Coordinated Regulation of the Orosomucoid-like Gene Family Expression Controls de Novo Ceramide Synthesis in Mammalian Cells*

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The orosomucoid-like (ORMDL) protein family is involved in the regulation of de novo sphingolipid synthesis, calcium homeostasis, and unfolded protein response. Single nucleotide polymorphisms (SNPs) that increase ORMDL3 expression have been associated with various immune/inflammatory diseases, although the pathophysiological mechanisms underlying this association are poorly understood. ORMDL proteins are claimed to be inhibitors of the serine palmitoyltransferase (SPT). However, it is not clear whether individual ORMDL expression levels have an impact on ceramide synthesis. The present study addressed the interaction with and regulation of SPT activity by ORMDLs to clarify their pathophysiological relevance. We have measured ceramide production in HEK293 cells incubated with palmitate as a direct substrate for SPT reaction. Our results showed that a coordinated overexpression of the three isoforms inhibits the enzyme completely, whereas individual ORMDLs are not as effective. Immunoprecipitation and fluorescence resonance energy transfer (FRET) studies showed that mammalian ORMDLs form oligomeric complexes that change conformation depending on cellular sphingolipid levels. Finally, using macrophages as a model, we demonstrate that mammalian cells modify ORM DL genes expression levels coordinately to regulate the de novo ceramide synthesis pathway. In conclusion, we have shown a physiological modulation of SPT activity by general ORMDL expression level regulation. Moreover, because single ORM DL3 protein alteration produces an incomplete inhibition of SPT activity, this work argues against the idea that ORM DL3 pathophysiology could be explained by a simple on/off mechanism on SPT activity.

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Orsomucoid-like (ORMDL) proteins belong to a family of transmembrane proteins that contains three members located in the endoplasmic reticulum (1). The association of ORM DL3 with childhood asthma in a genome-wide association study (2) stimulated renewed interest in the study of these proteins. Despite the absence of the single nucleotide polymorphism (SNP) identified in the genome-wide association study (rs 7216389) in a coding region of the gene, this first study correlated increased expression of ORM DL3 with the risk allele. Since then, several SNPs around the ORM DL3 gene that are associated with pathologies like inflammatory bowel disease, type I diabetes, and rheumatoid arthritis have been described (3–6).

The genome-wide association study approach to the diagnosis of genetic risk factors is not focused on precandidate genes, making it an excellent tool to identify new genes involved in diseases. However, sometimes the identified genes have ill defined functions as in the case of ORM DL3 at the time it was detected. To elucidate the pathophysiology associated with ORMDL3, several laboratories have been trying to understand the role of ORMDLs in cell physiology. Our laboratory has focused on the effects of ORMDL3 expression levels in calcium homeostasis, a likely connection between an endoplasmic reticulum-resident protein and immune system dysfunction. We have found that the expression levels of this protein are inversely correlated with the calcium content of the endoplasmic reticulum (1). The association of ORMDL3 with childhood asthma in a genome-wide association study (2) stimulated renewed interest in the study of these proteins. Despite the absence of the single nucleotide polymorphism (SNP) identified in the genome-wide association study (rs 7216389) in a coding region of the gene, this first study correlated increased expression of ORM DL3 with the risk allele. Since then, several SNPs around the ORM DL3 gene that are associated with pathologies like inflammatory bowel disease, type I diabetes, and rheumatoid arthritis have been described (3–6).

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(9, 10). In yeast and mammalian cells, the complete knockdown of ORMDLs releases serine palmitoyltransferase (SPT) activity and generates an increase in long chain bases and ceramides. The expression of any of the isoforms in this knockdown condition rescues the normal functioning of the pathway (10). This fact, together with the interaction between the yeast ORMDL isoforms (Orms) and the SPT enzyme, has established the idea that ORMDLs are the endogenous inhibitors of SPT. In addition, the SPT-Orm interaction is dependent on a phosphorylation reaction that disrupts an oligomeric complex of Orms and interferes with SPT-Orm interaction (10). The regulation of the pathway implicated in Orm phosphorylation and its sensitivity to ceramide cell content have been described in yeast (11). However, the role of this phosphorylation in SPT-ORMDL interaction is not clear in mammals because the N-terminal regulatory region described in yeasts is absent in mammalian ORMDLs. More remarkable is the lack of evidence that different ORMDL3 expression levels in mammalian cells alter SPT activity; this is a critical gap in understanding the pathophysiology associated with this gene.

We herein evaluate the role of mammalian ORMDLs in the ceramide synthesis context with three specific aims: (i) to explore the effect of ORMDL3 overexpression on SPT activity, (ii) to study the ORMDL-SPT complex interaction and its dependence on ceramide cell content, and (iii) to find a physiological context in which cells modify ORMDL expression levels to modulate SPT activity. For this purpose, we used HEK293 cells as the heterologous expression system and palmitate treatment to stimulate SPT activity. We performed coimmunoprecipitation studies between SPT-ORMDL complex elements and FRET studies to confirm and explore conformational changes. Moreover, we used the RAW264.7 monocytic cell line to study the regulation of ORMDL expression during de novo sphingolipid generation under the activation process.

**MATERIALS AND METHODS**

**Cell Culture and Transfection—**HEK293 and RAW264.7 cells were grown in DMEM (Sigma-Aldrich) supplemented with 10% heat-inactivated fetal calf serum, 100 units/ml penicillin, and 100 units/ml streptomycin. The cells were maintained in a 5% CO₂ environment at 37 °C. HEK293 cells were transiently transfected with the polycationic transfecting reagent polyethylenimine (PEI) (Polysciences), incubating cells with 6 eq of PEI/g of DNA for 5 h before changing to normal growing medium. Myriocin (10 μM) and C6-ceramide (10 μM) were both obtained from Sigma, and dimethyl sulfoxide (DMSO) was used as vehicle.

**Ceramide Quantification—**HEK293 cells were transfected with either pCDNA3 empty vector, pCDNA3-ORMDL1, pCDNA3-ORMDL2, pCDNA3-ORMDL3, pCDNA3-ORMDL3-D11–16, or the same final amount of all three ORMDL family members together. After 24 h of transfection, HEK293 cells were treated for 4 h with 500 μM palmitate (Sigma) complexed in 0.5% fatty acid-free BSA (Sigma), and BSA alone was used as a control. Cells were then washed twice with 1× PBS and centrifuged at 1800 rpm for 5 min at 4 °C, and the pellet was frozen in liquid nitrogen for ceramide quantification.

Lipid extraction and processing were performed as reported previously (12, 13). Lipid analysis was carried out by ultraperformance liquid chromatography coupled to time-of-flight (TOF) mass spectrometry in positive electrospray ionization mode. Instrument conditions were set as in previous studies (12, 13).

**In Vitro LPS Stimulation—**RAW264.7 cells were treated with 100 ng/ml *Escherichia coli* (055:B5) lipopolysaccharide (LPS) (Sigma-Aldrich) for the indicated time points. Total RNA or protein lysates were prepared from the cells and frozen at −80 °C until analysis. For ceramide quantification, RAW264.7 cells were washed twice in 1× PBS after LPS stimulation and centrifuged at 1800 rpm for 5 min at 4 °C, and the pellet was frozen in liquid nitrogen.

**Western Blot and Immunoprecipitation Assays—**HEK293 cells were transiently transfected with human ORMDL3-myc plus ORMDL1, ORMDL2, or ORMDL3 with different tags as indicated. In all conditions tested, cells were also transfected with a non-tagged ORMDL protein to ensure that all three proteins were present at the same ratio. After 24 h of transfection, cells were lysed with immunoprecipitation buffer (0.2% Triton X-100 plus protease inhibitor mixture in HEPES-buffered saline) and centrifuged at 100,000 × g to collect the total protein in supernatants. Then 1000 μg of total protein was incubated at 4 °C overnight with anti-myc antibody (Abcam) cross-linked with disuccinimidyl suberate (Pierce) to protein G-Sepharose beads. Immunocomplexes were washed with HEPES-buffered saline five times and eluted through a spin chromatography column with 0.2 M glycine, pH 2.5 before adding loading buffer and boiling for 5 min. Coimmunoprecipitation of ORMDL3-YFP, ORMDL1-YFP, or ORMDL2-YFP was detected by Western blot after separation by 4–12% gradient polyacrylamide gel electrophoresis and protein transfer to nitrocellulose membrane. Immunodetection was carried out using rabbit antibodies against ORMDL (1:300), SPTLC1 (1:300), and SPTLC2 (1:1000), all from Abcam. Secondary antibodies were horseradish peroxidase-conjugated anti-rabbit IgG (1:3000; GE Healthcare). The immunoreactive signal was detected by SuperSignal West Chemiluminescent substrate (Pierce) and visualized using the Molecular Imager Chemidoc XRS system (Bio-Rad).

In the case of RAW264.7 macrophages, after LPS stimulation, cells were washed with 1× PBS, lysed in 60 μl of lysis buffer (150 mM NaCl, 5 mM EDTA, 1% Triton X-100, 10 mM Tris-HCl, 1× Complete protease inhibitor) for 20 min on ice with agitation, and then centrifuged at 13,000 rpm at 4 °C for 30 min. Protein concentration in the supernatant was determined using the BCA Assay (Pierce). Equal amounts of protein (100 μg) were loaded into each lane and were separated on an SDS-polyacrylamide gel (12%).

**Quantitative Real Time PCR Analysis—**Total RNA of RAW264.7 cells was extracted using the Nucleospin RNA II kit (Macherey-Nagel) following the manufacturer’s instructions. Total RNA (1 μg) was reverse transcribed to cDNA using the SuperScript-RT system (Invitrogen). Quantitative RT-PCR was performed on an ABI Prism 7900HT (Applied Biosystems) with SYBR Green (SYBR Green Power PCR Master Mix, Applied Biosystems). Gene-specific mouse primers used were:
Coordinated ORMDL Expression Regulates SPT Activity

| Table 1: Ceramide content depending on ORMDL expression levels (pmol/1 × 10^6 cells) |
|----------------------------------|-----------|-----------|-----------|-----------|-----------|-----------|-----------|
| Control | ORMDL1 | ORMDL2 | ORMDL3 | ORMDL123 | siControl | si123 |
| C14 | 2.1 ± 0.3 | 2.8 ± 0.1 | 3.2 ± 0.5 | 3.2 ± 0.7 | 3.4 ± 0.5 | 0.8 ± 0.1 | 1.3 ± 0.0^a |
| C16 | 60.3 ± 9.0 | 58.4 ± 0.6 | 75.8 ± 14 | 75.5 ± 18 | 77.4 ± 7.6 | 14.9 ± 1.7 | 29.9 ± 2.4^a |
| C16:1 | 4.8 ± 0.8 | 5.3 ± 0.3 | 6.4 ± 1.6 | 6.5 ± 1.8 | 7.4 ± 0.9 | 1.4 ± 0.1 | 2.8 ± 0.2^a |
| C18 | 9.9 ± 1.9 | 6.7 ± 3.2 | 12.5 ± 2.5 | 10.4 ± 1.4 | 11.0 ± 0.3 | 2.7 ± 0.5 | 5.7 ± 0.1 |
| C20 | 3.8 ± 1.0 | 2.5 ± 0.1 | 3.9 ± 0.8 | 3.6 ± 0.2 | 3.0 ± 0.7 | 0.9 ± 0.2 | 2.3 ± 0.4 |
| C22 | 282.2 ± 4.9 | 260.0 ± 0.6 | 330.0 ± 7.4 | 276.2 ± 2.9 | 294.9 ± 0.9 | 3.7 ± 1.8 | 15.3 ± 1.5^a |
| C24 | 104.9 ± 16 | 98.4 ± 1.4 | 109.5 ± 22 | 89.1 ± 7.8 | 111.1 ± 6.8 | 25.0 ± 4.1 | 47.8 ± 3.3^b |
| Total | 214.1 ± 34 | 201.3 ± 3.8 | 244.3 ± 48 | 215.8 ± 33 | 242.8 ± 15 | 49.4 ± 4.8 | 105.7 ± 7.5^a |

*a p < 0.01.

mORMDL1, 5’-GATCCCTC TTGCGA GTTTT-3’ and 5’- CGGAGTCTC AAAAA CGGTT-3’; mORMDL2, 5’-GTCA- TCTACA AATGTCGCAAT-3’ and 5’-ACTGAGT CTCCAT- TCC-3’; mORMDL3, 5’-TGCCTGAGCATCCTCTT- TGT-3’ and 5’-CACGTTGTCGAGAAATGTT-3’; mβ-actin, 5’-TGGGATCC TCTGGCATCAGAAC-3’ and 5’-TAAA- ACGGCAGCTGAAACCTGG-3’; mSPTLC1, 5’-GTCCT- CTCCCTGAAACTGGTTA-3’ and 5’-CCCCATGTGCTG- GTGACT-3’; mSPTLC2, 5’-TGACAGCTCTGAGAGA- AAT-3’ and 5’-CAGGCAACCTTGGCAACAA-3’. PCR conditions in all cases were 95 °C for 5 min, 95 °C for 30 s, 60 °C for 30 s, 72 °C for 30 s, and 72 °C for 5 min with 40 cycles of amplification.

Fluorescence Resonance Energy Transfer Experiments—Proximity of the different tails of ORMDLs was assessed by FRET experiments as described previously (14). Cells were seeded on 25-mm glass coverslips and transfected using the indicated combinations. In addition to the YFP/CFP protein pair studied, 25-mm glass coverslips and transfected using the indicated

RESULTS
ORMDL Proteins Regulate de Novo Sphingolipid Synthesis—ORMDL proteins have been found to negatively regulate the activity of SPT, the enzyme that catalyzes the first rate-limiting step in de novo sphingolipid synthesis (10, 15). We used two different approaches, overexpression and silencing, to study how changes in ORMDL expression levels affect ceramide production and SPT activity. In overexpression experiments, monolayers of HEK293 cells were transiently transfected with either pCDNA3 empty vector (control), human pCDNA3-ORMDL1, -ORMDL2, -ORMDL3, or the combination of all three ORMDL proteins. For triple gene silencing, cells were treated with scrambled siRNA (control) and combined siRNA for ORMDL1, ORMDL2, and ORMDL3.

 Knocking down ORMDL1–3 significantly increased the cellular ceramide content compared with scrambled control siRNA (Table 1) in agreement with a previous report in HeLa cells (15). All ceramide species contributed to the total increase with C16, C18, C22, and C24 the most abundant. On the contrary, there was no alteration of specific or total ceramide amount after overexpressing the individual ORMDL isoforms separately or overexpressing all three ORMDL proteins simultaneously compared with control (Table 1).

We also evaluated the production of ceramides induced by addition of palmitate (PA), the substrate of the SPT enzyme. The increase of ceramide levels observed after 4-h incubation with 500 μM PA conjugated with BSA compared with control cells incubated with BSA alone was blocked in the presence of myriocin, a specific SPT inhibitor. This blockage demonstrated that the ceramide production using this protocol came mainly from the de novo pathway as it has been described previously (16). We normalized the value of the difference in ceramide content between cells treated with PA and control cells in the presence of myriocin or DMSO (Fig. 1B) and studied the contribution of the different ceramide species to this increase. After a 4-h incubation with PA, the newly formed ceramides contained mainly the C16 species, an early ceramide species from the de novo synthesis pathway (Fig. 1C). We then decided to explore, following the same protocol and normalization, the production of ceramides in cells knocked down for the three ORMDL isoforms with specific siRNAs (Fig. 1D). We observed a significant increase in total ceramide production, especially the C16 species, after a 4-h PA treatment in cells knocked down for ORMDLs compared with control cells with scrambled siRNA (Fig. 1, E and F). These results further demonstrated, together with the higher ceramide content in cells knocked down for the ORMDLs (Table 1), that the absence of ORMDL members promotes SPT activity. Conversely, cells overexpressing individual ORMDL isoforms did not show altered ceramide production under PA incubation (Fig. 1, G and H). A similar lack of effect was observed in cells expressing an ORMDL3 mutant lacking the first 16 amino acids of the N-terminal tail (ΔΝ1–16) that are essential for ORMDL3 calcium signaling function (8) (Fig. 1, G and H). Remarkably, only the overexpression of all three ORMDL proteins provoked a complete blockade in ceramide production after PA treatment in a manner similar to myriocin (Fig. 1, G and H). These results indicate that all three ORMDL isoforms are needed to efficiently block SPT activity. To date, there is no clear mechanistic explanation for the regulation of SPT by ORMDL proteins in mammals. Our observation that coordinated expression changes in all
ORMDL isoforms are required to modify SPT suggests the formation of a complex with close interaction among the three ORMDL members.

ORMDL Complex Formation and Rearrangement Depending on Lipid Environment—To evaluate possible interactions among ORMDL1, ORMDL2, and ORMDL3, immunoprecipitation assays were performed. HEK293 cells were transiently transfected with myc-ORMDL3 and either YFP-ORMDL1 or YFP-ORMDL2 to study hetero-oligomerization. For homo-oligomer formation studies of ORMDL3, cells were additionally transfected with YFP-ORMDL3. As a control, we used cells transfected with YFP-ORMDL3 in the absence of myc-ORMDL3. In all conditions tested, ORMDL isoforms without a tag were co-transfected at the same ratio to guarantee the presence of all members and to minimize artificial interactions due to the overexpression approach. Immunoprecipitation targeted the myc epitope in all cases, including the control condition. Coimmunoprecipitation was studied using an anti-ORMDL

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antibody, and the bands obtained migrated around 42 kDa, the expected size for the YFP-ORMDL constructs. As shown in Fig. 2A, mammalian ORMDL1 and ORMDL2 are able to directly interact with ORMDL3, forming heteromeric structures. These heteromeric structures showed the same extent of interaction (around 10%) as the homomeric structures of ORMDL3 as measured by band intensity analysis.

Regarding the formation of a complex between ORMDLs and SPT enzyme, we confirmed previous studies showing that ORMDL3 was able to interact with one subunit of the SPT complex, namely SPTLC1 (10). Moreover, we detected interaction between ORMDL3 and the catalytic subunit of the SPT complex, SPTLC2. The fraction of coimmunoprecipitation was around 4% with both subunits, demonstrating the existence of a functional ORMDL-SPT complex. Besides, we did not obtain coimmunoprecipitation with other endoplasmic reticulum membrane proteins such as STIM1 and calnexin (Fig. 2B). This result confirmed that the interactions we observed were not unspecific.

To further verify the observed interactions and extend the studies to all combinations of the three ORMDLs, the C terminus of ORMDL proteins was tagged with CFP/YFP FRET pairs. As in the previous experiments, the additional ORMDL isoforms were co-transfected at the same ratio without a tag to avoid artificial interactions. All tested combinations showed a positive FRET signal compared with the globular YFP control. Regarding the homomeric structures (ORMDL1-ORMDL1, ORMDL2-ORMDL2, and ORMDL3-ORMDL3), a similar FRET signal was obtained between the individual ORMDL isoforms (Fig. 2B). In the heteromerization studies, FRET signals for ORMDL1-ORMDL2 and ORMDL2-ORMDL3 were lower than those for ORMDL1-ORMDL3 (Fig. 2C). Considering that we observed no differences between homomeric and heteromeric interactions in the coimmunoprecipitation experiments,
a possible explanation for the lower FRET signals in the ORMDL1-ORMDL2 and ORMDL2-ORMDL3 combinations is that the final conformation places the C terminus of ORMDL2 more distant from ORMDL1 and ORMDL3.

Previous studies revealed that complex formation by yeast ORMDL orthologs is influenced by the amount of cellular sphingolipid composition. A decreased sphingolipid content not only reduces Orm oligomerization but also the Orm-Lcb1 binding (10). This release is supposed to allow for new sphingolipid generation. To explore the influence of sphingolipid content of mammalian ORMDL-SPTLC complex formation, we studied this process in both low and high sphingolipid content conditions. For a low sphingolipid environment, cells were treated with myriocin, a specific inhibitor of SPT; for a high sphingolipid environment, cells were incubated with C6-ceramide, a cell-permeable sphingolipid that has been shown to modulate SPT in an ORMDL-dependent manner (15). Our coimmunoprecipitation studies using a myc-ORMDL3 antibody showed that neither the ORM DL3-ORM DL3 interaction nor its binding to SPTLC1 or SPTLC2 was altered by changing the sphingolipidic environment (myriocin or C6-ceramide treatment) (Fig. 3A).

These data suggest that the interaction of ORMDL-SPT complexes is independent of the sphingolipidic environment. To evaluate putative conformational changes of ORMDL homo-oligomers due to the sphingolipid composition, FRET experiments were performed in both low and high sphingolipid content conditions. Interestingly, FRET efficiency was significantly reduced in a high sphingolipid environment (C6-ceramide treatment) for all three ORMDL isoforms compared with low sphingolipid content (myriocin treatment) (Fig. 3B). Taken together, these results indicate
the existence of a complex rearrangement between the ORMDL-SPTLC structures rather than a binding release mechanism.

**Role of ORMDLs in Macrophage Activation**—The importance of the ORMDL complex formation was further studied in cells of the innate immune response, a more physiological system. Previous studies have shown an increase in ceramide levels after activation of macrophages with bacterial LPS (16–19). In this process, the contribution of the de novo synthesis pathway at early time points is necessary for proper activation (19, 20), making it an interesting process for use in evaluating the role of the ORMDL complex in the regulation of ceramide levels. Therefore, RAW264.7 cells were treated with LPS, and the resulting changes in the sphingolipid content were monitored. Ceramide content increased steadily during the time of LPS activation, showing a peak after 8 h that decreases after long time activation (Fig. 4A). A similar trend was exhibited by sphinganine, another intermediate metabolite in the de novo ceramide synthesis pathway (Fig. 4B). These results agree with an early activation of the de novo pathway.

The endogenous expression of the ORMDLs after macrophage activation revealed a time-dependent regulation at both transcriptional and translational levels of all three ORMDL isoforms with a significant decrease at early time points (4 and 8 h) (Fig. 4C) followed by an increase at later times (24 h). To correlate the reduction of ORMDL expression with an increase in ceramide production, we compared the expression levels of ORMDLs with SPTLC2, the main SPT present in these cells. We observed a reduced ORMDL/SPTLC2 ratio during higher de novo ceramide production (Fig. 4D).

**DISCUSSION**

Complete absence of ORMDL protein expression in mammalian and yeast cells leads to an overproduction of ceramides through the de novo synthesis pathway (9, 10, 15). This fact, together with the positive coimmunoprecipitation of ORMDLs and SPT, supports the scenario in which ORMDLs are endogenous inhibitors of the first enzyme of de novo sphingolipid synthesis. Conversely, the rescue of normal growth on the Orm1/Orm2 double KO yeast strain by the expression of mammalian ORMDL3 isoform has generated the idea that all the ORMDL members have redundant functions and that this is a regulatory mechanism conserved throughout evolution (10). In addition to the different regulatory elements described in yeast, it has been shown that ORMDLs participate in the cascade that senses and inhibits SPT depending on the intracellular ceramide content (11, 15). However, the current knowledge regarding ORMDL function in controlling sphingolipid synthesis still has many blanks that hamper the understanding of the pathophysiological role of ORMDL3, a gene implicated as a risk factor in several diseases (2–6). The present study tries to cast light on the relationship between mammalian ORMDLs and SPT.

Until now it was not known whether single variations in ORMDL3 expression could cause changes in SPT function and cellular sphingolipid content. This information is important to understand whether SNPs that exert a cis regulation on ORMDL3 gene expression, claimed to be risk factors of several pathologies (2, 21), could be linked to the function of this protein in sphingolipid synthesis. Our results show that increased
expression levels of each different ORMDL or the three isoforms together do not affect the ceramide content of the cell. Interestingly, triple overexpression blocks SPT function when stimulating the de novo ceramide synthesis with palmitate. These findings can only be explained by additional regulatory mechanisms besides changes in expression. Thus, ORMDLs would not be allowed to lower the homeostatic content of ceramides in the cell, but their presence suffices to block SPT when ceramides increase to putatively deleterious levels. This effect is observed only in triple overexpression, meaning that all three members are needed for an effective sensing and blockage of the de novo pathway. Nevertheless, our results on the effect of ORMDL expression on SPT activity have to be cautiously interpreted as they may be partially dependent on the experimental settings, i.e. cell type and palmitate treatment. In this sense, we have modified exclusively ORMDL expression levels to mimic, in our in vitro model, as much as possible the pathophysiological context related to the diseases in which ORMDL3 has been implicated. Genetic analysis of ORMDL3-related inflammatory diseases only pointed to changes in ORMDL3 expression without information about possible changes in other elements participating in the SPT-ORMDL complex. In this context and based on the fact that overexpression of ORMDL3 alone did not modify ceramide content and palmitate-induced sphingolipid synthesis, our results do not support a pathophysiological mechanism based on the complete SPT inhibition. Besides, deletion of the first 16 amino acids of the ORMDL3 N-terminal tail has been shown to abolish its impact on calcium signaling (8). However, the lack of differences regarding ceramide content and production observed in the present work for overexpression of this mutant and wild type ORMDL3 argues against an explanation for the effect of ORMDL3 on calcium homeostasis based on its role in sphingolipid synthesis.

Another aspect we have addressed in this work is the dynamic regulation of ORMDL-SPTLC complexes that has been proposed in light of the experiments performed in yeasts (10). Our results demonstrate that the regulation based on interaction of the yeast ORMDL isoforms is lost in mammalian ORMDLs. The immunoprecipitation experiments did not support the occurrence of changes in the ORMDL-ORMDL or ORMDL-SPTLC interaction when the ceramide content of the cell was manipulated. Interestingly, another regulatory mechanism based on ORMDL phosphorylation via the TORC1 pathway that does not modify the interaction of ORMDL oligomers necessary for complex sphingolipid formation has been found in yeast (22). Whether this modulation is similar to the conformational changes observed in our experiments is a question that requires further investigation. It is also important to note that the regulation described in yeast is based on the phosphorylation of the N-terminal part of yeast Orms (10, 11). This domain is only present in yeast and plants, whereas vertebrate proteins cluster differently (1). Regarding the stoichiometry of the complex, we propose that ORMDLs are able to form homo- and hetero-oligomers with similar affinity because in our experiments we always overexpressed all the ORMDL members to avoid artificial interactions. Hetero-oligomerization likely results in a slightly different conformation based on the different FRET observed. Finally, our results also suggest that ORMDL regulation of SPT might be achieved by structural rearrangements depending on the intracellular ceramide content that do not involve changes in ORMDL-SPTLC binding affinity.

Finally, we demonstrate for the first time using physiological stimuli that mammalian cells modulate ceramide content by modifying expression levels of the ORMDL family. We have focused on macrophages where de novo ceramide synthesis has been characterized during LPS activation (16–18). The increase in ceramides through the SPT pathway is necessary for correct activation, and its blockage affects several processes such as autophagy and interleukin production (19, 20). In this context, we show that expression of the three ORMDL members is repressed at early time points in correlation with ceramide synthesis induction. The reduction in the ORMDL/SPTLC ratio suggests a regulatory scenario in which ceramide synthesis is induced by reducing the expression of the inhibitory subunits of the SPT complex. These results agree with the heterologous expression analysis where the triple knockdown of ORMDLs is necessary to release SPT activity (15).

Our work reinforces the idea that ORMDLs are negative modulators of SPT activity, providing physiological evidence for this role. However, increased individual expression of these proteins does not reduce the ceramide content in the cell, which conflicts with the idea that the pathophysiological link between disease-associated SNPs that change ORMDL3 expression is related to changes in cellular sphingolipid content due to a complete inhibition of the de novo pathway. This also implies that alternative mechanisms should be considered in determining the role of ORMDLs in immune/inflammatory diseases, i.e. the control of calcium homeostasis (7, 8) and the unfolded protein response (23, 24).

REFERENCES


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Coordinated ORMDL Expression Regulates SPT Activity


Lipids: Coordinated Regulation of the Orosomucoid-like Gene Family Expression Controls de Novo Ceramide Synthesis in Mammalian Cells

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