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ORMDL3 Transgenic Mice Have Increased Airway Remodeling and Airway Responsiveness Characteristic of Asthma

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Orosomucoid-like (ORMDL)3 is a gene localized to chromosome 17q21, which was initially linked to asthma in a genome-wide association study (1) with subsequent confirmation in multiple additional genome-wide association studies (2–4) and non– genome-wide association study genetic association studies in populations of diverse ethnic backgrounds (5–10). ORMDL3 has been linked to severe asthma (4, 9), childhood onset of asthma (1, 7, 8), exposure of children to environmental tobacco smoke, and risk of asthma (2, 10), as well as to rhinoviral wheezing illness and genetic risk of childhood onset of asthma (11), underscoring the importance of understanding its function. ORMDL3 is a member of the three member ORMDL gene family (ORMDL1, -2, -3), which encodes transmembrane proteins located at the endoplasmic reticulum (ER) (12). ORMDL1 (chromosome 2) (12) and ORMDL2 (chromosome 12) (12) are on different chromosomes from ORMDL3 (chromosome 17q21) (12) and have not been linked to asthma. Both humans and mice express the same three ORMDL family members, with ORMDL3 exhibiting 96% identity between these two species (12). ORMDL3 is a 153-aa protein with two predicted transmembrane domains (12). We recently demonstrated that in wild-type (WT) mice, ORMDL3 is an allergen and Th2 cytokine (IL-4 or IL-13) inducible gene localized to the ER and highly expressed in airway epithelial cells (13). Allergen challenge induced a 127-fold increase in ORMDL3 mRNA in bronchial epithelium in WT mice, with lesser 15-fold increases in ORMDL2 and no changes in ORMDL1 (13). We also demonstrated that transfection of ORMDL3 in human bronchial epithelial cells in vitro induced expression of CC chemokines (CCL20, also known as MIP-3α) (13), CXC chemokines (IL-8; CXCL10, also known as IFN-γ-induced protein 10 [IP-10]; CXCL11, also known as IFN-inducible T cell α [ITAC]) (13), metalloproteases (matrix metalloproteinase [MMP]-9; a disintegrin and metalloproteinase domain–containing protein 8 [ADAM8]) (13), and selectively activated transcription factor 6 (ATF6) (13), one of three ER unfolded protein response (UPR) pathway transcription factors (14) with subsequent regulation of sarco/endoplasmic reticulum Ca2+-ATPase (SERCA2b), which has been implicated in airway remodeling in asthma (15). Thus, these studies with bronchial epithelium in WT mice and in normal human bronchial epithelial cells suggest an important role for a pathway in which initial induction of ORMDL3 with subsequent activation of both ATF6-dependent pathways (i.e., SERCA2b) and/or ATF6-independent pathways.

ORSOMUCOID-LIKE (ORMDL)3; WT, wild-type.

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(MMP-9, ADAM8, CCL20, CXCL10, CXCL11) may contribute to the pathogenesis of asthma.

Although our previous studies demonstrated that Ormd3 is an allergen and Th2 cytokine–inducible gene that is dependent upon Stat6 for expression (13), these prior studies in WT mice did not determine which downstream pathways were regulated by ORMDL3 in vivo. To address this question we have generated ORMDL3 transgenic (Tg) mice, and in this study we demonstrate that Tg mice overexpressing human ORMDL3 (hORMDL3) spontaneously develop significantly increased levels of airway remodeling (smooth muscle, fibrosis, mucus) that precede the development of airway inflammation. Additionally, allergen challenge of ORMDL3 Tg mice resulted in enhanced OVA-specific IgE responses compared to OVA-challenged WT mice and was associated with increased major basic protein (MBP) positive peri-bronchial eosinophils and lung levels of IL-4. These studies in ORMDL3 Tg mice also provide evidence that the ER-localized ORMDL3 plays an important role in selective activation of one of the three UPR pathways in vivo (i.e., ATF6), and that expression of ORMDL3 in vivo regulates airway remodeling (smooth muscle, fibrosis, mucus) potentially through ATF6-independent genes (TGF-β1, ADAM8), which we detected at increased levels in the lungs of ORMDL3 Tg mice. ORMDL3 may therefore activate several pathways important to the pathogenesis of airway remodeling and asthma in vivo.

**Materials and Methods**

**Zp3-Cre mice**
Zp3-Cre mice (embryonic Cre expression) on a C57BL/6 background were acquired from The Jackson Laboratory.

**hORMDL3Zp3-Cre mouse generation**

All the mouse experimental protocols were approved by the University of California, San Diego Institutional Animal Care and Use Committee.

**Targeting plasmid construction.** The hORMDL3 Tg construct pCAGEN Lox red fluorescent protein (RFP)-H2B STOP Lox hORMDL3 was generated by cloning the 462-bp hORMDL3 open reading frame from pCMV6-AC-ORMDL3 (Origene) with AgeI/NotI into a construct previously developed and provided by A.J. Holland and D.W. Cleveland (Ludwig Institute for Cancer Research at the University of California, San Diego). RFP-StopZp3-LoxP-ORMLD3-Tg mouse generation. Spl/Pvul-linearized pCAGEN Lox RFP-H2B STOP Lox hORMDL3 (Fig. 1A, 1B) was microinjected into the pronuclei of fertilized mouse embryos and implanted into a pseudopregnant mouse (all on a C57BL/6 background) via an established protocol by the University of California, San Diego mouse Tg core. Progeny were then screened by PCR for the presence of the transgene. Subsequent mouse genotyping was performed using PCR with the following primers: F1-3530, 5'-GCA AGC TGC TGG TTA TGT TG-3'; F2-4009, 5'-CCC CCT GAA CCT GAA ACA TA-3'; R-4644, 5'-TAC AGC ACG ATG GGT GTG AT-3' (Fig. 1C). These RFP-StopZp3-hORMDL3-Tg mice (C57BL/6 background) were crossed with Zp3-Cre mice (C57BL/6 background) to generate hORMDL3Zp3-Cre mice (C57BL/6 background).

**Processing of lungs, bronchoalveolar lavage, and blood**

hORMDL3Zp3-Cre mice and littermate controls were euthanized at different ages (4, 8, and 26 wk) to quantitate levels of airway inflammation, airway remodeling, as well as expression of cytokines, chemokines, and remodeling genes. In addition to examining whole lung, purified populations of selected lung cell types (bronchial epithelium, bronchoalveolar lavage [BAL] macrophages) were also studied. Levels of IgE and other Igs were quantitated in peripheral blood.

**Lung processing.** Lungs were processed for protein and RNA extraction, as well as for immunohistochemistry (paraffin-embedded lung sections) as previously described in this laboratory (13, 16). For protein and RNA extractions, lungs were initially snap-frozen in liquid nitrogen and stored at −80 °C. For paraffin-embedded sections, lungs were equivalently inflated with an intratracheal injection of the same volume of 4% paraformaldehyde solution (Sigma-Aldrich, St. Louis, MO) to preserve the pulmonary architecture. Lung sections were processed for immunohistochemistry to detect MBP (anti-mouse MBP Ab was provided by Dr. James Lee, Mayo Clinic, Scottsdale, AZ), neutrophil elastase (anti-mouse neutrophil elastase Ab; Santa Cruz Biotechnology), F4/80 (anti-mouse F4/80 Ab; Santa Cruz Biotechnology), and CD4 (anti-mouse CD4 Ab; GeneTex). The number of individual cells staining positive for different cell types in the peribronchial space was counted using a light microscope. Results are expressed as the number of peribronchial cells staining positive per bronchiol with 150–200 μm internal diameter. At least 10 bronchiol were counted in each slide.

**BAL macrophages.** In selected experiments, purified populations of BAL macrophages (>98% purity) were obtained by adhesion by placing BAL cells in a 10-cm petri dish in complete media for 4 h at 37 °C as previously described in this laboratory (13). Pooled BAL macrophages from four mice per group were used for RNA and protein extraction.

**Isolation of bronchial epithelial cells.** The isolation of bronchial epithelial cells was performed as previously described in this laboratory (13, 17). Briefly, the epithelial brushing was performed using a sterile plastic feeding tube (Solomon Scientific) inserted into the right main and left main bronchi with gentle brushing and immediately placed in RNA STAT-60 (Tel-Test) for RNA extraction. Bronchial epithelial cells were of >95% purity as assessed by E-cadherin expression on FACS and histologic detection of ciliated epithelial cells (13, 17). Four mice per group were used for each data point.

**BAL fluid collection.** BAL fluid was collected by lavaging the lung with 1 ml PBS via a tracheal catheter as previously described (16). BAL fluid was centrifuged, and the supernatant frozen at −80 °C for subsequent cytokine analysis.

**Peripheral blood.** Peripheral blood was obtained from mice by cardiac puncture into tubes without anticoagulant added for quantitation of serum Ig levels.

**Detection of ORMDL3- and ORMDL3-regulated genes by quantitative RT-PCR**

Quantitative RT-PCR was performed as previously described in this laboratory (13). In brief, total RNA was extracted with RNA STAT-60 (Tel-Test) and reverse transcribed with oligo(dT) and SuperScript II kit (Life Technologies). Quantitative PCR was performed with TaqMan PCR Master Mix and ORMDL1, ORMDL2, ORMDL3 (human and mouse), SERCA2b, TGF-β1, ADAM8, MMP-9, ITAC, and IP-10 primers (all from Applied Biosystems). The relative amounts of transcripts were normalized to those of housekeeping gene (GAPDH) mRNA and compared between the different genes by the ΔΔCt method as previously described in this laboratory (13).

**Detection of airway remodeling**
Peribronchial smooth muscle layer. The thickness of the airway smooth muscle layer was measured by α-smooth muscle actin immunohistochemistry as previously described (16). Lung sections were immunostained with an anti-α-smooth muscle actin primary Ab (Sigma-Aldrich) to detect peribronchial smooth muscle. Species- and isotype-matched Abs were used as controls in place of the primary Ab. The area of peribronchial α-smooth muscle actin staining in paraffin-embedded lungs was outlined and quantified under a light microscope (Leica DMLS, Leica Microsystems) attached to an image analysis system (Image-Pro Plus, Media Cybernetics, Bethesda, MD) as previously described (16). Results are expressed as the area of peribronchial α-smooth muscle actin staining per micrometer length of basement membrane of bronchioles 150–200 μm internal diameter.

**Peribronchial trichrome staining.** The area of peribronchial trichrome staining in paraffin-embedded lungs was outlined and quantified under a light microscope (Leica DMLS, Leica Microsystems) attached to an image analysis system (Image-Pro Plus, Media Cybernetics) as previously described (16). Results are expressed as the area of trichrome staining per micrometer length of basement membrane of bronchioles 150–200 μm internal diameter.

**Lung collagen.** The amount of lung collagen was measured as previously described in this laboratory (16) with a collagen assay kit that uses a dye reagent that selectively binds to the [Gly-X-Y]n tripeptide sequence of mammalian collagens (BioColor, Newtonnabey, U.K.). In all experiments, a collagen standard was used to calibrate the assay.

**Airway mucus expression.** To quantitate the level of mucus expression in the airway, the number of periodic acid–Schiff (PAS)+ and PAS+ epithelial cells in individual bronchioles was counted as previously described in this
FIGURE 1. Generation of ORMDL3 Tg mice. (A) The human ORMDL3 Tg construct contains a CAG promoter for universal overexpression (light blue), H2B-RFP (RFP) (red), followed by a transcriptional stop site (orange), the human ORMDL3 open reading frame (green), and (Figure legend continues)
laboratory (16). At least 10 bronchioles were counted in each slide. Results are expressed as the percentage of PAS+ cells per bronchiole, which is calculated from the number of PAS+ epithelial cells per bronchus divided by the total number of epithelial cells of each bronchiole.

**Lung protein for ELISA and Western blot**

Lung tissue was homogenized in 500 μl lysis buffer and 100 mg stainless steel beads (Next Advance) for 5 min using the Bullet Blender homogenizer (Next Advance). The lysate was centrifuged at 13,000 rpm for 10 min at 4 °C. The supernatant was used for ELISA and Western blot.

**Quantification of lung cytokines**

BAL fluid and lung levels of selected Th2 cytokines (IL-4, IL-5, IL-13) and chemokines (eotaxin-1) was performed by ELISA according to the manufacturer’s instructions (R&D Systems). Levels of lung IL-4, IL-5, IL-13, and eotaxin-1 are expressed as the amount of cytokine/chemokine in picograms per milligram lung protein. Lung protein levels were quantitated by the bicinchoninic acid method (Thermo Scientific).

**Detection of SERCA2b by Western blot**

Lung protein was separated on a SDS-PAGE gel and transferred to a polyvinylidene difluoride membrane. Membranes were blocked in 5% (w/v) milk in 1× Tris-buffered saline with Tween 20 for 1 h and then incubated with the primary Ab overnight at 4°C. The primary Abs used in this study were mouse monoclonal anti-SERCA2b (Abcam) and rabbit monoclonal anti-GAPDH (Genetex).

**Quantitation of bronchial epithelial TGF-β1, ADAM8, SERCA2b and MMP-9 by image analysis**

Lung sections from hORMDL3zp3-Cre and WT mice were immunostained with either anti–TGF-β1, anti-ADAM8, SERCA2b, or anti–MMP-9 Abs. Levels of expression of either TGF-β1, ADAM8, SERCA2b, or MMP-9 by bronchial epithelial cells (outlined and visualized with a light microscope attached to an image analysis system) were quantitated by image analysis as described (18). The images were saved and analyzed with Image-Pro Plus 3 software (Media Cybernetics). The mean value of bronchial epithelial immunostaining intensity was normalized for the total area of bronchial epithelium.

**Quantitation of serum IgE, IgG, IgM, and IgA**

Serum IgE (IgGl1 and IgGl2a), IgM, and IgA were quantitated using an IgG ELISA kit (BD Biosciences). Serum IgE, IgM, and IgA were quantitated with a mouse IgG isotyping ELISA kit (BD Pharmingen) with results reported as OD at 450 nm per the manufacturer’s instructions. Serum OVA-specific IgE was quantitated with a mouse OVA-specific IgG kit (Biologend) with results reported as nanograms per milliliter. All ELISA plates were read with a Bio-Rad model 680 microplate reader.

**Activation of ATF6, Ire1, and PERK**

Purified populations of macrophages (>95% pure) were generated from hORMDL3zp3-Cre and WT mouse bone marrow cells cultured in complete DMEM media (Gibco) supplemented with stem cell growth factors (L Cells media, American Type Culture Collection) for 10 d as described (19). Activation of ATF6 was detected with immunofluorescence microscopy to detect nuclear localization of ATF6 using an ATF6 Ab (Regnaxx) as previously described (13, 20). Activation of ATF6, Ire1, and PERK was assessed by increased levels of phospho-eIF2α by Western blot using an Ab specific to the phosphorylated form of eIF2α (13, 20). In all UPR experiments, thapsigargin, a known activator of the UPR, was incubated with cells for 1 h before collecting cells for either RNA or protein analyses. Because insufficient numbers of bronchial epithelial cells were available for UPR Western blot studies, the UPR studies were performed on macrophages, a cell type that similar to epithelial cells expresses high levels of ORMDL3 (13).

**Acute OVA challenge model**

hORMDL3zp3-Cre and littermate control mice aged 12 wk were sensitized and challenged with OVA (Worthington, Lakewood, NJ) as previously described (13). In brief, mice were sensitized i.p. with 100 μg OVA and 2 mg aluminum hydroxide (Injekt Alum; Thermo Fisher Scientific, Waltham, MA) in a total volume of 200 μl PBS on days 0 and 10 and followed by intranasal administration of 200 μg OVA in 20 μl PBS on days 21, 23, and 25. Non-OVA-challenged ORMDL3 Tg and littermate groups of mice were sensitized and challenged with PBS only. Twenty-four hours after the last challenge, BAL fluid, lungs, and blood were collected as described above.

**Airway hyperreactivity to methacholine**

Airway responsiveness to methacholine was assessed in intubated and ventilated mice aged 12 wk (n = 8 mice/group) with the flexiVent ventilator (Scireq) anesthetized with ketamine (100 mg/kg) and xylazine (10 mg/kg) i.p. as previously described (16). The dynamic airway resistance and elastance were determined using Scireq software in mice exposed to nebulized PBS and methacholine (0, 3, 24, and 48 mg/ml). The following ventilator settings were used: tidal volume (10 ml/kg), frequency (150/ min), and positive end-expiratory pressure (3 cmH2O). Increased elastance values signal an increased stiffness of the lungs, and elastance is the inverse of compliance (Scireq).

**Statistical analysis**

All results are presented as means ± SEM. A statistical software package (GraphPad Prism; GraphPad Software, San Diego, CA) was used for the analysis. A p value <0.05 was considered statistically significant.

**Results**

**Generation of hORMDL3 Tg mice**

To perform studies in hORMDL3 Tg mice, we generated conditional hORMDL3 Tg floxed mice (RFP-Stop3lox ORMDL3-Tg mice) with the pCAGEN Iox RFP-H2B STOP lox hORMDL3 transgene construct (Fig. 1A, 1B), and crossed these with zona pellucida 3 (zp3) Cre mice, resulting in offspring with expression of the hORMDL3 transgene which prevents transcription of hORMDL3 (Fig. 1A). Thus, all cells in this RFP-StopFL hORMDL3-Tg mouse will not express hORMDL3 until crossed with a Cre-expressing mouse, which excises the transcriptional stop codon and protein kinase regulated by roman typefaces in red.
FIGURE 2. Airway remodeling in hORMDL3zp3-Cre mice. (A–C) Levels of peribronchial smooth muscle were quantitated by immunohistochemistry using an anti-α-smooth muscle actin Ab and image analysis. Results are expressed as the α-smooth muscle actin stained area (square micrometers) per micrometer length of basement membrane of bronchioles 150–200 μm internal diameter in (A) WT and (B, C) hORMDL3zp3-Cre mice at 4, 8, and 26 wk of age (n = 12 mice/group). (D–F) Levels of peribronchial trichrome staining were quantitated by image analysis. Results are expressed as the trichrome-stained area (square micrometers) per micrometer length of basement membrane of bronchioles 150–200 μm internal diameter in (D) WT and (E, F) hORMDL3zp3-Cre mice 4, 8, and 26 wk of age (n = 12 mice/group). (G) Levels of lung collagen were quantitated by a Sircol assay in WT and hORMDL3zp3-Cre mice at 4, 8, and 26 wk of age (n = 12 mice/group). (H–L) Levels of mucus were quantitated by PAS staining in (H) WT and (I, J) hORMDL3zp3-Cre mice at 4, 8, and 26 wk of age (n = 12 mice/group). Higher magnification view of PAS staining in (K) WT and (L) hORMDL3zp3-Cre mice. (A, B, D, E, H, and I) Original magnification ×200. (K and L) Original magnification ×400.
fluorescence prior to crossing to zp3 Cre mice. Crossing RFP-Stop\textsuperscript{FL}-hORMDL3-Tg mice to zp3 Cre mice results in the loss of RFP expression in cells that were designed to provide a rapid initial screen for successful expression of hORMDL3 (Fig. 1A, 1B). However, because levels of the RFP as detected by immunofluorescence microscopy varied widely in RFP-Stop\textsuperscript{FL}-hORMDL3-Tg mice, we utilized hORMDL3 PCR, rather than loss of red fluorescence, to detect hORMDL3 expression. hORMDL3\textsuperscript{zp3-Cre} mice overexpressing hORMDL3 constitutively in all cells were viable with no obvious developmental or morphologic defects (Fig. 1D), and their lung size and weights were similar to those of WT mice (Fig. 1E, 1F).

Levels of the human ORMDL3 transgene were highly expressed in hORMDL3\textsuperscript{zp3-Cre} mouse lung (Fig. 1G), bronchial epithelium (Fig. 1H), and BAL macrophages (Fig. 1I) as assessed by quantitative RT-PCR. In contrast, no expression of the human ORMDL3 transgene was detected in WT littermate mice (referred to as WT subsequently) (Fig. 1G–I). Levels of mouse ORMDL1, mouse ORMDL2, and mouse ORMDL3 in hORMDL3\textsuperscript{zp3-Cre} mice were not altered in mouse lung (Fig. 1J), bronchial epithelium (Fig. 1K), and BAL macrophages (Fig. 1L) as assessed by quantitative RT-PCR.

**Increased airway remodeling in hORMDL3\textsuperscript{zp3-Cre} mice**

hORMDL3\textsuperscript{zp3-Cre} mice have evidence of airway remodeling characteristic of asthma at 4 wk of age, which persisted through 26 wk of life (Fig. 2). Features of airway remodeling that are evident in hORMDL3\textsuperscript{zp3-Cre} mice include an increase in the area of peribronchial smooth muscle as assessed by immunostaining with an anti-α-smooth muscle actin Ab (Fig. 2A–C), an increase in peribronchial fibrosis as assessed by the area of peribronchial trichrome staining to detect lung collagen (Fig. 2D–F), and an increase in total lung collagen (Fig. 2G). Additionally, there was a significant increase in mucus expression detected by PAS staining (Fig. 2H–L). Although the levels of peribronchial smooth muscle (Fig. 2C) and peribronchial fibrosis as assessed by trichrome staining (Fig. 2F) remained stably increased from week 4 to week 26, levels of mucus continued to increase from week 4 to week 26 (Fig. 2J).

**Activation of ATF6 but not Ire1 or PERK in hORMDL3\textsuperscript{zp3-Cre} mice**

We have previously demonstrated that in vitro transfection of ORMDL3 activates only one of the three pathways of the UPR (i.e., activates the ATF6 pathway and not the Ire1 or PERK pathway) (13). To determine whether in vivo hORMDL3 activated the UPR, we cultured bone marrow–derived macrophages from WT and hORMDL3\textsuperscript{zp3-Cre} mice to have sufficient numbers of cells to perform a Western blot. hORMDL3\textsuperscript{zp3-Cre} mouse macrophages, but not WT macrophages, spontaneously activated the ATF6 pathway as assessed by translocation of ATF6 from being dispersed in the ER to the nucleus using immunofluorescence microscopy (Fig. 3A, 3B). In contrast, both WT and hORMDL3\textsuperscript{zp3-Cre} mouse macrophages did not activate either Ire1 (Fig. 3C) or PERK (Fig. 3D) pathways. Incubation of WT and hORMDL3\textsuperscript{zp3-Cre} mouse macrophages with thapsigargin, a known activator of the UPR, induced activation of the ATF6 (Fig. 3A, 3B), Ire1 (Fig. 3C), and PERK (Fig. 3D) pathways in both WT and hORMDL3\textsuperscript{zp3-Cre} mouse macrophages. Thus, bone marrow–derived macrophages...
from hORMDL3^{p3-Cre} mice, similar to cells transfected with ORMDL3 in vitro (13), exhibit selective activation of the ATF6 UPR pathway.

**Increased remodeling genes in lungs of hORMDL3^{p3-Cre} mice**

Because SERCA2b has been implicated in airway remodeling in asthma (15), we examined whether levels of SERCA2b were modulated in hORMDL3^{p3-Cre} mice. Our studies demonstrate that hORMDL3^{p3-Cre} mice have increased lung levels of SERCA2b as assessed by quantitative RT-PCR (Fig. 4A) and Western blot (Fig. 4B). Additionally, hORMDL3^{p3-Cre} mice have increased levels of bronchial epithelial expression of TGF-β1 (Fig. 4C) and ADAM8 (Fig. 4D), with a smaller increase in MMP-9 (Fig. 4E) as assessed by quantitative RT-PCR. Levels of TGF-β1 (Fig. 4F), ADAM8 (Fig. 4G), and SERCA2b (Fig. 4H), but not MMP-9 (data not shown), were increased in bronchial epithelium in

![Graphs and images showing quantitative analysis of SERCA2b, TGF-β1, ADAM8, MMP-9, CXCL10, and CXCL11 in hORMDL3^{p3-Cre} and WT mice.](http://www.jimmunol.org/)

**FIGURE 4.** hORMDL3^{p3-Cre} mice express SERCA2b, TGF-β1, ADAM8, MMP-9, CXCL10, and CXCL11. Levels of SERCA2b were quantitated in the lungs of hORMDL3^{p3-Cre} and WT mice by (A) quantitative PCR and (B) Western blot. Levels of (C) TGF-β1, (D) ADAM8, and (E) MMP-9 were quantitated by quantitative PCR in the lungs, bronchial epithelium, and BAL macrophages of hORMDL3^{p3-Cre} and WT mice. Levels of (F) TGF-β1, (G) ADAM8, and (H) SERCA2b were quantitated by immunohistochemistry and image analysis in bronchial epithelium of hORMDL3^{p3-Cre} and WT mice. Levels of (I) CXCL10 (IP-10) and (J) CXCL11 (ITAC) were quantitated by quantitative PCR in the lungs, bronchial epithelium, and BAL macrophages of hORMDL3^{p3-Cre} and WT mice (n = 12 mice/group).
hORMDL3<sup>2pr3-Cre</sup> mice as assessed by immunohistochemistry with quantitation by image analysis.

**Levels of selected CXC and CC chemokines in hORMDL3<sup>2pr3-Cre</sup> mice**

Because we had previously demonstrated that transfection of ORM DL3 in vitro induced high levels of expression of CXC chemokines (IL-8; CXCL10, also known as IP-10; CXCL11, also known as ITAC) (13) and lower levels of CC chemokines (CCL20, also known as MIP-3α) (13), we measured levels of these chemokines by quantitative PCR in the hORMDL3<sup>2pr3-Cre</sup> mice. hORMDL3<sup>2pr3-Cre</sup> mice had significantly increased levels of lung CXC chemokine mRNA, including CXCL10 in bronchial epithelial cells (Fig. 4I), as well as increased levels of CXCL11 mRNA in BAL macrophages (Fig. 4J), whereas levels of lung KC mRNA, the murine equivalent of IL-8, was not increased (data not shown).

Levels of selected CC chemokine mRNA (CCL20 and eotaxin-1) were also not different in lung, bronchial epithelium, or BAL macrophages in hORMDL3<sup>2pr3-Cre</sup> compared to WT mice (data not shown).

**Airway remodeling preceded airway inflammation in hORMDL3<sup>2pr3-Cre</sup> mice**

hORMDL3<sup>2pr3-Cre</sup> mice exhibit significant airway remodeling at 4 wk of age (Fig. 2), a time point at which there is no evidence of any increase in peribronchial CD4<sup>+</sup> cells (Fig. 5A), F4/80<sup>+</sup> macrophages (Fig. 5B), MBP<sup>+</sup> eosinophils (Fig. 5C), or neutrophil elastase<sup>+</sup> neutrophils (Fig. 5D) compared to WT mice. At 8 wk of age there is a small increase in F4/80<sup>+</sup> peribronchial macrophages (Fig. 5B) in the lungs of hORMDL3<sup>2pr3-Cre</sup> mice, with no change in CD4<sup>+</sup> cells, eosinophils, or neutrophils (Fig. 5A, 5C, 5D). At 26 wk of age there is a significant increase in peribronchial CD4<sup>+</sup> cells (Fig. 5A),
F4/80+ macrophages (Fig. 5B), eosinophils (Fig. 5C), and neutrophils (Fig. 5D) in hORMDL3zp3-Cre compared to WT mice.

Airway remodeling precedes increase in Th2 cytokines in hORMDL3zp3-Cre mice

Levels of Th2 cytokines IL-5 and IL-13 were not increased in either BAL or lung of hORMDL3zp3-Cre mice at 4 wk of age (Fig. 5E–H), a time point at which hORMDL3zp3-Cre mice exhibit significant airway remodeling. Levels of BAL IL-13 (Fig. 5E) and lung IL-13 (Fig. 5F) were significantly increased in hORMDL3zp3-Cre mice at 8 and 26 wk of age as assessed by ELISA. Levels of BAL IL-5 and lung IL-5 were only increased at 8 wk (Fig. 5H). In contrast, there was no increase in levels of lung IL-4 in hORMDL3zp3-Cre mice at 4, 8, or 26 wk of age (data not shown).

Increased IgE but not IgG, IgM, or IgA in hORMDL3zp3-Cre mice

Levels of IgE were significantly increased in hORMDL3zp3-Cre compared to WT mice at 4 wk of age, and this increase in IgE persisted at 8 and 26 wk (Fig. 6A). The increase in IgE was selective, as there was no increase in IgG (IgG1 or IgG2) (Fig. 6B, 6C), IgM (Fig. 6D), or IgA (Fig. 6E) in hORMDL3zp3-Cre mice.

Acute OVA allergen challenge enhances peribronchial eosinophilic inflammation, OVA specific IgE, and IL-4 in hORMDL3zp3-Cre mice

Acute OVA challenge induced a greater increase in peribronchial eosinophils (Fig. 7A–E) and OVA-specific IgE (Fig. 7F) in hORMDL3zp3-Cre mice compared to WT mice. This was associated with increased lung IL-4 levels in hORMDL3zp3-Cre mice compared to WT mice (Fig. 7G). Whereas acute OVA challenge induced increased levels of lung IL-5 (Fig. 7I), IL-13 (Fig. 7H), and eotaxin-1 (Fig. 7J) in hORMDL3zp3-Cre mice (OVA versus no OVA), these levels were not different from those detected in OVA-challenged WT mice.

Airway hyperreactivity to methacholine

hORMDL3zp3-Cre mice had spontaneously increased airway responsiveness to methacholine compared to WT mice at 12 wk of age (Fig. 8A). Additionally, hORMDL3zp3-Cre mice had increased lung elastance compared to WT mice (Fig. 8B).

Discussion

Although multiple genetic association studies have demonstrated that ORMDL3 is highly linked to asthma (1–10), the mechanism by which ORMDL3 may contribute to the pathogenesis of asthma in vivo is at present unknown. In this study using a mouse model in which the human ORMDL3 gene is overexpressed, we demonstrate the novel findings that expression of the human ORMDL3 transgene in vivo is associated with significantly increased airway remodeling, including increased airway smooth muscle, subepithelial fibrosis, and mucus. These airway remodeling changes in hORMDL3zp3-Cre mice were associated with the spontaneous development of increased airway responsiveness. Additionally, the remodeling changes were associated with an increased lung elas- tance (the inverse of lung compliance), which indicates an increased stiffness of the remodeled lungs. The mechanism of the increased airway remodeling did not appear to be dependent on increased airway inflammation, as significant airway remodeling was evident at 4 wk of age in the hORMDL3zp3-Cre mice, a time point not associated with an increased number of peribronchial eosinophils, neutrophils, macrophages, or CD4 cells. Increased levels of expression of genes associated with airway remodeling were detected in the lung (SERCA2b) and airway epithelium (TGF-β1, ADAM8, MMP-9) of hORMDL3zp3-Cre mice, suggesting that these pathways may contribute to airway remodeling detected in these mice. The importance of these genes that are highly expressed in hORMDL3zp3-Cre mice to airway remodeling...
and asthma is suggested from studies demonstrating expression of these pathways in the airways of human asthmatics (15, 22–27), induction of these mediators by allergen inhalation in asthmatics (TGF-β1, MMP-9) (28–30), and inhibition of asthma outcomes in mice deficient in these genes (ADAM8, Smad3, MMP-9) (31–35), or in mice treated with neutralizing Abs (TGF-β1) (36).

We have previously performed in vitro studies and demonstrated that transfection of ORMDL3 induces activation of one of the three pathways of the ER UPR, namely the ATF6 pathway (13). Using hORMDL3p3-Cre mice we have made the novel observation that the ATF6 pathway of the UPR is selectively activated by the human ORMDL3 transgene in vivo. ATF6 (consisting of the closely related ATF6α and ATF6β in mammals) (37) is a transcription factor known to regulate genes involved in ER protein folding (14), as well as expression of SERCA2b, which has been implicated in airway remodeling in asthma (15). In this study we

![FIGURE 7. Effect of acute OVA allergen challenge on airway inflammation, IgE, and Th2 cytokines in hORMDL3p3-Cre mice. hORMDL3p3-Cre and WT mice were sensitized to OVA and challenged with OVA allergen by inhalation. Compared to OVA-challenged WT mice, OVA-challenged hORMDL3p3-Cre mice had significantly increased levels of (A–E) lung MBP+ peribronchial eosinophils as assessed by immunohistochemistry, (F) OVA-specific IgE, and (G) lung IL-4. Levels of lung (H) IL-13, (I) IL-5, and (J) eotaxin-1 were similar in OVA-challenged WT and OVA-challenged hORMDL3p3-Cre mice (n = 12 mice/group). (A–D) Original magnification ×200.](http://www.jimmunol.org/)

![Lung eosinophils](http://www.jimmunol.org/)

![E](http://www.jimmunol.org/)

![F](http://www.jimmunol.org/)

![G](http://www.jimmunol.org/)

![H](http://www.jimmunol.org/)

![I](http://www.jimmunol.org/)

![J](http://www.jimmunol.org/)
demonstrate that hORMDL3zbp3-Cre mice exhibit both activation of ATF6 and increased levels of lung SERCA2b, suggesting that the ATF6 and SERCA2b pathways are downstream of ORMDL3 in vivo as previously demonstrated in vitro (13). In prior in vitro studies we have demonstrated that transfection of ORMDL3 induced activation of ATF6 and expression of SERCA2b, whereas knockdown of ATF6α inhibited SERCA2b expression (13). Activation of ATF6 was maximal, as further induction of the UPR upon treatment of the cells with thapsigargin, a well-characterized ER stress inducer, did not increase the nuclear localization of ATF6. Overall, these studies of hORMDL3zbp3-Cre mice provide in vivo evidence of ATF6a-dependent pathways (SERCA2b) and ATF6α-independent pathways (TGF-β1, ADAM8, MMP-9) through which the ER-localized ORMDL3 may be linked to airway remodeling and asthma. However, only future studies in which the ATF6 pathway is selectively inhibited will be able to determine the role played by the ATF6 pathway in any remodeling changes we have noted in hORMDL3zbp3-Cre mice.

hORMDL3zbp3-Cre mice also had an age-related increase in levels of airway inflammation, as well as an increase in lung cytokines and chemokines. The increased levels of airway inflammation were not evident in hORMDL3zbp3-Cre mice aged 4 wk, and only started to be evident at 8 wk (increase in peribronchial macrophages). At 26 wk hORMDL3zbp3-Cre mice had significantly increased levels of peribronchial CD4+ cells, eosinophils, macrophages, and neutrophils, suggesting that the airway inflammatory response in hORMDL3zbp3-Cre mice increased during the period from 8 to 26 wk of age. Because the inflammatory response in hORMDL3zbp3-Cre mice had evidence of both Th2-mediated inflammation (increased CD4+ cells and eosinophils), as well as increased neutrophils and macrophages, we investigated whether their lungs expressed cytokines and chemokines associated with Th2-mediated inflammation (IL-4, IL-5, IL-13, eotaxin-1) or chemokines known to be regulated by ORMDL3 in vitro (CXCL10, CXCL11, IL-8, CCL20) (13). Levels of lung Th2 cytokines (IL-5 and IL-13) were increased in hORMDL3zbp3-Cre mice at 8 and 26 wk of age. In contrast, there was no increase in IL-4 or eotaxin-1 at any time point. Allergen-challenged hORMDL3zbp3-Cre mice had significantly increased levels of peribronchial eosinophils and lung IL-4 compared to allergen-challenged WT mice. IL-4 can contribute to increased lung eosinophilic inflammation as evident in studies of IL-4 Tg mice (38). Our studies of hORMDL3zbp3-Cre mice confirmed that chemokines (CXCL10, CXCL11) we had observed to be highly expressed in vitro in ORMDL3-transfected cells (13) were highly expressed in hORMDL3zbp3-Cre mice bronchial epithelium (CXCL10) and alveolar macrophages (CXCL11). Although asthma has predominantly been associated with expression of CC chemokines, CXC chemokines have also been linked to asthma in studies in humans with asthma (39–42), as well as in studies in animal models (43, 44) (i.e., CXCL10 knockout mice have significant reduction in Th2-type allergic airway inflammation and airway responsiveness) (43).

In summary, these studies in hORMDL3zbp3-Cre mice provide evidence that ORMDL3 plays an important role in activation of the ATF6 UPR pathway in vivo, and that expression of ORMDL3 in vivo regulates airway remodeling (smooth muscle, fibrosis, mucus) potentially through ATF6 target genes such as SERCA2b and/or through ATF6-independent genes (TGF-β1, ADAM8, MMP-9) detected in airway epithelium, as well as through yet unidentified and/or uninvestigated pathways. In this regard sphingolipids are known to be regulated by ORMDL3, and sphingolipid pathways have been associated with asthma but not airway remodeling in mouse models (46–48). ORMDL3 also regulates ER-mediated calcium signaling (49) and lymphocyte activation in vitro (50). Because ORMDL3 is expressed in multiple cell types important to the pathogenesis of asthma (i.e., epithelial cells,
mature macrophages, eosinophils, T cells) (13, 51), in this study we have examined how increased expression of ORMDL3 in multiple cell types contributes to the pathogenesis of asthma. Future studies with selective overexpression of ORMDL3 in particular cell types will provide insight into the role of ORMDL3 in these individual cell types to the pathogenesis of asthma. Interestingly, increased levels of airway remodeling preceded increased levels of airway inflammation (CD4+ cells, macrophages, eosinophils, neutrophils) in the lungs of hORMDL33g-Cex mice, suggesting that airway remodeling can be dissociated from these pathways by activation of ORMDL3. As airway remodeling can be detected not only in adults with asthma, but also in childhood asthma (52), increased expression of ORMDL3 is dissociated from these pathways by activation of ORMDL3. As airway remodeling preceded increased levels of airway inflammation (52), evidence that severe airway remodeling in IL-5-deficient mice. Measurement in bronchoalveolar lavage fluid. Am. J. Respir. Crit. Care Med. 156: 642–647.


