples.

Lipid Extraction and Analysis. Lipid extracts were prepared using a modified Bligh/Dyer extraction procedure, In brief, membranes from organelle fractions (50 µL) were incubated with methanol and chloroform (2:1 ratio) at 0 °C for 5 min, and lipids were extracted by addition of KCl (1 M, 1:10 ratio) and centrifugation for 2 min at 7,500 × g at 4 °C. The organic phase was recovered, and extracted lipids were dried and stored at −80 °C. Lipid extracts were spiked with internal standards and analyzed by liquid chromatography (LC)/MS using multiple reaction monitoring (15–17). The internal standards used included 1,2-dioctanoyl-glycerol-3-phosphoinositol (PE; 16:0 ratio; Echelon Biosciences, Inc.), 1,2-dimyristoyl-glycerol-phosphoserine (PS; 28:0 ratio), 1,2-dimyristoyl-glycerol-3-phosphoethanolamine (PE; 28:0 ratio), 1,2-dimyristoyl-glycerol-3-phosphocholine (PC; 28:0 ratio), laur- yl-SM (C12-SM), N-heptadecanoyl-d-erythro-sphingosine [C17-CER], and N-glucosyl-β-1′-N-octanoyl-d-erythro-sphingosine [C8-glucosyl ceramide (Glu-CER)], which allowed the measurement of PI, PS, PE, phospholipid-PE, PC, ether-linked-PC, SM, CER, and Glu-CER, respectively. For LC separation, we followed a previously established method using a Luna silica column (3 µm, 2 mm × 150 mm; Phenomenex) with gradient elution of 100% chloroform/methanol/water/ammonia solution (90:9:5.0:3.2 vol/vol) changing to 100% chloroform/methanol/water/ammonia solution (50:48:2.0:3.2 vol by vol) over 50 min at 0.35 mL/min. Separated lipid classes were quantified via MRM mode on a triple-quadrupole instrument (API 3200; Applied Biosystems). PE/phospholipid-PE, PI, PS, and PA were detected in negative ionization, whereas CER, GluCER, PC/ether-linked-PC, and SM were detected in positive ionization. Diacylglycerides were analyzed using a modified version of reverse phase high pressure liquid chromatography/electrospray ionization/mass spectrometry (HPLC/ESI/MS) as described previously (15, 17). The remaining lipid classes were measured using previously reported MRM transition pairs and instrument settings (16, 17). EM. Lysosomal compartments pelleted after centrifugation were fixed in 2.5% glutaraldehyde in sodium cacodylate (SC) [100 mM SC (pH 7.4)] at room temperature for 45 min. The pellet was then rinsed in SC, postfixed in 1% osmium tetroxide in SC followed by 1% uranyl acetate, dehydrated through a graded series of ethanol, and embedded in LX112 resin (LADD Research Industries). Ultrathin sections were cut on a Reichert Ultratoc E, stained with uranyl acetate followed by lead citrate, and viewed on a JEOL 1200EX transmission electron microscope at 80 kV.

Measurement of Proteasomal Activity. Proteasomal activity was determined in liver homogenate. Briefly, liver pieces lyzed by sonication in 10 mM Tris-HCl (pH 7.8), 1 mM EDTA, 0.5 mM DTT, and 5 mM MgCl2 at 4 °C were centrifuged at 16,000 × g for 10 min, and the supernatant was used to measure proteasomal activity directly. The individual proteolytic activities of the proteasome were assayed as their ability to cleave fluorescent peptides specific for each of the activities as follows: 25 µg of liver homogenate was incubated at 37 °C with 1 mM N-succinyl-leu-leu-val-tyr-7-amido-4-methylcoumarin (Sigma–Aldrich) to measure peptidyl-glutamyl peptide-hydrolizing activity, with 1 mM N-CBZ-leu-leu-glu-β-naphthylamide (Sigma–Aldrich) to measure chymotrypsin-like activity, and with 1 mM Bz-val-gly-arg- AMC (BIOMOL International) to measure trypsin-like activity. Real-time fluorescence was measured at an excitation of 350 nm and an emission of 440 nm for 30 min. Proteasome inhibitor MG-115 and/or MG-132 was used as a control.

General Methods. Protein concentration was determined by the Lowry method, using BSA as a standard (18). For immunoblotting, the proteins recognized by the specific antibodies were visualized by chemiluminescence methods (Renaissance; NEN–Life Science). Lysosomal enzymatic hexosaminidase and cathepsin A activities were measured as reported previously (7). Quantification of cholesterol was performed using the Amplex Red cholesterol assay kit following the manufacturer’s instructions. GAPDH activity was measured with the GAPDH activity kit (ProteinSci).


Fig. S1. Effect of treatment with free-fatty acids on CMA activity in fibroblasts. (A) Mouse fibroblasts stably expressing a KFERQ-PS-CFP plasmid were exposed to the indicated concentrations of oleic acid in the presence or lack of leupeptin to block lysosomal degradation. Sixteen hours after photoswitching, cells were fixed and immunostained with anti-GFP. (Left) Representative fields. (Right) Number of fluorescent puncta per cell. (B) Mouse fibroblasts stably expressing a KFERQ-PA-mCherry plasmid were exposed to the indicated concentrations of palmitic acid. (Left) Representative images 24 h after photoactivation. (Right) Number of fluorescent puncta per cell. Values are the mean ± SEM of three different experiments with >50 cells quantified per experiment. *P < 0.05 compared with untreated cells. In A, the two-way ANOVA, followed by a Bonferroni posttest, showed no interaction between the leupeptin and oleic acid treatments.
Fig. S2. Effect of different inhibitors of lipid metabolism on intracellular lipid content. Mouse fibroblasts were maintained in the presence or absence of serum, treated with the indicated compounds, and stained with BODIPY and an antibody against LAMP-1. (A) Representative images. Individual channels and merged images are shown. (Bottom) Colocalization of both fluorophores is highlighted in white. (B) Quantification of the number of lipid droplets (LD) per cell after each treatment. (C) Percentage of colocalization between BODIPY and LAMP-1 after each treatment. Values are the mean ± SEM of three different experiments with >50 cells quantified per experiment. *P < 0.05 compared with untreated cells.
Fig. S3. Effect of lipid challenges on lysosomal proteolytic activity. (A) Lysosomes from livers of mice maintained on the RD, CHOL, or HFD were disrupted by a hypotonic shock and incubated with a pool of radiolabeled cytosolic proteins. Proteolysis is expressed as the percentage of acid-insoluble radioactivity transformed into acid-soluble radioactivity at the end of the incubation. Values are the mean ± SEM of three independent experiments. (B) Lysosomes with high (CMA+) or low (CMA−) activity for CMA were isolated from livers of mice maintained on the RD or HFD. Intact lysosomes (Left) or broken lysosomes disrupted by hypotonic shock (Right) were incubated with a pool of radiolabeled proteins, and proteolysis was calculated as in A. Values are the mean ± SEM of three experiments.
Fig. S4. Effect of high lipid content diets on CMA of different cytosolic proteins. (A) Immunoblot for GAPDH of cytosol from livers of mice maintained on the RD, HFD, or CHOL. (Right) Quantification of GAPDH levels normalized by actin. (B) Immunoblot for GAPDH of homogenates and lysosomes isolated from animals on the RD or HFD. LAMP-1 is shown as a loading control. (C) Specific activity of GAPDH (activity corrected for protein levels) in the cytosol of livers from RD-, HFD-, or CHOL-maintained mice. Values are the mean ± SEM of three experiments. (D) Lysosomes from mice on the RD or HFD were treated or not treated with a protease inhibitor (PI) as indicated and incubated with RNase A for 20 min at 37 °C in isotonic medium. Lysosomes collected at the end of the incubation by centrifugation were subjected to SDS/PAGE and immunoblotting for RNase A. Uptake was calculated as the amount of RNase A associated with lysosomes treated with protease inhibitors (association) after discounting the amount associated with untreated lysosomes (binding) for each experiment. Values are the range of two independent experiments. *P < 0.05 compared with the RD group. INP, input.
Fig. S5. Effect of lipid challenges on CMA in young and old mice. Lysosomes from 3- and 22-mo-old mice maintained on the RD or 3-wk CHOL were treated or not treated with a protease inhibitor (PI) as indicated and incubated with GAPDH for 20 min at 37 °C in isotonic medium. Lysosomes collected at the end of the incubation by centrifugation were subjected to SDS/PAGE and immunoblotting for GAPDH. (A and B) Representative immunoblots. (C) Uptake was calculated as the amount of GAPDH associated with lysosomes treated with protease inhibitors (association) after discounting the amount associated with untreated lysosomes (binding) for each experiment. Values are the mean ± SEM of three different experiments. inp, input.

Fig. S6. Changes in the association of LAMP-2A with membrane microdomains in response to high-content lipid diets. Lysosomes from livers of fed mice or mice starved for 24 h maintained on the RD or CHOL were extracted with 1% Triton X-114 and then subjected to flotation in discontinuous sucrose density gradients. Four aliquots collected from the detergent-resistant (DR) to detergent-soluble (DS) region of the gradient were subjected immunoblotting for LAMP-2A (L2A) or dot blot analysis for GM1 using cholera toxin and an antibody against this toxin. (A) Representative immunoblots and immunodot blots. (B) Densitometric quantification of LAMP-2A (Left) and GM-1 (Right). Values are expressed as a percentage of the total lysosomal levels and are means of four independent experiments. *P < 0.05 compared with fed mice. STV, starved.
Fig. S7. Morphological and biochemical characterization of the group of lysosomes active for CMA. (A) Ultrastructure of lysosomes active for CMA (Left), lysosomes inactive for CMA (Center), and autophagolysosomes (Right). (Insets) Higher magnification images. (B) Immunoblot for the indicated proteins in fractions isolated from livers of mice starved for 24 h. Cyt, cytosol; ER, endoplasmic reticulum; Hom, homogenate; Lys, lysosomes with high (+) and low (−) CMA activity; Mito, mitochondria. (C) Immunoblot for mannose-6-phosphate receptor (M6PR) in homogenates (Hom) and CMA-active lysosomes (Lys+) isolated from animals maintained on the RD, CHOL, or HFD. (Scale bar: 500 nm.)
Fig. S8. Glycosphingolipid changes in the lysosomal membrane in response to dietary lipids and aging. Lysosomal membranes from livers of mice maintained on the RD, CHOL (A–D), or HFD (E–H) or from 3- and 22-mo-old mice (I–L) were subjected to comparative lipidomic analysis. Graphs show the percentage of saturated SM, unsaturated SM, CER, and glucoceramide (GC) relative to the total SM, CER, and GC in the samples. Values are the mean ± SEM from five different experiments. *P < 0.05 compared with the RD-maintained (A–H) or 3-mo-old (I–L) mice.
Fig. S9. PC changes in the lysosomal membrane in response to dietary lipids and aging. Lysosomal membranes from livers of mice maintained on the RD, CHOL (A–D), or HFD (E–H) or from 3- and 22-mo-old mice (I–L) were subjected comparative lipidomic analysis. Graphs show the percentage of saturated, unsaturated, and different acyl chain-length PC relative to the total PC in the samples. Values are the mean ± SEM from five different experiments. *P < 0.05 compared with RD-maintained (A–H) or 3-mo-old (I–L) mice.
Fig. S10. Changes induced by dietary lipids in the ubiquitin/proteasome system. (A) Immunoblot for the subunits of the 19S and 20S proteasomes in livers of mice maintained on the RD, CHOL, and HFD. (Lower) Densitometric quantification of three immunoblots as the one shown here. Values are expressed in arbitrary units (a.u.) and are corrected for actin. (B) Catalytic activities of the 20S proteasome measured with specific fluorescent substrates in the same livers. CTL, chymotrypsin-like; PGPH, peptidyl-glutamyl peptide hydrolyzing activity; TL, trypsin-like. (C) Immunoblot for ubiquitin in the same livers. (Lower) Densitometric quantification of three immunoblots as the one shown here. Values are expressed in arbitrary units and are corrected for actin. Poly-ub, poly-ubiquitin. Values are the mean ± SEM from three different experiments. *P < 0.05 compared with the RD.