

This information is current as
of November 17, 2013.

Epithelial NF- κ B Orchestrates House Dust Mite –Induced Airway Inflammation, Hyperresponsiveness, and Fibrotic Remodeling

Jane E. Tully, Sidra M. Hoffman, Karolyn G. Lahue, James D. Nolin, Vikas Anathy, Lennart K. A. Lundblad, Nirav Daphtary, Minara Aliyeva, Kendall E. Black, Anne E. Dixon, Matthew E. Poynter, Charles G. Irvin and Yvonne M. W. Janssen-Heininger

J Immunol published online 13 November 2013
<http://www.jimmunol.org/content/early/2013/11/12/jimmunol.1301329>

-
- Supplementary Material** <http://www.jimmunol.org/content/suppl/2013/11/13/jimmunol.1301329.DC1.html>
- Subscriptions** Information about subscribing to *The Journal of Immunology* is online at:
<http://jimmunol.org/subscriptions>
- Permissions** Submit copyright permission requests at:
<http://www.aai.org/ji/copyright.html>
- Email Alerts** Receive free email-alerts when new articles cite this article. Sign up at:
<http://jimmunol.org/cgi/alerts/etoc>

Epithelial NF- κ B Orchestrates House Dust Mite–Induced Airway Inflammation, Hyperresponsiveness, and Fibrotic Remodeling

Jane E. Tully,* Sidra M. Hoffman,* Karolyn G. Lahue,* James D. Nolin,* Vikas Anathy,* Lennart K. A. Lundblad,[†] Nirav Daphtary,[†] Minara Aliyeva,[†] Kendall E. Black,[†] Anne E. Dixon,[†] Matthew E. Poynter,[†] Charles G. Irvin,[†] and Yvonne M. W. Janssen-Heininger*

NF- κ B activation within the epithelium has been implicated in the pathogenesis of asthma, yet the exact role of epithelial NF- κ B in allergen-induced inflammation and airway remodeling remains unclear. In the current study, we used an intranasal house dust mite (HDM) extract exposure regimen time course in BALB/c mice to evaluate inflammation, NF- κ B activation, airway hyperresponsiveness (AHR), and airway remodeling. We used CC10-I κ B α _{SR} transgenic mice to evaluate the functional importance of epithelial NF- κ B in response to HDM. After a single exposure of HDM, mRNA expression of proinflammatory mediators was significantly elevated in lung tissue of wild-type (WT) mice, in association with increases in nuclear RelA and RelB, components of the classical and alternative NF- κ B pathway, respectively, in the bronchiolar epithelium. In contrast, CC10-I κ B α _{SR} mice displayed marked decreases in nuclear RelA and RelB and mRNA expression of proinflammatory mediators compared with WT mice. After 15 challenges with HDM, WT mice exhibited increases in inflammation, AHR, mucus metaplasia, and peribronchiolar fibrosis. CC10-I κ B α _{SR} transgenic mice displayed marked decreases in neutrophilic infiltration, tissue damping, and elastance parameters, in association with less peribronchiolar fibrosis and decreases in nuclear RelB in lung tissue. However, central airway resistance and mucus metaplasia remained elevated in CC10-I κ B α _{SR} transgenic mice, in association with the continued presence of lymphocytes, and partial decreases in eosinophils and IL-13. The current study demonstrates that following airway exposure with an asthma-relevant allergen, activation of classical and alternative NF- κ B pathways occurs within the airway epithelium and may coordinately contribute to allergic inflammation, AHR, and fibrotic airway remodeling. *The Journal of Immunology*, 2013, 191: 000–000.

The NF- κ B pathway is a critical regulator of both innate and adaptive immune responses in a wide variety of cell types. Upon stimulation, the I κ B kinase (IKK) signalosome, consisting of IKK β , IKK α , and IKK γ , is activated, leading to IKK β -mediated phosphorylation of I κ B α . Phosphorylation of I κ B α in turn leads to its subsequent ubiquitination and degradation by the 26S proteasome, thus allowing for the transcription factor RelA to translocate to the nucleus. This event results in RelA-dependent transcription of genes important in cell survival, proliferation, and inflammation (1, 2). A wide variety of agonists

can activate the classical NF- κ B pathway in lung epithelial cells and the resultant release of proinflammatory mediators crucial in the recruitment and activation of dendritic cells, lymphocytes, neutrophils, and many other cells in the lung (3). In addition, an alternative NF- κ B pathway exists, which requires activation of NF- κ B-inducing kinase (NIK) and subsequent phosphorylation of IKK α . IKK α in turn phosphorylates p100, leading to its partial processing to p52. This allows subsequent nuclear translocation of RelB/p52 and transcriptional activation of a subset of NF- κ B-dependent genes (4, 5). It was originally thought that the alternative NF- κ B pathway played a predominant role in lymphocyte activation and lymphoid organ development (6). However, recent work from our laboratory demonstrated that both classical and alternative NF- κ B pathways are activated in lung epithelial cells in response to diverse proinflammatory stimuli and that both pathways coordinately regulate proinflammatory responses (7).

Activation of the classical NF- κ B pathway within the airway epithelium has been demonstrated to play a critical role in acute inflammation and allergic airways disease. CC10-I κ B α _{SR} transgenic mice, which are refractory to I κ B α degradation and NF- κ B activation in the lung epithelium, were demonstrated to be strongly protected from airway inflammation induced by LPS (8). Following i.p. sensitization and challenge with OVA, CC10-I κ B α _{SR} transgenic mice showed a marked diminution of airway inflammation compared with WT littermate controls, although OVA-induced airway hyperresponsiveness (AHR) was unaffected in CC10-I κ B α _{SR} transgenic mice compared with controls (9). A similar protection against OVA-induced allergic inflammation and

*Department of Pathology, University of Vermont College of Medicine, Burlington, VT 05405; and [†]Department of Medicine, University of Vermont College of Medicine, Burlington, VT 05405

Received for publication May 20, 2013. Accepted for publication October 11, 2013.

This work was supported by National Institutes of Health Grants T32 HL076122, T32 ES07122, NIGMS P30, GM103532, and R01 HL060014, and by an American Thoracic Society unrestricted grant and a Parker B. Francis Fellowship (to V.A.).

Address correspondence and reprint requests to Dr. Yvonne M.W. Janssen-Heininger, Department of Pathology, University of Vermont Health Sciences Research Facility, Room 216A, Burlington, VT 05405. E-mail address: Yvonne.janssen@uvm.edu

The online version of this article contains supplemental material.

Abbreviations used in this article: AHR, airway hyperresponsiveness; BAL, bronchoalveolar lavage; CCL20, chemokine (C-C motif) ligand 20; G, tissue damping (parameter); H, elastance (parameter); HBE, human bronchial epithelial (cell); HDM, house dust mite; IKK, I κ B kinase; KC, keratinocyte-derived chemokine; NIK, NF- κ B-inducing kinase; PAS, periodic acid–Schiff; Rn, Newtonian resistance (parameter); α -SMA, α -smooth muscle actin; WT, wild-type.

Copyright © 2013 by The American Association of Immunologists, Inc. 0022-1767/13/\$16.00

peribronchiolar fibrosis has been observed in mice following epithelial-specific ablation of IKK β (10).

It remains unclear to date whether activation of NF- κ B within epithelial cells plays a role in the orchestration of inflammatory responses in vivo to an asthma-relevant allergen following sensitization via the airways. It also remains unknown whether both NF- κ B pathways are activated following exposure to an Ag. House dust mite (HDM) is a multifaceted allergen to which up to 85% of asthmatics are allergic (11). HDM has been shown to signal through the classical NF- κ B pathway in human bronchial epithelial cells in vitro (12). Therefore, the goal of the current study was to determine the activation of classical and alternative NF- κ B in epithelial cells in vivo in response to HDM, and to address its effect on HDM-triggered airway inflammation, remodeling, mucus, and AHR. Our results demonstrate the functional importance of epithelial NF- κ B in HDM-induced acute inflammatory responses, AHR, and airway remodeling. We also demonstrate activation of both classical and alternative NF- κ B pathways in response to HDM. These findings illustrate the complexity of activation of the NF- κ B pathways in settings of allergic airways disease and suggest a broader role for epithelial NF- κ B in lung disease pathogenesis.

Materials and Methods

Animal studies

CC10-IkB α _{SR} mice were generated as previously described (8) and backcrossed onto the BALB/cJ ($n = 10$) (The Jackson Laboratory, Bar Harbor, ME) background. Transgene-negative littermates were used as a control. All experiments were approved by the University of Vermont Institutional Animal Care and Use Committee.

Cell culture

A human bronchial epithelial (HBE) cell line was kindly provided by Dr. Albert van der Vliet (University of Vermont) and cultured as described previously (13, 14), and primary human nasal epithelial cells were cultured as described previously (15). Human cell lines were exposed to either PBS or 25 μ g HDM (Greer, Lenoir, NC). All protocols that use primary human nasal epithelial cells were approved by the Institutional Review Board.

HDM exposure

BALB/cJ mice (The Jackson Laboratory) were subjected to daily intranasal instillation with 50 μ g HDM (35 endotoxin units/mg, normalized to protein content) extract resuspended in PBS, or PBS alone as a vehicle control. Briefly, mice were either instilled with 1 dose of HDM and euthanized 2 h, 6 h, or 24 h later or instilled with 3 doses of HDM on 3 consecutive days and euthanized 24 h thereafter. In addition, mice were exposed to HDM 5 d/wk for 1, 2, or 3 wk (5, 10, or 15 instillations, respectively) and harvested 72 h after the final exposure (Fig. 1A).

Assessment of AHR

Mice subjected to 5, 10, or 15 administrations of HDM were anesthetized with an i.p. injection of pentobarbital sodium (90 mg/kg), tracheotomized, and mechanically ventilated at 200 breaths per minute and assessed in response to increasing doses of methacholine (0, 3.125 mg, 12.5 mg, and 25 mg). Respiratory mechanics were assessed with a forced oscillation technique on a computer-controlled small animal ventilator (Flexi-vent, SCIREQ, QC, Canada), as previously described (16, 17), and the parameters Newtonian resistance (Rn), tissue damping (G), and elastance (H) were calculated.

Serum IgG1 and IgE

Following euthanization, blood was collected by heart puncture and immediately spun through a microtainer, and serum was separated. Analysis of serum IgG1 and IgE was performed via ELISA methods, using 1 μ g/ml HDM to coat the 96-well plates.

Bronchoalveolar lavage

Following euthanization, bronchoalveolar lavage (BAL) was collected using 1 ml PBS. Cell counts were determined (Advia 120 Automated Hematology Analyzer), and differential cells were analyzed by the Hema3 kit (Fisher Scientific, Kalamazoo, MI) by counting a minimum of 300 cells per mouse, as previously described (17).

mRNA/protein analysis

Right lung lobes were flash frozen and pulverized for protein and mRNA. Total mRNA was isolated using the RNeasy kit (QIAGEN, Valencia, CA). Next, 1 μ g mRNA was used to generate cDNA, followed by quantitative PCR using SYBR Green (Bio-Rad, Hercules, CA) to assess expression of Muc5ac, keratinocyte-derived chemokine (KC), MIP-2, GM-CSF, IL-6, chemokine (C-C motif) ligand 20 (CCL20), and IL-33. Primers were as follows: KC (Forward: 5'-GCTGGATTCACCTCAAGAA-3', Reverse: 5'-TGGGGACACCTTTTAGCATC-3'), IL-6 (Forward: 5'-CTGATGCTGGTGACAACCAC-3', Reverse: 5'-CAGAATTGCCATTGCACAAAC-3'), CC-L20 (Forward: 5'-AAGACAGATGGCCGATGAAG-3', Reverse: 5'-AGCCCTTTTCAACCAGTTCT-3'), Muc5ac (Forward: 5'-CAGTGAATTCTGGAGGCCAACAAAGGTAGAG-3', Reverse: 5'-AGCTAAGCTTAGATCTGTGGGACAGCAGC-3'), MIP-2 (Forward: 5'-AGTGAAGTGGCTGTCAATG-3', Reverse: 5'-TTCAGGGTCAAGGCCAACTT-3'), GM-CSF (Forward: 5'-GGCCTTGAAGCATGTAGAGG-3', Reverse: 5'-GGA-GAACTCGTTAGAGACGACTT-3'), and IL-33 (Forward: 5'-GCTGCGTCTGTTGACACATT-3', Reverse: 5'-GACTTGCAGGACAGGGAGAC-3'). The expression of all genes was normalized to the housekeeping gene cyclophilin (Forward: 5'-TTCCTCCTTTCACAGAATTATTCCA-3', Reverse: 5'-CCGCCAGTGCCATTATGG-3') and calculated via the ddCt method. Abs for Western blot analysis were RelB and NIK (Santa Cruz Biotechnology, Santa Cruz, CA); histone H3, p52, and phospho RelA536 (Cell Signaling Technology, Danvers, MA); and β -actin (Sigma-Aldrich, St. Louis, MO). ELISAs were performed according to the manufacturer's instructions (R&D Systems, Minneapolis, MN) on either BAL or homogenized lung extracts. Protein was equalized before analysis.

Histopathology/immunofluorescence/ α -smooth muscle actin immunohistochemistry

Following euthanization, the left lobe was inflated with 4% paraformaldehyde and mounted in paraffin-embedded 5- μ m sections. H&E and periodic acid-Schiff (PAS) imaging was all performed with three small bronchioles ($\times 20$ magnification) per animal. Mucus metaplasia was assessed with a blinded scoring system by two independent investigators with the following scale: 0, no reactivity; 1, minimal staining; 2, moderate staining; and 3, prominent staining. Scores were averaged according to treatment group. Immunofluorescence was performed as previously described to detect nuclear RelA and RelB in situ (18). RelA and RelB Abs were purchased from Santa Cruz Biotechnology. Staining for α -smooth muscle actin (α -SMA) was performed on lung sections following incubation of slides for 20 min in 0.01 M sodium citrate, pH 6.0 at 95°C. Slides were then blocked with 2% normal goat serum for 30 min, incubated with mouse mAb against α -SMA (1:5000 dilution; Sigma Aldrich) overnight at 4°C, and then incubated in biotinylated anti-mouse IgG for 30 min at room temperature. Subsequently, the slides were incubated in avidin-biotin complex-alkaline phosphatase (Vectastain ABC-AP, Vector Laboratories, Burlingame, CA) for another 30 min at room temperature. After rinsing the sections in PBS, the substrate Vector Red (Vector Laboratories) was added for 20 min; this substrate reacts with the bound alkaline phosphatase, thus producing an intense red color. The slides were then counterstained in hematoxylin and imaged ($\times 20$ magnification) for analysis.

Assessment of collagen content

Collagen was assessed from the upper right lobe of the lung after an overnight digestion with 10 mg pepsin in 0.5 M acetic acid. Quantification was performed by the Sircol Assay according to the manufacturer's instructions (Accurate Chemical and Scientific, Westbury, NY). Masson's trichrome reactivity was evaluated in three bronchioles ($\times 20$ magnification) per animal, and analyzed by two blinded investigators with the following scale: 0, no reactivity; 1, minimal staining; 2, moderate staining; and 3, prominent staining.

Statistical analysis

All data were evaluated using Graphpad Prism 6 Software (Graphpad, San Diego, CA). A one-way ANOVA was used with Bonferroni corrections to adjust for multiple comparisons, and all p values < 0.05 were considered statistically significant. Histopathological/ α -SMA immunohistochemistry scores were analyzed using the Kruskal-Wallis test and Dunn multiple comparison post hoc tests.

Results

Inflammatory response in BALB/c mice exposed to HDM

Because of the well-known role of NF- κ B in the orchestration of inflammation, we first sought to assess the extent and kinetics of

the HDM-induced inflammatory response through a time-course analysis illustrated schematically in Fig. 1A. Neutrophils in BAL were significantly increased 6 h following a single intranasal challenge with HDM, and remained elevated until 24 h post three challenges, compared with PBS controls (Fig. 1B). It is important to note that neutrophils were elevated 24 h following exposure to the vehicle PBS, potentially indicative of a nonspecific response to the intranasal instillation. Neutrophils were also detectable in BAL 72 h post 5, 10, or 15 challenges, with statistically significant increases occurring in response to HDM 72 h post 15 challenges. Although macrophages in BAL tended to increase in response to HDM, these trends were not statistically significant. No eosinophils or lymphocytes were detected in BAL up to 24 h post three challenges. In contrast, robust increases in eosinophils and lymphocytes occurred in BAL 72 h post 10 and 15 challenges with HDM, compared with PBS controls (Fig. 1B). Although some fluctuations in the total number of macrophages in BAL were observed throughout these time points, these were not statistically significant (Fig. 1B).

Increases in nuclear localization of RelA and RelB in airway epithelium in response to HDM

We next evaluated a potential role for epithelial NF- κ B in response to HDM. We investigated kinetics of activation of both classical and alternative pathways in the bronchiolar epithelium via the assessment of nuclear presence of RelA and RelB. Nuclear pres-

ence of RelA and RelB (indicated by yellow staining) within the bronchiolar epithelium was increased 2 and 6 h following a single challenge with HDM, compared with PBS controls in the nuclei (Fig. 2). Nuclear RelA decreased to control reactivity by 24 h, but increased again 24 h post 3 challenges with HDM, indicative of a bimodal activation pattern (Fig. 2). In contrast, increases in nuclear RelB were apparent throughout the time course evaluated in this study (Fig. 2). No clear evidence for nuclear localization of RelA or RelB within the airway epithelium was apparent 72 h after 5, 10, or 15 challenges with HDM compared with PBS (data not shown).

Proinflammatory mediator expression following acute HDM exposures

To investigate the early inflammatory response after HDM exposure, we evaluated mRNA expression of IL-33, KC, GM-CSF, MIP-2, CCL20, IL-6, IL-25, and thymic stromal lymphopoietin, proinflammatory mediators shown to be produced by epithelial cells. Expression levels of all genes analyzed were significantly increased following 2-h exposure to HDM, with the exception of IL-25 and thymic stromal lymphopoietin, whose expression did not change (data not shown). mRNA expression of CCL20, GM-CSF, MIP-2, KC, and IL-6 remained elevated 6 h after a single administration of HDM and tended to decrease toward control levels thereafter (Supplemental Fig. 1).

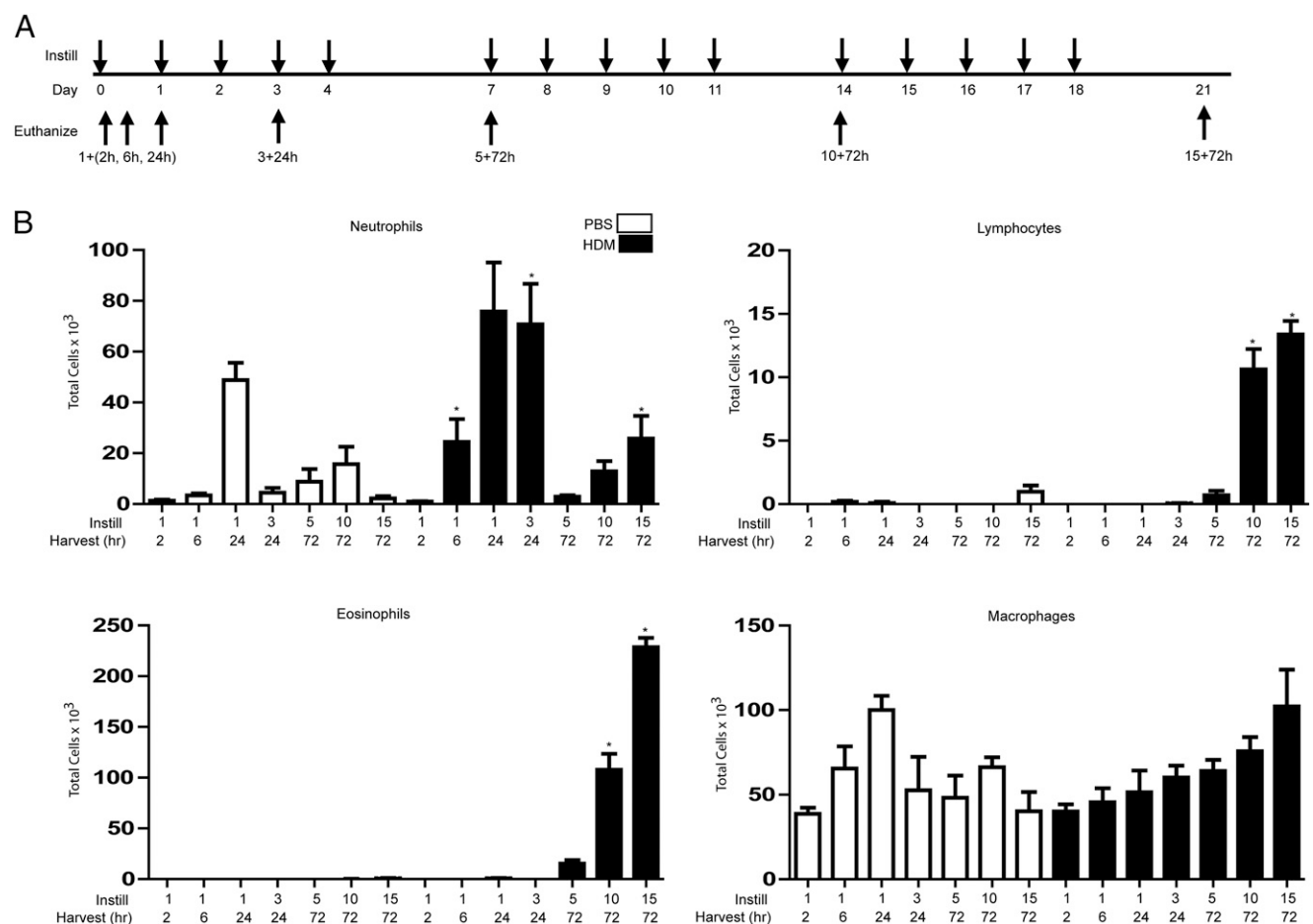


FIGURE 1. Analysis of cell totals and differentials in BAL of mice exposed to HDM. (A) Schematic depicting the time course of HDM exposure and euthanization. For instillation, 50 μ g HDM or PBS as the vehicle control was administered intranasally once at the days indicated via downward arrows. Mice were euthanized 2 h, 6 h, or 24 h post a single challenge, 24 h after 3 challenges, or 72 h after 5, 10, or 15 challenges, indicated by the upward arrows. (B) Mice ($n = 5$ per group per time point) were exposed to PBS or HDM according to the schematic in (A), and BAL was assessed for neutrophils, eosinophils, lymphocytes, and macrophages. * $p < 0.05$ (ANOVA) compared with the PBS group at the same time point.

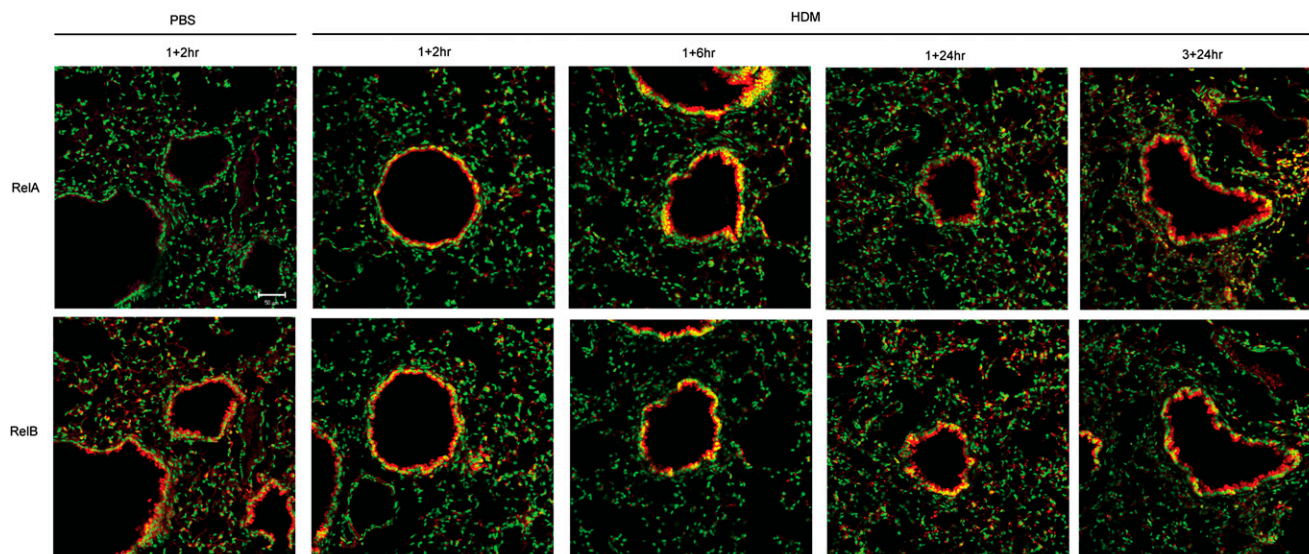


FIGURE 2. Assessment of nuclear localization of RelA and RelB in airway epithelium in response to HDM. BALB/c mice were exposed to PBS and HDM and harvested as indicated. Deparaffinized lung sections were incubated with Abs directed against RelA or RelB, followed by fluorophore-conjugated secondary Abs, and the nuclei were counterstained with Sytox Green. Images were captured by laser scanning confocal microscopy using identical instrument settings in all groups. Images are representative results of two independent experiments; $n = 5$ per group per time. RelA/RelB: red, DNA: green, nuclear RelA/RelB: yellow. Scale bar, 50 μ m.

Role of epithelial NF- κ B in proinflammatory cytokine expression induced by HDM

We next sought to address the role of epithelial NF- κ B in HDM-induced proinflammatory responses using CC10-I κ B α _{SR} transgenic mice. Although nuclear RelA and RelB content increased in the bronchiolar epithelium in WT mice in response to HDM, these increases were not observed in CC10-I κ B α _{SR} mice exposed to HDM (Fig. 3A). HDM-mediated increases in mRNA expression of CCL20, GM-CSF, MIP-2, KC, IL-33, and IL-6 in WT mice were strongly attenuated in CC10-I κ B α _{SR} mice (Fig. 3B), demonstrating the importance of NF- κ B activation in the bronchiolar epithelium in the orchestration of the acute proinflammatory responses to HDM.

Effects of repeated HDM exposure on AHR, mucus metaplasia, and remodeling

We next subjected mice to 5, 10, and 15 challenges of HDM. Serum content of HDM-specific IgG1 and IgE showed no apparent increases 72 h following 5-d challenge. Marked increases in HDM-specific IgG1 and IgE were apparent following 10 and 15 challenges with HDM, compared with PBS controls, indicative of activation of adaptive immune responses (Supplemental Fig. 2A). AHR, a feature of allergic airways disease, was assessed via forced oscillation mechanics using ascending doses of methacholine. Rn increased significantly over PBS controls after 10 and 15 challenges with HDM, whereas no changes were apparent in mice subjected to 5 challenges with HDM (Supplemental Fig. 2B). G, which is indicative of tissue resistance and small airway dysfunction, was increased following 5 challenges with HDM, and remained increased throughout 10 and 15 challenges. H was also increased following 5, 10, and 15 challenges of HDM. Increases in H were most prominent after 10 challenges of HDM and tended to decrease after 15 challenges (Supplemental Fig. 2B). Histopathological evaluation revealed a robust inflammatory response to HDM, with prominent peribronchiolar and perivascular cellular infiltrates being apparent following 10 and 15 challenges with HDM (Supplemental Fig. 2C). In addition, mucus metaplasia was apparent following 5, 10, and 15 challenges of HDM, based upon

staining with PAS reagent and Muc5ac expression (Supplemental Fig. 2C, 2D). Another hallmark of severe allergic airways disease is peribronchiolar fibrotic remodeling. Following 15 challenges with HDM, overall collagen content increased in the lung tissue (Supplemental Fig. 2E), consistent with increases in peribronchiolar collagen deposition evaluated via Masson's trichrome staining (Supplemental Fig. 2C). These results collectively demonstrate that the HDM exposure regimen used in this study induces a number of the hallmark features of allergic airways disease.

Role of epithelial NF- κ B in HDM-induced inflammation

We next determined the role of epithelial NF- κ B in the inflammatory response induced following 15 challenges of HDM. HDM-mediated increases in total cell counts in BAL observed in WT mice were significantly decreased in CC10-I κ B α _{SR} mice (Fig. 4A). BAL eosinophils were significantly decreased in CC10-I κ B α _{SR} transgenic mice exposed to HDM, whereas BAL lymphocytes remained elevated in HDM-exposed CC10-I κ B α _{SR} transgenic mice compared with WT littermates (Fig. 4B). Total macrophage number in BAL remained unchanged in CC10-I κ B α _{SR} mice (Fig. 4B). In contrast, HDM-mediated increases in BAL neutrophils in WT mice were completely attenuated in CC10-I κ B α _{SR} mice. Consistent with these observations, increases in BAL content of the neutrophil chemoattractant KC in WT mice exposed to HDM were absent in CC10-I κ B α _{SR} mice (Fig. 4C). HDM-specific IgG1 and IgE levels in serum were decreased slightly in CC10-I κ B α _{SR} HDM-exposed mice, although not significantly, compared with littermate controls, but remained elevated over PBS controls (Fig. 4D). Levels of IFN- γ , IL-13, and IL-17A in homogenized lung tissue were significantly decreased in CC10-NF- κ B_{SR} mice exposed to HDM, compared with WT animals (Fig. 4E), collectively suggesting an attenuation of Th-mediated responses to HDM following epithelial-specific inhibition of NF- κ B. We next sought to assess inhibition of nuclear RelA and RelB in lung tissue in CC10-NF- κ B_{SR} mice exposed to HDM. After 15 challenges with HDM, no consistent increases in nuclear RelA or RelB were detected in the bronchiolar epithelium (data not shown), in contrast to earlier time points at which increases in

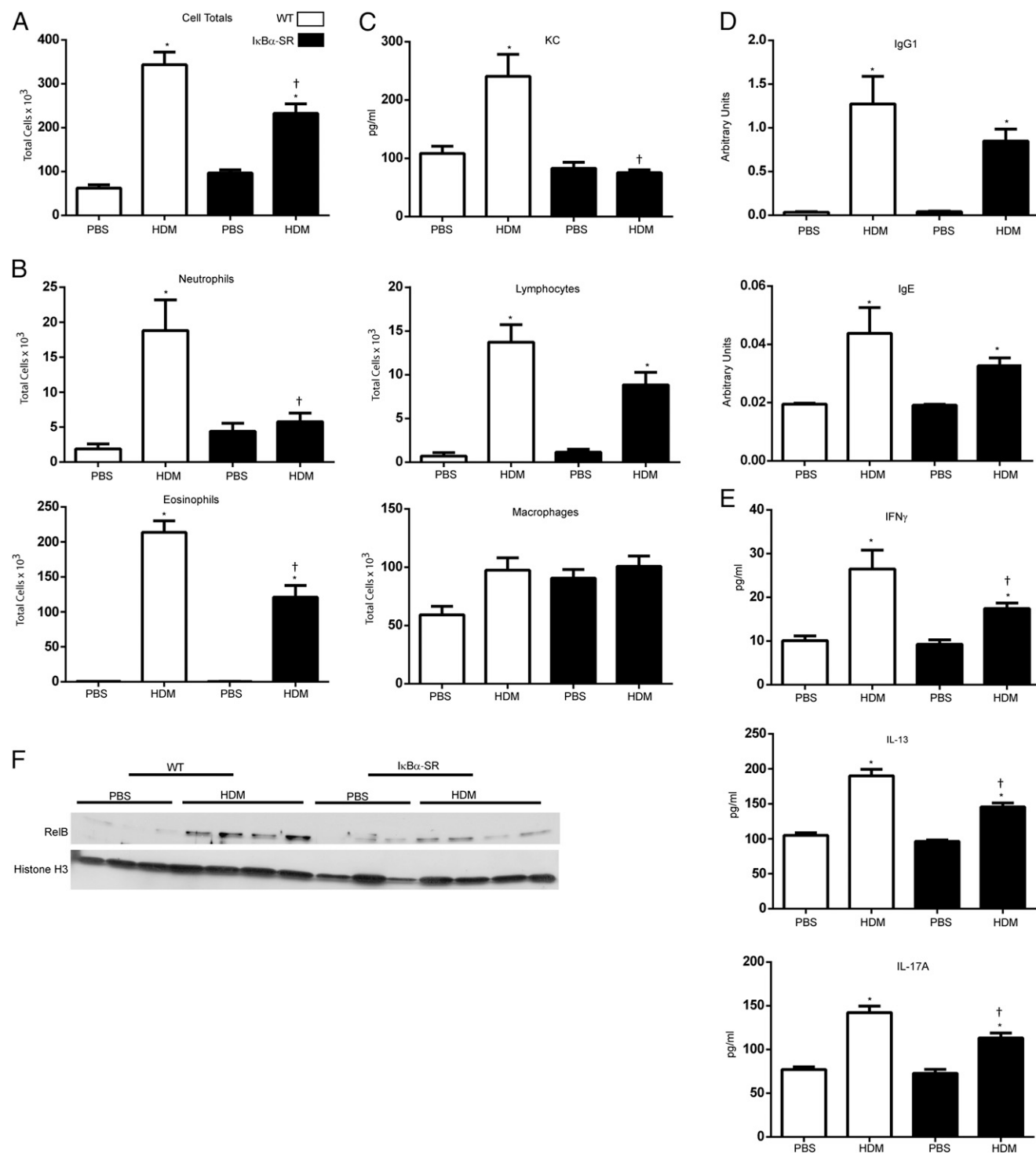


FIGURE 4. Assessment of HDM-induced inflammation, immunoglobulins, T cell cytokines, and nuclear RelB in WT or CC10-NF- κ B_{SR} mice following 15 challenges with HDM. Assessment of total cells (**A**) and differential cell counts (**B**) and KC levels (**C**) in BAL. (**D**) Total IgG1 and IgE in serum. (**E**) IFN- γ , IL-13, and IL-17A in homogenized tissue by ELISA. (**F**) Nuclear content of RelB in homogenized lung tissue. Histone H3 is shown as a loading control. Data represent eight mice per group per time point. * $p < 0.05$ (ANOVA) compared with the PBS group at the same time point; [†] $p < 0.05$ (ANOVA) compared with WT mice exposed to HDM.

mice, which were absent in the CC10-IkB α _{SR} transgenics (Fig. 4F).

Role of epithelial NF- κ B in HDM-induced AHR and remodeling

WT and CC10-IkB α _{SR} mice subjected to 15 challenges with HDM demonstrated equivalent increases in Rn. In contrast, HDM-mediated increases in G and H were significantly decreased in

CC10-IkB α _{SR} mice compared with littermate controls (Fig. 5A). Consistent with attenuated inflammatory cell profiles in BAL, peribronchiolar infiltrates were attenuated in CC10-IkB α _{SR} mice compared with controls (Fig. 5B). In contrast, mucus metaplasia was not observed to be decreased in HDM-exposed CC10-IkB α _{SR} mice, compared with WT animals (Fig. 5B, 5D). These findings are consistent with a lack of attenuation of HDM-induced Muc5AC mRNA (Fig. 5C) and modest decreases in IL-13 con-

tent (Fig. 4E). Biochemical analysis of collagen demonstrated that HDM-mediated increases in WT mice were significantly decreased in CC10-I κ B α _{SR} transgenic mice (Fig. 5E), consistent with less peribronchiolar collagen detected via histopathological analysis (Fig. 5B, 5F). Furthermore, α -SMA, a known marker of airway thickening and remodeling, was assessed via immuno-

histochemistry. Peribronchiolar α -SMA staining was significantly increased following HDM administration in WT mice, as compared with PBS controls. In contrast, no increases in peribronchiolar α -SMA reactivity were detected in CC10-I κ B α _{SR} mice exposed to HDM, in comparison with PBS controls (Fig. 5G, 5H).

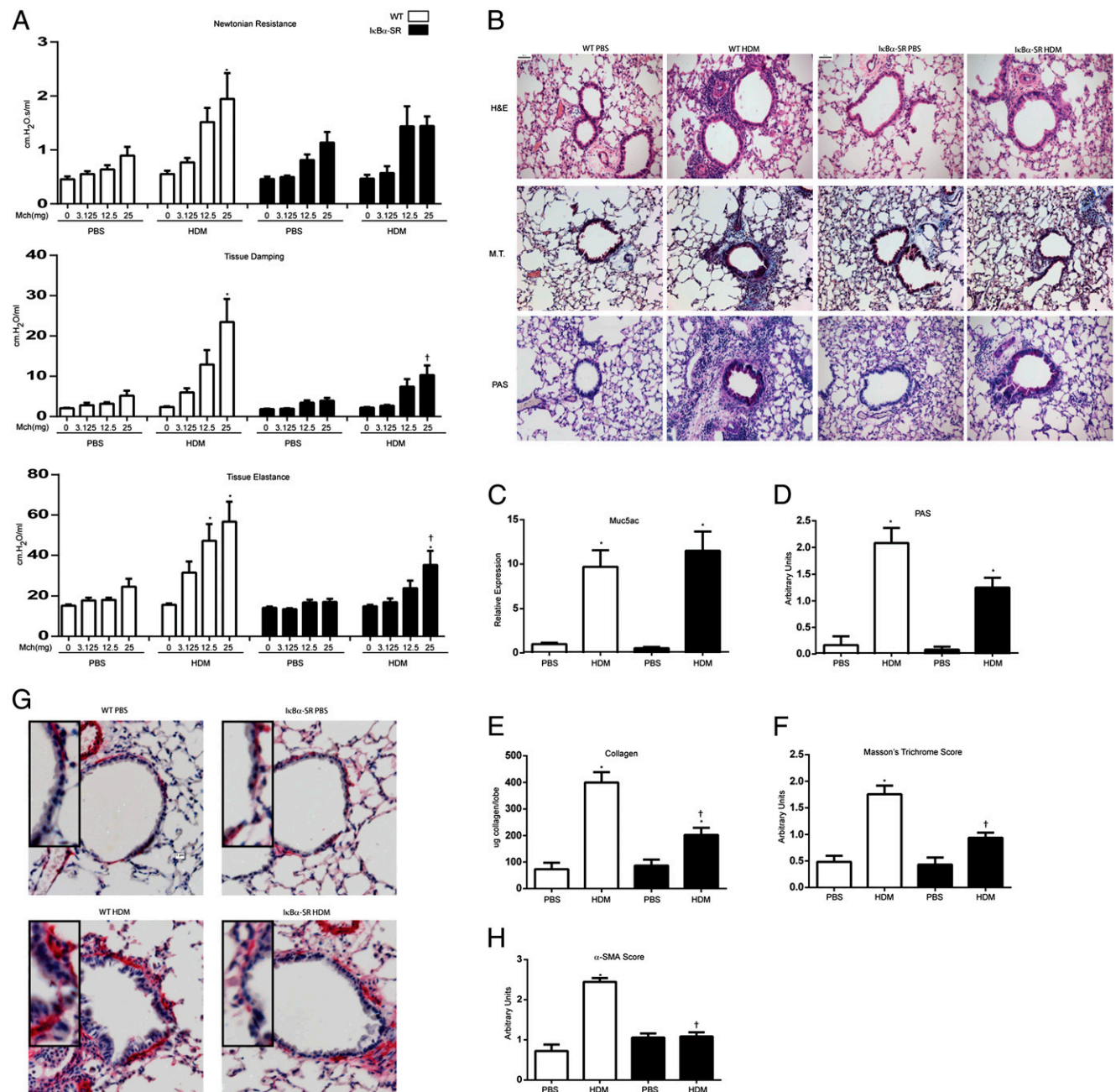


FIGURE 5. Assessment of HDM-induced alterations in respiratory mechanics and airway remodeling in WT or CC10-I κ B α _{SR} mice following 15 challenges with HDM. **(A)** Evaluation of AHR via forced oscillation mechanics in response to 3.125, 12.5, and 25 mg methacholine. Shown are the parameters Rn, G, and H assessed via forced oscillation **(B)** Histopathological evaluation of tissue inflammation (H&E), mucus metaplasia (PAS), and peribronchiolar collagen deposition (Masson's trichrome). Scale bar, 50 μ m. **(C)** Analysis of Muc5ac mRNA expression in homogenized lung tissue. Data were normalized to cyclophilin and are presented as relative expression. **(D)** Quantification of mucus metaplasia. Bronchioles of similar size ($n = 3$) were analyzed in each mouse by two blinded scorers, and average scores were presented as average units. **(E)** Assessment of collagen from the upper right lobe. **(F)** Quantification of peribronchiolar collagen deposition. Bronchioles of similar size ($n = 3$) were analyzed in each mouse by two blinded scorers, and average scores presented as average units. **(G)** Assessment of α -SMA immunohistochemistry. Red, α -SMA; scale bar, 1 μ m. **(H)** Quantification of α -SMA immunoreactivity. Bronchioles of similar size ($n = 3$ per mouse) were analyzed by two independent blinded scorers, and average scores were presented as average units. Data represent eight mice per group per time point. * $p < 0.05$ (ANOVA) compared with the PBS group at the same time point; † $p < 0.05$ (ANOVA, Kruskal–Wallis) compared with WT mice exposed to HDM.

HDM-induced activation of the classical and alternative NF- κ B pathways in human nasal and HBEs

Because we demonstrated increases in nuclear RelB in the bronchial epithelium in response to HDM administration, we sought to determine whether the alternative NF- κ B pathway could be activated directly by HDM in human epithelial cells. Results in Fig. 6A demonstrate increased levels of NIK and p52 in response to HDM in primary human nasal epithelial cells obtained from two independent donors, and similar increases were observed (Fig. 6B) in HBE cells, albeit less robust, demonstrating activation of the alternative NF- κ B pathway. In addition, phosphorylation of RelA at serine 536, a separate parameter of NF- κ B activation, was also increased in response to HDM (Fig. 6A, 6B), demonstrating that in addition to the known ability of HDM to activate the classical NF- κ B pathway, it also induces activation of the alternative NF- κ B pathway in lung epithelial cells.

Discussion

NF- κ B is a regulator of inflammation and immunity, and its role in the pathogenesis in asthma has been suggested on the basis of evidence of its activation in the bronchiolar epithelium from asthmatics (19) and from studies in mouse models of allergic airways disease (18). Notably, studies previously performed in our laboratory have revealed a crucial role for lung epithelial NF- κ B in the ALUM/OVA model of allergic airways disease (9). Moreover, transgenic expression of constitutively active IKK β in lung epithelial cells was sufficient to cause neutrophilic inflammation and AHR, as well as enhanced sensitization to an inhaled Ag (17, 20). Despite these prior observations, the more generalized importance of epithelial NF- κ B in allergic airways disease following sensitization/exposure to a relevant allergen via the airways has yet to be determined. Results presented in this article describe a critical role for nonciliated airway epithelial NF- κ B in promoting inflammation, AHR, and fibrotic remodeling following extended challenges of HDM, whereas differences in mucus metaplasia did not appear affected.

Results of the current study somewhat contrast our previous work using the ALUM/OVA model of i.p. sensitization followed by

challenges of aerosolized OVA. Notably, CC10-I κ B α _{SR} mice subjected to the ALUM/OVA protocol were strongly protected against the development of eosinophilic inflammation and mucus metaplasia, but were not protected against OVA-induced AHR (9). The reasons for these discrepant findings remain unclear. It is likely that the route of sensitization of an allergen dictates the nature of the subsequent immunological and pathophysiological response. In support of the latter is a previous report demonstrating that the i.p. sensitization regimen with ALUM/OVA triggers eosinophilic, Th2-driven inflammation (21). In contrast, sensitization with OVA via the airways, along with LPS as the adjuvant, led to neutrophilic-dependent AHR and Th17-associated inflammation (22).

In the current study, we demonstrated that CC10-I κ B α _{SR} mice did not display increases in airway neutrophils following 15 challenges with HDM compared with PBS controls, whereas neutrophils were robustly increased in BAL of WT mice. In contrast, HDM-induced eosinophilia was only partially attenuated in CC10-I κ B α _{SR} mice, whereas BAL lymphocytes were not significantly affected. IL-5, a potent eosinophil chemoattractant in mice and humans, was not detectable at any experimental time points (data not shown); however, we did demonstrate a strong decrease in KC (human IL-8) in CC10-I κ B α _{SR} mice in comparison with WT mice exposed to HDM. These findings demonstrate that following inhibition of epithelial NF- κ B in the setting of HDM-induced disease, the inflammatory process is not uniformly inhibited but preferentially affects neutrophils. This potential differential effect of NF- κ B inhibition on the inflammatory process could explain the impact of airway remodeling and AHR. Consistent with the continued presence of eosinophils, HDM-specific IgG1 and IgE, and IL-13 in CC10-I κ B α _{SR} mice exposed to HDM, mucus metaplasia and Muc5AC expression remained elevated in these animals. Previous studies demonstrated that IL-13 can activate epithelial cells to produce Muc5ac by an NF- κ B-independent mechanism (23). In light of those observations, the lack of an impact of CC10-I κ B α _{SR} mice in HDM-induced mucus metaplasia is, therefore, perhaps, not surprising.

The role of neutrophils in the pathogenesis of asthma remains unclear. HDM exposure in mice has been associated with mixed

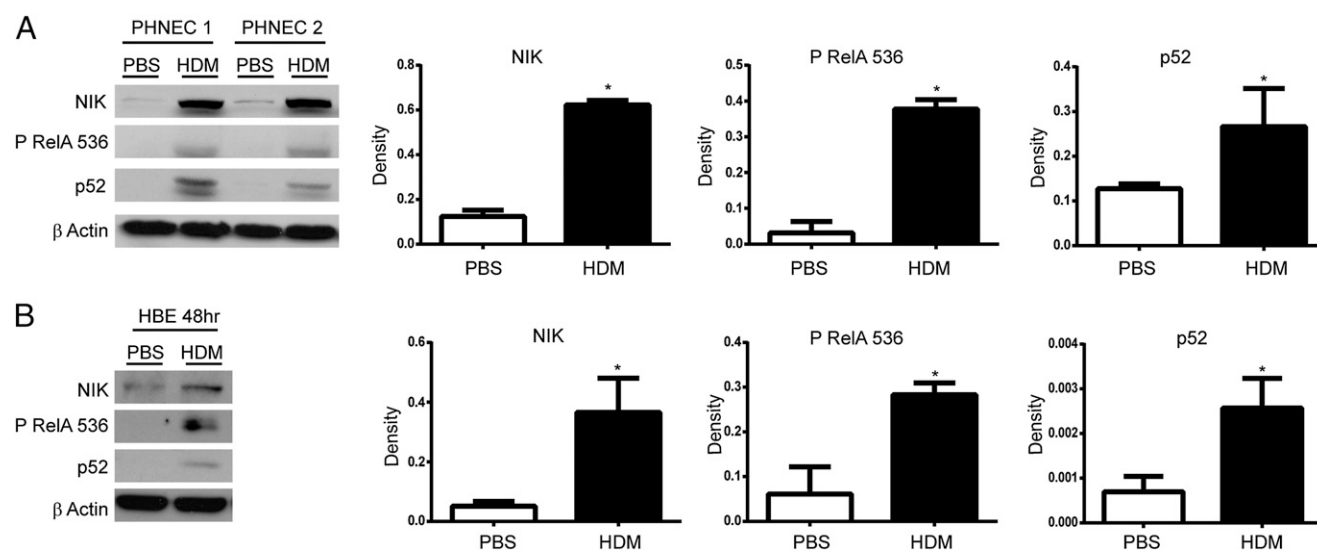


FIGURE 6. Assessment of activation of classical and alternative NF- κ B pathways in human lung epithelial cells exposed to HDM. (A) Primary human nasal epithelial (PHNE) cells or (B) HBE cells were exposed to PBS or 25 μ g HDM once a day for consecutive days and harvested thereafter at either 72 h (PHNE) or 48 h (HBE) following first HDM exposure. Cells were lysed for evaluation of NIK, p52, and phospho RelA 536, by Western blot analyses. β -Actin (loading control). Right panels: Densitometric evaluation of Western blots shown in (A) PHNE ($n = 3$ patients) or (B) HBE ($n = 3$ experimental repeats). Results are expressed as arbitrary density and were normalized to corresponding β -actin bands.

neutrophilic/eosinophilic, Th2/Th17-linked inflammation, as well as production of IL-13 and IL-17A (24). In this study, the preferential diminution of neutrophils following inhibition of epithelial NF- κ B was associated with normalization of G and H parameters toward values observed in controls; in contrast, Rn, reflective of the central airways, was unaffected. The disparate effects of the CC10-I κ B α_{SR} transgene on G and H, compared with Rn, are puzzling. Previous studies have suggested that inflammation contributed to enhanced leakage of fibrin to the airway surface, leading to decreased stability of surfactant proteins and increased surface tension, and facilitated contractility of smooth muscle. These mechanisms have been linked to increases in the closure of distal airways (25–27). It is plausible that decreases in inflammation, along with decreased α -SMA in bronchioles, account for decreases in G and H, which were observed in CC10-I κ B α_{SR} transgenic mice. This putative explanation would need to be addressed with additional studies. Our data also demonstrate that KC, a potent neutrophil chemokine, was not increased in CC10-I κ B α_{SR} HDM-exposed mice in comparison with PBS controls, indicating a potential mechanism whereby neutrophils are decreased in CC10-I κ B α_{SR} mice. Altogether, these findings are suggestive of a role for neutrophils in promoting increases in peribronchiolar collagen deposition and associated changes in G and H. Alternatively, it is also plausible that other mediators control neutrophil trafficking to the airways, promote peribronchiolar remodeling, and produce changes in AHR. In this regard, IL-17A, which we also demonstrate to be decreased in the tissue of CC10-I κ B α_{SR} mice, has been shown to stimulate production of chemokines important in neutrophil recruitment in a KC-dependent manner (28) and has been implicated in the pathogenesis of pulmonary fibrosis (29); its functional contribution, however, remains to be determined. In addition, eosinophils expressing TGF- β 1 have been shown to be important in allergen-induced peribronchial fibrosis (30); hence it is possible that TGF- β 1 also plays a role in peribronchial fibrosis downstream of activation of NF- κ B. Additional studies will be required to formally address these putative scenarios.

In addition to demonstrating a role for epithelial NF- κ B in promoting inflammation, AHR, and fibrosis following 15 challenges of HDM, we also established a contributing role for epithelial NF- κ B in regulating proinflammatory gene expression following a single exposure of HDM. Numerous proinflammatory mediators have been implicated in the development of allergic airways disease (3), and our studies demonstrate a putative role for epithelial NF- κ B in regulating expression of several of these molecules in response to HDM. GM-CSF and CCL20 have been shown to be important in the recruitment/activation of dendritic cells, which are required for T cell activation and recognition of Ags. Notably, exposure of human asthmatic bronchial epithelial cells to the HDM component, Derp1, was shown to be important in dendritic cell recruitment in a CCL20-dependent manner (31). In addition, IL-33 has been shown to be important in the activation of a variety of cell types crucial to the development of asthma, such as T helper type 2 cells, eosinophils, dendritic cells, and mast cells (32). As previously mentioned, the activation and infiltration of neutrophils is emerging as a potential phenotype in severe, steroid-resistant asthma. Cytokines KC and MIP-2 have both been shown to be important in the recruitment of neutrophils and production of HDM-specific IgE (33). Altogether, our data indicate a likely role for epithelial NF- κ B in the recruitment/activation of several cell types important for the development of HDM-induced asthma. This observation suggests a mechanism whereby epithelial NF- κ B activation is the crucial step between contact with an allergen and downstream manifestations of asthma.

In addition to the role of NF- κ B in airway epithelium demonstrated in this study, it is plausible that activation NF- κ B in other cell types contributes to the pathophysiology of allergic airways disease. Notably, our findings demonstrate increased immunofluorescence of nuclear RelA and RelB in parenchymal regions following exposure to HDM (Fig. 3). Unraveling the cell types in which NF- κ B is activated and their functional contribution to allergic airways disease would require additional cell-specific labeling and targeting strategies, which were beyond the scope of the current study. For example, neutrophil elastase-induced secretion of TGF- β 1 from smooth muscle cells was shown to be dependent on NF- κ B activation (34), and the smooth muscle contractile force in response to IL-17A was dependent on NF- κ B activation (24), suggesting a putative role of NF- κ B activation in smooth muscle cells in AHR and remodeling. Secretion of eotaxin, a potent eosinophil-activating factor, by fibroblasts was also shown to be dependent on NF- κ B activation (35). One notable finding of the current study is that HDM activates both the classical and the alternative NF- κ B pathway within the bronchiolar epithelium, as evidenced by the increases in the nuclear presence of both RelA and RelB in mice exposed to HDM. Similar to observations in this study, we recently demonstrated increases in nuclear RelA and RelB in the parenchymal regions following administration of LPS, a component of HDM (7), suggesting that TLR4 activation by HDM may be contributing to the observed increases in RelA and RelB observed. Increases in nuclear RelA and RelB within the epithelium occurred rapidly and were sustained for at least 24 h after three challenges. However, 72 h post 5, 10, or 15 challenges, there was no clear evidence of increased RelA or RelB in the bronchiolar epithelium, possibly owing to the timing of tissue analysis after the last administration of HDM, which was 72 h, a time when NF- κ B activation may have resolved. However, we did observe sustained increases in nuclear RelB in lung tissue homogenates 72 h after 15 challenges with HDM in WT mice, suggestive of NF- κ B activation in other cell types. Increases in nuclear RelB were attenuated in CC10-I κ B α_{SR} mice, suggesting a putative role for RelB in the orchestration of HDM-mediated AHR and fibrotic remodeling. In addition to the demonstration that RelB was increased in the bronchiolar epithelium following HDM exposure, we also demonstrated that HDM directly activated both NF- κ B pathways in both human bronchial and nasal epithelial cells. Although extensive studies have been conducted to unravel the molecular regulation and pathophysiological relevance of the classical NF- κ B pathway, far less information is available for the alternative pathway. The latter pathway was originally thought to play a role in adaptive immune responses and in development of lymphocytes and lymphoid organs. However, emerging studies have pointed to a coordinate function of both classical and alternative NF- κ B in the orchestration of proinflammatory responses. For example, exposure of lung epithelial cells with TNF- α , polyinosinic acid, LPS, IL-17A, lipoteichoic acid, and CD40L, agonists that signal through distinct families of receptors, led to a coordinate activation of classical and alternative NF- κ B pathways and subsequent proinflammatory responses (7). In contrast, adenovirus-mediated delivery of RelB afforded protection against cigarette smoke-induced neutrophilic inflammation (36), suggesting potentially complex roles for the alternative NF- κ B pathway in the regulation of proinflammatory and immune responses. The classical NF- κ B pathway previously has been shown to be important in HDM-mediated proinflammatory responses in HBE cells *in vitro* (12), and Derp1, a component of HDM, was shown to be important in the activation of NF- κ B in human asthmatic bronchial epithelial cells (37). Patients with allergic asthma demonstrate increased classical NF- κ B

activation in nasal epithelial cells in response to HDM in comparison with healthy controls (38). Intriguingly, the HDM component, β -glucan, was shown to be crucial for activation of allergic rhinitis in nasal epithelial cells via TLR2, in contrast to another HDM component, LPS, which is important in promoting allergic airways disease via activation of TLR4 in bronchial epithelial cells (39). Despite these previous studies and data presented in this study, additional studies are needed to better understand the timing and locale of activation of classical and alternative NF- κ B pathways, the components of HDM that are required to activate either pathway, as well as the relative contributions of the classical and alternative NF- κ B pathways in eliciting HDM-triggered allergic airways disease.

In summary, we demonstrate in the current study the importance of NF- κ B activation within the bronchiolar epithelium in HDM-induced inflammation, AHR, and fibrotic airway remodeling. We also demonstrate that both classical and alternative NF- κ B pathways are activated by HDM. Data presented in this study showed that in the setting of HDM-induced allergic airways disease, inhibition of epithelial NF- κ B plays a more prominent role in attenuating neutrophilia, AHR, and remodeling, in comparison with eosinophilia, IgE, and mucus metaplasia. Therefore, it is plausible that therapeutic approaches that target NF- κ B via interference with degradation of I κ B α may have a stronger impact on asthmatic patients with predominant neutrophilia in contrast to patients with predominant eosinophilia. Corticosteroids, the most common therapy for asthma, inhibit NF- κ B, but have many off-target effects. Current therapies that are being developed for asthma are aimed at inhibiting the proinflammatory effects of NF- κ B signaling in the lung (40) and are focused on inhibition of IKK β , the dominant kinase in the classical NF- κ B pathway (40). The small-molecule IKK β inhibitor, IMD-0354, attenuated HDM-induced eosinophilia, goblet cell hyperplasia, subepithelial fibrosis, smooth muscle cell hypertrophy, and lung resistance using an i.p. sensitization model (41). The disparate findings of the latter study relative to the present findings may relate to the different sensitization route, the mechanism of inhibition of NF- κ B, and the cell types wherein NF- κ B inhibition occurred. Furthermore, i.v. administration of RelA antisense oligonucleotides prior to OVA challenge resulted in dampened responses of inflammation, AHR, and TH2 responses in mice (42). On the basis of these collective findings, therapeutics designed to inhibit both facets of the NF- κ B pathway may hold larger therapeutic potential for allergic asthma and other allergic diseases of the lung.

Acknowledgments

We thank the University of Vermont Microscopy Imaging Center and the Vermont Lung Center, Burlington, VT, for assistance with these studies.

Disclosures

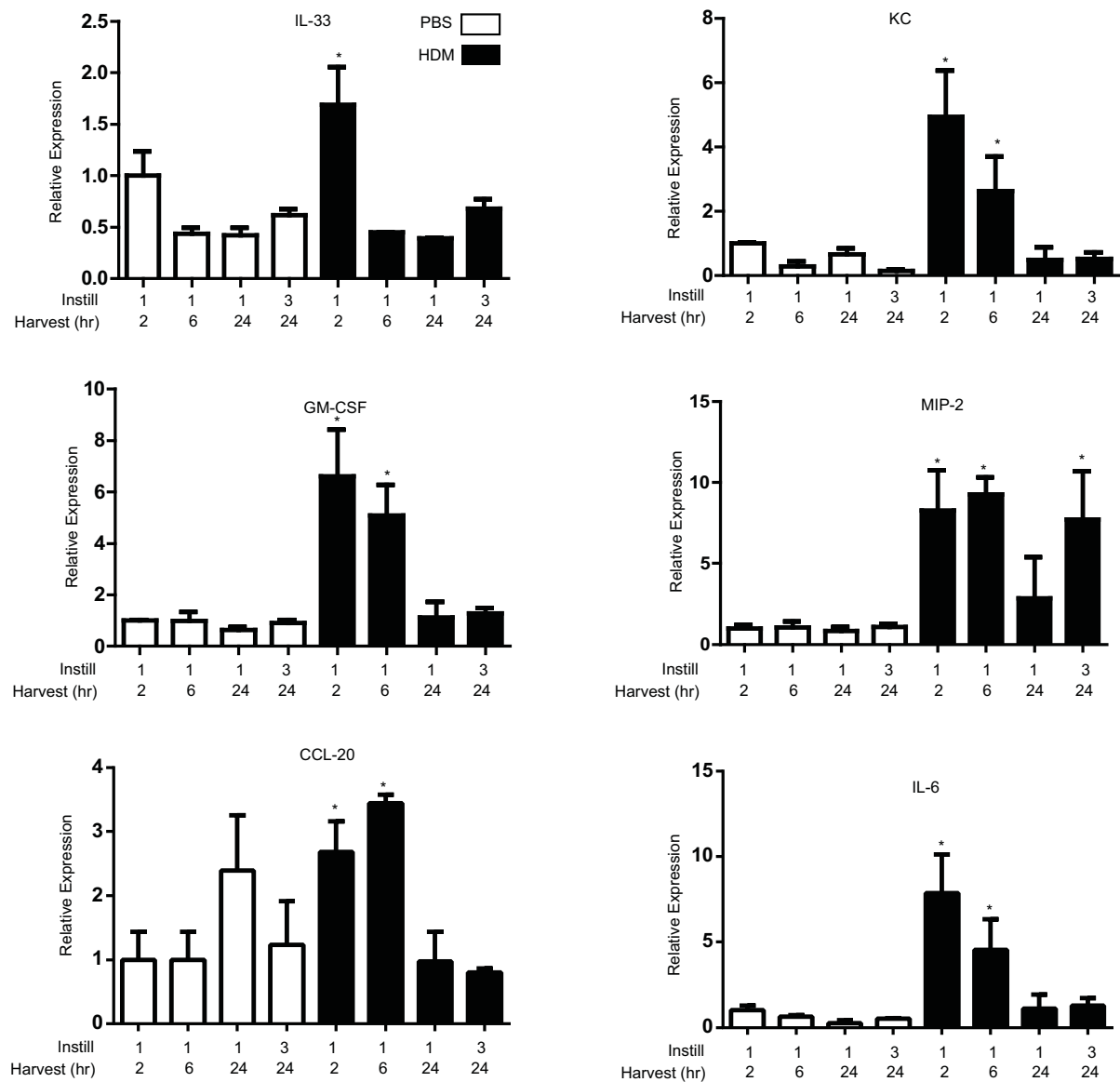
The authors have no financial conflicts of interest.

References

- Hayden, M. S., and S. Ghosh. 2004. Signaling to NF-kappaB. *Genes Dev.* 18: 2195–2224.
- Scheidereit, C. 2006. IkappaB kinase complexes: gateways to NF-kappaB activation and transcription. *Oncogene* 25: 6685–6705.
- Swamy, M., C. Jamora, W. Havran, and A. Hayday. 2010. Epithelial decision makers: in search of the 'epimunome'. *Nat. Immunol.* 11: 656–665.
- Senftleben, U., Y. Cao, G. Xiao, F. R. Greten, G. Krähn, G. Bonizzi, Y. Chen, Y. Hu, A. Fong, S. C. Sun, and M. Karin. 2001. Activation by IKK α of a second, evolutionary conserved, NF-kappa B signaling pathway. *Science* 293: 1495–1499.
- Oeckinghaus, A., M. S. Hayden, and S. Ghosh. 2011. Crosstalk in NF- κ B signaling pathways. *Nat. Immunol.* 12: 695–708.
- Xiao, G., A. B. Rabson, W. Young, G. Qing, and Z. Qu. 2006. Alternative pathways of NF-kappaB activation: a double-edged sword in health and disease. *Cytokine Growth Factor Rev.* 17: 281–293.
- Tully, J. E., J. D. Nolin, A. S. Guala, S. M. Hoffman, E. C. Roberson, K. G. Lahue, J. van der Velden, V. Anathy, T. S. Blackwell, and Y. M. Janssen-Heininger. 2012. Cooperation between classical and alternative NF- κ B pathways regulates proinflammatory responses in epithelial cells. *Am. J. Respir. Cell Mol. Biol.* 47: 497–508.
- Poynter, M. E., C. G. Irvin, and Y. M. Janssen-Heininger. 2003. A prominent role for airway epithelial NF-kappa B activation in lipopolysaccharide-induced airway inflammation. *J. Immunol.* 170: 6257–6265.
- Poynter, M. E., R. Cloots, T. van Woerkom, K. J. Butnor, P. Vacek, D. J. Taatjes, C. G. Irvin, and Y. M. Janssen-Heininger. 2004. NF-kappa B activation in airways modulates allergic inflammation but not hyperresponsiveness. *J. Immunol.* 173: 7003–7009.
- Broide, D. H., T. Lawrence, T. Doherty, J. Y. Cho, M. Miller, K. McElwain, S. McElwain, and M. Karin. 2005. Allergen-induced peribronchial fibrosis and mucus production mediated by IkappaB kinase beta-dependent genes in airway epithelium. *Proc. Natl. Acad. Sci. USA* 102: 17723–17728.
- Gregory, L. G., and C. M. Lloyd. 2011. Orchestrating house dust mite-associated allergy in the lung. *Trends Immunol.* 32: 402–411.
- Osterlund, C., H. Grönlund, N. Polovic, S. Sundström, G. Gafvelin, and A. Bucht. 2009. The non-proteolytic house dust mite allergen Der p 2 induce NF-kappaB and MAPK dependent activation of bronchial epithelial cells. *Clin. Exp. Allergy* 39: 1199–1208.
- Olson, N., A. K. Greul, M. Hristova, P. F. Bove, D. I. Kasahara, and A. van der Vliet. 2009. Nitric oxide and airway epithelial barrier function: regulation of tight junction proteins and epithelial permeability. *Arch. Biochem. Biophys.* 484: 205–213.
- Wu, R., Y. H. Zhao, and M. M. Chang. 1997. Growth and differentiation of conducting airway epithelial cells in culture. *Eur. Respir. J.* 10: 2398–2403.
- Jaspers, I., J. M. Cieniewicz, W. Zhang, L. E. Brighton, J. L. Carson, M. A. Beck, and M. C. Madden. 2005. Diesel exhaust enhances influenza virus infections in respiratory epithelial cells. *Toxicol. Sci.* 85: 990–1002.
- Riesenfeld, E., G. B. Allen, J. H. Bates, M. E. Poynter, M. Wu, S. Aimian, and L. K. Lundblad. 2012. The temporal evolution of airways hyperresponsiveness and inflammation. *J. Allergy Ther.* 1: 1–7.
- Pantano, C., J. L. Ather, J. F. Alcorn, M. E. Poynter, A. L. Brown, A. S. Guala, S. L. Beuschel, G. B. Allen, L. A. Whittaker, M. Bevelander, et al. 2008. Nuclear factor-kappaB activation in airway epithelium induces inflammation and hyperresponsiveness. *Am. J. Respir. Crit. Care Med.* 177: 959–969.
- Poynter, M. E., C. G. Irvin, and Y. M. Janssen-Heininger. 2002. Rapid activation of nuclear factor-kappaB in airway epithelium in a murine model of allergic airway inflammation. *Am. J. Pathol.* 160: 1325–1334.
- Zhao, S., Y. Qi, X. Liu, Q. Jiang, S. Liu, Y. Jiang, and Z. Jiang. 2001. Activation of NF-kappa B in bronchial epithelial cells from children with asthma. *Chin. Med. J. (Engl.)* 114: 909–911.
- Ather, J. L., S. R. Hodgkins, Y. M. Janssen-Heininger, and M. E. Poynter. 2011. Airway epithelial NF-kB activation promotes allergic sensitization to an innocuous inhaled antigen. *Am. J. Respir. Cell Mol. Biol.* 44: 631–638.
- Kung, T. T., H. Jones, G. K. Adams, III, S. P. Umland, W. Kreutner, R. W. Egan, R. W. Chapman, and A. S. Watnick. 1994. Characterization of a murine model of allergic pulmonary inflammation. *Int. Arch. Allergy Immunol.* 105: 83–90.
- Wilson, R. H., G. S. Whitehead, H. Nakano, M. E. Free, J. K. Kolls, and D. N. Cook. 2009. Allergic sensitization through the airway primes Th17-dependent neutrophilia and airway hyperresponsiveness. *Am. J. Respir. Crit. Care Med.* 180: 720–730.
- Whittaker, L., N. Niu, U. A. Temann, A. Stoddard, R. A. Flavell, A. Ray, R. J. Homer, and L. Cohn. 2002. Interleukin-13 mediates a fundamental pathway for airway epithelial mucus induced by CD4 T cells and interleukin-9. *Am. J. Respir. Cell Mol. Biol.* 27: 593–602.
- Kudo, M., A. C. Melton, C. Chen, M. B. Engler, K. E. Huang, X. Ren, Y. Wang, X. Bernstein, J. T. Li, K. Atabai, et al. 2012. IL-17A produced by α β T cells drives airway hyper-responsiveness in mice and enhances mouse and human airway smooth muscle contraction. *Nat. Med.* 18: 547–554.
- Hohlfeld, J. M. 2002. The role of surfactant in asthma. *Respir. Res.* 3: 4.
- Wagers, S. S., R. J. Norton, L. M. Rinaldi, J. H. Bates, B. E. Sobel, and C. G. Irvin. 2004. Extravascular fibrin, plasminogen activator, plasminogen activator inhibitors, and airway hyperresponsiveness. *J. Clin. Invest.* 114: 104–111.
- Yager, D., R. D. Kamm, and J. M. Drazen. 1995. Airway wall liquid. Sources and role as an amplifier of bronchoconstriction. *Chest* 107(3, Suppl):105S–110S.
- Laan, M., Z. H. Cui, H. Hoshino, J. Lötvall, M. Sjöstrand, D. C. Gruenert, B. E. Skoogh, and A. Lindén. 1999. Neutrophil recruitment by human IL-17 via C-X-C chemokine release in the airways. *J. Immunol.* 162: 2347–2352.
- Hasan, S. A., B. Eksteen, D. Reid, H. V. Paine, A. Alansary, K. Johansson, C. Gwozd, K. A. Goring, T. Vo, D. Proud, and M. M. Kelly. 2013. Role of IL-17A and neutrophils in fibrosis in experimental hypersensitivity pneumonitis. *J. Allergy Clin. Immunol.* 131: 1663–1673.
- Cho, J. Y., M. Miller, K. J. Baek, J. W. Han, J. Nayar, S. Y. Lee, K. McElwain, S. McElwain, S. Friedman, and D. H. Broide. 2004. Inhibition of airway remodeling in IL-5-deficient mice. *J. Clin. Invest.* 113: 551–560.
- Pichavant, M., A. S. Charbonnier, S. Taront, A. Bricet, B. Wallaert, J. Pestel, A. B. Tonnel, and P. Gosset. 2005. Asthmatic bronchial epithelium activated by the proteolytic allergen Der p 1 increases selective dendritic cell recruitment. *J. Allergy Clin. Immunol.* 115: 771–778.
- Borish, L., and J. W. Steinke. 2011. Interleukin-33 in asthma: how big of a role does it play? *Curr. Allergy Asthma Rep.* 11: 7–11.
- McKinley, L., J. Kim, G. L. Bolgos, J. Siddiqui, and D. G. Remick. 2005. CXC chemokines modulate IgE secretion and pulmonary inflammation in a model of allergic asthma. *Cytokine* 32: 178–185.

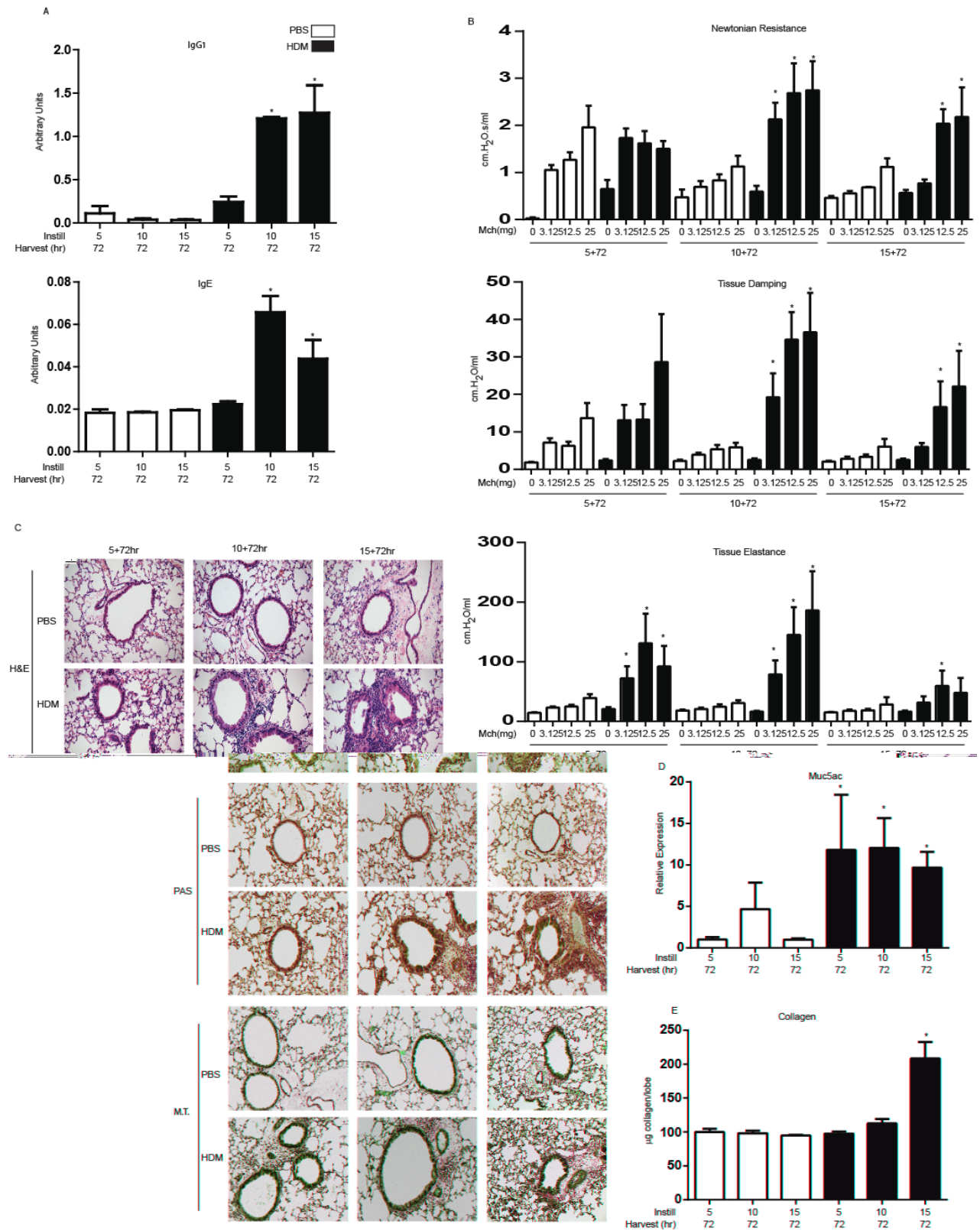
34. Lee, K. Y., S. C. Ho, H. C. Lin, S. M. Lin, C. Y. Liu, C. D. Huang, C. H. Wang, K. F. Chung, and H. P. Kuo. 2006. Neutrophil-derived elastase induces TGF- β 1 secretion in human airway smooth muscle via NF- κ B pathway. *Am. J. Respir. Cell Mol. Biol.* 35: 407–414.
35. Rokudai, A., Y. Terui, R. Kuniyoshi, Y. Mishima, Y. Mishima, E. Aizu-Yokota, Y. Sonoda, T. Kasahara, and K. Hatake. 2006. Differential regulation of eotaxin-1/CCL11 and eotaxin-3/CCL26 production by the TNF- α and IL-4 stimulated human lung fibroblast. *Biol. Pharm. Bull.* 29: 1102–1109.
36. McMillan, D. H., C. J. Baglione, T. H. Thatcher, S. Maggirwar, P. J. Sime, and R. P. Phipps. 2011. Lung-targeted overexpression of the NF- κ B member RelB inhibits cigarette smoke-induced inflammation. *Am. J. Pathol.* 179: 125–133.
37. Stacey, M. A., G. Sun, G. Vassalli, M. Marini, A. Bellini, and S. Mattoli. 1997. The allergen Der p1 induces NF- κ B activation through interference with IkappaB alpha function in asthmatic bronchial epithelial cells. *Biochem. Biophys. Res. Commun.* 236: 522–526.
38. Vroeling, A. B., M. J. Jonker, S. Luiten, T. M. Breit, W. J. Fokkens, and C. M. van Drunen. 2008. Primary nasal epithelium exposed to house dust mite extract shows activated expression in allergic individuals. *Am. J. Respir. Cell Mol. Biol.* 38: 293–299.
39. Ryu, J. H., J. Y. Yoo, M. J. Kim, S. G. Hwang, K. C. Ahn, J. C. Ryu, M. K. Choi, J. H. Joo, C. H. Kim, S. N. Lee, et al. 2013. Distinct TLR-mediated pathways regulate house dust mite-induced allergic disease in the upper and lower airways. *J. Allergy Clin. Immunol.* 131: 549–561.
40. Edwards, M. R., N. W. Bartlett, D. Clarke, M. Birrell, M. Belvisi, and S. L. Johnston. 2009. Targeting the NF- κ B pathway in asthma and chronic obstructive pulmonary disease. *Pharmacol. Ther.* 121: 1–13.
41. Ogawa, H., M. Azuma, S. Muto, Y. Nishioka, A. Honjo, T. Tezuka, H. Uehara, K. Izumi, A. Itai, and S. Sone. 2011. IkB kinase β inhibitor IMD-0354 suppresses airway remodelling in a *Dermatophagoides pteronyssinus*-sensitized mouse model of chronic asthma. *Clin. Exp. Allergy* 41: 104–115.
42. Choi, I. W., D. K. Kim, H. M. Ko, and H. K. Lee. 2004. Administration of antisense phosphorothioate oligonucleotide to the p65 subunit of NF- κ B inhibits established asthmatic reaction in mice. *Int. Immunopharmacol.* 4: 1817–1828.

Figure 1S



Supplemental Figure 1: Pro-inflammatory mediator mRNA expression analysis following HDM exposure. BALB/c mice were exposed to a single dose of either PBS or HDM and harvested 2, 6, or 24 h following challenge. Additionally, mice were challenged for three consecutive days and harvested 24 h following the 3rd challenge. mRNA isolated from homogenized lung tissue was analyzed for expression of IL-33, GM-CSF, CCL20, KC, MIP-2, and IL-6. Data represent 5 mice/group/time point. * $p < 0.05$ (ANOVA) compared to the PBS group at the same time point.

Figure 2S



Supplemental Figure 2: Effects of Repeated HDM Exposure on HDM-specific IgG1 and IgE, AHR and airways remodeling: BALB/c mice were exposed to HDM and harvested at the indicated time points. (A) Assessment of total IgG1 and IgE in serum. (B) Assessment of airway hyperresponsiveness in response to increasing doses (3.125, 12.5, and 25 mg) of methacholine via forced oscillation mechanics. Parameters measured were Newtonian Resistance (Rn), Tissue Damping (G) and Tissue Elastance (H). (C) Histopathological analysis of inflammation (H&E), mucus metaplasia (PAS) and peri-bronchiolar collagen deposition (M.T.) in mice exposed to HDM for the indicated times. scale bar=50 μ m. (D) Analysis of Muc5ac mRNA expression in homogenized lung tissue. Data were normalized to cyclophilin and are presented as relative expression. (E) Quantification of collagen from the upper right lobe. Data represent 5 mice/group/time point. * $p < 0.05$ (ANOVA) compared to the PBS group at the same time point.