Chronic Alcohol Induces M2 Polarization Enhancing Pulmonary Disease Caused by Exposure to Particulate Air Pollution

Paul Thevenot, Jordy Saravia, Joseph Giaimo, Kyle I. Happel, Tammy R. Dugas, and Stephania A. Cormier

Background: Chronic alcohol consumption causes persistent oxidative stress in the lung, leading to impaired alveolar macrophage (AM) function and impaired immune responses. AMs play a critical role in protecting the lung from particulate matter (PM) inhalation by removing particulates from the airway and secreting factors which mediate airway repair. We hypothesized AM dysfunction caused by chronic alcohol consumption increases the severity of injury caused by PM inhalation.

Methods: Age- and sex-matched C57BL/6 mice were fed the Lieber-DeCarli liquid diet containing either alcohol or an isocaloric substitution (control diet) for 8 weeks. Mice from both diet groups were exposed to combustion-derived PM (CDPM) for the final 2 weeks. AM number, maturation, and polarization status were assessed by flow cytometry. Noninvasive and invasive strategies were used to assess pulmonary function and correlated with histomorphological assessments of airway structure and matrix deposition.

Results: Co-exposure to alcohol and CDPM decreased AM number and maturation status (CD11c expression), while increasing markers of M2 activation (interleukin [IL]-4Rα, Ym1, Fizz1 expression, and IL-10 and transforming growth factor [TGF]-β production). Changes in AM function were accompanied by decreased airway compliance and increased elastance. Altered lung function was attributable to elevated collagen content localized to the small airways and loss of alveolar integrity. Intranasal administration of neutralizing antibody to TGF-β during the CDPM exposure period improved changes in airway compliance and elastance, while reducing collagen content caused by co-exposure.

Conclusions: Combustion-derived PM inhalation causes enhanced disease severity in the alcoholic lung by stimulating the release of latent TGF-β stores in AMs. The combinatorial effect of elevated TGF-β, M2 polarization of AMs, and increased oxidative stress impairs pulmonary function by increasing airway collagen content and compromising alveolar integrity.

Key Words: Alcohol, Particulate, Macrophage, Pollution, Environmentally Persistent Free Radicals.

The most common causes of alcohol-related morbidity and mortality are organ damage and susceptibility to infection (Nelson and Kolls, 2002). In the lungs, chronic alcohol consumption has a profound and negative impact on immune cell function and the development of immune defenses (Happel and Nelson, 2005). In particular, alveolar macrophage (AM) dysfunction has been observed in chronic alcoholics (Mehta and Guidot, 2012). Specifically, chronic alcohol intake impairs AM terminal differentiation through the production of oxidative stress, leading subsequently to deficits in surfactant clearance, phagocytosis, and cytokine production (Brown et al., 2009; Joshi and Guidot, 2007).

AMs, in addition to controlling immune function, also participate in tissue remodeling and repair dependent upon their polarization status (Gibbons et al., 2011). In response to stimuli such as interleukin (IL)-1β/lipopolysaccharide (LPS), IL-4/IL-13, and transforming growth factor (TGF)-β/IL-10, AMs attain distinct activation/polarization states (reviewed in Gordon and Martinez, 2010). Polarizing stimuli subsequently alter expression of cytokine and scavenger receptors, costimulatory molecules and major histocompatibility complex, cytokine/chemokine production, and production of reactive oxygen/nitrogen species and enzymes. In the local tissue compartment, polarization thus influences recruitment of cytotoxic verses helper T-cells, oxidative stress, as well as fibroblast proliferation and matrix deposition. Thus, AM polarization status dictates repair and remodeling processes in airways and alveoli and influences
pulmonary function. Macrophage polarization has been well studied with focus on specific disease states such as bacterial infection, cancer, and asthma; however, research into how alcohol intake modulates polarization states in vitro and in vivo is in its infancy. Recent literature shows that M2 polarization occurs in AMs cultured in ethanol (EtOH)-containing media (Brown and Brown, 2012). M2 (or alternatively)-activated macrophages have been shown to play major roles in airway remodeling and pulmonary fibrosis, due to polarization by IL-4/IL-13 and the production of TGF-β/IL-10 (Gibbons et al., 2011; Homer et al., 2011; Sun et al., 2011). Although in vivo evidence suggests AM polarization may be impaired as a result of alcohol intake (Brown et al., 2009), how alcohol consumption may influence polarization in the immature AM pools has yet to be defined.

Limited epidemiological studies suggest an association between chronic alcohol consumption and increased airflow obstruction independent of smoking history (Emirgil and Sobol, 1977; Sisson et al., 2005), a disease state characterized by small airway remodeling and fibrosis accompanied by destruction of the alveoli. Although little is known regarding the relationships between chronic alcohol intake and pulmonary function (Sisson, 2007), chronic M2 macrophage polarization appears to play a prominent role in alcohol morbidity and mortality by stimulating remodeling/fibrosis (altering pulmonary function). However, links between the chronic M2 macrophage polarization in alveolar lungs and damage to the airways/alveolar space are not well understood.

Exposure to airborne particulate matter (PM) derived from cigarette smoke, industrial/urban, or automotive sources has been consistently linked with altered pulmonary function (Balakrishna et al., 2011; Gauderman et al., 2007; Moshammer et al., 2006) and predisposition or exacerbation of pulmonary diseases such as chronic obstructive pulmonary disease (COPD) and pulmonary fibrosis (Kelly and Fussell, 2011). Although the risks of PM exposure to pulmonary health are clear, limited epidemiological studies have examined chronic alcoholism as a comorbidity to PM exposure (Sisson, 2007). The relationships between alcoholism and PM exposure are particularly concerning given an estimated 18 million people in the United States with alcohol use disorders (Grant et al., 2004) and the high percentage of individuals who heavily consume alcohol and smoke (Batel et al., 1995). Although comorbid studies of alcohol and smoking are limited with conflicting observations, combinatorial effects have been observed in 2 clinical studies showing increased glutathione oxidation (Yeh et al., 2007) and exacerbation of COPD (Saric et al., 1977) in smoking alcoholics.

Our group has shown that combustion-derived PM (CDPM), such as that found in cigarette smoke and airborne PM derived from industrial treatment and refinery processes and consumption of biomass fuels, produces oxidative stress in biological environments (Balakrishna et al., 2011; Thevenot et al., 2013). Inhalation of CDPM leads to pulmonary oxidative stress, airway remodeling, increased airways resistance, and decreased compliance. As chronic alcohol ingestion increases expression of TGF-β and IL-13 (Brown and Brown, 2012), which are markers for M2 polarization, we hypothesized that M2 polarization and TGF-β production along with injury induced by CDPM inhalation would lead to fibrotic remodeling of the Airways. We tested this hypothesis in C57BL/6 mice fed a 5.3% liquid alcohol diet or isocaloric control diet for 8 weeks, an alcohol feeding period sufficient to induce clinical indicators of chronic alcoholism including liver microsteatosis, altered aspartate aminotransferase/alanine aminotransferase (ALT) ratio levels, and an increased pulmonary ratio of oxidized to reduced glutathione and elevated 8-isoprostanes. During the last 2 weeks of the protocol, mice were exposed to a laboratory-generated, endotoxin-free CDPM (referred to as DCB-230; Balakrishna et al., 2011; Thevenot et al., 2013) or filtered air. DCB-230 exposure at the concentration 200 μg/m³, exposure period, and duration employed in this study induces airway hyperresponsiveness analogous to epidemiological reports (Banauch et al., 2003).

MATERIALS AND METHODS

Mouse Model of Chronic Alcohol Ingestion

Female C57BL/6 mice (8 weeks of age; Harlan, Indianapolis, IN) were fed with Lieber-DeCarli liquid diet (32% calories from EtOH, 5.3% by volume) or nonalcohol isocaloric control diet for 8 weeks, an alcohol feeding period sufficient to induce clinical indicators of chronic alcoholism as previously described (Lomnicki et al., 2008). Briefly, copper oxide was reacted with amorphous cab-o-sil (<200 nm in diameter). These particles were then exposed to 1,2 dichlorobenzene (DCB) vapor at 230°C chemisorbing DCB onto the CuO-silica matrix. Particles were then cooled, dried, and stored under vacuum until use. DCB-230 had an aerodynamic diameter of 0.2 μm and was free of endotoxin.

DCB-230 Exposure Model

DCB-230 (DCB-230)—Synthesis and Dispersion

The environmentally persistent free radical containing PM₀.2<sub>DCB-230</sub>, was used as a model CDPM (Balakrishna et al., 2011; Thevenot et al., 2013). DCB-230 synthesis and chemical characterization were performed in the laboratories of Drs. Barry Dellinger and Slawomir Lonnicki (Louisiana State University, Baton Rouge, LA) as previously described (Lomnicki et al., 2008). Briefly, copper oxide was reacted with amorphous cab-o-sil (<200 nm in diameter). These particles were then exposed to 1,2 dichlorobenzene (DCB) vapor at 230°C chemisorbing DCB onto the CuO-silica matrix. Particles were then cooled, dried, and stored under vacuum until use. DCB-230 had an aerodynamic diameter of 0.2 μm and was free of endotoxin.

Invasive Measurements of Pulmonary Function

Twenty-four hours after the last exposure, lung function was assessed using the forced oscillation technique (FlexiVent; SCIREQ). Mice were anesthetized by intraperitoneal injection of...
Chemical exposure by inhalation was performed as previously described (Podowski et al., 2012) and was confirmed by qualitative real-time PCR. Quantitative real-time PCR on the ABI PRISM 7700 Sequence Detector System (PE Applied Biosystems, Foster City, CA) as previously described (Alcorn et al., 2003). Primers were synthesized with gene-specific intron-spanning primers (Yml, Arg1,Fizz1, Nos2, Tgf-β1). Single-species amplicons for each primer pair were verified by dissociation curves. Relative expression was calculated using the delta–delta C_t method and normalized to the reference gene Hprt1.

**Histology and Immunohistochemistry**

Blood was removed from the lungs by retrograde saline perfusion. A cannula was inserted into the trachea, and lungs were inflated to a pressure of 25 cm H2O, Zn-formalin-fixed, and paraffin-embedded. Sections (4 μm) were then taken of lung tissues and stained with hematoxylin and eosin (H&E; to visualize morphology and structure) or Masson’s trichrome (to visualize collagen and fibrosis). Immunohistochemistry was performed as previously described (Podowski et al., 2012) and quantified using a hydroxyproline assay kit (Biovision, Mountain View, CA) according to the manufacturer’s instructions.

**Flow Cytometry**

Perfused lungs were minced and enzymatically digested with collagenase and DNase to obtain a single-cell suspension using the gentleMACS Octo Dissociator (Miltenyi, Germany). Red blood cells were removed with RBC lysis buffer (eBioscience, San Diego, CA). Live cells were determined by trypan blue staining. Cells were stained with surface antibodies to F4/80-Pacific Blue, CD11b-APC, Live cells were determined by trypan blue staining. Cells were stained with antibodies to F4/80-Pacific Blue, CD11b-APC, and paraaffin-embedded. Sections (4 μm) were then taken of lung tissues and stained with hematoxylin and eosin (H&E; to visualize morphology and structure) or Masson’s trichrome (to visualize collagen and fibrosis). Immunohistochemistry was performed as previously described (Podowski et al., 2012) and quantified using a hydroxyproline assay kit (Biovision, Mountain View, CA) according to the manufacturer’s instructions.

**Bronchoalveolar Lavage, AM Isolation, and Ex Vivo Polarization and DCB-230 Exposure**

Bronchoalveolar lavage (BAL) was performed by inflating the lungs via the trachea with 1 ml of phosphate buffered saline (PBS; 3% bovine serum albumin [BSA]). As chronic alcohol ingestion alters AM polarization, maturation, and function, we hypothesized that chronic alcohol ingestion would alter AM numbers in the lungs after exposure to CDPM. DCB-230 exposure in control-fed mice led to a significant increase in the total number of AMs in the lungs; however, the total number of AMs in the lungs of alcohol-fed, DCB-230-exposed mice was significantly lower with respect to mice fed a control diet and exposed to DCB-230 (Fig. 1A). Depressed AM levels in the alcohol-fed, DCB-230-exposed mice were accompanied by a dramatic decrease in glutathione availability (GSH/GSSG; Fig. 1B). AM maturation status was assessed by flow cytometry via mean fluorescent intensity of CD11c and CD11b on gated AMs. CD11c expression was significantly decreased on AMs from alcohol-fed, DCB-230-exposed mice compared to control-fed mice. Overall, these results suggest that chronic alcohol ingestion impairs AM maturation and function after exposure to CDPM.

**RESULTS**

**Chronic Alcohol Ingestion Impairs AM Maturation and Exposure to CDPM Increases Oxidative Stress and Suppresses Pulmonary AM Numbers**

As chronic alcohol ingestion alters AM polarization, maturation, and function, we hypothesized that chronic alcohol ingestion would alter AM numbers in the lungs after exposure to CDPM. DCB-230 exposure in control-fed mice led to a significant increase in the total number of AMs in the lungs; however, the total number of AMs in the lungs of alcohol-fed, DCB-230-exposed mice was significantly lower with respect to mice fed a control diet and exposed to DCB-230 (Fig. 1A). Depressed AM levels in the alcohol-fed, DCB-230-exposed mice were accompanied by a dramatic decrease in glutathione availability (GSH/GSSG; Fig. 1B). AM maturation status was assessed by flow cytometry via mean fluorescent intensity of CD11c and CD11b on gated AMs. CD11c expression was significantly decreased on AMs from alcohol-fed, DCB-230-exposed mice compared to control-fed mice. Overall, these results suggest that chronic alcohol ingestion impairs AM maturation and function after exposure to CDPM.
alcohol-fed, DCB-230-exposed mice compared with alcohol or DCB-230 exposure alone (Fig. 1C), indicating impaired terminal maturation. Conversely, CD11b expression was significantly higher on AMs from alcohol-fed, air-exposed mice compared with mice fed a control diet and exposed to air (Fig. 1D). This increase was absent in alcohol-fed, DCB-230-exposed mice compared with mice fed a control diet and exposed to DCB-230, suggesting a comparatively more immature phenotype due to chronic alcohol ingestion that is decreased with DCB-230 exposure. Interestingly, an increase in the percentage of CD11b hi AMs (F4/80+, CD11c+) was observed in only the alcohol-fed mice exposed to DCB-230. This indicates that, while overall expression of CD11b in AMs is not changed in this group, a larger population of very highly expressing immature cells are present (Fig. 1E).

**Chronic Alcohol Ingestion Induces M2 Polarization Which is Exacerbated Following Exposure to CDPM**

We hypothesized that chronic alcohol ingestion would prime AMs for M2 polarization and that exposure to CDPM would further enhance M2 polarization. Expression of IL-10 and IL-4Rx was used to assess M2 polarization by flow cytometry, following antibody optimization using IL-4-stimulated MH-S macrophages in vitro. Polarization status of gated AMs from whole lungs was analyzed for each treatment group (Fig. 2). IL-10 and IL-4Rx levels were unchanged on AMs from the lungs of mice fed control diet alone or exposed to DCB-230 (Fig. 2A). However, both IL-10 and IL-4Rx were significantly upregulated in mice fed the alcohol diet compared with the control diet, with a further significant increase in alcohol-fed mice exposed to DCB-230.

M2 polarization was further confirmed by expression of Ym1 and Fizz1 in AMs obtained from the BAL (Fig. 2B). Both markers were significantly increased following alcohol ingestion and exposure to DCB-230 compared with both alcohol and DCB-230 exposure alone, suggesting that DCB-230 exposure increased M2 polarization following alcohol ingestion. As no change in M2 markers was observed following DCB-230 exposure in control-fed mice, we examined expression of the M1 marker inducible nitric oxide synthase 2 (Nos2) and the M2 marker Arginase 1 (Arg1; Fig. 2C). Arg1 levels were not statistically different among treatment groups; however, increased Nos2 expression and decreased Arg1:Nos2 ratios are consistent with an M1 phenotype. Nos2 expression was significantly elevated in control-fed, DCB-230-exposed mice resulting in a significant decrease in the Arg1:Nos2 ratio compared with mice fed an alcohol diet and exposed to DCB-230, indicating that chronic alcohol ingestion alters macrophage polarization associated with exposure to CDPM.

**Chronic Alcohol Ingestion During Exposure to CDPM Increase Active TGF-β Levels in the Lung**

To assess TGF-β-driven pulmonary fibrosis, we examined levels of TGF-β in the lungs and expression and production of TGF-β in AMs (Fig. 3) from the lungs of mice fed an alcohol diet and exposed to DCB-230. In the lung, total TGF-β was elevated in mice fed an alcohol diet and exposed to DCB-230, with respect to both alcohol and DCB-230 alone (Fig. 3A). To determine whether AMs were contributing to the increased levels of TGF-β in the lung, Tgf-β1 mRNA expression was quantified in AMs obtained from BAL (Fig. 3B). Tgf-β1 expression was significantly elevated in AMs from mice fed an alcohol diet and exposed to DCB-230 with respect to all other groups of mice. To determine whether chronic alcohol ingestion increased latent levels of TGF-β, we performed intracellular staining for LAP in AMs and analyzed by flow cytometry. Chronic alcohol ingestion resulted in a significant increase in the percentage of LAP+ cells (Fig. 3C). This suggests chronic alcohol intake increases latent levels of TGF-β in AMs and subsequent exposure to CDPM results in elevated expression of TGF-β and increased levels of active TGF-β in the lung.

**Exposure of Chronic Alcohol-Fed Mice to CDPM Reduces Lung Function in a Manner Consistent with Fibrosis**

As DCB-230 exposure of mice fed a chronic alcohol diet led to increased TGF-β production, we hypothesized...
that co-exposure might result in pulmonary fibrosis worsening pulmonary function. Pressure-volume loops were collected to derive quasi-static compliance (Cst). Exposure to DCB-230 resulted in a significant decrease in Cst (associated with fibrosis), with no additional exacerbation due to chronic alcohol ingestion (Fig. 4A). Newtonian resistance (Rn; Fig. 4B) was significantly reduced only in mice fed a chronic alcohol diet and exposed to DCB-230. Collectively, functional parameters in DCB-230-exposed mice trend toward fibrosis, due to loss of compliance and increased elastance (H; Fig. 4C) regardless of diet. However, DCB-230 exposure in chronic alcohol-fed mice appeared to elicit additional obstructive components as suggested by decreased Newtonian or central airway resistance.

Chronic Alcohol Ingestion and Exposure to CDPM Increases Airway Collagen

Pulmonary fibrosis was assessed by histological examination of collagen deposition in the lungs and quantification of lung collagen content using the hydroxyproline assay (Fig. 5). Histological quantification resulted in an insignificant increase in airway collagen content only in the mice fed a chronic alcohol diet and exposed to DCB-230 (Fig. 5A,B). However, hydroxyproline levels were significantly elevated only in the lungs of mice fed a chronic alcohol diet and exposed to DCB-230 (Fig. 5C). These data suggest that the decreased compliance observed in mice fed an alcohol diet and exposed to DCB-230 is at least partially due to increased airway collagen deposition.
CDPM Exposure in Chronic Alcohol-Fed Mice Causes Loss of Alveolar Integrity

We hypothesized that elevations in oxidative stress caused by chronic alcohol ingestion and exposure to DCB-230 may be accelerating parenchymal tissue destruction and emphysema compared with mice exposed to DCB-230 and fed a control diet. Therefore, alveolar integrity was examined by calculating space percentage and mean linear intercept (Fig. 6). Mice on a chronic alcohol diet exposed to DCB-230 exhibited significant increases in both the percentage of alveolar space (Fig. 6B) and mean linear intercept (Fig. 6C), compared with mice exposed to DCB-230 and fed a control diet. This suggested that chronic alcohol ingestion and exposure to CDPM increase alveolar tissue damage and emphysema.

Neutralizing Antibody to TGF-β Improves Pulmonary Function and Reduces Collagen Deposition Due to Chronic Alcohol Ingestion and Exposure to CDPM

We hypothesized that increased TGF-β production by M2-polarized AMs was central to increased collagen deposition and reduced pulmonary function indicative of fibrosis in the mice fed a chronic alcohol diet and exposed to DCB-230. To test this, we administered TGF-β-neutralizing antibody i.n. immediately following DCB-230 exposure on days 0, 2, 6, 10, and 14 of the exposure period. Pulmonary function and lung collagen levels were then analyzed (Fig. 7). In mice fed an alcohol diet and exposed to DCB-230, TGF-β neutralization did not statistically effect Cst or Rn (Fig. 7A,B). However, TGF-β neutralization significantly reduced tissue elastance in alcohol-fed, DCB-230-exposed mice (Fig. 7C). Additionally, neutralization of TGF-β significantly decreased collagen content in the lungs of mice fed an alcohol diet and exposed to DCB-230 (Fig. 7D). In total, these data suggest that aberrant TGF-β production in mice is responsible for the pulmonary dysfunction observed following DCB-230 exposure in mice chronically fed an alcohol diet.
Chronic alcohol intake leads to tissue destruction and organ damage (Guidot and Hart, 2005), which in the lungs manifest as deficits in pulmonary function consistent with airflow obstruction (Emirgil and Sobol, 1977; Sisson et al., 2005). Alcohol mediates these effects by altering immune cell function (Happel and Nelson, 2005; Joshi and Guidot, 2007), increasing TGF-β production (Bechara et al., 2004; Guidot and Hart, 2005), and increasing oxidative stress (Moss et al., 2000; Yeh et al., 2007). Pathologically, severe obstructive and restrictive lung diseases are also associated with oxidative stress and elevated TGF-β production, which stimulates remodeling, matrix deposition, and loss of alveolar structure (Bergeron et al., 2003; Podowski et al., 2012; Takizawa et al., 2001). Macrophages obtained from the lungs of patients with mild to moderate COPD (Shaykhiev et al., 2009) and fibrotic diseases (Furuhashi et al., 2010; Gibbons et al., 2011; Mathai et al., 2010; Murray et al., 2010) are M2 polarized, and several studies suggest the development of pulmonary fibrosis is directly related to elevated IL-4/IL-13 and TGF-β leading to persistent and/or exaggerated M2 polarization (Gibbons et al., 2011; Martinez et al., 2009; Pulichino et al., 2008; Yogo et al., 2009). In addition to stimulating remodeling and fibrosis, M2s increase TGF-β and IL-10 production (Mosser and Edwards, 2008). AM depletion or reversing M2 polarization greatly diminishes collagen deposition in many models of fibrosis (Gibbons et al., 2011; Homer et al., 2011).

As chronic alcohol intake induces oxidative stress and TGF-β production in lungs, we hypothesized that exposure of the alcoholic lung to CDPM (at concentrations found in urban locales) would further increase M2 polarization and activation exacerbating deficits in pulmonary function and tissue destruction. We investigated this hypothesis using C57BL/6 mice using nose-only inhalation of a laboratory-generated CDPM (DCB-230) in mice chronically fed a liquid alcohol diet.

We previously showed that nose-only exposure to PM in rodents induces inflammatory responses including increased AMs into the lung (Balakrishna et al., 2011; Thevenot et al., 2013). However, chronic alcohol appears to interrupt AM...
influx by suppressing myeloid development (Melvan et al., 2012) and reducing the maturation status of AMs (Brown et al., 2009; Joshi et al., 2005). Therefore, it was not surprising that AM levels in lungs of DCB-230-exposed mice chronically fed alcohol were reduced compared with mice fed a control liquid diet and exposed to DCB-230. The AM maturation markers, CD11c and CD11b, were also decreased in DCB-230-exposed, alcohol-fed mice compared with controls, demonstrating decreased AM maturation. These decreases could be the result of impaired differentiation from interstitial macrophages in the lung or from circulating monocyte pools, as AMs may derive from both.

As AMs of chronic alcohol-fed mice were less mature, we initially hypothesized M2 polarization in this setting would be impaired. However, steady-state AM turnover in the lung is slow, with data suggesting that a mere 15% of AMs turnover at 3 months and at most 40% at 1 year (Janssen et al., 2011). Our data demonstrate that resident BAL-derived AMs, whether obtained from mice fed a control or alcohol diet, are capable of polarizing to the M2s during extended culture in media containing alcohol. We therefore formulated the refined hypothesis that chronic alcohol intake increases M2 polarization of resident AMs.

IL-10 expression in AMs is used as a marker of M2 polarization, and IL-4Rα expression has also been a factor strongly correlated with M2. Chronic alcohol ingestion caused increased expression and production of these M2-associated factors in AMs consistent with an M2 phenotype. Exposure to DCB-230 in alcohol-fed mice led to further elevations in IL-10 and IL-4Rα with respect to alcohol intake alone as well as increased expression of the M2 genes Ym1 andFizz1, demonstrating increased capacity for AMs to polarize into M2s. Conversely, DCB-230 exposure in mice fed a control liquid diet led to an increase in Nos2 expression, indicative of M1 polarization. Several conflicting observations with regard to the effects of CDPM on AM polarization have been reported, in which AM phenotype is likely attributable to route of exposure, chemical/endotoxin contamination, and perhaps most influenced by the degree and duration of exposure (Brown et al., 2011; Miyata and van Eeden, 2011). Our results suggest that alcohol predisposes AMs to the M2 phenotype and that is exacerbated upon CDPM exposure.

Mounting evidence suggests AMs, specifically M2-polarized AMs, are the predominant source of TGF-β and depleting M2-polarized macrophages during the progressive phase of fibrosis reduces collagen deposition by limiting TGF-β production (Gibbons et al., 2011). TGF-β alone has been shown to predispose to M2 polarization by increasing expression of IL-4Rα expression (Homer et al., 2011). Increased levels of TGF-β contribute to the immature phenotype observed in AMs in this and previous chronic alcohol ingestion models (Joshi and Guidot, 2007). We observed elevated levels of total TGF-β and IL-4Rα in the lungs of alcohol-fed mice, and this was enhanced upon exposure to DCB-230.

With chronic alcohol, we observed increases in latent TGF-β in AMs with no increase in active TGF-β (similar to others; Bechara et al., 2004). CDPM exposure in chronic alcohol-fed mice led to elevated TGF-β levels in lung digests, elevated TGF-β1 transcripts in AMs, and decreased GSH/GSSG ratios and increased collagen content in the lungs. Oxidative stress alone has been shown to play an important role in the development of fibrosis through activation of latent TGF-β (reviewed in Liu and Gaston Pravia, 2010). In agreement with clinical data from smoking chronic alcoholics (Yeh et al., 2007), we found that co-exposure to CDPM and chronic alcohol induced oxidative stress. Therefore, it is plausible that increased oxidative stress activates latent TGF-β in addition to increasing TGF-β production through increased M2 activation.

Using pulmonary functional assessments and morphometric analyses of lung sections, we demonstrate that CDPM exposure decreased lung function in alcohol-fed mice by increasing fibrosis and disrupting alveolar integrity. Concurrently, recent studies on alcohol and lung function demonstrate that alcohol consumption alone decreases airway smooth muscle contraction (Oldenburg et al., 2009, 2012). We have previously shown that DCB-230 exposure reduced compliance and increased tissue elastance regardless of diet (Balakrishna et al., 2011), indicating an increase in stiffness. As anticipated, increased collagen deposition in the perivascular regions of the airways was observed in mice fed an alcohol diet and exposed to DCB-230 and correlated with a significant increase in lung hydroxyproline content. Histological examination of the alveolar spaces showed significant loss of alveolar tissue in alcohol-fed, DCB-230-exposed mice. Collectively, function and histological data illustrate how increased M2 polarization coupled with elevations in TGF-β increased disease severity in alcohol-fed mice exposed to CDPM.

Coupling our data with the work of others, we propose the following sequence of events which ultimately lead to enhanced fibrosis and lung dysfunction in CDPM-exposed chronic alcohol-consuming mice. AMs in chronic alcohol-fed mice are primed to develop into M2s capable of producing TGF-β. Subsequent pulmonary disturbances demonstrated here with CDPM further increase pulmonary oxidative stress and enhance M2 polarization/activation, leading to significant elevations in total active TGF-β. Similar to the phenotypes observed in pulmonary bleomycin and TGF-β overexpression models, M2 polarization and increased TGF-β lead to fibrosis, which here appears localized to the airways, in addition to destruction of the alveolar space. Collectively, these tissue architectural changes lead to decrements in pulmonary function. We hypothesized that excess TGF-β was critical in manifesting deficits in pulmonary function in alcohol-fed and DCB-230-exposed mice and show that decreasing TGF-β by administration of neutralizing antibody effectively reduced collagen content in the lungs and improved pulmonary function. This suggests targeting M2 polarization may be of therapeutic value to inhibiting...
tissue damage in chronic alcoholics exposed to CDPM. In conclusion, our data demonstrate alcohol consumption increases M2 predisposition in the lungs and leads to increased tissue damage and lung dysfunction.

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SUPPORTING INFORMATION

Additional Supporting Information may be found in the online version of this article:

Fig. S1. AM gating strategy.