Respiratory Infections Cause the Release of Extracellular Vesicles: Implications in Exacerbation of Asthma/COPD

Eltom, S., Dale, N., Raemdonck, K. R. G., Stevenson, C. S., Snelgrove, R. J., Sacitharan, P. K., ... Birrell, M. A. (2014). Respiratory Infections Cause the Release of Extracellular Vesicles: Implications in Exacerbation of Asthma/COPD. PloS one, 9(6), [e101087]. DOI: 10.1371/journal.pone.0101087

Published in:
PloS one

Document Version:
Publisher's PDF, also known as Version of record

Queen's University Belfast - Research Portal:
Link to publication record in Queen's University Belfast Research Portal

Publisher rights
© 2014 The Authors
This is an open access article distributed under the terms of the Creative Commons Attribution License (https://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited

General rights
Copyright for the publications made accessible via the Queen's University Belfast Research Portal is retained by the author(s) and / or other copyright owners and it is a condition of accessing these publications that users recognise and abide by the legal requirements associated with these rights.

Take down policy
The Research Portal is Queen's institutional repository that provides access to Queen's research output. Every effort has been made to ensure that content in the Research Portal does not infringe any person's rights, or applicable UK laws. If you discover content in the Research Portal that you believe breaches copyright or violates any law, please contact openaccess@qub.ac.uk.
Respiratory Infections Cause the Release of Extracellular Vesicles: Implications in Exacerbation of Asthma/COPD

Suffwan Eltom¹, Nicole Dale¹, Kristof R. G. Raemdonck¹, Christopher S. Stevenson²*, Robert J. Snelgrove³, Pradeep K. Sacitharan¹, Chiara Recchi⁴, Silene Wavre-Shapton⁵, Daniel F. McAuley⁶, Cecilia O’Kane⁶, Maria G. Belvisi⁴⁵, Mark A. Birrell¹⁷†

¹Respiratory Pharmacology, Imperial College, London, United Kingdom, ²Novartis, Horsham, United Kingdom, ³Leukocyte Biology, Imperial College, London, United Kingdom, ⁴Molecular Medicine, Imperial College, London, United Kingdom, ⁵Institute of Ophthalmology, University College London, London, United Kingdom, ⁶Centre for Infection and Immunity, Queen’s University of Belfast, Belfast, United Kingdom, ⁷MRC-Asthma UK Centre in Allergic Mechanisms of Asthma, Imperial College, London, United Kingdom

Abstract

**Background:** Infection-related exacerbations of respiratory diseases are a major health concern; thus understanding the mechanisms driving them is of paramount importance. Despite distinct inflammatory profiles and pathological differences, asthma and COPD share a common clinical facet: raised airway ATP levels. Furthermore, evidence is growing to suggest that infective agents can cause the release of extracellular vesicle (EVs) in vitro and in bodily fluids. ATP can evoke the P2X₇/caspase 1 dependent release of IL-1β/IL-18 from EVs; these cytokines are associated with neutrophilia and are increased during exacerbations. Thus we hypothesized that respiratory infections causes the release of EVs in the airway and that the raised ATP levels, present in respiratory disease, triggers the release of IL-1β/IL-18, neutrophilia and subsequent disease exacerbations.

**Methods:** To begin to test this hypothesis we utilised human cell-based assays, ex vivo murine BALF, in vivo pre-clinical models and human samples to test this hypothesis.

**Results:** Data showed that in a murine model of COPD, known to have increased airway ATP levels, infective challenge causes exacerbated inflammation. Using cell-based systems, murine models and samples collected from challenged healthy subjects, we showed that infection can trigger the release of EVs. When exposed to ATP the EVs release IL-1β/IL-18 via a P2X₇/caspase-dependent mechanism. Furthermore ATP challenge can cause a P2X₇ dependent increase in LPS-driven neutrophilia.

**Conclusions:** This preliminary data suggests a possible mechanism for how infections could exacerbate respiratory diseases and may highlight a possible signalling pathway for drug discovery efforts in this area.

Introduction

Asthma and Chronic Obstructive Pulmonary Disease (COPD) are respiratory diseases with ever-increasing global prevalence [1–3] that represent a social and economic burden for industrialised and developing countries [4]. The World Health Organization currently states the number of patients suffering from asthma is 300 million and predicts this figure to rise to 400 million by 2025 [5], whereas there are 600 million COPD sufferers worldwide and the disease is predicted to be the third ranked leading cause of death by 2020 [5].

Exacerbations are common events in the lives of patients with asthma and COPD [6–8]. These episodes are often associated with infections by viruses or bacteria [9] and cause worsening of symptoms, which can be fatal. Often these heightened symptoms are far less responsive to normal treatments and are associated with increased health care costs and societal impact [10]. Increases in inflammatory status, particularly IL-1β and neutrophilia, in the airway are evident during exacerbations of both diseases [9,11–16]. Furthermore, there is increasing evidence to suggest that the exacerbations accelerate the progressive decline in lung function [8,11]. Therefore there is an urgent need to understand the mechanisms driving exacerbations and identify novel therapeutic interventions to target this cohort of patients.

Extracellular vesicles (EV) such as exosomes and microvesicles have been shown to be released from a diverse range of cell types...
in response to infective agents/pathogens and are believed to primarily function in immune surveillance and host defence (recently reviewed [17–19]. These vesicles contain proteins, lipids, mRNA and microRNA; they typically range from 30 nm to 1 μm in size and are found in many biological fluids. Recent cell-based studies have shown that ATP-stimulated EVs release IL-1β and IL-18 via the P2X7/caspase-1 axis [20–23] and it is known that these cytokines are involved in airway neutrophilia, activation of macrophages and the maintenance of a chronic inflammatory response [4]. Furthermore, it has been reported that ATP levels are increased in the airways of patients with asthma and COPD [24–27]. Indeed, despite distinct inflammatory and pathological patterns, raised ATP levels in asthma and COPD represents one common clinical attribute. Therefore our hypothesis is that exacerbations of asthma and COPD during respiratory infections are due to ATP (a known danger associated molecular pattern) activating the P2X7/caspase-1 axis within EVs resulting in the release of IL-1β and IL-18, and subsequently increasing neutrophilia and worsening of symptoms that may accelerate disease pathogenesis.

**Materials and Methods**

**Demonstration that bacterial mimetic (LPS)-induced release of EVs can enhance IL-1β and neutrophil levels and change disease phenotype in model known to have increased levels of ATP**

To begin to investigate our hypothesis we first determined if we could model the exacerbated inflammatory response to inhaled infective agents. Recently, it has been shown that inhaled bacterial mimetic, LPS, causes greater inflammation if rats have been exposed to cigarette smoke [22]. We, and others, have shown that cigarette smoke causes an increase in airway ATP levels [21,23], indeed it seems likely that the ATP, acting on P2X7 receptor, that is driving the inflammation after exposure to cigarette smoke [24,25]. Thus it is conceivable that the exacerbated neutrophilia reported by Hardaker et al. [22] could be through the ATP-EV-IL-1β axis. As much of the recent work performed by our group and our collaborators is performed using mice, we first wanted to repeat the study reported by Hardaker et al. (2010) in mice. All animal procedures were approved by the British Home Office, under the United Kingdom Animal (Scientific Procedures) Act 1986 (Project license 70/7212). Male adult C57bl/6 mice (18–25 gm) were purchased from Harlan UK Ltd and maintained in cages under controlled temperatures (19–23°C) and lighting (12-h day/12-h night) conditions. Animals were acclimatized to the laboratory conditions for a minimum of one week prior to the commencement of experiments.

Figure 1. Demonstration that LPS-induced release of EVs can enhance IL-1β and neutrophil levels and change disease phenotype in model known to have increased levels of ATP. Mice (n = 8 per treatment group) were exposed to either room air (control) or CS (3R4F cigarettes) using a negative pressure system. Mice were subjected to 2 periods of CS exposure (500 ml/minute) per day (4 hours apart) for 3 consecutive days. On the morning of the third challenge day, the mice were exposed to aerosolised vehicle of endotoxin free saline or LPS (1 mg/ml) in Perspex chambers for 30 minutes. Animals were culled and BALF and lung tissue samples were collected 24 hours after LPS treatment. IL-1β levels were measured in the BALF and neutrophil numbers were determined in the BALF and lung tissue. In separate BALF samples collected from parallel smoke or LPS driven challenges ATP levels were measured (Panel A). Data shown as mean ± S.E.M. (A: ATP #: = P = 0.0023, Mann-Whitney; B: IL-1β #: = P = 0.0009, Mann-Whitney; C: BALF neutrophil number, #: = P = 0.0431, Students T test; D: lung tissue neutrophil number; #: = P = 0.0006, Mann-Whitney).

doi:10.1371/journal.pone.0101087.g001

Airway Infections Triggers EV Release

PLOS ONE | www.plosone.org 2 June 2014 | Volume 9 | Issue 6 | e101087
light, 12-h dark cycle, lights on at 07:00 hrs). All mice had *ad libitum* access to water and chow (RM1 diet; Special Diet Services, Devon, UK).

The protocol for this study was designed specifically to examine the effect of the combination of the two stimuli. We focused not on the optimum time for smoke induced inflammation but when we knew the ATP levels were increased. Mice (*n* = 6 per treatment group) were exposed to either room air (control) or CS (3R4F cigarettes; Tobacco Health Research Institute, University of Kentucky, Lexington, KY, USA) using a negative pressure system as previously described [28,29]. Mice were subjected to 2 periods of CS exposure (500 ml/minute) per day (4 hours apart) for 3 consecutive days. On the morning of the third day, the mice were challenged with aerosolised vehicle of endotoxin-free saline (Fresenius Kabi Ltd, Warrington, UK) or LPS (1 mg/ml; to mimic bacterial infection) in Perspex chambers for 30 minutes (as previously described, [28,30]). Animals were culled and BALF and lung tissue samples were collected 24 hours after LPS treatment. IL-1β levels were measured in the BALF and neutrophil numbers determined in the BALF and lung tissue digest as described in Birrell *et al.* and Eltom *et al.* [28,31].

BALF ATP levels were determined in samples from separate parallel smoke and LPS driven studies using an ATPlite luminescence assay.

**Demonstration of the concept that an infective insult can cause EV release**

To confirm that it was appropriate to use ATP induced IL-1β as a biological marker of functional EV presence we used a cell based system [23]. THP-1 (Human acute monocytic leukemia cell line) monocytes were obtained from the European Collection of Cell Cultures (Wiltshire, UK), cultured in Roswell Park Memorial Institute medium (RPMI) 1640 medium plus GlutaMAX-I and supplemented with 10% foetal calf serum and 1% Penicillin/Streptomycin/Amphotericin B. Cells were maintained under sterile conditions in a humidified atmosphere of 37 °C containing 95% air, 5% (v/v) CO₂. Cells were cultured 10⁶ per T-75 culture flasks (Corning Inc, NY, USA). Trypan Blue exclusion was performed to determine cell viability. Cells were then incubated for one hour at 37 °C in a humidified atmosphere (95% air; 5% (v/v) CO₂) in order to settle. Flasks were treated with RPMI (vehicle) or LPS (0.1 μM, final concentration of *Escherichia coli* serotype 0111:B4 (Sigma-Aldrich Co)) and incubated overnight at 37 °C (95% air; 5% (v/v) CO₂). Eighteen hours after treatment, samples were collected and centrifuged at 900 g for 10 minutes to remove the cells. The supernatants were collected and split into two equal fractions. One fraction (EV-rich) was stored at −80 °C without further processing. The other fraction was ultracentrifuged at 100,000 × g for 2 hours (4 °C) to remove the EVs. The supernatant (EV-deficient) was stored at −80 °C for further experiments. Supernatant samples from both fractions were thawed at room temperature and divided into four separate fractions: non-ultracentrifuged (EV-rich – left side) and ultracentrifuged (EV-deficient – right side). The samples were pre-treated with vehicle (DMSO, 0.1%, V/V) or P2X7 antagonist (AZ 11645373; 10⁻⁷ M). Samples were incubated for one hour and then treated with vehicle (PBS) or exogenous ATP ((10⁻³ M). The samples were then incubated for a further 4 hours prior to ELISA assessment for cytokines (A: IL-1β, B: IL-18, C: TNFα, D: MMP-9). The data is shown as mean ± S.E.M. doi:10.1371/journal.pone.0101087.g002
temperature and 100 μl of each sample was pre-treated with vehicle (DMSO, final concentration 0.1%, V/V) or P2X7 antagonist (AZ 11645373; 10^{-7} M; AstraZeneca Pharmaceuticals PLC (AZ), UK – concentration established in Eltom et al. 2011). Samples were incubated for one hour (37°C; 95% air; 5% (v/v) CO₂) and then subsequently treated with RPMI (vehicle) or Adenosine 5’-[y-thio]triphosphate tetralithium salt (ATPγS; Sigma-Aldrich Co; 10^{-2} M – concentration established in Eltom et al. 2011). Thereafter, the samples were incubated for 4 hours (37°C; 95% air; 5% (v/v) CO₂) prior to ELISA assessment for cytokines linked to the P2X7-inflammasome axis, IL-1β and IL-18, and control non-inflammasome markers such as TNFα and MMP-9. ELISAs used were purchased from R&D Systems Europe Ltd, Oxfordshire, UK.

Determining if a bacterial mimic can cause the release of EVs in the lung

Having demonstrated the concept that a bacterial mimic insult can cause the release of EVs and IL-1β is useful biological marker of their presence, we then wanted to determine if a range of infective mimics/agents would cause the release of EVs in the airway. To do this we challenged mice with bacterial and viral mimetics, collected the lavage fluid and examined ATP-induced EV release of IL-1β (IL-1β was not measured in all experiments because of the limited sample obtained and the cost of the assays).

Bacterial mimic: LPS

Mice (n = 6 per group) were challenged with the aerosolised vehicle or LPS as described above. Animals were sacrificed and BALF samples were obtained 6 hours after challenge.

Visualisation of extracellular vesicles by electron microscopy

BALF was collected from a parallel set of mice described above, the white cells and debris were removed by centrifugation (3000 g) and then the EVs collected via ultracentrifugation (100,000 g). A drop of the resuspended EVs was placed on Formvar carbon-coated copper electron microscopy grids. Samples were subsequently fixed in 4% paraformaldehyde/1% gluteraldehyde in phosphate buffer, washed 10 times with ddH₂O, stained with 1% methyl cellulose/2% uranyl acetate pH 4 on ice and visualised with a JEOL 1010 Transmission Electron Microscope (Welwyn Garden City, UK). Images were taken with a Gatan Orius SC100B charge-coupled device camera.

Visualisation of extracellular vesicles by Nanosight Technology

The Nanosight (Amsebury, UK) system has recently been suggested to be the most appropriate way of assessing the size profile of EVs in biological fluids [32,33]. It was used under the guidance of one of their trained technicians and made possible through a grant from The Rosetrees Trust, (A523).

Quantitating extracellular vesicles by measuring ATP induced cytokine release

Samples were treated with RPMI (vehicle) or exogenous ATPγS (Sigma-Aldrich; 10^{-3} M, concentration established previously,[28]) and incubated for a further 4 hours (37°C; 95% air; 5% (v/v) CO₂). Samples were then collected and stored for cytokine analysis by specific ELISA. Levels of IL-1β, IL-1α and IL-18 were determined.

Assessing the role of P2X7 and caspase in the release of IL-1β

Samples were pre-treated with inhibitors (P2X7 antagonist A438079 (10^{-6} M); Abbot Laboratories and caspase-1 inhibitor VX-765 (10^{-7} M; Vertex Pharmaceuticals, Oxfordshire, UK) and incubated for 1 hour. For these experiments we used the
Abbott compound because we have previously found that, unlike the AZ compound, it is effective antagonist of the murine P2X7 receptors [28]. Thereafter, RPMI (vehicle) or exogenous ATPcS (Sigma-Aldrich; 10^{-3} M) were added to the samples and incubated for a further 4 hours (37°C; 95% air; 5% (v/v) CO2). Samples were then collected and stored for cytokine analysis by ELISA. TNFa was measured as a negative control.

LPS challenge in healthy subjects

To translate our murine findings we repeated some of the assessment in BALF samples collected from healthy human subjects challenged with vehicle (n = 5) or LPS (n = 10). Samples were generated as described in a study by Shyamsundar [34]. Shyamsundar et al. reported that “The study was approved by the local research ethics committee, and written informed consent was obtained from all subjects before enrolment in the study”. Briefly, LPS (Escherichia coli serotype O26:B6) was dissolved in endotoxin-free sterile 0.9% saline and inhaled via an automatic inhalation-synchronized dosimeter nebulizer (Spira). The total dose of inhaled LPS was 50 mg. Bronchoalveolar lavage (BAL) was performed 6 hours after vehicle of LPS challenge. Cells were removed via centrifugation (900 g) and the BALF supernatant processed as described above.

BALF ATP levels were determined in samples from separate parallel smoke and LPS driven studies using an ATPlite luminescence assay.

Live bacterial challenge: H. influenzae

To show that live bacterial challenge causes the release of EVs, mice were infected with Haemophilus influenzae (H. influenzae). Briefly, Hib Eagan strain was a kind gift from P. Langford (Faculty of Medicine, St Mary’s Hospital, Imperial College London). Bacteria were cultured at 37°C in 5% CO2 in Brain heart Infusion broth (OXOID) supplemented with 10 μg/ml of both Hemin (10ug/ml) and Nicotinamide adenine dinucleotide (NAD) (Sigma-Aldrich, UK) or on BHI agar (OXOID) supplemented with 4% Levinthals when agar was ≤50°C. Levinthals was made by adding 50% horse blood (TCS Biosciences) to BHI broth and heating to 70°C for 45 minutes. On cooling to 50°C, 0.7 mg/ml NAD was added and the supernatant stored at −80°C in 10% glycerol as single use aliquots.

Groups of mice (n = 4–5 per group) were infected i.n. with 1×10^7 colony forming units of H. influenzae serotype b (strain Eagan) in sterile phosphate buffered saline (PBS). Terminal anaesthesia was induced at 6, 24 and 72 hours after challenge. Mice were culled and BALF samples were collected at 6, 24 and 48 hours. Presence of EVs was assessed as detailed above.

Figure 5. Determining if a bacterial mimetic (LPS) can cause the release of EVs in the lung – Cytokine release. Mice were challenged with the aerosolised vehicle of endotoxin-free saline or LPS (1 mg/ml) in Perspex chambers for 30 minutes. Animals were sacrificed and BALF obtained 6 hours after challenge. The samples were centrifuged (900 g) to remove the white blood cells and debris, and then treated with vehicle (PBS) or ATPcS (10^{-3} M), and incubated for a further 4 hours and subsequent cytokine release was analysed by ELISA. Data shown as mean +/- S.E.M. (A: IL-1β, B: IL-18, C: IL-1α).

doi:10.1371/journal.pone.0101087.g005
Viral mimic: Poly IC

To determine if viral challenge would cause EV we challenged mice with vehicle or the viral mimic of Polynosinic: polycytidylic acid (Poly IC) as previously described [35]. Briefly, mice \( n = 6 \) per group were anesthetized with isoflurane and intranasally administered with 50 \( \mu l \) (approximately 25 \( \mu l \) per nostril) of either saline (vehicle) or a 0.6 mg/ml solution of Poly IC (Invitrogen Ltd, UK). Mice were culled and BALF samples were collected at 2, 6 and 24 hours. Presence of EVs was assessed as described above.

Determining whether the ATP/P2X7 axis is central to the exacerbation response \textit{in vivo}

Finally, experiments were performed to show that the ATP/P2X7 axis can exacerbate responses to inhaled bacterial mimic \textit{in vivo}. We were not able to use the smoke plus LPS exacerbation model systems because we, and others, have shown that ATP and P2X7 are central to the smoke only model, thus interpretation of any agents designed to modulate ATP/P2X7 would be difficult [27,28]. Previously, we have shown that the inflammation in the basic LPS model is not modulated by a P2X7 inhibitor [22], thus it was ideal to use as a mechanistic model for testing the hypothesis that the P2X7 receptor was involved in the ATP-driven exacerbation. Briefly, the mice were challenged with vehicle or LPS, and four hours later intranasally dosed with saline (2 ml/kg) or ATP\( \gamma \)S (0.001 mg/kg, Sigma, UK) whilst under light anaesthesia (4% isoflurane in oxygen). The mice received oral vehicle or P2X7 inhibitor, A38079 (1000 mg/kg), 30 minutes prior to the ATP challenge, 4 hours after the challenge and 1 hour prior to cull. Twenty Four hours after the LPS exposure the mice were culled and lavaged. Neutrophils numbers were measured in the BALF as described in Eltom et al. [28].

Data analysis

All data are expressed as mean \(+/-\) standard error of mean (S.E.M) of \( n \) observations. Experiments were statistically analysed utilising the Students unpaired t-test for parametric data and the Mann-Whitney U-test for non-parametric data with independent groups compared with their specific controls or time-matched controls. For multiple comparison tests, statistical analysis was performed by applying One-way analysis of variance (ANOVA) followed by a Bonferroni’s multiple comparison post-test for parametric data or a Kruskal-Wallis test incorporating a Dunn’s multiple comparison post-test for non-parametric data. A \( P \) value of \(<0.05\) was accepted as significant. Data analysis was performed using GraphPad Prism 5.0.

Results

LPS-induced release of EVs exacerbates ATP associated airway inflammation

LPS challenge caused an increase in BALF IL-1\( \beta \) levels, neutrophils and lung tissue neutrophils. Utilising mice that had been previously exposed to cigarette smoke, and thus had increased ATP levels in the BALF, the response to LPS was significantly exacerbated (Figure 1B–D). This suggests that the mouse is an appropriate species to model the exacerbation responses and thus can be used for further investigations into infection induced release of EVs.

BALF ATP levels were increased in samples from a separate parallel smoke driven study but not after LPS challenge (Figure 1A).

Demonstration of the concept that an infective insult can cause EV release

Before using the \textit{in vivo} model systems, we initially wanted to use a cell based system to demonstrate the concept that an infective insult can cause the release of EVs and that subsequent challenge of these EVs with ATP will drive the P2X7-dependent release of IL-1\( \beta \)/IL-18. Treating cultured THP-1 cells with a bacterial mimic, LPS, caused an increase in supernatant levels of IL-1\( \beta \), IL-18, TNF\( \alpha \) and MMP-9 (Figure 2). As expected the levels of these mediators were not altered by ultracentrifugation, nor were they altered when the P2X7 inhibitor was added to the cell-free supernatants (Figure 2). Interestingly, it would appear that cultured cells release some EVs under “basal” conditions. As can be seen in Figure 2, ATP triggers some P2X7-dependent IL-1\( \beta \) release. This could imply that cultured THP-1 cells release EVs constantly or that it is a consequence of culturing procedures.

Figure 6. Determining if a bacterial mimic (LPS) can cause the release of EVs in the lung – Signalling. Mice \( n = 6 \) per group) were challenged with the aerosolised vehicle of endotoxin-free saline or LPS (1 mg/ml) in Perspex chambers for 30 minutes. Animals were sacrificed and BALF obtained 6 hours after challenge. The samples were then centrifuged (900 g) to remove the white blood cells and then pre-treated with inhibitors (P2X7 antagonist A 438079 (10 M)) and incubated for 1 hour. Samples were then treated with vehicle (PBS) or ATP\( \gamma \)S (10 M), and incubated for a further 4 hours and subsequent cytokine release was analysed by ELISA. Data shown as mean \(+/-\) S.E.M. (A: IL-1\( \beta \), B: TNF\( \alpha \)). * = \( P = 0.0138 \) (One way ANOVA followed by a Bonferroni’s Multiple Comparison test). doi:10.1371/journal.pone.0101087.g006
Treatment of the cell-free supernatants with ATP increased the levels of IL-1β and IL-18 but not TNFα and MMP-9 in the EV-rich (non-ultracentrifuged) LPS challenged samples (Figure 2). This increase was attenuated when the P2X7 inhibitor, which suggests this receptor is central to the response. Significantly, when the samples were ultracentrifuged to remove the EVs, the ATP-induced increase in IL-1β/IL-18 was lost (Figure 2, right side of panels). Together this data shows that LPS can cause the release of EVs which in turn can release IL-1β/IL-18 after ATP stimulation in a P2X7 receptor dependent manner. Furthermore, importantly this data suggests that measuring ATP induced IL-1β in cell free medium is an appropriate surrogate marker of the presence of viable/functional EVs.

Determining whether infective agents can cause the release of EVs in the lung

Having demonstrated the concept that an infective insult can cause the release of EVs in a cell culture system, we then wanted to determine if this occurred in vivo. To do this we challenged mice with LPS, collected the lavage fluid and performed EM, Nanosight assessment and examined ATP-induced IL-1β production. The inflammatory response after LPS challenge was typical of that previously described [28,30]. When we prepared the EV fractions from mice challenged with the bacterial mimic, they appeared to contain more EV-like structures when compared to the saline challenged controls (Examples of the images are in Figure 3). Furthermore, assessing the cell-free BALF samples using Nanosight technology showed that the mice challenged with LPS had more particles around the size of EVs in their airways (20–1000 nM, Figure 4).

LPS challenging the mice caused an increase in IL-1β, IL-18 and IL-1α in the BALF as expected (Figure 5). Stimulating the cell-free BALF with ATP resulted in a marked increase in IL-1β and IL-18, but not IL-1α (Figure 5). Pharmacological assessment of this increase showed that the ATP induced exacerbation was dependent on P2X7 and caspase 1 (Figure 6). The levels of TNFα, a cytokine not thought to be linked to the ATP-P2X7-Caspase 1 axis, were not altered as expected (Figure 6).

In an attempt to translate our pre-clinical findings, we collected BALF samples from healthy human subjects challenged with LPS, removed the white cells and debris and then spiked the BALF with vehicle or ATP. As can be seen from Figure 7, ATP significantly increased IL-1β, but not TNFα, in the samples from the subjects challenged with LPS. Similarly, we measured an increase in IL-18 but not IL-1α after ATP spike in the BALF from LPS challenged subjects (Figure 7 and data not shown). Like the murine model samples, we did not detect any change in ATP levels in the human BALF after LPS challenge (data not shown).

To demonstrate that actual live bacteria could trigger EV release, we used BALF samples from a H. influenzae bacterial
challenge model. The inflammatory response observed in the model was typical of that reported previously [36]. As can be seen from Figure 8 (left panels) ATP spiking the samples increased the levels of IL-1β, and not TNFα, suggesting there appears to be functional EVs present in the airways of bacterially infected animals.

Similarly, in a model that uses a viral mimetic, poly I:C, which resulted in a typical inflammatory response [35], ATP spiking increased levels of IL-1β (Figure 8, right panels). Thus we are confident that bacterial and viral infections cause the release of EVs into the airway, and if there are high levels of ATP present in the airway (like in asthma and COPD), it could trigger the release of IL-1β and IL-18.

Determining whether the ATP/P2X7 axis is central to the exacerbation response in vivo

Finally, an experiment was performed to determine if ATP could trigger P2X7 dependent exacerbation of LPS induced airway inflammation in vivo. Mice were challenged with LPS +/-/− ATP in the presence and absence of a P2X7 inhibitor. LPS challenge caused an increase in BALF IL-1β and neutrophilia; this was exacerbated in the mice that received ATP (Figure 9). Whilst the P2X7 inhibitor, A438079, had no effect on neutrophil numbers after just LPS challenge (as shown before [28]), the ATP exacerbated response was attenuated by the inhibitor (Figure 9).

Discussion

Exacerbations of both asthma and COPD can be fatal and represent a growing concern to patients and healthcare providers [6,8]. These episodes are usually associated with viral and bacterial infections which enhance the existing inflammatory status, and in turn compromise lung function [9,11]. We propose that these exacerbations could be caused by the increased levels of ATP in the lungs of these patients. Increased levels of ATP could activate the P2X7/caspase 1 axis in EVs resulting in the release of IL-1β / IL-18, and subsequent increase in inflammatory and disease status. To investigate this, we wanted to demonstrate that infective mimetics/agents could cause the release of EVs in the airway and that these EVs are functional/viable i.e. they can be triggered into releasing IL-1β and IL-18 in response to ATP.

It had been previously shown that exposure to smoke increases ATP levels in the BALF from mice [27] and that co exposure with smoke and the bacterial mimetic, LPS, led to an enhanced neutrophilic response which was similar in profile to that in exacerbating asthma and COPD sufferers [12,13,37,38]. Together this suggested that the LPS exposure could be causing the release
of EVs and the smoke-induced ATP could be causing the enhanced neutrophilia response via the release of enhanced levels of IL-1β/IL-18. To investigate this hypothesis we repeated the experiment using our murine model systems. The data clearly shows that combination of smoke and LPS leads to an exacerbated inflammatory response with IL-1β and neutrophilia levels enhanced. The ability to repeat the published data in a second species gave us confidence in the observation but more importantly indicated to us that mice are an appropriate species to use and enabled us to utilise our murine model systems. Furthermore, whereas we routinely observe an increase in ATP levels in our smoke model, as can be seen from Figure 1A we do not measure an increase in our LPS model. In corroboration, in these models we only observe an increase in caspase 1 activity (a downstream marker of ATP activity on the P2X7 receptor) in the smoke model [28]. The lack of LPS induced increase in ATP levels allows us to interpret the combination and ATP spiking data more readily.

Before further studying the possibility that respiratory infections can lead to the release of functional EVs in the airway, we first wanted to demonstrate the phenomenon in a cell based system. We found that the supernatant from cells treated with a bacterial mimetic had measurable levels of IL-1β, IL-18, MMP-9 and TNFα but only the levels of IL-1β and IL-18 were further increased in the presence of ATP. Furthermore, data showed that this enhanced release was P2X7-dependent. The role of EVs in this exacerbated response was confirmed when we compared supernatants that were ultracentrifuged, to remove the EVs (EV-deficient), with non-ultracentrifuged samples (EV-rich). Whereas the addition of ATP increased IL-1β and IL-18 levels in the EV-rich samples, we did not see a similar increase in the same samples when EVs were removed. These findings are similar to that observed by others which gave us confidence in our assessments and the use of IL-1β as a marker of the existence of functional/viable EVs [17,20–23,39,40].

To begin to investigate whether respiratory infections caused the release of EVs we challenged mice with an inhaled bacterial mimetic. Assessment of the cell free BALF gave EM images that suggested an increase in particles after LPS challenge. This finding was quantified using Nanosight technology which gave a clear increase in particles in the samples harvested from LPS challenged mice. The size of these particles appeared to be in the 100–400 nm range, which is reminiscent of microvesicles, rather than exosomes [32,33]. However, more detailed studies are required to establish the EV profile and, importantly, to determine the cellular source. While imaging data is useful it lacks the information required of these proof of concept studies i.e. that EVs are functional/viable, can release cytokines upon ATP stimulation via a P2X7-caspase 1 axis. To do this we used the bioassay as described. LPS challenge in mice caused an increase in the BALF IL-1α, IL-1β, IL-18 and TNFα levels as previously found [28]. The levels of IL-1β and IL-18, but not TNFα and IL-1α, were further increased when the cell-free BALF was spiked with exogenous ATP. This increase appeared to be dependent on both P2X7 and caspase 1 suggesting this axis was integral to the response. Whilst we do not know the exact mechanism that triggers the release of IL-1β and IL-18 (and not TNFα); the fact that it can be modulated by P2X7 and caspase 1 inhibitors suggests it is not simply through lysis. Together however, this data package does indicated that EVs are released and they are functional. Furthermore, using ATP induced IL-1β release was an appropriate surrogate marker for the biological fluid we were studying. By using this bioassay we were able to show that inhaled LPS can trigger the release of EVs in humans. This suggests that the data obtained in our murine model systems is translatable. Thus additional experiments were performed on other murine airway infection systems. We were able to show that as well as a bacterial mimetic a live bacteria challenge (H. influenzae, a gram-negative bacterium) also caused the release of EVs. Furthermore a viral mimetic had measurable levels of IL-1β and IL-18, MMP-9 and TNFα but only the levels of IL-1β and IL-18 were further increased in the presence of ATP. Furthermore, data showed that this enhanced release was P2X7-dependent. The role of EVs in this exacerbated response was confirmed when we compared supernatants that were ultracentrifuged, to remove the EVs (EV-deficient), with non-ultracentrifuged samples (EV-rich). Whereas the addition of ATP increased IL-1β and IL-18 levels in the EV-rich samples, we did not see a similar increase in the same samples when EVs were removed. These findings are similar to that observed by others which gave us confidence in our assessments and the use of IL-1β as a marker of the existence of functional/viable EVs [17,20–23,39,40].

Finally, to determine whether the ATP/P2X7 axis is central to the exacerbation seen in vivo, we used a mechanistic model. The reason was that data in the disease model systems has shown that modulation of ATP levels or attenuation of P2X7 inhibits the inflammatory response [27,28,41], making interpretation of the exacerbation part of the model system difficult. Previously we have shown that the inflammation in the standard LPS model is not...
modulated by a P2X7 inhibitor [28], making it ideal to use as a mechanistic model for testing the importance of the P2X7 receptor in the ATP-driven exacerbation. The data generated clearly show that exogenous ATP caused a P2X7-dependent exacerbation of airway neutrophilia. The limitation of this in vitro assessment is that we can only infer from the in vivo/ex vivo data that the IL-1β released after ATP challenge is coming from EVs, there are other cellular sources in the airway like macrophages which we cannot rule out.

In our systems ATP alone did not cause airway neutrophilia; this is different to that reported by others [42]. Whilst we do believe that ATP can be a driving factor in airway inflammation, a good example is our published data in the smoke model [28], we think that ATP alone is not sufficient to cause cellular inflammation, other co-factors such as chemoattractants are required. Thus we are not surprised when the ATP alone did not cause neutrophilia, we cannot however explain the data published by Cicco et al. [42], it could perhaps be due to experimental design, health status of animals, etc. Another aspect of our findings that would appear to contradict published data is the lack of effect of the P2X7 receptor inhibitor on the LPS-only inflammation. Moncao-Ribeiro et al. published that P2X7 KO mice were protected from LPS challenge, with all aspects of the inflammatory response reduced in the GM mice compared to the wild type controls [43]. We have now repeated this study with litter matched controls and in agreement with our previous published findings [28], we find that in our model system the P2X7 receptor is not involved in the inflammation (data not shown). To us this result makes sense as it is what one would expect given there is no increase in ATP or the downstream marker, caspase 1 activity, in our model and it is well know that the LPS response triggers NF-κB driven inflammation. Again we cannot explain the difference in data but suggest it could perhaps be due to experimental design.

In summary, the results of this study indicate that respiratory infections, bacterial and viral, can trigger the release of functional EVs in mice and man. Furthermore, upon ATP activation, the EVs released IL-1β and IL-18 in a P2X7/caspase-1 axis dependent manner resulting in exacerbated neutrophilia. We suggest EVs and this signalling pathway could prove to be a viable therapeutic target in the future to improve the clinical outcomes of the exacerbations of asthma and COPD.

Author Contributions
Conceived and designed the experiments: SE MGB CSS RJS DFM MAB. Performed the experiments: SE ND KR CSS RJS SW-S DFM CO MAB. Analyzed the data: SE ND KR CSS RJS SW-S MAB. Wrote the paper: SE MGB MAB.

References
1. Maest M, Fabian D, Holt S, Beasley R, Program Gi for A (GINA) (2004) The global burden of asthma: executive summary of the GINA Disinfection Committee report. Allergy 59: 469–478.
2. Mannino DM, Buist AS (2007) Global burden of COPD: risk factors, prevalence, and future trends. Lancet 370: 765–773.
3. Parer E, Am-Khaled N, Beasley R, Mallol J, Keil U, et al. (2007) Worldwide trends in the prevalence of asthma symptoms: phase III of the International Study of Asthma and Allergies in Childhood (ISACA). Thorax 62: 758–766.
4. Barnes PJ (2008) Immunology of asthma and chronic obstructive pulmonary disease. Nat Rev Immunol 8: 193–202.
5. Organisation WH (2007) World Health Organisation. Global surveillance, prevention and control of chronic respiratory diseases: a comprehensive approach. ISBN 978 92 4. 12–36.
6. Jackson DJ, Sykes A, Mallia F, Johnson SL (2011) Asthma exacerbations: origin, effect, and prevention. J Allergy Clin Immunol 120: 1165–1174.
7. O’Byrne PM (2009) Allergen-induced airway inflammation and its therapeutic intervention. Allergy Asthma Immunol Res 1: 5–9.
8. Wedzicha JA, Seemungal TA (2007) COPD exacerbations: defining their cause and prevention. Lancet 370: 786–796.
9. Pauwels RA (2004) Similarities and differences in asthma and chronic obstructive pulmonary disease exacerbations. Proc Am Thorac Soc 1: 73–76.
10. Schwenklenck M, Lowy A, Andrehub H, Seacs TD (2003) Costs of asthma in a cohort of Swiss adults: associations with exacerbation status and severity. Value Health 6: 75–83.
11. O’Byrne PM, Pedersen S, Larum C, Tan WJ, Busse W (2009) Severe allergic asthma exacerbations: origin, effect, and prevention. J Allergy Clin Immunol 120: 1165–1174.
12. Botelho FM, Bauer CMT, Finch D, Nikota JK, Zavitz CCJ, et al. (2011) IL-1a/ IL-1R1 Expression in Chronic Obstructive Pulmonary Disease and Mechanistic Relevance to Smoke-Induced Neutrophilia in Mice. PLoS One 6: e20457– e20457.
13. Bafadhel M, McKenna S, Terry S, Mistry V, Reid C, et al. (2011) Acute experimental infection with extracellular vesicles in mice. Respir Res 9: 121–130.
14. Kersul AL, Iglesias A, Rı ´os A´ , Noguera A, Forteza A, et al. (2011) Molecular mechanisms of exacerbations of chronic obstructive pulmonary disease. Arch Bronconeumol 47: 176–183.
15. Marin A, Garcia-Aymerich J, Sauleda J, Belda J, Millares L, et al. (2012) Effect of bronchial colonisation on airway and systemic inflammation in stable COPD. COPD 9: 121–130.
16. Goyberg B, Szabó TG, Pasztói M, Pal Z, Miják P, et al. (2011) Membrane vesicles, current state-of-the-art: emerging role of extracellular vesicles. Cell Mol Life Sci 68: 2667–2680.
17. Robbins PD, Morelli AE (2014) Regulation of immune responses by extracellular vesicles. Int Immunopharmacol 16: 290–298.
18. Dragovic RA, Gardiner C, Brooks AS, Tannetta DS, Ferguson DJP, et al. (2011) Exhaled breath condensate cytokine patterns in chronic obstructive pulmonary disease. Arch Bronconeumol 47: 176–183.
19. Gyo¨rgy B, Szabo´ TG, Pa´szto´i M, Pa´l Z, Misja´k P, et al. (2011) Membrane vesicles, current state-of-the-art: emerging role of extracellular vesicles. Cell Mol Life Sci 68: 2667–2680.
20. Robbins PD, Morelli AE (2014) Regulation of immune responses by extracellular vesicles. Int Immunopharmacol 16: 290–298.
34. Shyamsundar M, McKeown STW, O’Kane CM, Craig TR, Brown V, et al. (2009) Simvastatin decreases lipopolysaccharide-induced pulmonary inflammation in healthy volunteers. Am J Respir Crit Care Med 179: 1107–1114.
35. Harris P, Sridhar S, Peng R, Phillips JE, Cohn KG, et al. (2013) Double-stranded RNA induces molecular and inflammatory signatures that are directly relevant to COPD. Mucosal Immunol 6: 474–484.
36. LeVine AM, Whitsett JA, Gwozdz JA, Richardson TR, Fisher JH, et al. (2000) Distinct effects of surfactant protein A or D deficiency during bacterial infection on the lung. J Immunol 165: 3934–3940.
37. Hardaker EL, Freeman MS, Dale S, Bahra P, Raza F, et al. (2010) Exposing rodents to a combination of tobacco smoke and lipopolysaccharide results in an exaggerated inflammatory response in the lung. Br J Pharmacol 160: 1985–1996.
38. De Kluijver J, Grünberg K, Pons D, de Klerk EP, Dick CR, et al. (2003) Interleukin-1beta and interleukin-1ra levels in nasal lavages during experimental rhinovirus infection in asthmatic and non-asthmatic subjects. Clin Exp Allergy 33: 1415–1418.
39. Qu Y, Ramachandra L, Mohr S, Franchi L, Harding CV, et al. (2009) P2X7 receptor-stimulated secretion of MHC class II-containing exosomes requires the ASC/NLRP3 inflammasome but is independent of caspase-1. J Immunol 182: 5052–5062.
40. Wang J-G, Williams JC, Davis BK, Jacobson K, Doerschuk CM, et al. (2011) Monocytic microparticles activate endothelial cells in an IL-1β-dependent manner. Blood 118: 2366–2374.
41. Lucattelli M, Cicko S, Müller T, Lommatzsch M, De Cunto G, et al. (2011) P2X7 receptor signaling in the pathogenesis of smoke-induced lung inflammation and emphysema. Am J Respir Cell Mol Biol 44: 423–429.
42. Cicko S, Lucattelli M, Müller T, Lommatzsch M, De Cunto G, et al. (2010) Purinergic receptor inhibition prevents the development of smoke-induced lung injury and emphysema. J Immunol 185: 688–697.
43. Monção-Ribeiro LE, Cagido VR, Lima-Murad G, Santana PT, Riva DR, et al. (2011) Lipopolysaccharide-induced lung injury: role of P2X7 receptor. Respir Physiol Neurobiol 179: 314–325.