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Pattern Recognition Scavenger Receptor A/CD204 Regulates Airway Inflammatory Homeostasis Following Organic Dust Extract Exposures

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Abstract

Exposure to agriculture organic dusts, comprised of a diversity of pathogen-associated molecular patterns, results in chronic airway diseases. The multi-functional class A macrophage scavenger receptor (SRA)/CD204 has emerged as an important class of pattern recognition receptors with broad ligand binding ability. Our objective was to determine the role of SRA in mediating repetitive and post-inflammatory organic dust extract (ODE)-induced airway inflammation. Wild-type (WT) and SRA knockout (KO) mice were intra-nasally treated with ODE or saline daily for 3 wk and immediately euthanized or allowed to recover for 1 wk. Results show that lung histopathologic changes were increased in SRA KO mice as compared to WT following repetitive ODE exposures marked predominately by increased size and distribution of lymphoid aggregates. After a 1-wk recovery from daily ODE treatments, there was significant resolution of lung injury in WT mice, but not SRA KO animals. The increased lung histopathology induced by ODE treatment was associated with decreased accumulation of neutrophils, but greater accumulation of CD4⁺ T-cells. The lung cytokine milieu induced by ODE was consistent with a T_H1/T_H17 polarization in both WT and SRA KO mice. Overall, our data demonstrate that SRA/CD204 plays an important role in the normative inflammatory lung response to ODE as evidenced by the enhanced dust-mediated injury viewed in the absence of this receptor.

Keywords

lung; inflammation; Toll-like receptor; farm; lymphocyte; T_H17

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Introduction

The major macrophage scavenger receptor class A (SRA/CD204), expressed preferentially on myeloid lineage cells but also on airway epithelial cells, is a multi-functional receptor with broad ligand binding potential (de Winther et al., 2000; Wyatt et al., 2005; Seimon et al., 2006; Dieudonne et al., 2012; Canton et al., 2013). SRA is well recognized for its role in atherosclerosis as it binds endogenous ligands including oxidized and acetylated low-density lipoproteins as well as apoptotic cells (de Winther et al., 2000). SRA has emerged as an important pattern recognition receptor because it recognizes several pathogen-associated molecular patterns (PAMPs) including endotoxin, lipoteichoic acid, fungal glucans, bacterial CpG, and double-stranded RNA. SRA has also been found to contribute to Toll-like receptor (TLR)-signaling pathways, and in particular, SRA may limit the inflammatory response to systemic endotoxin through its interaction with TLR4 (Seimon et al., 2006; Areschoug and Gordon, 2009; Yu [X] et al., 2011; Yu [H] et al., 2012).

Consistent with this protective role for SRA, absence of SRA has been associated with worsening outcomes following infections with *Listeria* (Ishiguro et al., 2001), *Neisseria* (Pluddemann et al., 2009), and pneumococcal (Arredouani et al., 2007) infections in mice. In contrast, others have reported the opposite: SRA knockout (KO) mice were protected from surgically induced polymicrobial sepsis (Ozment et al., 2012) and *Cryptococcus neoformans* infections (Qiu et al., 2013). In addition to its role in modulating infectious insults, SRA has been implicated in regulating lung consequences following exposure to environmental particles such as crystalline silica and titanium dioxide (Beamer and Holian, 2005; Thakur et al., 2008). Namely, SRA KO animals were protected from the development of silica-induced lung fibrosis, but yet, lung inflammatory cell accumulation was enhanced (Beamer and Holian, 2005). These data suggest that SRA could play an important role in regulating lung inflammatory outcomes following exposure to occupational organic dusts, which are complex mixtures of particulate matter and a wide-diversity of microbial components.

Chronic exposure to occupational agricultural organic dusts or bioaerosols results in significant airway inflammatory diseases. The chronic airway inflammatory response, which is marked by influx of inflammatory cells (neutrophils, macrophages, and lymphocytes) and pro-inflammatory cytokine/chemokine production, leads to progressive decline in lung function over time (May et al., 2012). Although wearing personnel protective equipment is recommended, respirator compliance is poor and treatment options are limited for symptomatic persons, even after removal from the exposure (May et al., 2012). Recent advances in understanding the host response have focused on acute and sub-chronic exposures; however, there is little information about the recovery phase of repetitive organic dust induced airway inflammation, which may be a more relevant target area for improving disease outcomes once injury has occurred. Important etiologic factors identified within the complex organic dusts include Gram-negative endotoxins, Gram-positive peptidoglycans, fungal glucans, and bacterial CpG DNA (Poole and Romberger, 2012). Moreover, important roles for Toll-like receptor (TLR)-2, TLR4, TLR9, and MyD88 have been reported to mediate organic dust-induced airway inflammatory consequences in mice and humans

(Charavaryamath et al., 2008; Senthilselvan et al., 2009; Poole et al., 2011; Bauer et al., 2013; Gao et al., 2013).

Because SRA/CD204 is a multi-functional receptor that recognizes a diversity of ligands that are predominate within complex agricultural environmental organic dust samples, we hypothesized that SRA would play an important functional role in modulating organic dust extract-induced airway inflammation. We also aimed to investigate the lung response following a 1-wk post-exposure recovery phase, focusing on a potential regulatory role for SRA. Our studies demonstrated that lung histopathologic changes were enhanced with increased numbers of CD4⁺ T-cells following repetitive organic dust extract exposure in SRA KO mice. Increased inflammatory consequences remained pronounced at 1 wk post-exposure in SRA KO mice compared to WT mice. These studies suggest that SRA plays an important role in regulating airway inflammation and the post-inflammatory lung tissue homeostasis with organic dust extract exposures.

Materials and Methods

Animals

C57BL/6 wild-type (WT) mice and SRA (CD204) knockout (KO, *Msr1^{tm1Cskl}/-*) mice on C57BL/6 background were purchased from The Jackson Laboratory (Bar Harbor, ME). All mice were housed in specific pathogen-free facilities maintained at a set-point of $\approx 22^{\circ}\text{C}$ (range of 21–24°C) and a range of 30–70% relative humidity and a 12-hr light:dark cycle. All mice had *ad libitum* access to standard rodent chow and filtered water throughout the course of the studies. All experimental animal procedures were conducted according to the NIH guidelines for the use of rodents, and the University of Nebraska Medical Center Institutional Animal Care and Use Committee approved all procedures.

Organic dust extract (ODE)

Aqueous organic dust extract (ODE) was prepared from settled dust collected from horizontal surfaces (~3 feet above floor) of local swine confinement feeding operations (housed ≈ 400 –600 animals), and dust extracts were batch prepared utilizing previously described methods (Poole et al., 2009). Briefly, dust (10 g) was placed into sterile Hank's Balanced Salt Solution (10 ml; Sigma, St. Louis, MO), incubated at room temperature for 1 hr, and then centrifuged twice for 20 min at 2000g. The final supernatant was filter-sterilized (0.22 μm), a process that removes coarse particles and also yields inhalable particle sizes (equal to and less than 0.22 μm size particles). Stock ODE was aliquoted and frozen until use in experimental assays.

For use in the experiments, stock ODE was diluted to a final concentration of 12.5% (vol/vol) in sterile phosphate-buffered saline (PBS; pH 7.4 [diluent]). The 12.5% ODE has been previously shown to elicit optimal and reproducible lung inflammation in mice and is well-tolerated by these hosts (Poole et al., 2009). Prior work to understand the components within these organic dust extracts have been previously published (Poole et al., 2010) and these previous studies demonstrated the presence of high concentration of muramic acid (component of peptidoglycan) and 3-hydroxy fatty acid (component of endotoxin) and low

concentration of ergosterol (component of fungi) by gas chromatography-tandem mass spectrometry. In these studies, independent batches of ODE were prepared and tested, but due to the biologic complexity of the samples, inherent variability in the samples is inevitable. The range of diluted 12.5% ODE contained 2.91–3.88 mg/ml of total protein as measured by Nanodrop spectrophotometry (NanoDrop Technologies, Wilmington, DE) and confirmed by Bradford protein assay. Endotoxin levels in the 12.5% ODE ranged from 22.1 to 91.1 EU/ml as assayed using the limulus amoebocyte lysate assay according to manufacturer's instruction (Sigma). Peptidoglycan levels were not quantitated, since commercially available measurement assays do not exist, to the best of our knowledge.

Murine Model

Per established protocol (Poole et al., 2009), mice received intra-nasal inhalation of 50 μ l of sterile saline (PBS) or 12.5% ODE. For experimental studies, mice were treated daily for 3 wk (with week-ends excluded) and immediately euthanized (repetitive exposure), or allowed to recover for 1 wk without treatments (recovery studies). No mice exhibited respiratory distress or weight loss throughout the treatment periods.

Bronchoalveolar lavage fluid cell analysis

Bronchoalveolar lavage (BAL) was achieved using 3 \times 1-ml instillates/recoveries of PBS. After pooling the three lavages, the total number of cells/mouse was enumerated and differential cell counts performed using cytopsin-prepared slides (Cytopro Cyto centrifuge, Wescor Inc, Logan, UT) stained with DiffQuick (Siemens, Newark, DE).

Histopathology

Following lung lavage, the whole lungs were excised and slowly inflated (and hung under 20 cm H₂O pressure) with 10% formalin (Sigma) for 24 hr. Next, by routine histology processing, the fixed lung tissue was embedded in paraffin and entire lung sections cut (4–5 μ m) and stained with hematoxylin and eosin (H&E). Each slide was entirely reviewed at scanning magnifications (2X, 4X, and 10X objectives; Nikon Eclipse Model E600 microscope, Nikon, Tokyo, Japan) and semi-quantitatively assessed for the degree and distribution of lung inflammation by a pathologist (W.W.W.), blinded to the treatment conditions, utilizing a previously published scoring system (Poole et al., 2009).

Flow cytometry phenotyping of whole lung cells

Cells were isolated from whole lungs as previously described (Poole et al., 2012a,b). Briefly, 24 hr after the final treatment condition, animals were euthanized, and lung lavage performed. Next, after exposing the chest cavity, the right ventricle was infused with 10 ml sterile PBS to remove blood from the pulmonary vasculature and the lungs were harvested and subjected to an automated dissociation procedure using a gentleMAC Dissociator instrument (according to manufacturer instructions [Miltenyi Biotech, Auburn, CA]) in a solution containing collagenase type I (324 U/mL; Fisher, Pittsburgh, PA), bovine DNase (75 U/ml), porcine heparin (25 U/ml) and PBS with Ca²⁺ and Mg²⁺ (pH 7.4). The resulting solution was passed through a nylon mesh (40 μ m; Thermo Fisher Scientific, Waltham, MA) to remove any large fragments, red blood cells were lysed using a 0.84% (w/v)

ammonium chloride treatment (5 min at 4°C), and after centrifugation at 422g, the remaining cells were re-suspended in PBS and final cell counts obtained using a hemocytometer. Viability of the final cell preparations was assessed by trypan blue exclusion. A LIVE/DEAD Fixable Violet Dead Cell Stain kit (Life Technologies, Carlsbad CA) was also used to assess cell viabilities; ultimately, < 1% of (gated) macrophages and lymphocytes were seen to be dead, with no differences noted between the saline and ODE-treated groups (data not shown). Whole lung cells from each animal were then incubated with anti-CD16/32 (F_c Block, BD Biosciences, San Jose, CA) to minimize non-specific antibody staining, and then stained with monoclonal antibodies (mAb) directed against CD11c, CD11b, IA-b, CD80, CD86, Ly-6G, CD3, CD4, and CD8; parallel cell preparations were treated with appropriate isotype controls (BD Biosciences). Compensation was performed with antibody capture beads (BD Biosciences) stained separately with each individual mAb used in test samples.

The gating strategy for CD11c⁺ macrophages, lymphocytes, and Ly6G⁺ neutrophils are depicted in Supplemental Figure 1 and follow that outlined in previously published reports (Osterholzer et al., 2011; Poole et al., 2012b). Briefly, initial gating on CD45⁺ lung leukocytes excludes debris, and the percentage of respective population (i.e., macrophage, lymphocyte, neutrophil) was determined from CD45⁺ leukocytes. This percentage (determined by flow cytometry) is multiplied by the original hemocytometer count of total cells recovered for each animal. All populations were gated by characteristic forward and side-scatter properties and antibody-specific staining fluorescence intensity using a FACSAria cell sorter system and associated software (BD Biosciences). In each case, a minimum of 25,000 CD45⁺ events/mouse sample was acquired for analysis.

Lung homogenate cytokine assays

Cytokine profiles were also characterized in total lung homogenates. Briefly, total lung homogenates were prepared by homogenizing lung samples in 500 µl sterile PBS, and cell-free homogenates were quantitated for cytokine contents using a cytometric bead array protein multiplex immunoassay array (interleukin [IL]-17A, IL-4, IL-10; BD Biosciences) or by ELISA kits (IL-1β and IL-12p40; R&D Systems, Minneapolis, MN).

Statistical methods

Data are presented as mean and standard error of mean (SEM). Statistical significance was assessed by one-way analysis of variance (ANOVA) and a two-tailed Mann-Whitney test, where appropriate. GraphPad (Version 5.02, La Jolla, CA) software was used. Significance was accepted at p-values < 0.05.

Results

SRA KO mice display increased lung inflammatory histopathology following repetitive ODE treatment

We have previously reported that repetitive ODE treatment induces lung inflammation in C57BL/6 WT animals (Poole et al., 2009); in this study, we sought to determine the role of SRA in regulating ODE-induced lung inflammation. WT and SRA KO mice were

intranasally treated in side-by-side experiments with saline or ODE for 3 wk whereupon whole lung sections were processed and reviewed as described in the Methods. By microscopic review there was a notable increase in inflammatory cells and the size and distribution of lymphoid aggregates in the SRA KO mice as compared to WT mice following repetitive ODE exposures (Figures 1A–1D). The range of dust-induced histopathologic changes were semi-quantitatively assessed by reviewer (pathologist) blinded to treatment conditions; there were significant increases in the graded distribution of lung alveolar inflammation and lymphoid aggregates, but not bronchiolar inflammation, induced by repetitive ODE treatment in SRA KO mice as compared to WT mice (Figure 1E). Thus, these studies demonstrated enhanced pathologic injury in SRA KO mice following repetitive ODE treatment, suggesting a role for SRA in mediating lung inflammation consequences following ODE exposures.

Lung inflammation in SRA KO mice remained increased following a 1-wk recovery after repetitive ODE exposures

In the next set of experiments, we investigated lung inflammatory histopathology following a 1-wk recovery phase (no treatment for one week) following the 3-wk repetitive ODE treatment in WT and SRA KO mice. Though decreased compared to the effects from the 3-wk repetitive ODE treatment, there remained notable evidence of inflammatory cells within the bronchiolar and alveolar compartments with a decreased number of lymphoid aggregates in WT mice at the end of the 1-wk recovery period (Figures 2A–2D). Interestingly, as compared to WT mice, SRA KO animals demonstrated increased lung inflammation at 1 wk post-repetitive ODE treatments (Figures 2A–2C). Significant differences were demonstrated in the range of inflammatory scores of lung lymphoid aggregates, bronchiolar inflammation, and alveolar inflammation between the exposed KO and WT treatment groups (Figure 2D). These studies show that the resolution of lung inflammation was slowed in SRA KO mice following ODE exposures, and suggest that SRA may play a significant role in post-inflammatory homeostasis.

Differences in cell recruitment in lavage fluid demonstrated after daily repetitive ODE and following a 1-wk recovery in SRA KO mice

In these experiments, cellular influx in lavage fluid was determined in WT and SRA KO mice after daily repetitive ODE (or saline) and following a 1-wk recovery. There were significant increases in total cells, neutrophils, macrophages, and lymphocytes after repetitive ODE treatments as compared to saline in the WT and KO mice (Figure 3). Following the 1-wk recovery, total cell influx marked by macrophages and lymphocytes, but not neutrophils, remained elevated as compared to in the saline control groups (Figure 3). In comparing ODE-treated WT animals with SRA KO mice (Figure 3), there was no significant difference in total cell influx at either time point, but there were slight, but significant, differences observed with the numbers of macrophages, neutrophils, and lymphocytes. Namely, SRA KO mice demonstrated less neutrophil influx as compared to WT mice after the repetitive exposures, yet increased macrophages and lymphocytes after the 1-wk recovery. These data suggest the role of SRA may be more important to specific cells (macrophages and lymphocytes) as opposed to all inflammatory cells in the airways.

Total lung cells and lung neutrophils following repetitive ODE and 1-wk recovery in SRA KO mice

To further delineate the inflammatory cellular effect observed with SRA KO mice treated with ODE as compared to with WT hosts, whole lung cells following treatment conditions were dissociated and analyzed as described in the Methods; overall gating strategy is shown in Supplemental Figure 1. Repetitive ODE treatment resulted in increased total whole lung cell influx and increased neutrophil influx as compared to saline treatment (Figure 4). Although trends for decreased total lung cells were observed in WT mice following the 1-wk recovery as compared to after the repetitive ODE treatment, this did not reach statistical significance. There was also no difference in total lung cells between WT and SRA KO ODE treatment groups. Neutrophil numbers were significantly decreased ($p < 0.001$) after the 1 wk recovery. The data suggest there were no pronounced differences in total lung cells between the WT and SRA KO mice following ODE treatments, but that neutrophil counts were diminishing after the 1-wk post-exposure respite.

Exudative CD11c^{hi}/CD11b^{hi} macrophages are the most prevalent macrophage population in lungs of WT and SRA KO mice following repetitive ODE exposure and following a 1-wk recovery

In previous studies, we found that repetitive ODE treatment induces the influx of lung exudative CD11c^{hi}/CD11b^{hi} macrophages (Poole et al., 2012b). Because SRA was initially described as a macrophage receptor, we had hypothesized that it would play an important role in ODE-induced lung inflammation via its macrophage immunomodulatory properties. Consistent with that prior work, repetitive ODE treatment resulted in increased levels of lung exudative CD11c^{hi}/CD11b^{hi} macrophages; however, in this current study, we report that the exudative macrophages remain significantly increased following a 1-wk recovery (Figure 5). Further, there were no significant differences in the number of exudative CD11c^{hi}/CD11b^{hi} macrophages between the WT and SRA ODE repetitive and recovery treated groups. The numbers of alveolar macrophages (CD11c^{hi}/CD11b^{lo} macrophages) were decreased in WT animals as compared to SRA KO mice after the 1-wk recovery. We also investigated cell surface expression of MHC Class II (murine I-A^b) and co-stimulatory CD80 and CD86 molecules on these alveolar and exudative macrophages (Supplemental Figure 2; the numbers of exudative CD11c^{hi}/CD11b^{hi} macrophages in saline-treated animals were unsurprisingly too low for adequate flow cytometric analysis). Consistent with prior studies, repetitive ODE treatment was associated with increased CD80, CD86, and I-A^b cell surface expression on exudative CD11c^{hi}/CD11b^{hi} macrophages in WT animals (data not shown), and at one week recovery CD80, CD86, and I-A^b expression remains enhanced (Supplemental Figure 2). Though there were subtle differences in I-A^b expression (less robust cell surface expression in SRA KO mice), there were no significant differences overall in CD80, CD86, and I-A^b surface expression on exudative CD11c^{hi}/CD11b^{hi} macrophages between the WT and SRA ODE repetitive and recovery treatment groups. Collectively, these data did not support our hypothesis that the macrophage was significantly dysregulated in SRA KO animals as an explanation for the enhanced lung inflammation these hosts displayed following ODE treatments (Figures 1 and Figures 2).

CD4⁺ lymphocyte recruitment is increased in SRA KO mice following repetitive ODE exposure and remains elevated following a 1-wk recovery

Because histopathology demonstrated enhanced lymphoid aggregates (Figures 1 and 2) following ODE treatments, infiltrating lung T-cell phenotypes were investigated by flow cytometry. Consistent with prior reports (Ivanov et al., 2005; Poole et al., 2012a), repetitive ODE treatments resulted in increased levels of CD3⁺CD4⁺ T-cells; moreover, levels of these cells remained increased after 1 wk of recovery (Figure 6). Interestingly, SRA KO mice demonstrated significantly increased numbers of CD3⁺CD4⁺ T-cells following ODE repetitive and recovery treatment periods as compared to WT animals (Figure 6). Although CD3⁺CD8⁺ T-cells were slightly increased following ODE as compared to saline treatment, this did not reach statistical significance in WT mice. There were more CD3⁺CD8⁺ T-cells in SRA KO mice following the 1-wk recovery as compared to in the WT mice. These studies suggest that infiltrating CD4⁺ T-cells may play an important role in explaining the enhanced lung inflammatory response in SRA KO mice following ODE treatment.

ODE exposure induces a T_H17/T_H1-polarized cytokine response in the whole lung

Next we investigated the cytokine milieu in lung homogenates, and found evidence of both a T-helper (T_H)-17 and T_H1 cell responses induced by repetitive ODE treatment in both WT and SRA KO animals (Figure 7). There were no significant differences in the cytokine milieu between groups with the exception that IL-17A remained significantly elevated in SRA KO mice as compared to WT following one week recovery from ODE treatments (Figure 7).

Discussion

Our studies demonstrate that SRA plays a significant role in regulating complex organic dust extract-induced lung injury, and that SRA KO mice recover less effectively as compared to control animals, implicating a role for SRA in post-inflammatory homeostasis. SRA KO mice repetitively treated with ODE demonstrated increased lung histopathology changes marked by increased lymphoid aggregates and CD4⁺ T-cell infiltration. This lung inflammatory process remained significantly increased in SRA KO mice following one week of recovery post exposure. The T-cell polarization, consistent with a T_H1/T_H17 skewing, was similar between SRA and WT animals following the ODE treatments. Collectively, these are the first studies to establish a potential protective functional role for SRA in repetitive organic dust extract-induced airway inflammation, and moreover, propose its role in regulating post-inflammatory recovery.

Recent studies of occupational/agricultural organic dusts and dust extract exposure-induced airway disease have led to an appreciation for the presence of a wide diversity of microbial components, particularly Gram-positive as well as Gram-negative bacterial cell wall components (May et al., 2012). Although most studies focus on swine confinement facility environments due to its strong pro-inflammatory airway response and associated lung disease manifestations, other industrial animal farm environments such as dairy farming have also been associated with a high predominance of Gram-positive and Gram-negative bacterial cell wall components (Poole et al., 2010). Roles for TLR2 that recognizes

lipoteichoic acids and peptidoglycans, and TLR4 that recognizes endotoxin have been described (Charavaryamath et al., 2008; Senthilselvan et al., 2009; Poole et al., 2011; Gao et al., 2013). Namely, TLR2 and TLR4 KO mice have blunted airway inflammatory responses following acute and sub-chronic swine facility dust/dust extract exposures (Charavaryamath et al., 2008; Poole et al., 2011; Bauer et al., 2013), and polymorphisms in the TLR2 and TLR4 have been associated with lung function consequences in exposed swine workers (Senthilselvan et al., 2009; Gao et al., 2013). However, the ODE-induced airway inflammatory response is not completely abrogated in TLR2 and TLR4 KO mice (Charavaryamath et al., 2008; Poole et al., 2011; Bauer et al., 2013), and moreover, direct comparisons with lipopolysaccharide at concentrations found in ODE did not replicate the magnitude of lung inflammatory responses induced by ODE (Dusad et al., 2013). Thus, there may not be one critical ligand responsible for eliciting complex organic dust extract-induced disease. Indeed, MyD88, the common adaptor protein for all TLR pathway signaling responses (except TLR3), appears to be the central pathway mediating ODE-induced airway inflammation because there was a near absent airway response following acute dust challenge in MyD88 KO mice (Bauer et al., 2013). However, targeting MyD88 may not be a viable potential preventative or therapeutic option due to its critical role in overall host defense.

Thus, due to its broad ligand binding affinity, SRA might be an important pattern recognition receptor to target in organic dust-induced airway inflammatory diseases because the dust is noted to contain an abundance of pathogen-associated molecular patterns (PAMPs) (Canton et al., 2013). Although there are reports that knocking out SRA results in improved outcomes following polymicrobial sepsis (Ozment et al., 2012) and *Cryptococcus neoformans* lung infection (Qiu et al., 2013), several studies have suggested the opposite. Namely, several studies have reported that SRA acts as an immunosuppressive molecule because infectious consequences were worsened in the absence of SRA (Seimon et al., 2006; Areschoug and Gordon, 2009; Yi et al., 2009; Ohnishi et al., 2011; Yu X et al., 2011; Yu H et al., 2012). The data here are entirely consistent with these later observations in that SRA may play an important role in the limiting airway inflammatory responses. We interpret our findings to be mixed in comparison to results demonstrated in a non-infectious, silica exposure model (Beamer and Holian, 2005). Specifically, absence of SRA was protective against silica-induced lung fibrosis development but not against inflammatory cell infiltration. It is likely that these differences are not explained by one critical ligand, but instead by the complexity of occupational/agricultural organic dusts and dust extracts that contain a wide diversity of PAMPs and particulate matter.

The remarkable finding was the enhanced histopathologic changes following repetitive ODE exposures that remained elevated after a 1 wk recovery phase in the SRA KO mice. The dominant cell type to explain this difference was CD4⁺ T-cells (Figure 6). Yi et al. (2012) also showed that SRA KO mice developed more robust CD4⁺ T-cell responses after OVA immunization, with a concomitant increase in IL-12 production. Lung CD4⁺ T-cell accumulation has also been demonstrated following cryptococcal lung infections in SRA KO animals (Qiu et al., 2013). Our group has previously reported that repetitive ODE exposure results in a cytokine milieu consistent with a T_H1/T_H17 polarized lung response,

and in this study, a T_H1/T_H17 response was confirmed in the WT animals and also extended to the SRA KO mice, but no predominant difference in the lung cytokine milieu was found between groups. Neutrophil infiltration was not enhanced in SRA KO mice, but rather, neutrophil recruitment was diminished. Others have observed no effect on neutrophil recruitment in SRA KO mice challenged with infectious insults (Ozment et al., 2012). SRA is not expressed on neutrophils (Yamamoto et al., 1999), and therefore, regulation of neutrophil functions are likely not directly dependent on SRA.

Exudative macrophages (i.e., $CD11c^{hi}CD11b^{hi}$) (Osterholzer et al., 2009, 2011) with an increased expression of co-stimulatory molecules (CD80 and CD86) and MHC Class II expression are normal, protective responses following infectious and organic dust-induced inflammatory insults in the lung (Poole et al., 2012b). Others have found that expression markers (co-stimulatory molecules) are up-regulated on classically activated lung macrophages following lung infections as a potential explanation for improved host defense in SRA KO mice (Ozment et al., 2012). Here, the number of lung exudative macrophages did not differ significantly in SRA KO mice treated with ODE as compared to WT animals, and the pattern of expression markers also did not differ. These results might suggest that the mechanism by which the macrophage SRA/CD204 receptor alters organic dust-induced inflammatory responses is indirectly through its modulation of T-cell recruitment instead of direct changes to the lung macrophage response.

Prior research investigating effects of occupational exposure to agricultural organic dusts or bioaerosols has focused on acute and repetitive dust exposure scenarios. In contrast, information regarding post-inflammatory recovery events is lacking but represents a novel area to potentially intervene, as symptomatic workers are more apt to seek medical care after disease has established. For these reasons, we investigated the normative post-inflammatory response to repetitive ODE exposures in WT mice and compared that response to SRA KO animals. At 1 wk after the repetitive exposures, the inflammatory lung consequences in WT mice persisted marked by significant lung histopathology and increased infiltration of $CD4^+$ T-cells and exudative $CD11c^{hi}CD11b^{hi}$ macrophages. However, there was clear evidence of a resolving neutrophil response, and IL-17 (but not IL-12) levels were also diminishing at 1 wk post-exposure. This association of IL-17 and neutrophil responses might be expected because T_H17 responses have been closely associated with neutrophilic inflammation (Korn et al., 2009). Although subtle, it is noted that neutrophil response was slightly lower in the ODE-treated SRA KO mice despite higher levels of IL-17A (Figures 4 and 7), suggesting that the IL-17 response is not the sole driver of neutrophil inflammation following ODE exposures. The inflammatory differences between WT and SRA KO mice appeared stronger in the post-recovery experiments. In the SRA KO mice, resolution of neutrophil infiltration was observed, but levels of $CD4^+$ T-cells, $CD8^+$ T-cells, and macrophages remained strikingly elevated. These data support the role of involvement of SRA in immune homeostasis.

In summary, this study demonstrated several new features important in understanding organic dust-induced airway inflammation. First, SRA appeared to serve a functional role in regulating repetitive organic dust-induced airway inflammation as evident by worsening lung histopathology in the absence of SRA. As opposed to an overall global increase in

inflammatory cellular infiltration; the enhanced response was associated with a robust CD4⁺ T-cell influx. Even more pronounced was the enhanced inflammatory response following a post-exposure, 1-wk recovery in SRA KO mice. Collectively, SRA/CD204 is a functionally diverse pattern recognition receptor that might be exploited to impact health consequences in organic dust exposed workers. There also may be merit in future studies to examine if genetic polymorphisms in SRA contribute to airway responses in exposed workers.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Declaration of interest

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Abbreviations

AHR	airway hyper-responsiveness
BALF	bronchoalveolar lavage fluid
COPD	chronic obstructive pulmonary disease
IL	interleukin
KO	knockout
ODE	swine confinement organic dust extract
PAMP	pathogen-associated molecular pattern
SRA	scavenger receptor A, CD204
TLR	toll-like receptor
TNFα	tumor necrosis factor- α
WT	wild-type

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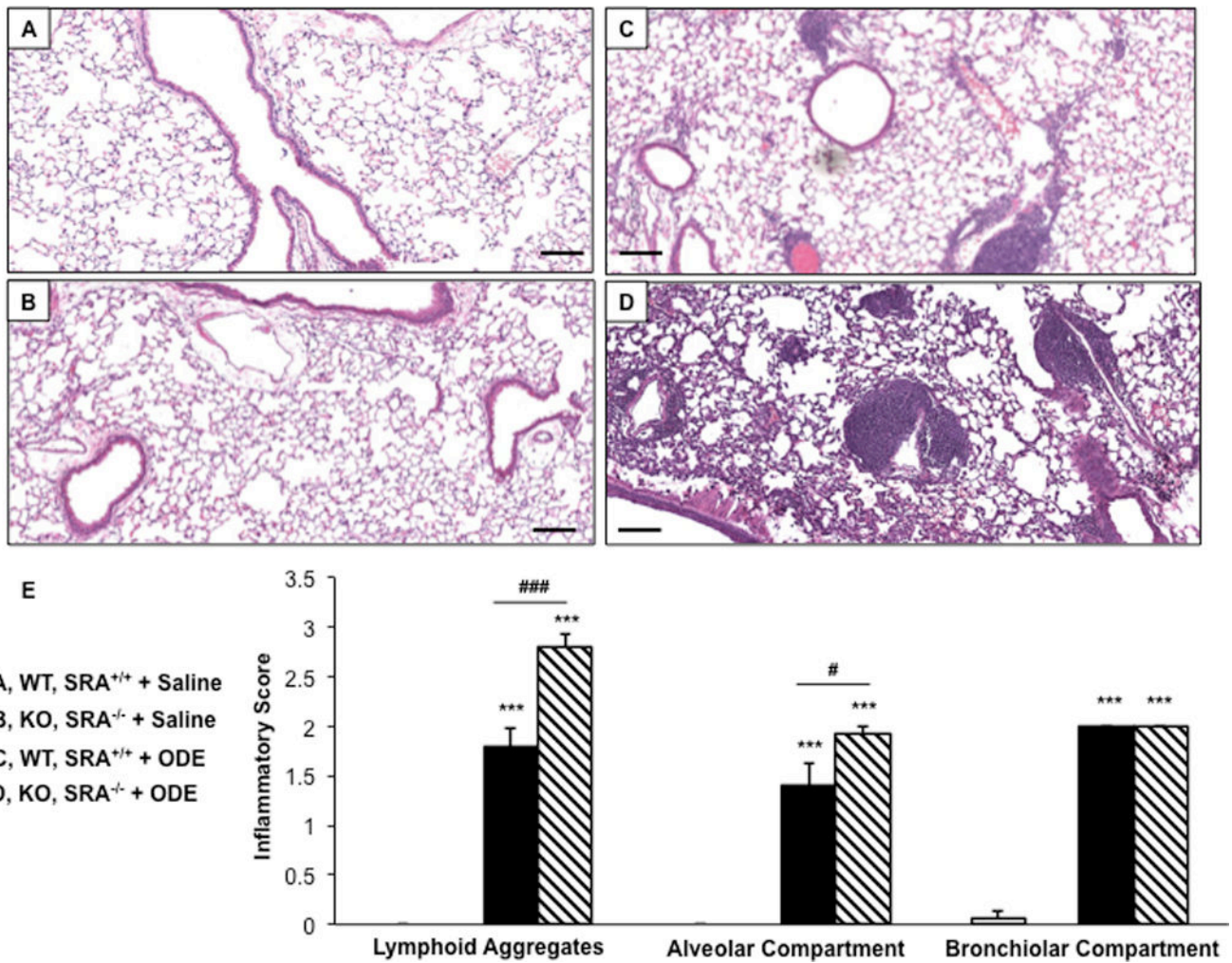


Figure 1. Lung inflammation in SRA knockout mice after repetitive ODE exposures
Mice were intra-nasally treated with saline or organic dust extract (ODE) daily for 3 wk. A representative 4–5- μ m thick section (H&E stained) of one mouse per treatment group is shown (10 \times magnification). (A) Wild-type (WT) SRA^{+/+} + saline. (B) Knockout (KO) SRA^{-/-} + saline. (C) WT + ODE. (D) KO + ODE. (E) Mean semi-quantitative distribution of inflammatory scores of lung lymphoid aggregates, alveolar inflammation and bronchiolar inflammation in mice (n = 4–5 mice/group). Error bars represent SE. Statistically significant (***) $p < 0.001$ vs. saline. # $p < 0.05$ and ### $p < 0.001$: significant differences between groups as indicated (1-way ANOVA with Tukey’s *post-hoc* comparison). Line scale represents 100 μ m.

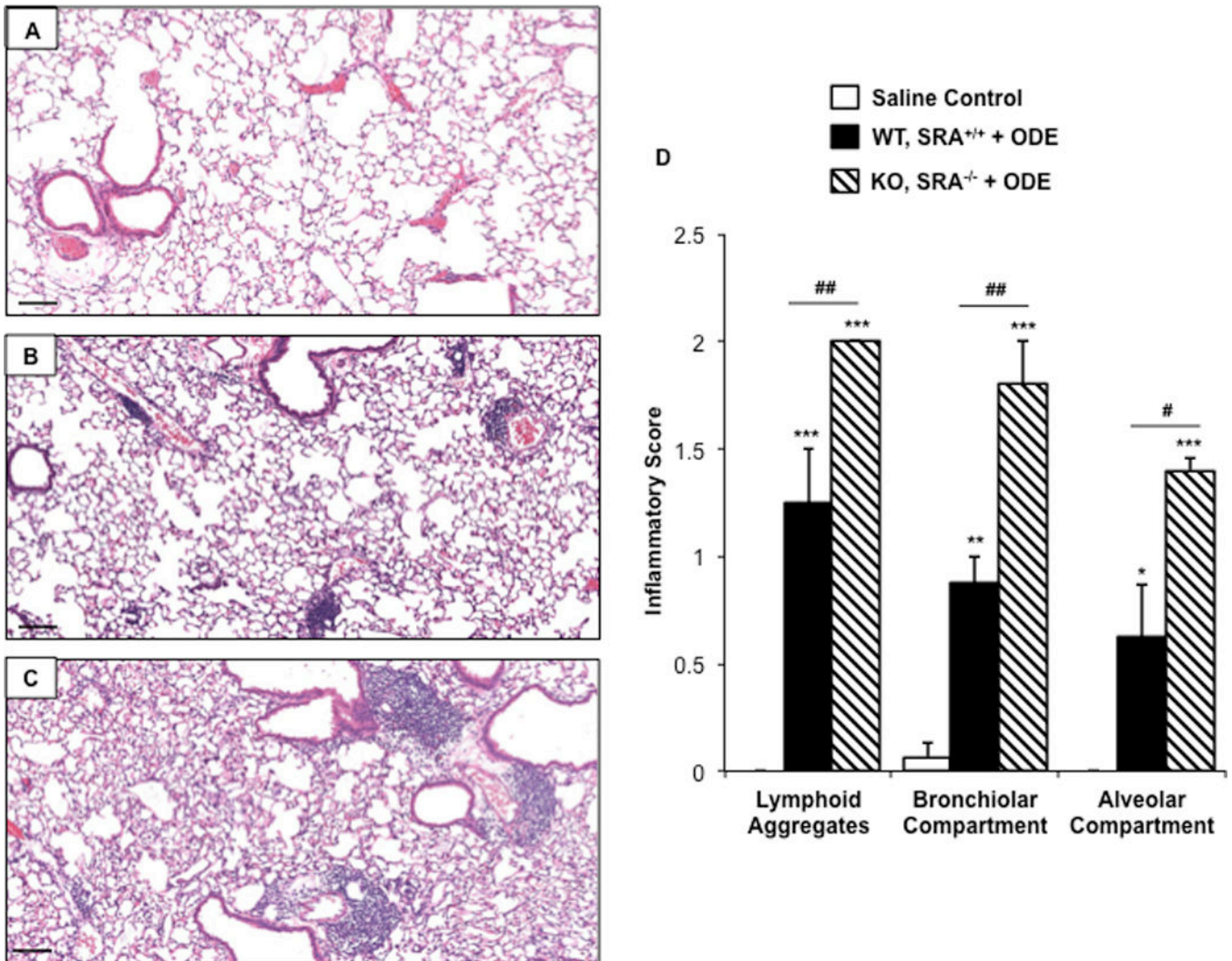


Figure 2. Lung inflammation in SRA KO following 1 wk recovery after repetitive ODE exposures

Mice intra-nasally treated with saline or ODE daily for 3 wk and subsequently rested (no intra-nasal treatments) for a 1 wk recovery phase. A representative 4–5- μ m thick section (H&E stained) of one mouse per treatment group is shown (10 \times magnification). (A) Saline. (B) Wild-type (WT) SRA^{+/+} treatment group. (C) KO SRA^{-/-} treatment group. (D) Mean semi-quantitative distribution of inflammatory scores of lung lymphoid aggregates, bronchiolar inflammation and alveolar inflammation in mice (n = 4–5 mice/group). Error bars represent SE. Statistically significant (*p < 0.05, **p < 0.01, ***p < 0.001) vs. saline. #p < 0.05 and ###p < 0.001: significant differences between groups as indicated (1-way ANOVA with Tukey's *post-hoc* comparison). Line scale represents 100 μ m.

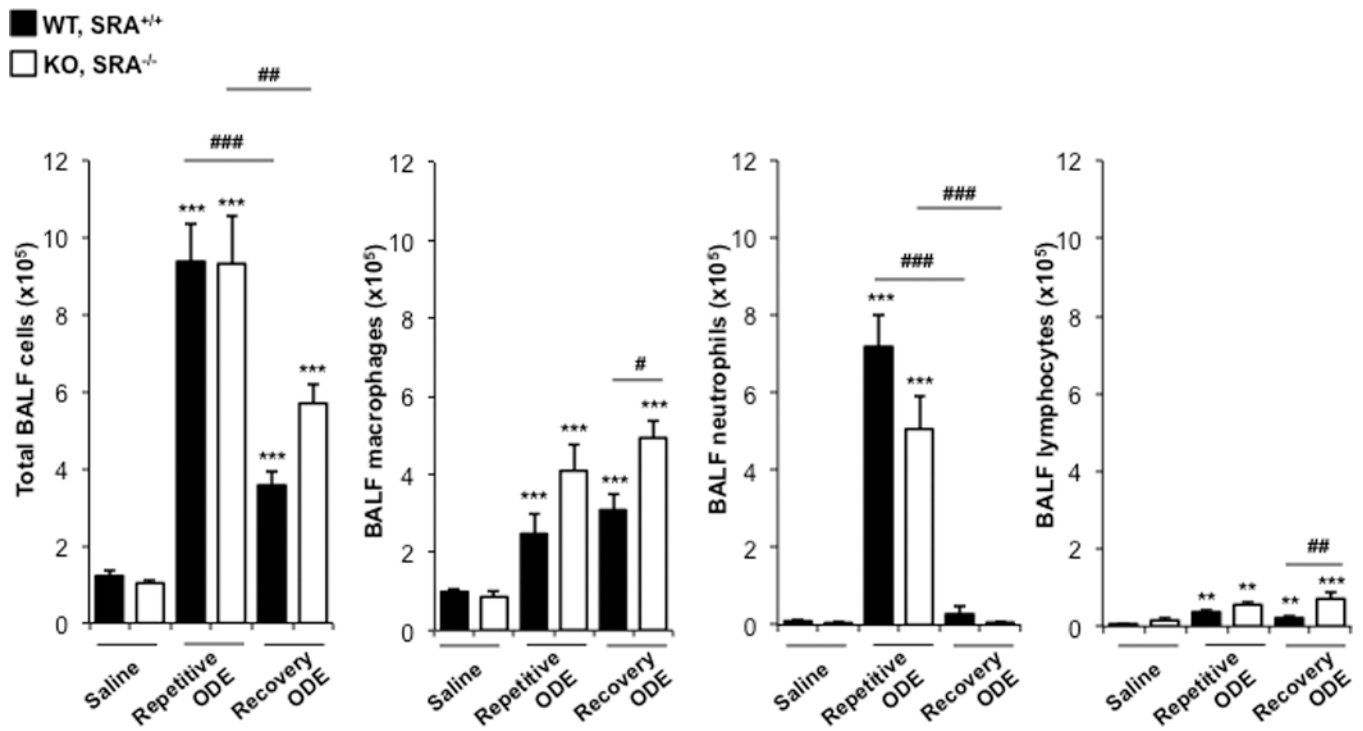


Figure 3. Lavage fluid cell (type) levels indicative of recruitment after repetitive ODE treatments and following the 1 wk recovery

Mice (WT and SRA KO) were intra-nasally treated with saline or ODE daily for 3 wk (repetitive ODE) or ODE daily for 3 wk followed by no treatments for 1 wk (recovery ODE). Results shows are means (\pm SEM) bronchoalveolar lavage fluid counts of total cells, macrophages, neutrophils and lymphocytes as determined by hemocytometer (n = 4–8 mice/group). Statistically significant (**p < 0.01, ***p < 0.001) vs. saline. #p < 0.05, ###p < 0.01, ####p < 0.001: significant differences between groups as indicated by line (one-way ANOVA with Tukey’s *post-hoc* comparison).

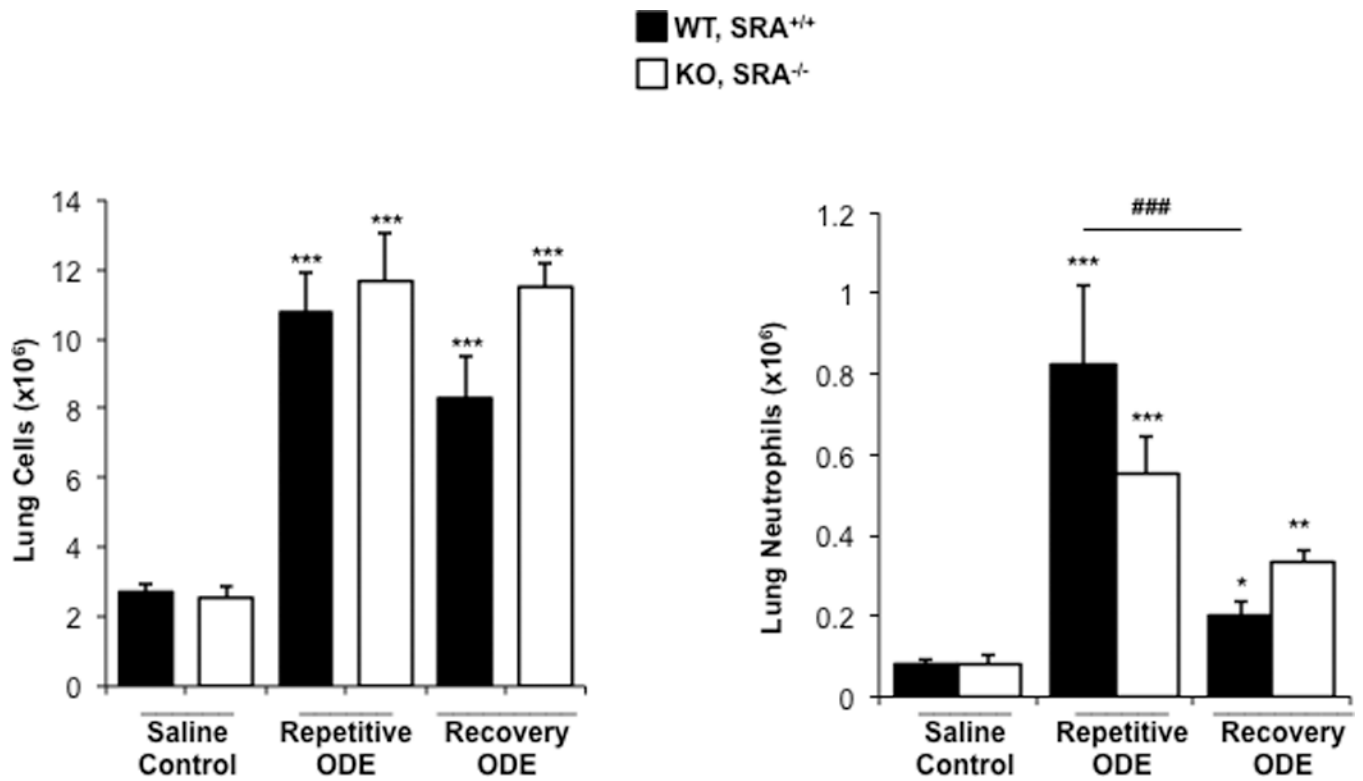


Figure 4. Total lung cells and lung neutrophils following repetitive ODE treatments and 1 wk recovery in SRA KO mice

Mice were intra-nasally treated with saline or ODE daily for 3 wk (repetitive ODE) or treated daily for 3 wk followed by no treatment for 1 wk (recovery ODE) whereupon mice were euthanized, lavage fluid removed, pulmonary vasculature perfused, and lung cells dissociated. **(A)** Mean total lung cells as determined by hemocytometer. **(B)** Mean total neutrophils (% LY6G⁺ neutrophils (see Supplemental Figure 1) of CD45⁺ cells [as determined by flow cytometry] multiplied by total lung cells) in WT and SRA KO mice. Error bars represent SE. Statistical significance (*p < 0.01, **p < 0.01, ***p < 0.001) vs. saline. ###p < 0.001: significant difference between WT repetitive ODE and WT recovery ODE (1-way ANOVA with Tukey’s *post-hoc* test). N = 5–8 mice/group.

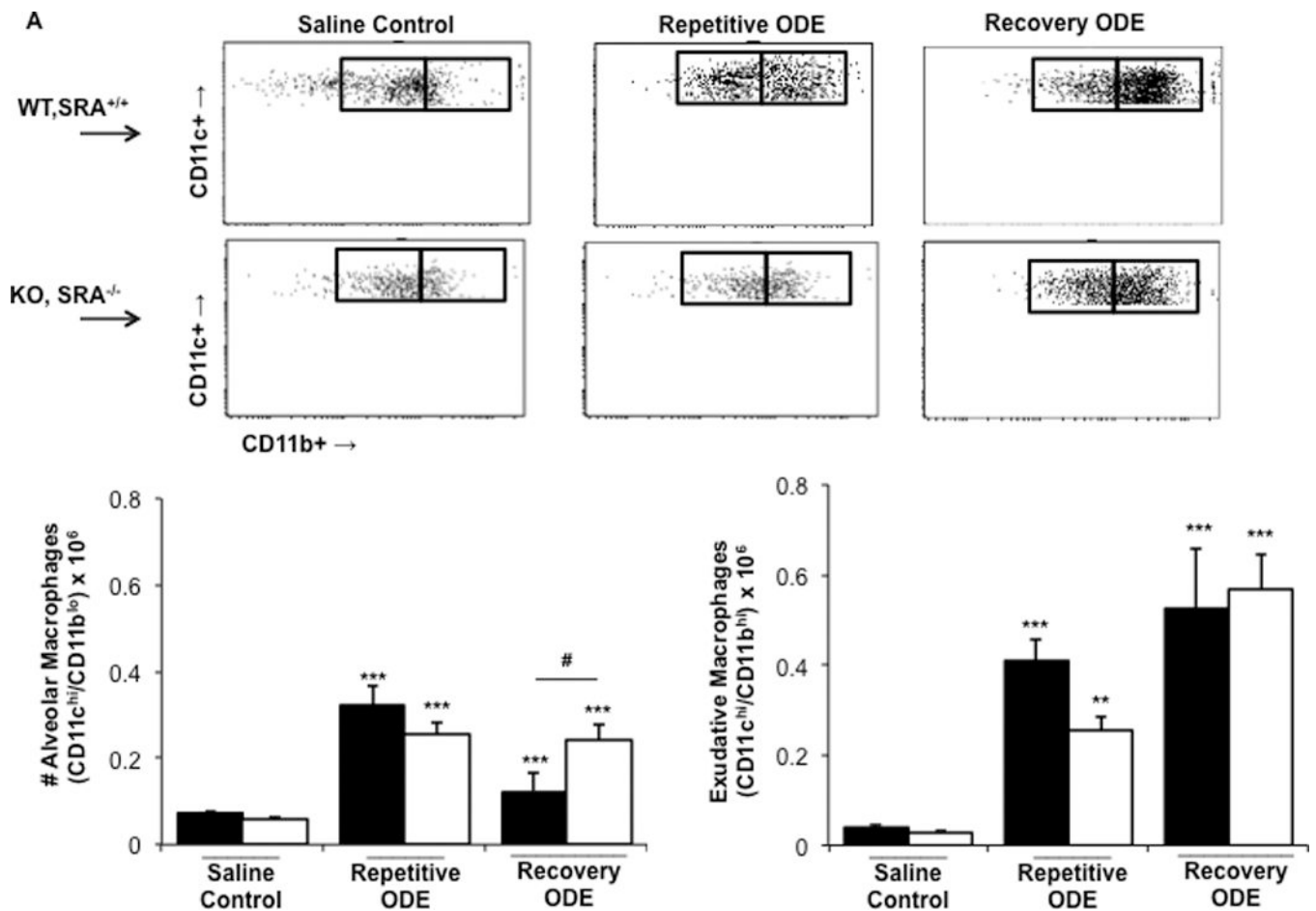


Figure 5. Exudative CD11c^{hi}/CD11b^{hi} macrophages in lungs of WT and SRA KO mice after repetitive ODE exposure and following the 1 wk recovery
Mice were intra-nasally treated with saline control or ODE daily for 3 wk (repetitive ODE) or treated daily for 3 wk followed by no treatment for 1 wk (recovery ODE) whereupon mice were euthanized, lavage fluid removed, pulmonary vasculature perfused, and lung cells dissociated and analyzed by flow cytometry. Initial gating on CD45⁺ lung leukocytes excludes debris, lymphocytes, and neutrophils, and a plot identified CD11c vs. FSC identifies CD11c⁺ cells (see Supplemental Figure 1). Next, a plot of CD11c vs. CD11b distinguishes alveolar macrophages and exudative macrophages (**top panel, A**). **B (bottom panel)**: numbers of macrophages were calculated by multiplying the % of macrophages in respective gate (% of all CD45⁺ cells, as analyzed by FACS) multiplied by total lung cells (determined using a hemocytometer) for each mouse. Data shown are means ± SEM of 5–8 mice/group. Statistically significant (**p < 0.01, ***p < 0.001) vs. saline. #p < 0.05: significant difference between SRA deficient mice (1-way ANOVA with Tukey’s *post-hoc* test).

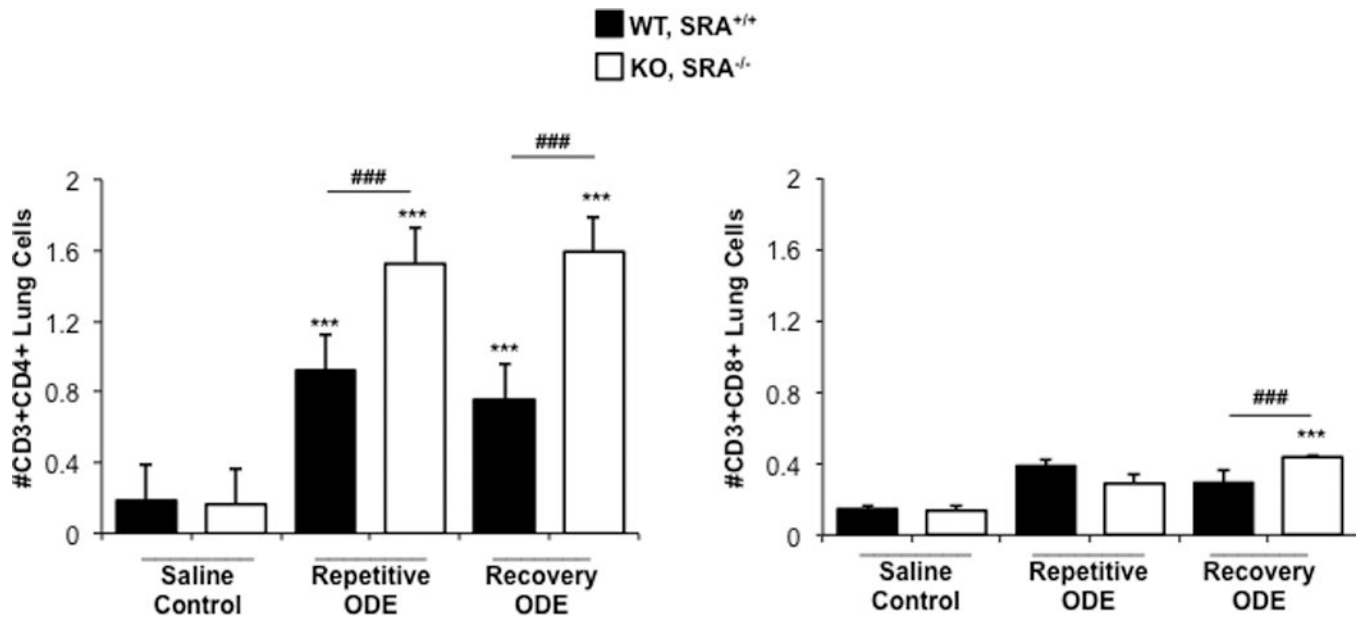


Figure 6. CD4⁺ lymphocyte recruitment in SRA KO mice following repetitive ODE exposure and after the 1 wk recovery
Mice were treated and process as outlined in Figure 5 legend. Lymphocytes identified by CD45⁺ leukocytes excluding debris and characteristic FSC and SSC properties of lymphocytes (see Supplemental Figure 1) followed by staining for CD3, CD4, and CD8. Total numbers of each lymphocyte population were determined by multiplying the frequency of CD3⁺CD4⁺ cells or CD3⁺CD8⁺ cells (among the CD45⁺ leukocytes) by the total number of cells (determined by hemocytometer) for each mouse. Data shown are means ± SEM of 5–8 mice/group. Statistically significant (***) $p < 0.001$ vs. saline. ### $p < 0.001$: significant difference between SRA KO and WT mice (1-way ANOVA with Tukey’s *post-hoc* test).

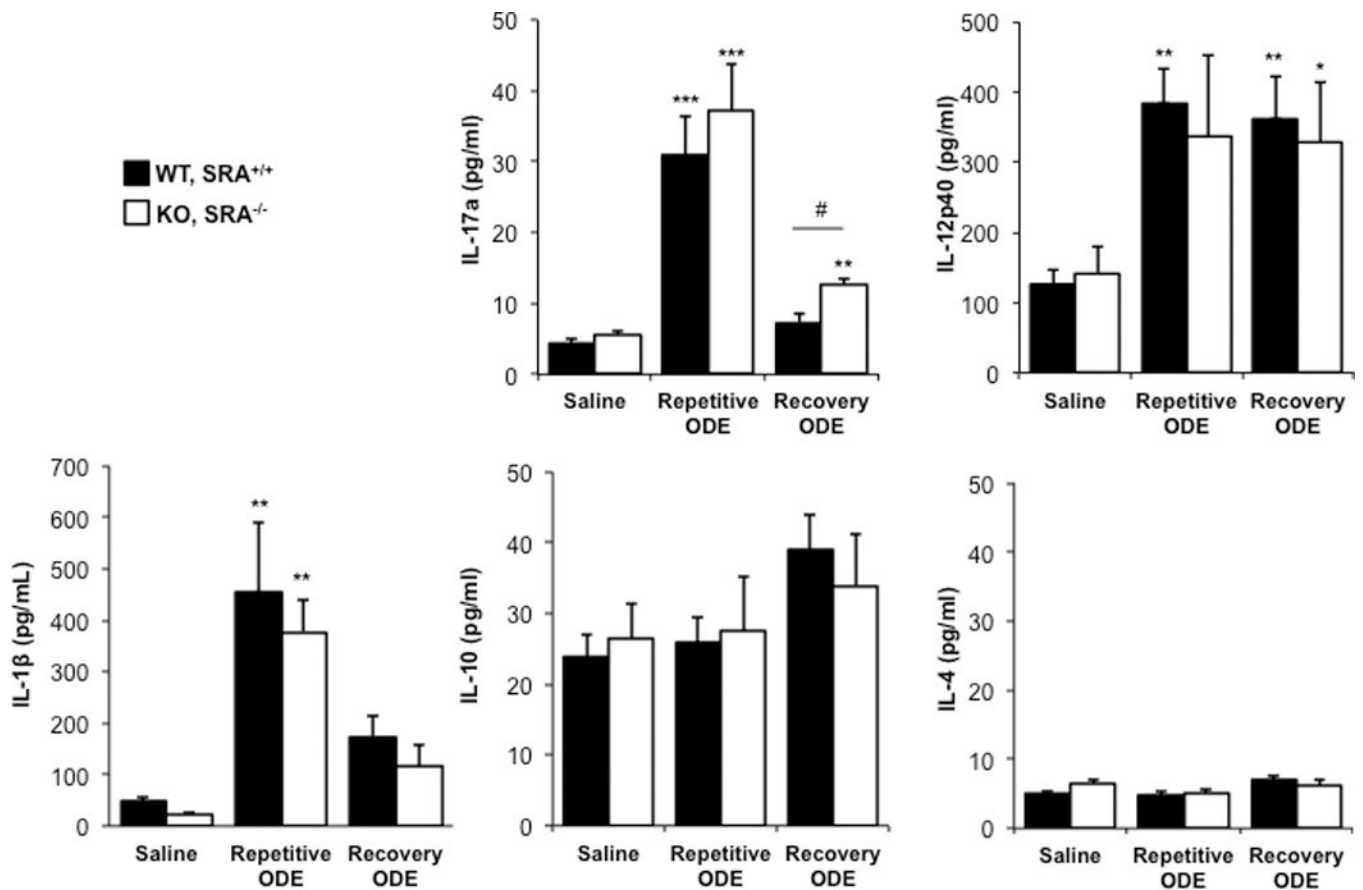


Figure 7. ODE exposure-induced T_H17/T_H1-polarized cytokine response in lung

Mice were intra-nasally treated with saline or ODE daily for 3 wk (repetitive ODE) or treated daily for 3 wk followed by no treatment for 1 wk (recovery ODE) whereupon they were euthanized, lavage fluid removed, pulmonary vasculature perfused, and half of their lung tissue was homogenized. Cytokine profiles (associated with T-cell subsets) were analyzed from cell-free supernatants of the homogenates using protein multiplex immunoassay and ELISA. Results shown are means ± SEM of 5–8 mice/group. #p < 0.05: significant difference between SRA-deficient and WT mice (1-way ANOVA with Tukey’s *post-hoc* test).