Efferocytosis impairs pulmonary macrophage and lung antibacterial function via PGE$_2$/EP2 signaling

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The lung is highly susceptible to injury from systemic (e.g., sepsis) or inhalational (e.g., acid aspiration) exposures, and acute lung injury, manifesting clinically as adult respiratory distress syndrome (ARDS), exacts a huge human toll (1). ARDS is characterized by apoptosis of pulmonary epithelial and endothelial cells as well as of infiltrating neutrophils recruited during the inflammatory response to injury (2). The surface binding/ingestion of ACs during macrophage efferocytosis has been shown to trigger the release of molecules such as transforming growth factor β, interleukin-10 (IL-10), nitric oxide, and prostaglandin E$_2$ (PGE$_2$). Although the antiinflammatory actions of these mediators may contribute to the restoration of homeostasis after tissue injury, their potential impact on antibacterial defense is unknown. The lung is highly susceptible to diverse forms of injury, and secondary bacterial infections after injury are of enormous clinical importance. We show that ACs suppress in vitro phagocytosis and bacterial killing by alveolar macrophages and that this is mediated by a cyclooxygenase–PGE$_2$–E prostanoid receptor 2 (EP2)–adenylyl cyclase–cyclic AMP pathway. Moreover, intrapulmonary administration of ACs demonstrated that PGE$_2$ generated during efferocytosis and acting via EP2 accounts for subsequent impairment of lung recruitment of polymorphonuclear leukocytes and clearance of Streptococcus pneumoniae, as well as enhanced generation of IL-10 in vivo. These results suggest that in addition to their beneficial homeostatic influence, antiinflammatory programs activated by efferocytosis in the lung have the undesirable potential to dampen innate antimicrobial responses. They also identify an opportunity to reduce the incidence and severity of pneumonia in the setting of lung injury by pharmacologically targeting synthesis of PGE$_2$ or ligation of EP2.

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AMs ingest and kill pathogens, thereby serving a critical function in immune defense of the delicate gas-exchanging region of the lung. One such receptor that plays an important role in antimicrobial responses (8) and that has been extensively investigated (9) is the Fcγ receptor (FcR) for IgG antibodies. To determine if efferocytosis suppresses in vitro FcR-mediated phagocytosis, rat AMs were preincubated with different ratios of ACs and then challenged with IgG-opsonized erythrocytes (RBCs) or *Escherichia coli* for 90 min. As a source of ACs, we used Jurkat T cells treated with camptothecin using a protocol resulting in 25.6% of cells in early apoptosis with only 3.1% contamination by late apoptotic or necrotic cells (Fig. 1 A). Because Fadok et al. (3) previously demonstrated that PGE2 production by human macrophages occurred after 90 min of incubation with ACs, we initially used this pretreatment interval. Microscopic visualization (unpublished data) indicated that AMs bound and ingested ACs, as previously reported (10). Preincubation of AMs for 90 min with various ratios of ACs dose-dependently inhibited subsequent FcR-mediated phagocytosis of both RBCs and *E. coli* (Fig. 1 B), with ≥50% inhibition being observed at AC/AM ratios of 3:1. The inhibition by ACs (3:1) of FcR-mediated ingestion of both targets was also time dependent over a 15–90-min pretreatment interval, and a 16-h pretreatment resulted in near complete suppression (Fig. 1 C). Similar inhibitory effects were obtained when rat thymocytes, rat PMNs, or RLE-6TN rat lung epithelial cells were used as the source of ACs (unpublished data). Preincubation with either viable or necrotic cells had no effect on subsequent FcR-mediated phagocytosis (Fig. 1 D). After their ingestion, macrophages must kill bacteria. Preincubation with ACs (3:1) for 90 min significantly enhanced the intracellular survival of phagocytosed bacteria, reflecting an impairment of AM microbiocidal activity against IgG-opsonized *Klebsiella pneumoniae* (Fig. 1 E). Together, these results demonstrate that preexposure to ACs markedly impairs the ability of AMs to carry out two crucial functions involved in immune defense against bacterial pneumonia: microbial phagocytosis and killing.

We next assessed whether the inhibition of FcR-mediated phagocytosis by efferocytosis in AMs was dependent on soluble mediators. Pretreatment of naive AMs with cell-free supernatant harvested from parallel AM cultures incubated for 90 min with ACs (3:1) inhibited subsequent FcR-mediated phagocytosis to the same degree as did direct addition of
Inasmuch as the interaction with ACs leads human macrophages to also secrete PGE$_2$ (3), and we have reported that endogenously produced PGE$_2$ inhibits FcR-mediated phagocytosis by AMs (14), we evaluated the role of endogenous prostanoids in the suppression of FcR-mediated phagocytosis. Pretreatment with the cyclooxygenase (COX) inhibitors indomethacin and aspirin completely abrogated the inhibition of FcR-mediated phagocytosis by ACs, and such inhibition was reproduced by addition of exogenous PGE$_2$ (Fig. 2 A). Indeed, AMs secreted PGE$_2$ in response to ACs and this was

Figure 2. PGE$_2$ mediates the suppressive effects of efferocytosis on AM antimicrobial functions via EP2. (A) AMs were pretreated with culture supernatant derived from parallel incubations of ACs/AMs (3:1), with 5 μM PGE$_2$, or with 3:1 ACs in the absence or presence of 6 μg/ml of anti–TGF-β blocking antibody or 5 μM of the COX inhibitors indomethacin (Indo) and 200 μM of aspirin (Asp). They were subsequently challenged with IgG RBCs and phagocytosis was determined. (B) AMs were incubated with medium alone or with ACs in the absence or presence of aspirin. PGE$_2$ in supernatant was quantitated by immunoassay after 30 min. (C) AM phagocytosis of IgG RBCs was determined after a 90-min pretreatment with medium alone or with ACs (3:1) in the absence or presence of 100 μM of the EP2 antagonist AH-6809. (D) AMs from EP2$^{-/-}$ or WT control mice were preincubated with or without apoptotic thymocytes (5:1) for 90 min before challenge with IgG RBCs and phagocytosis was determined. Results represent the mean ± SEM from three independent experiments, each performed in quintuplicate (A–C) or the mean ± SEM of quintuplicate values from one experiment representative of three independent experiments (D). *, P < 0.05 versus control; #, P < 0.05 versus AC.
We have reported that the suppressive effects of PGE\(_2\) on AM antimicrobial functions are primarily mediated by its binding to the EP2 receptor, a G\(_\text{s}\)-coupled receptor that activates adenylyl cyclase activity with resulting cAMP formation (14, 17, 18). To assess the specific contribution of PGE\(_2\), among other prostanoids, to the efferocytosis-induced inhibition of FcR-mediated phagocytosis and to specifically dissect the participation of EP2 in this process, AMs were pretreated with ACs in the presence or absence of the EP2 antagonist AH-6809 and then challenged with IgG RBCs. Inhibition of FcR phagocytosis by ACs was completely abrogated by the EP2 antagonist (Fig. 2 C). Moreover, AC pretreatment of AMs harvested from mice genetically deficient in the EP2 receptor was unable to significantly inhibit FcR-mediated phagocytosis as it was in cells from WT mice (Fig. 2 D). The suppressive effect of ACs on AM phagocytosis was also abrogated when these cells were pretreated with the adenylyl cyclase inhibitor SQ 22536 (Fig. 3 A). In addition, inhibition of phagocytosis by ACs directly correlated with increased intracellular levels of cAMP, which was also abrogated by aspirin and SQ 22536 (Fig. 3 B). Bactericidal capacity in the context of efferocytosis was likewise markedly augmented by coincubation with a COX inhibitor, an EP2 antagonist, or an adenylyl cyclase inhibitor (Fig. 3 C). These results demonstrate that PGE\(_2\), acting via an EP2–adenylyl cyclase–cAMP pathway, mediates the suppressive effects of efferocytosis on FcR-mediated antimicrobial functions by AMs.

To confirm the biological significance of these in vitro results, we modeled a secondary lung infection after initial pulmonary exposure to ACs such as what would be seen in acute lung injury. For these experiments, ACs were generated by dexamethasone treatment of murine thymocytes for 6 h, which yielded 40.3% early apoptotic and 4.93% late apoptotic plus necrotic cells (Fig. 4 A). Initially, different numbers of ACs were coadministered intratracheally in C57BL/6 WT mice along with a standard inoculum of the important respiratory pathogen *Streptococcus pneumoniae*, and 48 h after challenge the bacterial burdens in lung homogenates were evaluated. As shown in Fig. 4 B, there was no difference in pulmonary bacterial clearance of mice that were infected and simultaneously exposed to ACs using this protocol compared with those infected alone. Because our in vitro results indicated that the efferocytosis–induced inhibition of FcR-mediated phagocytosis was time dependent and reached a level of 89% when phagocytic target challenge was performed after an interval of 16 h (Fig. 1 C), we devised a second model in which *S. pneumoniae* was administered intratracheally 16 h after various numbers of apoptotic thymocytes were instilled intranasally. As \(\sim55\%\) of the total numbers of cells obtained after treatment of thymocytes with dexamethasone remain viable, viable cells were also administrated intranasally as an experimental control. Results showed that pretreatment with ACs, but not viable cells, using this protocol dose-dependently impaired pulmonary bacterial clearance (Fig. 4 C) and also led to the dissemination of *S. pneumoniae* into the bloodstream (Fig. 4 D) 48 h after infection. Finally, to test the role
of the EP2 receptor in this impairment of in vivo pulmonary defense against *S. pneumoniae* by AC pretreatment, we compared the lung and the bloodstream bacterial burdens in WT versus EP2−/− mice. In contrast with WT mice, the pulmonary bacterial burden in AC-pretreated EP2−/− mice was no greater than in non–AC-pretreated controls but was 2.5 logs lower than in AC-pretreated WT mice (Fig. 4E). In addition, EP2−/− mice exhibited no bacteremia (Fig. 4F). Because these mice lack preexisting antibodies against *S. pneumoniae*, bacterial recognition and clearance by phagocytes in the in vivo model is likely independent of FcR, indicating that the PGE2/EP2/cAMP axis also suppresses innate defense.

Figure 4. Intrapulmonary administration of ACs impairs host defense in a mouse model of pneumococcal pneumonia. (A) Thymocytes were incubated with 1 μM dexamethasone for 6 h and ACs were detected by AnnexinV-FITC/PI and analyzed by flow cytometry. Early ACs comprise 40.3% of total cells. (B) 10^6 CFU of *S. pneumoniae* and varying numbers of apoptotic thymocytes were coadministered intratracheally in WT mice. Lung homogenates were assessed for bacterial CFUs 48 h later. (C) Indicated numbers of apoptotic or viable thymocytes were instilled intranasally in WT mice and, 16 h later, 10^6 CFU *S. pneumoniae* were administered intratracheally. Lung homogenates were assessed for bacterial CFUs 48 h after *S. pneumoniae* challenge. (D) Bacterial CFUs were determined in blood obtained 48 h after *S. pneumoniae* challenge from the same WT mice studied in C. (E) WT and EP2−/− mice were subjected to intranasal administration of apoptotic thymocytes 16 h before intratracheal challenge with *S. pneumoniae* as described in C. Lung homogenate CFUs 48 h after bacterial challenge are presented. (F) Bacterial CFUs were determined in blood obtained 48 h after *S. pneumoniae* challenge from the same EP2−/− mice studied in E. Results represent the mean ± SEM of one experiment representative of two. The number of animals analyzed in each group is indicated above each bar. ND, none detected. *, P < 0.05 versus control; #, P < 0.05 versus AC.
Immune responses when bacterial recognition proceeds via other relevant recognition receptors such as toll-like receptors (19), collectins (20), or scavenger receptors (21).

We next sought to address the impact of PGE2/EP2 signaling on lung levels of antiinflammatory mediators. TGF-β has previously been implicated as an important antiinflammatory mediator in the lung in vivo (6). We verified that lung homogenate TGF-β and IL-10 levels were indeed dose-dependently increased 16 h after the administration of thymocytes in uninfected WT mice. In contrast, levels of NO3− (the stable oxidized derivative of nitric oxide) were unchanged (unpublished data). As expected, PGE2 levels in lung homogenates of infected animals were increased after administration of ACs (Fig. 5 A). S. pneumoniae infection in WT mice increased lung homogenate levels of TGF-β and NO3− but not IL-10 (unpublished data). In this experimental context, administration of ACs had no effect on lung levels of either NO3− or TGF-β (Fig. 5 B) but significantly increased levels of IL-10 (Fig. 5 C). This increase in IL-10 generation after AC administration was not seen in EP2−/− animals (Fig. 5 C). These studies show that PGE2/EP2 signaling drives the enhanced IL-10 production associated with efferocytosis in vivo.

In view of the contribution of PMN recruitment to antibacterial defense of the lung, we next assessed whether efferocytosis influenced PMN influx to the lungs of infected mice. Intrapulmonary instillation of apoptotic thymocytes had no impact on total and differential cell counts in bronchoalveolar lavage fluid (BALF) obtained 16 h later from noninfected WT mice (unpublished data). However, pretreatment with ACs 16 h before infection with S. pneumoniae dose-dependently reduced the numbers of total cells (not depicted) and PMNs (Fig. 5 D), but not mononuclear cells (not depicted), in BALF harvested 48 h after infection. EP2−/− mice infected with S. pneumoniae exhibited significantly lower numbers of total BALF cells and PMNs than WT animals, an effect likely related to reduced alveolar capillary permeability (unpublished data). However, in contrast to their WT counterparts, AC pretreatment failed to attenuate PMN recruitment in EP2−/− mice (Fig. 5 D). These results suggest that PGE2/EP2 signaling also mediates the impairment in PMN recruitment to the infected lung, which likely contributes to defective bacterial clearance.

In a mouse model of Chagas disease, administration of ACs was found to enhance Trypanosoma cruzi parasitemia, an effect blocked by COX inhibitors; however, this study did not use the natural route of inoculation via the skin (12). Ours is the first in vivo study to use a natural route of microbial inoculation, the first to address antibacterial defenses, and

![Figure 5. PGE2/EP2 signaling impairs PMN recruitment and promotes in vivo generation of IL-10 in a mouse model of pneumococcal pneumonia. 10^6 apoptotic thymocytes were instilled intranasally in WT and EP2−/− mice and, 16 h later, 10^6 CFU S. pneumoniae were administered intratracheally. (A–C) PGE2 (A), total TGF-β (B), and IL-10 levels (C) were quantified in the supernatant of lung homogenates from animals studied in Fig. 4 (C and E). (D) PMNs in BALF from WT and EP2−/− mice were counted. Results represent the mean ± SEM of one experiment representative of two (A–C) or of one experiment (D). The number of animals analyzed in each group is indicated above each bar. *, P < 0.05 versus control; #, P < 0.05 versus AC.]
the first to examine a pulmonary infection. As World Health Organization statistics document that respiratory infections account for more life years lost around the world than any other category of disease (22), the implications of these findings for global health may be substantial. Although ARDS is the most conspicuous example of such a scenario, chronic lung diseases such as cystic fibrosis (23) and emphysema (24) also involve ongoing apoptosis and confer a high risk of infection (25, 26). The capacities of AMs to bind and ingest ACs (10, 27), as well as to generate prostanoids including PGE$_2$ (28), are known to be reduced relative to those of peritoneal macrophages. For this reason, it was not obvious that either eff erocytosis itself or PGE$_2$ generated under such conditions would mediate suppression of antimicrobial defense in AMs in vitro or in the lung in vivo. Nevertheless, the studies reported here demonstrate that eff erocytosing AMs generate sufficient PGE$_2$ as well as sufficient EP2-mediated signaling responses to this lipid mediator to largely account for impairment of subsequent antibacterial responses in vitro and in vivo. As a variety of COX inhibitors are currently available, and as EP2 antagonists are under active development, our findings suggest that therapeutic targeting of this pathway has the potential to augment innate immunity in the lung under conditions characterized by apoptosis.

MATERIALS AND METHODS

Reagents. RPMI 1640 and penicillin/streptomycin/amphotericin B solution were purchased from Invitrogen. Camptothecin and dexmethylone were obtained from EMD. Trypsin soy broth was supplied by BD. Aspirin, cytochalasin D, indomethacin, o-phenylenediamine dihydrochloride, and SDS were obtained from Sigma-Aldrich. AH-8989, PGE$_2$, and NO$_2^-$ colorimetric assay kits were obtained from Cayman Chemical, and PGE$_2$ and cAMP EIA kits were obtained from assay designs. SQ 22536 was obtained from BIOMOL International. Anti-TGF-$\beta_1$ antibody was obtained from R&D Systems. TGF-$\beta$ and IL-10 ELISAs were obtained from BD. Compounds requiring reconstitution were dissolved in DMSO.

Animals. EP2$^{-/-}$ mice on a C57BL/6 background (29) (Ono Pharmaceutical), WT C57BL/6 mice (The Jackson Laboratory), and Wistar rats (Charles River Laboratories) were treated according to National Institutes of Health guidelines for the use of experimental animals with the approval of the University of Michigan Committee for the Use and Care of Animals.

Cell isolation and culture. Resident AMs from mice and rats were obtained via lung lavage and cultured as previously described (14).

Apoptotic cells. Jurkat T cells (American Type Culture Collection) and glycogen-elicited PMNs (30) were incubated with 8 $\mu$g/ml camptothecin for 5 and 3 h, respectively. Rat or mouse thymocytes were incubated with dexmethylone for 6 h (31), and RLE-6TN rat lung epithelial cells (American Type Culture Collection; gift of V. Thannickal, University of Michigan, Ann Arbor, MI) were serum starved overnight to induce apoptosis. Apoptotic cells were detected by AnnexinV-FITC/PI staining (BD) and analyzed using a FACSCalibur (BD). Jurkat cells were rendered $\sim$85% AnnexinV$^+/P^+$ by freeze thawing (necrotic cells).

Phagocytosis and bacterial killing assays. Phagocytosis of IgG RBCs or IgG E. coli was assessed as previously described (14). The ability of K. pneumoniae to survive intracellularly after phagocytosis was assessed as previously described (32, 33).

Measurement of cAMP, PGE$_2$, IL-10, TGF-$\beta$, and NO$_2^-$ levels. Intracellular cAMP levels in AM lysates (34, 35), and PGE$_2$ (35), IL-10, and TGF-$\beta$ levels in culture supernatants or lung homogenates were quantified by ELISA. NO$_2^-$ (the stable oxidized derivative of nitric oxide) was determined using the Greiss reaction (36).

In vivo experiments. WT and EP2$^{-/-}$ mice were subjected to intratracheal or intranasal administration of ACs or viable cells in PBS. Either simultaneously or 16 h thereafter, 10$^6$ CFU S. pneumoniae were administered intratracheally. Lung homogenates and blood were assessed for bacterial CFUs 48 h after S. pneumoniae challenge (S. pneumoniae provided by P. Mancuso, University of Michigan, Ann Arbor, MI). In another set of experiments, BAL was performed with 3 ml HBSS 16 h after instillation of ACs or 46 h after infection with S. pneumoniae. Cell counts and differentials were determined by light microscopy.

Statistical analysis. Data are presented as the mean  ±  SEM. Comparisons among groups were assessed with ANOVA followed by Bonferroni analysis. Differences were considered significant if P-values were <0.05.

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