A Role for Mast Cells in Alcohol-Induced Tissue Damage and Remodeling

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Abstract

Introduction: Alcohol continues to be one of the most frequently abused drugs in the world. While low levels of alcohol consumption may have health benefits, chronic abuse of alcohol deleteriously impacts most body systems and contributes to or exacerbates over sixty disease conditions. The mechanisms of organ and tissue damage in response to alcohol abuse include altered metabolic pathways, accumulation of reactive oxygen species and depressed immune function. Mast cells are multi-functional cells that have been classically described for their role in hypersensitivity reactions. More recently, roles for these cells have been elucidated in innate immunity and tissue remodeling. Mast cells perform these functions primarily through the secretion of a plethora of mediators that include histamine, heparin, serine proteases, cytokines and others. The specific factors that are produced and secreted at any time by mast cells depend in part on the tissue microenvironment providing the basis for extensive plasticity of these cells. Recent studies are beginning to define the role of mast cells in mediating the deleterious effects of chronic alcohol abuse. For instance, alcohol-induced damage to the gastrointestinal mucosa is at least in part mediated by activation of mast cells. Pharmacological inhibition of mast cell degranulation attenuates the increased permeability of the gastrointestinal epithelium associated with alcohol abuse.

Conclusion: Mast cells and their secretory products have been implicated in promoting a number of disease conditions. Recent studies have suggested an important role for these cells in alcohol-induced tissue remodeling. These cells and their specific secretory mediators may provide novel therapeutic targets in prevention or reversal of alcohol-induced tissue damage.

Keywords: Mast cells; Remodelling; Alcohol abuse

Introduction

Despite the clear health ramifications of chronic alcohol abuse, this continues to be one of the most frequently abused chemicals in the world. Approximately fourteen million Americans chronically abuse alcohol (National Institute of Alcohol Abuse and Alcoholism) and excessive alcohol use is the third leading cause of preventable death in the United States [1]. Alcohol abuse affects most organ systems and causes or exacerbates over sixty diseases including heart disease, Alzheimer’s, stroke, liver disease, diabetes mellitus and others [2-6]. The pathological effects of chronic alcohol abuse involve a number of mechanisms including altered metabolic, immunological, signaling and inflammatory pathways. Studies in several organ systems have illustrated that chronic consumption of high levels of alcohol alters immune and inflammatory responses including impairment of antimicrobial and antiviral immunity. Excessive consumption of alcohol is associated with increased incidence of pneumonia, tuberculosis, hepatitis C infection, enhanced susceptibility to HIV and increased propensity for some tumors [7-9]. These effects have been attributed to suppression of immune function by alcohol abuse. Many questions remain regarding the cellular and molecular mechanisms of these effects; however, substantial advances have recently been made in this area. Exposure to ethanol affects cytokine production by various immune cells [10-12]. Among the cells shown to be affected by chronic alcohol exposure, mast cells fill a particularly important role in innate immunity and deleterious tissue remodeling.

Mast cells

Mast cells are bone marrow-derived effector cells found abundantly in connective tissues situated at the interface of the body and the external environment including the integument and mucous membranes [13,14]. They are also found in connective tissue components of many organs, particularly in association with blood vessels, nerves and mucous glands [15]. Mast cells were originally described based upon their unique histological staining of large cytoplasmic granules [16]. The initial view that these cells were involved in nutrition of surrounding tissue gradually evolved as it was discovered that mast cell granules contain histamine and a functional relationship between histamine and anaphylaxis was developed [17,18]. These and other discoveries set the stage for the classical view of mast cells functioning in the early phases of immediate hypersensitivity reactions. In this process, antigens react with immunoglobulin E bound to the surface of mast cells promoting degranulation and release of mast cell mediators that are responsible for a number of the features typically associated with allergies including mucous secretion, bronchoconstriction and increased vascular permeability [19,20].

As research into mast cell function has continued, it has become increasingly clear that these cells are multifunctional and their specific
roles are likely regulated by the tissue microenvironment. Mast cells are involved in a variety of physiological processes including tissue repair, wound healing, angiogenesis and likely adaptive immunity [21]. Mast cells also play roles in defense against parasitic and bacterial infections [22]. More recently, dysregulation of mast cells and their secretory mediators has been implicated in playing a causal role in deleterious tissue remodeling, autoimmune disorders and cancers [23].

Mast cells produce and secrete a variety of biochemical mediators. Indeed, the large array of mediators produced by mast cells contributes to their functional plasticity [24]. These mediators are subdivided into three general groups: 1) preformed mediators, 2) neoformed or lipid mediators and 3) newly synthesized mediators. Preformed mediators are stored in cytoplasmic granules which are released upon mast cell activation (degranulation). Mast cells store a wide variety of preformed mediators including histamine, heparin, proteases and select cytokines such as tumor necrosis factor-α [21]. The release of preformed mediators typically occurs within seconds to minutes of stimulation providing an advantage for the involvement of these cells in immune surveillance [25]. One of the first and most extensively studied mast cell preformed mediators is histamine [18]. Histamine is a biogenic amine with diverse functions associated with allergic reactions including vasodilation, increased capillary permeability and bronchoconstriction [26-29]. More recent studies have described novel roles for histamine including involvement in autoimmune diseases and regulation of dendritic cell function [30]. Mast cell granules also contain active enzymes of the mast cell protease family, which includes chymases, tryptases and carboxypeptidase A [31]. Mast cells can be stimulated to degranulate by cross-linking of immunoglobulin E receptors, mechanical force, electrical activity and by various chemical activators [32]. The neoformed mediators are of the eicosanoid family and are synthesized as required from the fatty acids of membrane phospholipids [33,34]. Two families of enzymes catalyze fatty acid oxygenation to produce eicosanoids from arachidonic acid - cyclooxygenases and lipooxygenases. The mast cell-produced eicosanoids, prostaglandins and leukotrienes, have a variety of functions including stimulation of leukocyte migration, smooth muscle contraction, mucus production and others [15,35,36]. Members of the final group of mediators are newly synthesized when the mast cell is activated and are regulated in a stimulus-specific manner. These include diverse cytokines, chemokines and growth factors. Unlike release of preformed mediators during degranulation, production of newly synthesized mediators is a slower process more in line with the responses seen in other inflammatory and immune cells.

Mast cells exhibit extensive plasticity and heterogeneity in the production and release of mediators that directly reflect their microenvironment and the stimulus encountered. Multiple phenotypes of mast cells have been described based upon their localization and array of biochemical mediators produced. In rodents, two subtypes have been described that vary in the proteases produced. In mice, mucosal mast cells typically reside in the mucosa of the lung and gastrointestinal tract and produce mouse mast cell proteases 1 and 2. In contrast, connective tissue mast cells are located in intestinal submucosa, peritoneum and skin and are characterized by the production of mouse mast cell proteases 4,5 and 6 [37,38]. In humans, mature mast cells have also been divided into two subsets based upon their protease content. The mast cell tryptase (MCT) subset store tryptases in their granules while the mast cell tryptase/chymase (MCTC) subset store tryptases, chymases and carboxypeptidases in their granules [39-40]. While these characterizations provide a simple scheme for classifying mast cells, it is clear that these phenotypes can vary between tissues or even between cells in the same tissue. This is likely modulated by the biochemical milieu of the local environment as particular cytokines have been shown to promote the differentiation of specific mast cell phenotypes [15,41,42].

Roles for mast cells in the response to alcohol
While mast cells have begun to receive more attention regarding their role in normal tissue physiology and in diverse disease conditions, very few studies have focused on the contribution of these cells to alcohol-induced tissue damage. However, accumulating evidence suggests a role for these cells in pathogenesis associated with alcohol abuse. These studies will be summarized below.

Ethanol, mast cells and asthma
While controversial for some time, there is now compelling evidence that mast cells play a substantial causative role in the consequences of asthma [43]. Mast cells secrete mediators that can stimulate mucus secretion, bronchoconstriction and mucosal edema, which are all features of asthma. Bronchial mast cells exhibit features of ongoing activation in patients with asthma [44]. This is associated with alterations in the microlocalization of mast cells within respiratory tissues [45]. Most of the mast cells in the airway smooth muscle are characterized by expression of both tryptase and chymase. Experimental studies have illustrated that delivery of exogenous tryptase induces bronchoconstriction in dog and sheep models [46]. This response is blocked by anti-histamine suggesting that the tryptase effect on bronchoconstriction is mediated by histamine. Thus, mast cell activation appears to be an important mediator of asthma. Indeed, prostaglandin 2 has a protective effect against asthma and this may be in part due to prevention of mast cell activation [47]. Despite these advances, the relationship of mast cells to alcohol-induced respiratory conditions has not been extensively investigated.

The incidences of asthma and binge drinking are on the rise, particularly in young adults [48]; however, their interactions have remained relatively unexplored. A number of clinical studies have illustrated that alcohol use exacerbates asthmatic symptoms [49,50]. A recent study in allergen-sensitized mice illustrated that ethanol treatment induced mast cell degranulation and exacerbated the symptoms of asthma including airway constriction and mucous production [48]. Similarly, acetaldehyde, the first metabolite of ethanol, induces degranulation and histamine release from airway mast cells resulting in constriction of bronchi [51,52]. To date, perturbation studies have not been carried out to conclusively define the functional role of mast cells and their mediators in the effects of alcohol on asthmatic conditions.

Ethanol-induced gastric damage
Exposure of the gastrointestinal tract to ethanol results in vascular leakage in mucosal capillaries and postcapillary venules [53]. This contributes to hemorrhage and the development of necrotic lesions in the gastric wall [54]. Several studies have illustrated a correlation between the number of mast cells and the appearance of gastric lesions [55,56]. Compounds that promote mast cell degranulation (compound 48/80, for instance), induce gastric damage that appears very similar to the damage elicited by ethanol exposure [57]. A hallmark of the mast cell product histamine is its effects on the vasculature and induction of vascular leakage. The mechanism of this effect is the production of small gaps between endothelial cells in capillaries and small venules
Exposure of the gastrointestinal tract to alcohol also increases permeability of the mucosal epithelium [65,66]. This appears to be due to the metabolism of ethanol to acetaldehyde as ethanol itself has little direct effect on the epithelium. Acetaldehyde has been demonstrated to disrupt epithelial tight junctions [67]. Disruption of the epithelial barrier results in increased blood endotoxin levels, which in turn has substantial effects on other organs, particularly the liver. Endotoxin is a potent activator of liver Kupffer cells and appears to be a prerequisite for cirrhosis [68,69]. Mast cells themselves can induce disruption of the epithelial barrier suggesting that they may be involved in ethanol-induced alterations in this barrier [70,71]. Studies have illustrated that gastrointestinal bacteria are able to metabolize ethanol into acetaldehyde and inhibition of this with antibiotics reduces the disruption of the epithelial barrier by alcohol [72]. In addition, stabilization of mast cells with doxantrazole reduces the disruption of the epithelial barrier function by ethanol exposure. These studies have established a paradigm whereby ethanol is metabolized by endogenous gut bacteria and epithelial function is reduced in part by activation of mucosal mast cells by acetaldehyde.

Alcohol, mast cells and cancer

While the impact of alcohol abuse on a number of diseases has been investigated, its impact on cancer has not been as widely appreciated. Chronic alcohol use, in fact, promotes several epithelial cancers including esophageal, liver, breast and colorectal cancers [73-75] and results in poor prognosis of those. The underlying mechanisms mediating the effects of ethanol on cancer susceptibility and progression are not well understood but may include the formation of DNA adducts, lipid peroxidation, oxidative stress, inflammation or other processes. In the colon, mast cells participate in innate immunity to protect against microbial pathogens and parasites and to modulate the permeability of the gut epithelium. However, mast cells also respond to oncogenic signals in ways that appear to contribute to cancer progression [76-78]. Recent studies illustrated that chronic alcohol consumption induces an increase in intestinal polyps in the APC min/+ mouse model [79]. There was a corresponding increase in the number of mast cells in the stroma of polyps in the ethanol-treated animals including both tryptase-positive and chymase-positive cells. The functional significance of these alterations remain to be determined.

Mast cells and alcoholic cardiomyopathy

Mast cells and their secretory products have been implicated in mediating cardiac remodeling associated with a number of pathological conditions including hypertension, myocarditis, myocardial infarction and heart transplantation [80-84]. Increased cardiovascular load as occurs in hypertension results in myocardial hypertrophy and fibrosis. Concurrent with these changes is an increase in mast cell density in the myocardium [85]. Studies in which mast cells were stabilized with nedocromil illustrated a causal role for mast cells in myocardial remodeling and fibrosis of spontaneously hypertensive rats [84]. In these studies, mast cell stabilization also normalized the cytokine expression profiles associated with chronic hypertension. Mast cells have also been implicated in damage resulting from myocardial ischemia/reperfusion. In response to myocardial ischemia/reperfusion, mast cells release renin, which activates the local renin-angiotensin system [86,87]. This culminates in the local formation of angiotensin II, increased sympathetic activity and arrhythmias [87,88]. Interestingly, ischemic preconditioning inhibits subsequent ischemia/reperfusion-induced mast cell release of renin and activation of the local renin-angiotensin system [89]. Collectively these studies illustrate an important role for mast cells in diverse cardiovascular diseases and suggest these cells may be an important therapeutic target.
rapidly increased in the hearts of mice on the alcohol-containing diet (Figure 1). Further studies will be required to determine the functional role of mast cell activation in the progression of alcohol-induced myocardial remodeling; however, it is tempting to speculate that activation of these cells contributes to the progression of alcoholic cardiomyopathy similar to other cardiovascular conditions.

Cellular and molecular mechanisms of the alcohol effects

While it is becoming increasingly clear that mast cells contribute to the progression of alcohol-induced pathology; many questions remain regarding the molecular mechanisms of their effects. Alcohol exposure may impact a number of parameters of mast cell physiology including differentiation, activation/degranulation, gene expression, proliferation, migration and other processes. The response of mast cells and other inflammatory cells to alcohol will likely depend upon the local microenvironment. To date, few studies have focused on the direct effects of alcohol on mast cells.

As mentioned previously, chronic alcohol abuse impairs the immune response resulting in increased susceptibility of alcoholics to infectious diseases. Several studies have illustrated that alcohol treatment of animals inhibits expression of proinflammatory cytokines by macrophages and other cell types [7,95]. Pretreatment of isolated bone marrow-derived mast cells with ethanol in vitro results in attenuated IgE-mediated degranulation [12]. Similarly, long-term exposure to ethanol in an inhalation model, results in diminished mast cell degranulation following treatment with compound 48/80 [96]. These studies also illustrated that treatment of the HMC-1 human mast cell line with ethanol, inhibits tumor necrosis-alpha (TNF-α) and interleukin-8 production. In contrast to this, other studies have illustrated that treatment of mast cells with low doses of ethanol or acetaldehyde results in increased expression of cytokines including TNF-α, transforming growth factor-beta and interleukin-6 [97]. These effects were attenuated by pretreatment of the cells with mitogen-activated protein kinase (MAPK) inhibitors. The differences in these results may reflect a dose-dependent response of mast cells to ethanol and contribute to a potential explanation for the beneficial effects of low doses of alcohol consumption versus harmful effects of alcohol abuse.

Mast cells produce and store exceptionally high levels of serine proteases collectively termed mast cell proteases. These proteases include tryptase, chymase and carboxypeptidase A. Mast cell proteases may account for up to 25% of the total protein content in mast cells [98]. A diverse array of functions has been attributed to these enzymes including roles in the clearance of parasites and bacteria, cleavage of extracellular matrix components, activation of matrix metalloproteases and others. Studies have illustrated that relatively high doses of acetaldehyde stimulates the production of chymase by mast cells [99]. We have performed experiments to evaluate the effects of long-term (1 week) exposure of the rat basophil/mast cell line (RBL 2H3 cells) on mast cell protease expression. Following culture for 1 week in ethanol, the expression of mRNAs for mast cell proteases was evaluated. Mast cell protease 4, also known as β chymase, mRNA levels were significantly increased in a dose-dependent manner in these experiments (Figure 2). The mechanisms of this response and the functional significance of enhanced chymase production in alcohol-induced tissue remodeling remain to be determined.

![Figure 2](Image)

Figure 2: This graphically illustrates the relative expression of mast cell protease-4 (MCPT-4) by RBL 2H3 cells following treatment for seven days with the indicated ethanol concentration. Cells were treated continually with the respective ethanol concentration. Medium was changed daily during the treatment period. MCPT-4 expression was assayed by semiquantitative polymerase chain reaction. Expression of MCPT-4 was normalized to acidic ribosomal binding protein and the experimental treatments are expressed relative to the no ethanol control. Significance was determined by one way Anova comparison to the no ethanol control (*<0.05; n=3 independent cultures).

Mast cells are generated from bone marrow-derived precursors and mature in response to c-kit ligand, stem cell factor and the cytokine/growth factor milieu of the tissue. Increased numbers of mast cells in tissues can arise by proliferation of resident mast cells or recruitment of additional cells. Few studies have examined the effects of alcohol on either of these aspects of mast cell biology. Treatment of bone marrow-derived mast cells and the human mast cell line, HMC-1, with ethanol inhibits proliferation [12,100]. This was not accomplished by treatment of mast cells with acetaldehyde suggesting that this is a direct effect of ethanol on these cells [100]. Due to the effects of alcohol abuse on neurological development, a number of studies have evaluated the modulation of proliferation in the developing nervous system by ethanol. Exposure to high doses of ethanol inhibits proliferation of multiple cell types including glial cells [101,102], dermal fibroblasts [103] and neural crest cells [104]. Studies with a binge alcohol exposure model in adolescent rats illustrated an alteration of cell cycle kinetics in hippocampal progenitor cells [105]. Mechanistic studies evaluating inhibition of hepatic cell proliferation by ethanol have suggested dysfunction of iron metabolism as a contributing factor [106]. In neural stem cells, inhibition of proliferation appears to be dependent on alteration of extracellular signal-regulated kinase and phospholipase D by ethanol exposure [107]. In contrast to other cell types, ethanol exposure results in enhanced cell growth and proliferation of human embryonic stem cell lines [108]. Further studies are required to conclusively determine the effects of alcohol on proliferation of mast cells and their progenitors and to elucidate the mechanisms of these effects.

Ethanol exposure induces apoptosis of various cells including liver Hep G2 cells, macrophages, embryonic neural crest cells and cardiac myocytes [90,109,110]. Recent studies have also illustrated that...
treatment of human mast cells in vitro with ethanol decreases viability of the cells [100]. This was true of both the HMC-1 cell line and primary bone marrow-derived mast cells. Ethanol treatment promoted apoptosis of mast cells, as determined by TUNEL staining and activation of caspase-3. The fact that exposure to ethanol is thought to decrease proliferation of mast cells and promote apoptosis appears contradictory to the increase numbers of mast cells in tissues following alcohol consumption. Other processes that would increase mast cell number in tissues include enhanced migration of precursor cells or the promotion of mast cell differentiation by alcohol.

As mentioned previously, chronic alcohol consumption results in impaired immune function. Long-term ethanol exposure in an inhalation model results in reduced neutrophil and eosinophil migration [96]. Similarly, ethanol exposure reduces migratory capacity of cutaneous dendritic cells [111] and natural killer cells [112]. In the former studies, exposure to ethanol prevented the transition of the dendritic cells to a migratory phenotype with concomitant changes in gene expression that promote migration [113]. Other studies have illustrated that ethanol exposure alters microtubule assembly, which may impact migration, proliferation and other cellular processes [114]. In contrast to this, ethanol exposure appears to enhance the migratory ability of some cancer cells. Recent studies have illustrated that exposure to alcohol results in enhanced epithelial to mesenchymal transition including increased expression of matrix metalloproteases required for migration and metastasis [115]. Alcohol also has effects on the tumor microenvironment including alterations in the extracellular matrix that may enhance migration through the matrix [116]. The effect of alcohol on migration of mast cells has not been evaluated, but is an important aspect of mast cell-mediated responses that needs to be explored.

Conclusions

Mast cells play substantial roles in a variety of disease processes. Several recent studies have indicated a role for these cells in mediating the deleterious effects of chronic alcohol abuse making mast cells an intriguing therapeutic target. Ethanol or acetaldehyde treatment in vivo and in vitro has been shown to impact a variety of parameters important to mast cell function including gene expression, differentiation, degranulation/secretion, migration, proliferation and even survival. However, the molecular and cellular mechanisms whereby mast cells affect alcohol-induced disease are not well established. Further research in this area will be critical to development of specific therapeutic approaches with few side effects. Global inhibition of mast cell degranulation has been shown to prevent or slow the progression of several pathological conditions; however, this is likely to have significant detrimental effects as these cells play important physiological roles. Identification of individual factors involved in mast cell activation or secretory products of mast cells involved in specific processes will provide more specific targets for therapeutic purposes.

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References


