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## Heme Oxygenase-1 Promotes Granuloma Development and Protects Against Dissemination of Mycobacteria

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### Abstract

Non-tuberculous mycobacterial (NTM) infections occur in both immunocompromised and immunocompetent hosts and are an increasingly recognized cause of morbidity and mortality. The hallmark of pulmonary mycobacterial infections is the formation of granuloma in the lung. Our study focuses on the role of heme oxygenase-1 (HO-1), a cytoprotective enzyme, in the regulation of granuloma development and maturation following infection with *Mycobacterium avium*. We examined the role of HO-1 in regulating monocyte chemoattractant protein-1 (MCP-1) and chemokine receptor 2 (CCR2), two molecules involved in monocyte-macrophage cell trafficking after infection. We showed that RAW 264.7 mouse monocytes exposed to *M. avium* expressed HO-1 and MCP-1. Inhibition of heme oxygenase by zinc protoporphyrin-IX led to inhibition of MCP-1 and increased expression of CCR2, its cognate receptor. HO-1<sup>-/-</sup> mice did not develop organized granuloma in their lungs, had higher lung colony forming unit of *M. avium* when infected with intratracheal *M. avium*, and had loose collections of inflammatory cells in the lung parenchyma. Mycobacteria were found only inside defined granulomas but not outside granuloma in the lungs of HO-1<sup>+/+</sup> mice. In HO-1<sup>-/-</sup> mice, mycobacteria were also found in the liver and spleen and showed increased mortality. Peripheral blood monocytes isolated from GFP<sup>+</sup> mice and given intravenously to HO-1<sup>+/+</sup> mice localized into tight granulomas, while in HO-1<sup>-/-</sup> mice they

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remained diffusely scattered in areas of parenchymal inflammation. Higher MCP-1 levels were found in bronchoalveolar lavage fluid (BALF) of *M. avium* infected HO-1<sup>-/-</sup> mice and CCR2 expression was higher in HO-1<sup>-/-</sup> alveolar macrophages when compared to HO-1<sup>+/+</sup> mice. CCR2 expression localized to granuloma in HO-1<sup>+/+</sup> mice but not in the HO-1<sup>-/-</sup> mice. These findings strongly suggest that HO-1 plays a protective role in the control of *M. avium* infection.

## Keywords

Heme oxygenase-1 (HO-1); Chemokine receptor 2 (CCR2); Chemoattractant protein-1 (MCP-1); *Mycobacterium avium*; Granuloma; Disseminated lung infection

Non tuberculous mycobacterial (NTM) infection is a growing problem in the United States and remains poorly diagnosed in developing countries. An increase in NTM infections has been observed in the Southeastern United States<sup>1-5</sup>. Disseminated mycobacterial infection is one of the major causes of morbidity and mortality in patients who are immunocompromised such as those with HIV and AIDS. Recently, there has been an increase in pulmonary NTM infections in immunocompetent patients; mostly in older women and in patients with cystic fibrosis<sup>1,2</sup>.

*Mycobacterium avium* (*M. avium*) is a NTM that can cause infection in patients with underlying conditions that compromise local lung or systemic immunity. It is increasingly recognized as a major human pathogen, primarily following the emergence of the AIDS epidemic<sup>6</sup>. A hallmark of disseminated *M. avium* infections is the presence of mycobacteria in multiple organs including the liver, spleen, lymph nodes, bone marrow, and lung<sup>1,3,6</sup>. Granulomas are formed as a consequence of chronic antigen persistence and their formation involves the interaction between the infectious organism and host immune cells, including macrophages, and T-cells, as well as immune effectors such as chemokines and cytokines<sup>6,7</sup>. Mature granulomas include fibroblasts and extracellular matrix, which surround and isolate the granulomas from other tissues<sup>8</sup>. Importantly, organisms are not always eliminated from the granuloma, but can become dormant, resulting in latent infection<sup>9</sup>.

Heme oxygenase-1 (HO-1) is a cytoprotective enzyme which breaks down heme to produce carbon monoxide, iron and biliverdin<sup>10</sup>. HO-1 is induced by multiple stimuli including oxidative stress, pro-inflammatory cytokines and has been shown to be upregulated in lungs following mycobacterial infection<sup>10-12</sup>. While HO-1 derived carbon monoxide can induce the DosR dormancy regulon in mycobacteria leading to latency and survival of the organism inside host granuloma<sup>13</sup>, it is not clear whether HO-1 regulates the key host response of granuloma formation. Monocyte chemoattractant protein-1 (MCP-1/CCL2), a C-C chemokine, along with its receptor chemokine receptor 2 (CCR2) on monocytes-macrophages is responsible for the recruitment of mononuclear cells from peripheral blood to sites of inflammation<sup>14,15</sup>. However, a link between *M. avium* induced granuloma formation and HO-1 has not yet been established.

We evaluated the regulatory role of HO-1 in the recruitment of monocyte-macrophages and found that the activation of the MCP-1/CCR2 axis by *M. avium* infection was impaired by

inhibition of heme oxygenase (HO) activity. HO-1<sup>+/+</sup> mice showed mature, organized granuloma formation in lung tissue following *M. avium* infection without dissemination. In contrast, HO-1<sup>-/-</sup> mice had diffused, unorganized collections of mononuclear cells in the lungs with mycobacteria in the spleen and liver as evidence of dissemination of infection.

## Materials and Methods

### Mouse monocyte culture

RAW 264.7 cells were obtained from American Type Culture Collection (Manassas, VA) and maintained in Dulbecco's Modified Eagle's Medium (DMEM) as previously described<sup>16</sup>.

### Treatment of RAW 264.7 cells with *M. avium* and Zinc protoporphyrin-IX (ZnPP-IX)

Cells were plated on 60 mm culture dishes (Corning, Lowell, MA) at a concentration of  $1 \times 10^6$  cells and treated with *M. avium* ( $50 \times 10^6$  bacteria per plate) in serum free medium (SFM). Alternatively, cells were pretreated with 10  $\mu$ M ZnPP-IX (Frontier Scientific, Logan, UT) for 30 minutes in SFM and incubated at 37° C for different time points.

### Quantitative PCR analysis

Total RNA from cultured cells was purified by a commercial kit (RNeasy Mini Kit, Qiagen Science, Maryland). The quantitative analysis of MCP-1 and CCR2 receptor genes were assessed by PCR as described<sup>17,18</sup>. The mRNA levels of HO-1, MCP-1, and CCR2 were quantified using the mouse HO-1 forward - 5'-CACGCATATACCCGCTACCT-3', reverse - 5'-AAGGCGGTCTTAGCCTCTTC-3'; mouse MCP-1 forward - 5'-GGCTCAGCCAGATGCAGTTAA-3', reverse - 5'-CCTACTCATTGGGATCATCTTGCT-3'; and mouse CCR2 forward - 5'-CAACTCCTTCATCAGGCACAR-3', reverse - 5'-GGAAAGAGGCAGTTTGCAAAG-3' respectively.

### HO-1 knockout mouse model

We used the HO-1<sup>-/-</sup> mice generated by Poss and Tonegawa, and Kapturczak *et al.*<sup>19,20</sup>. Age-matched wild-type (HO-1<sup>+/+</sup>) littermates were used as controls. The protocol was approved by the Institutional Animal Care and Use Committee (IACUC) at the University of Florida.

### Mouse model of *Mycobacterium avium* infection

*M. avium* subspecies avium Chester (ATCC# 15769) was maintained in ATCC medium 90 Lowenstein Jenson medium and grown in Lowenstein-Jensen Medium Slants (BD Biosciences, San Jose, CA) according to manufacturer's instructions. HO-1<sup>+/+</sup> and HO-1<sup>-/-</sup> mice were infected with  $1 \times 10^7$  *M. avium* cells in PBS via intratracheal route once per week for three weeks. An additional group of wild type (HO-1<sup>+/+</sup>) control mice were inoculated with equal volume of PBS. After 6 months the mice were euthanized and lung tissue was harvested. Tissue was fixed in 4% paraformaldehyde at room temperature for 24 hours and processed for immunohistochemistry.

### Determination of Mycobacterial Colony Forming Units (CFU) in the lung

To assess mycobacterial growth, the lungs were removed aseptically at specified time points. The lungs were cut into small pieces, and homogenized. Viable mycobacteria in the lung homogenates were then assessed as CFU by performing serial dilutions from the lung homogenate and plating onto 7H11 agar in 6-well plates in duplicates. The plates were incubated under 100% humidity, 5% CO<sub>2</sub>, at 37 °C for two to three weeks, and colonies were counted. The plates were again incubated for an additional two weeks to decrease the possibility of failure to detect slower growing *M. avium* strains. No difference in the number of colonies was observed in re-incubated plates.

### Estimation of HO-1 protein by Western blot analysis

Lung tissues were weighed and lysed in RIPA buffer and 20 µg of total proteins were resolved by SDS-polyacrylamide gel electrophoresis and transferred to a nitrocellulose membrane. Proteins were detected with antibodies against HO-1 (1:1000), and β-actin (1:2000) (Santa Cruz Biotechnology, CA). Blotted antibody was developed by horseradish peroxidase-conjugated secondary antibody and enhanced chemiluminescence detection system. β-Actin was probed as an internal loading control.

### Quantitative morphometric examination

The effect of *M. avium* infection on pleural macrophage recruitment and granuloma formation as measured by the number and size of granuloma on the pleural surface and monocyte infiltration in the sub mesothelial area on light microscopy morphometric examinations. A quantitative method was used to evaluate the number and size of granuloma on gross pathology as well as light microscopy. Following sacrifice of the animal the chest cavity was opened and the image captured onto a Macintosh computer screen. Imaging data for granuloma is expressed as number of granuloma/mm<sup>2</sup>. The lung section were fixed and stained with hematoxylin and eosin. A bar grid scale set at 1-1000 prm is superimposed on the photomicrograph to permit quantification of numbers of macrophages and size of granuloma. Areas were randomly selected and include a maximum of 10 grids. The light microscopic images were captured onto the computer screen via an Olympus MT-2 microscope.

### Tracking GFP<sup>+</sup> monocytes in HO-1<sup>+/+</sup> and HO-1<sup>-/-</sup> mice infected with *M. avium*

Peripheral blood monocytes (PMBC) from GFP-transgenic C57BL/6 mice (monocytes in these mice constitutively express GFP) were isolated by density gradient centrifugation and incubated with serum free media either with IL-2 (200 units/ml based on preliminary observation), or IFN-γ (500 U/ml). 3 × 10<sup>6</sup> PMBC were inoculated intravenously via caudal vein into HO-1<sup>-/-</sup> mice with established pulmonary parenchymal infection by *M. avium*. The mice were euthanized after 24 hours and the lungs were perfused with heparinized Hanks' Balanced Salt Solution (HBSS). The lungs were flash frozen in OCT compound and 4 micron thick sections were sectioned. The sections were viewed under a fluorescence microscope at 465 excitation and 535 emission spectra (Zeiss Axiovert 200M) to localize the GFP positive monocytes in the lung section. The granuloma formation, bacterial clearance and mortality were estimated as previously described <sup>21</sup>.

### **MCP-1 expression by ELISA**

Mouse whole blood was collected through the abdominal aorta using a 25 gauge needle, centrifuged, aliquoted and stored at -80°C. Bronchoalveolar lavage fluid (BALF) was collected by ligating a catheter into the airway using thread and injecting 800-1000 µL of (HBSS) into the lung. The BALF was centrifuged, the supernatant was aliquoted and stored at -80° C and the BALF pulmonary alveolar macrophages (PAM) were collected for flow cytometry studies for analysis of CCR2 receptor expression. Supernatants were tested for MCP-1 using ELISA (R&D Systems, Minneapolis MN).

### **CCR2 expression by flow cytometry**

PBMC were isolated by density gradient centrifugation, and SFM either with IL-2 (200 units/ml based on preliminary observation), or IFN-γ (500 U/ml based on previously published (59) work) for 2 to 48 hours at 37°C in 5% CO<sub>2</sub> atmosphere. The PBMC were washed three times in PBS with 5% BSA and 5 mM sodium azide and incubated for 45 minute at 4°C either in the presence of rabbit anti-mouse CCR2 (1 µg/10<sup>6</sup> cells) monoclonal antibody or isotype Ab (IgG1). Cells were washed three times and labeled with goat anti-rabbit IgG1 FITC-conjugate to detect the antibody bound to the antigens. After incubations the cells were washed three times and fixed in 4% paraformaldehyde. The fluorescence associated with the PBMC's and PAM were analyzed by flow cytometry using a FACS star (Becton-Dickinson Immunocytometry Systems, Mountain View, CA). Fluorescence data were collected on log scale and the relative fluorescence intensity was reported by comparing their light scatter characteristics to those of normal cells analyzed in the same experiment.

### **Immunohistochemistry**

Paraffin embedded tissues were stained with hematoxylin and eosin (H&E). Sections were stained for CCR2 using anti mouse primary antibody (Capralogics, Hardwick, MA) and ABC staining kit (Vector laboratories, Burlingame, CA) according to manufacturer's recommendations. Alternatively, sections were stained with primary antibody for HO-1 (Calbiochem, San Diego, CA). Sections were also stained with TB Fluorescent Stain Kit M (BD Biosciences, Sparks, MD, USA) to visualize *M. avium* in lung tissue.

### **Statistical Analysis**

Data were analyzed by the unpaired student t-test using Statview software version 5.0, SAS Inst. INC. (Cary, NC). All experiments were done in triplicate. Differences were considered significant at P < 0.05 level.

## **Results**

### ***M. avium* induces HO-1 expression**

To address the role of HO-1 in *M. avium* infection, we exposed RAW 264.7 cells to *M. avium* at 50 multiplicity of infection (MOI). One day post infection, the cells were fixed and stained for HO-1. HO-1 was induced following 24 hours infection with *M. avium* as compared with untreated cells (Figure1 A, B). To substantiate the above observation, we

analyzed mRNA levels of HO-1 in RAW 264.7 cells treated with *M. avium* or untreated as previously described in methods section. Real-time PCR results showed that *HO-1* relative gene expression was significantly increased in a time dependent manner in RAW 264.7 cells following *M. avium* treatment (Figure 1C).

### **MCP-1 levels increase in RAW 264.7 cells treated with *M. avium***

In order to test the relationship between HO-1 and MCP-1, RAW 264.7 cells were pre-treated with 10  $\mu$ M ZnPP-IX, a competitive inhibitor of HO enzymatic activity for 30 minutes and then incubated with *M. avium* or SFM. After 4, 8 and 24 hours of incubation, cells were harvested and mRNA levels of *MCP-1* were analyzed by RT-PCR. MCP-1 protein levels were also measured in the supernatant of treated and control cells. We found that *MCP-1* mRNA levels were significantly increased in a time dependent manner when cells were treated with *M. avium* as compared to untreated cells (Figure 2A). However, as expected, these levels decreased significantly when we pre-treated the cells with the HO inhibitor ZnPP-IX (Figure 2A).

### **CCR2 mRNA expression is increased in RAW 264.7 cells after pretreatment with ZnPP-IX**

To determine whether *M. avium* infection and HO inhibition affect gene expression of *CCR2*, mRNA levels of *CCR2* were measured in RAW 264.7 cells following the same protocol as explained above. Our results indicated that *CCR2* gene expression was significantly increased following pretreatment with ZnPP-IX and *M. avium* infection. However no significant increase in *CCR2* gene expression was found in cells treated with *M. avium* as compared with non-treated cells (Figure 2B).

### **HO-1<sup>-/-</sup> mice failed to form mature, organized granulomas following *M. avium* infection**

Our *in vitro* results suggested that HO-1 appeared to play a role in the regulation of both MCP-1 and CCR2 following infection with *M. avium*. To evaluate the role of HO-1 in granuloma formation *in vivo*, we used a mouse model of HO-1 deficiency and compared granuloma formation in HO-1<sup>+/+</sup> mice and HO-1<sup>-/-</sup> mice following *M. avium* infection. We examined the lungs of HO-1<sup>+/+</sup> and HO-1<sup>-/-</sup> mice, six weeks, sixteen weeks and at six months post *M. avium* infection and compared the results with lungs from control mice inoculated with saline at the same time period. We examined the lungs of HO-1<sup>+/+</sup> and HO-1<sup>-/-</sup> mice six months post *M. avium* infection and compared the results with lungs from control mice inoculated with saline at the same time period. At the end of the experimental period, mice were euthanized and the lungs, spleen and liver were harvested for analysis. Lungs from HO-1<sup>+/+</sup> infected mice showed organized, mature granulomas whereas, lungs from HO-1<sup>-/-</sup> infected mice presented unorganized, loose collections of mononuclear cells (Figure 3, B, D).

To confirm granuloma formation, we analyzed H&E stained lung sections from the same mice. As shown in Figure 3, panel E-H, it was clear that HO-1<sup>+/+</sup> infected mice developed organized granulomas as compared to HO-1<sup>-/-</sup> mice which displayed loose, unorganized lesions in the lungs after exposure to *M. avium*. We used HO-1<sup>+/+</sup> and HO-1<sup>-/-</sup> mice inoculated with saline as a negative control for *M. avium* infection and observed no significant damage in lung tissue and no histological presence of granulomas (Figure 3A, C,

E, G). The inability of HO-1<sup>-/-</sup> mice to form mature, organized granulomas led to disseminated infection with more inflammation and lesions covering a greater area of the lungs (Figure 3 panel D, H). Disseminated infection was also evident in infected HO-1<sup>-/-</sup> mice, which presented diffuse collections of mononuclear cells as well as positive cultures for *M. avium* in the liver and spleen and displayed increased mortality (Table 1).

### **HO-1<sup>-/-</sup> mice failed to form mature, organized granulomas with dispersed *M. avium* in lung tissue**

Using the TB auramine stain M (BD Biosciences, Sparks, MD, USA), we were able to visualize fluorescently labeled *M. avium* in mouse lung tissue and in individual cells. We compared our H&E stained tissue (Figure 4, D-F) with the corresponding Auramine stained tissue (Figure 4, A-C). We observed a diffuse presence of mycobacteria in lung tissue in HO-1<sup>-/-</sup> mice following infection (Figure 4, panel C, F). However, mycobacteria were contained inside mature granulomas in HO-1<sup>+/+</sup> mice following infection (Figure 4, panel B, E). Saline controls did not show any mycobacteria in lung tissue (Figure 4, panel A, D).

### **HO-1<sup>+/+</sup> PBMC (from GFP<sup>+</sup> mice) home to the granuloma in HO-1<sup>+/+</sup> but not HO-1<sup>-/-</sup> mice**

In order to evaluate if reconstitution of HO-1<sup>-/-</sup> mice with HO-1<sup>+/+</sup> monocytes resulted in the recruitment of GFP<sup>+</sup> monocytes to the lung, with increased granuloma formation, increased clearance of mycobacteria and decreased mortality, monocytes were isolated from GFP<sup>+</sup> transgenic C57BL/6 mice and were inoculated intravenously via caudal vein into HO-1<sup>-/-</sup> mice with established *M. avium* infection. The GFP-PBMCs were disseminated diffusely in the lung tissue in HO-1<sup>-/-</sup> mice (Figure 5 C), pointed by arrows) and not organized into the granuloma at 6 weeks through six months. The GFP-PBMCs were disseminated diffusely in the lung tissue in HO-1<sup>-/-</sup> mice (Figure 5 C), pointed by arrows) and not organized into the granuloma. In HO-1<sup>+/+</sup> mice the GFP-PBMC were localized into granulomas and were better organized (Figure 5F, pointed by arrow) compared to HO-1<sup>-/-</sup> mice. These data demonstrate that in HO-1<sup>+/+</sup> mice, PBMC were recruited into lung granulomas suggesting modulates monocyte recruitment into granulomas.

### ***M. avium* CFU counts are higher in HO-1<sup>-/-</sup> lung tissue than in HO-1<sup>+/+</sup> tissue**

To test whether the failure of HO-1<sup>-/-</sup> mice to form mature granulomas had any effect on their ability to control and contain the infection, we harvested lung tissue from infected HO-1<sup>+/+</sup> and HO-1<sup>-/-</sup> mice and counted CFU in the tissue. *M. avium* were recovered from liver and spleen culture (Table-1). In HO-1<sup>+/+</sup> mice there was no dissemination of infection. There was no mortality (n=8) in this group even at 16 weeks following intratracheal infection of *M. avium*. In contrast, HO-1<sup>-/-</sup> mice infected with *M. avium* had persistent infection in the liver and spleen at all-time points (Table 1). Notably, HO-1<sup>-/-</sup> mice which failed to form organized, mature granulomas, also had significantly higher average *M. avium* CFU counts in their lung tissue as compared with HO-1<sup>+/+</sup> mice (Figure 6). It is possible that the increased mortality seen in the HO-1<sup>-/-</sup> is secondary to the dissemination of the organisms to other vital organs.

### **MCP-1 levels are significantly increased in BALF and serum of HO-1<sup>-/-</sup> mice following *M. avium* infection**

Since MCP-1 is the soluble ligand for CCR2, we analyzed the MCP-1 levels in BALF and serum samples from the HO-1<sup>+/+</sup> and HO-1<sup>-/-</sup> mice infected with *M. avium* or treated with saline. We observed a significant increase of MCP-1 levels in the BALF of infected HO-1<sup>+/+</sup> mice as compared to non-infected controls, as well as increased levels in HO-1<sup>-/-</sup> *M. avium* infected mice as compared to HO-1<sup>-/-</sup> non-infected controls (Figure 7A). Additionally, MCP-1 levels were higher in HO-1<sup>-/-</sup> control mice as compared with HO-1<sup>+/+</sup> control mice. Similar results were found when we analyzed the MCP-1 levels in the corresponding serum samples. HO-1<sup>-/-</sup> mice infected with *M. avium* showed higher levels of the cytokine as compared to HO-1<sup>+/+</sup> samples (Figure 7A). Furthermore, the levels of MCP-1 in sera of HO-1<sup>-/-</sup> mice infected with *M. avium* were significantly higher than HO-1<sup>+/+</sup> and non-infected HO-1<sup>-/-</sup> mice (Figure 7A). Although the MCP-1 profile was the same between BALF and serum samples for each condition, the amount of MCP-1 was up to 1000 fold higher in BALF than serum.

### **CCR2 expression was increased in PBMC of *M. avium* infected HO-1<sup>+/+</sup> and HO-1<sup>-/-</sup> mice and in infected pulmonary alveolar macrophages (PAM) of HO-1<sup>-/-</sup> mice**

CCR2 expression was significantly higher in PAM from *M. avium* infected HO-1<sup>-/-</sup> mice compared to PAM from *M. avium* infected HO-1<sup>+/+</sup> mice. In HO-1<sup>+/+</sup> mice infected with *M. avium*, the PBMC CCR2 expression was higher than compared to PAM CCR2 expression. These data demonstrate that in the absence of HO-1, CCR2 expression perpetuates a lack of retention of monocyte-macrophages at the site of infection (Figure 7B).

### **HO-1 protein levels increase in mouse lung tissue following *M. avium* infection**

In accordance with the HO-1 results found in RAW 264.7 cells treated with *M. avium*, we analyzed expression of this protein in the lung tissue of HO-1<sup>+/+</sup> and HO-1<sup>-/-</sup> infected and non-infected mice. We detected that HO-1 was expressed and co-localized in granuloma in HO-1<sup>+/+</sup> lungs infected with *M. avium* as compared to non-infected mice (Figure 8, A and B). Furthermore, we stained lung tissue from HO-1<sup>-/-</sup> infected and non-infected mice to confirm that these mice did not express HO-1 (data not shown). Western blot analysis confirmed that HO-1 protein levels were upregulated in lung tissue from *M. avium* infected mice as opposed to uninfected controls (Figure 8C).

### **CCR2 co-localizes to granulomas in HO-1<sup>+/+</sup> infected lung tissue but not in HO-1<sup>-/-</sup> infected lung tissue**

To confirm previous *in vitro* results that showed that HO-1 regulates the expression of CCR2, we examined CCR2 expression in the lung tissue of HO-1<sup>+/+</sup> and HO-1<sup>-/-</sup> *M. avium* infected and non-infected mice. We observed that HO-1<sup>-/-</sup> non-infected mice had higher basal levels of CCR2 in lung tissue compared to HO-1<sup>+/+</sup> non-infected mice (Figure 9, A and C). When we analyzed CCR2 levels in *M. avium* infected mice, we found that this receptor was up-regulated and co-localized to the granulomas in the lung tissue from HO-1<sup>+/+</sup> mice (Figure 9B). On the other hand, CCR2 expression in infected HO-1<sup>-/-</sup> lung



tissue showed a diffuse pattern of expression in agreement with loose collections of mononuclear cells that did not organize into granuloma (Figure 9D).

## Discussion

In this study, we tested the interaction and regulation of three proteins: HO-1, MCP-1 and CCR2 in the process of granuloma formation in a mouse model. HO-1, an inducible, cytoprotective protein has been implicated in many inflammatory diseases as well as mycobacterial infections<sup>10</sup>. Our results demonstrated that HO-1 plays a role in granuloma formation and maturation by regulating MCP-1 and CCR2 expression in mouse monocytes and therefore regulating the cellular recruitment of these immune cells from the periphery to the site of infection and injury. Our findings may also have substantial implications for understanding the role of HO-1 in the control of *M. tuberculosis* infection.

Since the advent of HIV and the AIDS epidemic, NTM infections have become a major source of disease around the world<sup>1,22-26</sup>. *M. avium* complex (MAC) infections have been on the rise in both immune competent patients as well as the immunocompromised<sup>1</sup>. The formation of granuloma is a vital part of the body's immune defense against invading pathogens. After infection with *M. avium*, failure to develop organized granulomas results in the dissemination of the infection and greater injury to the host<sup>6</sup>. The regulated recruitment of immune cells including monocyte-macrophages, T-cells, dendritic cells and B-cells to the site of infection and injury is essential for the host to contain and eliminate invading pathogens<sup>27</sup>. In the course of granuloma maturation, the recruitment of phagocytes and lymphocytes is driven by various cytokines and chemokines that are initially produced by mycobacteria-infected macrophages<sup>8</sup>.

MCP-1 is the chemokine responsible for creating a gradient for the chemotaxis of immune cells from the periphery to the site of infection and CCR2 is the cell surface receptor which is bound by MCP-1<sup>28</sup>. MCP-1 and CCR2 are both involved in the process of cellular recruitment to granuloma and in this study we demonstrate that HO-1 is a key regulator of these cytokines and therefore the immune recruitment process and granuloma formation.

From our *in vitro* experiment using RAW 264.7 mouse monocytes, we observed that HO-1 is induced by *M. avium* as early as 4 hours post treatment (data not shown) and stayed highly expressed up to 24 hours (Figure 1). Once activated by *M. avium*, these monocytes begin to express higher levels of both MCP-1 (Figure 2 A) and CCR2 (Figure 2 B). However, when the cells were treated with ZnPP-IX, an inhibitor for HO enzyme activity, a decreased expression of MCP-1 and an increase in expression of CCR2 were observed, suggesting that HO-1 levels are directly correlated with MCP-1 expression and inversely correlated with CCR2 expression.

Previous studies suggest conflicting evidence in terms of the correlation between HO-1, MCP-1 and CCR2. MCP-1 was shown to be downregulated at both mRNA and protein level with the inhibition of HO-1<sup>29-31</sup>. Zampetaki et al<sup>32</sup> showed that despite high mRNA levels, significantly lower levels of cytokine protein were found in BALF of mice over-expressing HO-1. These findings suggest that HO-1 overexpression can suppress the induction of

cytokine mRNA levels. Our *in vitro* and *in vivo* results, however, demonstrate that inhibiting HO enzyme activity led to altered MCP-1 protein levels in BALF and serum (Figure 7A), suggesting that HO-1 directly regulates MCP-1 expression in mouse monocytes. Since we observed an induction of HO-1 and increased levels of MCP-1 in the lung tissue, BALF and serum of infected HO-1<sup>+/+</sup> mice, it is possible that an MCP-1 gradient was established in order to recruit monocytes from the peripheral blood to the lung. This corresponds to a normal immune response in which cells expressing CCR2 are recruited from the periphery and move along an MCP-1 gradient towards the site of infection in order to form granulomas<sup>13</sup>.

This phenomenon of regulated expression of CCR2 is abnormal in HO-1<sup>-/-</sup> mice. However, in spite of high MCP-1 levels in the lung, there is poor recruitment of monocytes to the granuloma in the lung parenchyma. Chemokine receptor expression, the other critical component of the ligand-receptor interaction is affected since we found that in HO-1<sup>-/-</sup> mice CCR2 was not down regulated in the pulmonary alveolar macrophages (PAM). When testing MCP-1 protein levels in infected mice, we observed higher basal levels of MCP-1 in sera and BALF in control HO-1<sup>-/-</sup> mice as compared to wild type controls (Figure 7A). These results support previous findings<sup>19,32-34</sup> and suggest that HO-1<sup>-/-</sup> mice are in a chronic state of inflammation. This chronic inflammatory state could lead to the dysregulation of the MCP-1 chemotactic gradient, which would then disrupt cellular recruitment to the site of granuloma.

Previous findings along with our results suggest that different cell types in various organ systems may react differently to altered levels of HO-1. This may be due to direct regulation by HO-1 on MCP-1 gene expression via the NFκB signaling pathway. Jadhav *et al.* have shown that HO-1 and more specifically, hemin therapy, can reduce NFκB mRNA levels by 87.1%, which would therefore lead to a reduction in cytokine production<sup>35</sup>. It has been reported that when HO-1 enzymatic activity is inhibited by ZnPP-IX, the levels of CCR2 increase<sup>36,37</sup> suggesting that HO-1 regulates the expression of CCR2 on the surface of monocytes, macrophages and other immune cells. Our results show that HO-1<sup>-/-</sup> mice had increased levels of CCR2 on monocyte/macrophage cell surfaces as seen by relative gene expression (Figure 2B) and immunohistochemistry (Figure 9). Other studies<sup>38-40</sup> suggest that the lack of HO-1 causes a pro apoptotic milieu which leads to loss of an adequate monocyte-macrophage that can lead to decreased granuloma formation.

Our findings also suggest that HO-1 normally regulates the expression of the CCR2 during infection and allows directed migration of monocyte-macrophages along an MCP-1 gradient towards the organism. When we analyzed the results from our *in vivo* mouse model, we observed that the loss of HO-1 expression led to loose or unorganized collections of mononuclear cells in mice following *M. avium* infection. HO-1<sup>-/-</sup> mice failed to form mature, organized granulomas and this led to diffuse inflammation throughout the lungs and to dissemination of the infection to the liver and spleen. The significant increase of MCP-1 in the BALF of HO-1<sup>-/-</sup> mice (Figure 7A) following *M. avium* infection may lead to increased or unregulated recruitment of CCR2 expressing cells into the lung parenchyma and could also explain the diffuse, widespread inflammation in the lung tissue of these mice. These factors taken together could explain why these mice failed to form mature granulomas

and therefore, failed to contain the infection. An increase in colony forming units (CFU) was also observed in the lung issue of HO-1<sup>-/-</sup> mice as compared with HO-1<sup>+/+</sup> mice following *M. avium* infection (Figure 6). It is possible that the increased numbers of organisms combined with the dissemination to other vital organs contribute to the increased mortality in these mice. Based on these results, it seems that HO-1 plays a critical role in the regulation of cellular recruitment to the site of infection and in organized granuloma formation following infection with *M. avium*.

HO-1 is recognized to affect not just host responses but also mycobacterial expression of specific genes. CO a breakdown product of heme metabolism causes mycobacteria to express the dormancy (Dos) regulon<sup>13</sup>.

In conclusion, HO-1 is not only involved, but plays an essential role in the processes involved in cellular recruitment of monocytes from the peripheral blood to the sites of infection with mycobacteria. It is possible that other diseases in which granuloma formation is a vital host defense response, HO-1 may play a regulatory role in the response. The HO-1<sup>-/-</sup> mouse model provides a useful tool in the study of granuloma formation following *M. avium* infection. Our findings further substantiate the role of HO-1 in the regulation of MCP-1 and CCR2 and its key involvement in innate and adaptive immune responses following *M. avium* infection.

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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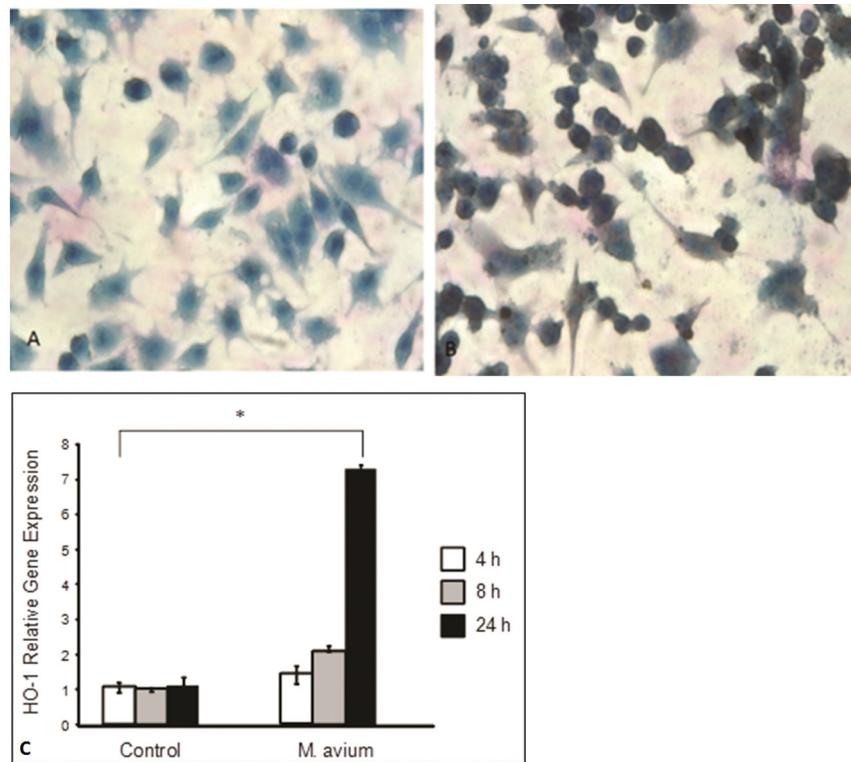
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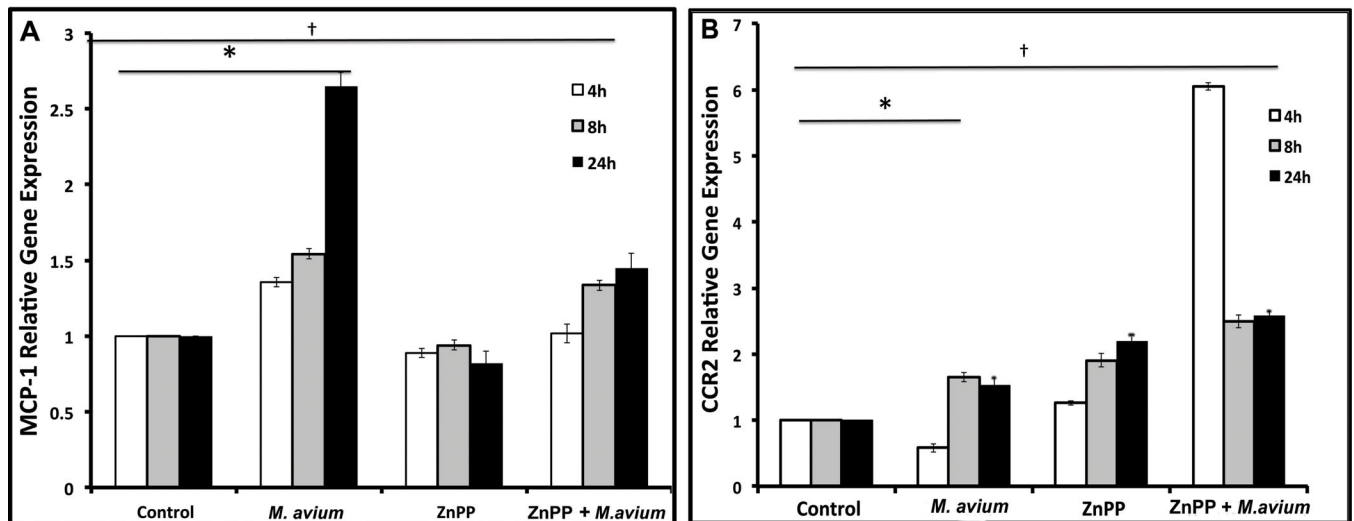
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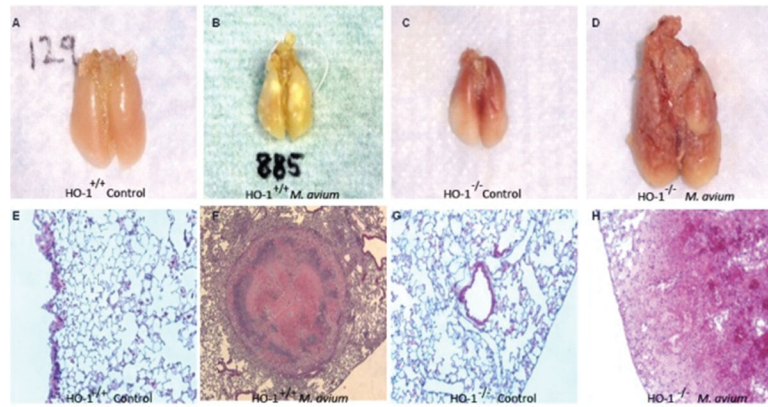


**Figure 1. *M. avium* infection induces HO-1 in RAW 264.7 monocyte-macrophages**  
**(A, B)** Representative light micrograph showing immunocytochemical detection of HO-1 in RAW 264.7. Panel A shows the control cells were incubated in SFM while panel B shows the cells infected with *M. avium* (50 MOI) for 24 hours. Cells were fixed and stained with anti-HO-1 antibody and visualized by optical microscopy at 400 $\times$  magnification. The infected cells stained for HO-1 expression (brown) after 24 hours as compared to control cells. Blue color shows counter-stain with hematoxylin. The photomicrograph is the representative of three individual staining. **(C)** HO-1 relative mRNA expression significantly increases in RAW 264.7 cells after infection with *M. avium*. Control cells were incubated in SFM and treated cells were incubated with *M. avium*. For 4, 8 and 24 hours. HO-1 relative gene expression was analyzed by RT-PCR. HO-1 levels significantly increased in cells treated with *M. avium* as compared to nontreated cells in a time dependent manner (\*P < 0.001). Values represent the mean of three different experiments.



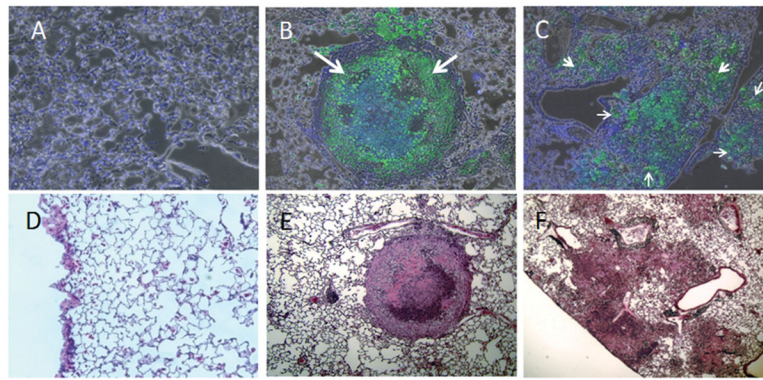
**Figure 2. HO inhibition decreases MCP-1 while increase CCR2 relative gene expression *in vitro* following the infection by *M. avium***

RAW 264.7 cells were pretreated with SFM (control) or ZnPP-IX (HO inhibitor) for 30 minutes and then incubated with *M. avium* for 4, 8 and 24 hours. MCP-1 and CCR2 relative gene expression was analyzed by RT-PCR. **(A)** MCP-1 levels were higher in *M. avium* treated cells as compared to controls (\* $P < 0.001$ ) and were significantly down-regulated († $P < 0.001$ ) when cells were pretreated with ZnPP-IX, for 8 and 24 hours as compared to *M. avium* alone for the same time points. No significant differences were observed at the 4 hour time-point. Values represent the mean of three different experiments. **(B)** CCR2 gene expression was significantly increased as compared with controls at all time points (\* $P < 0.05$ ). CCR2 relative gene expression was higher in cells pre-treated with ZnPP-IX as compared with cells treated with *M. avium* alone († $P < 0.05$ ). Values represent the mean of three different experiments.

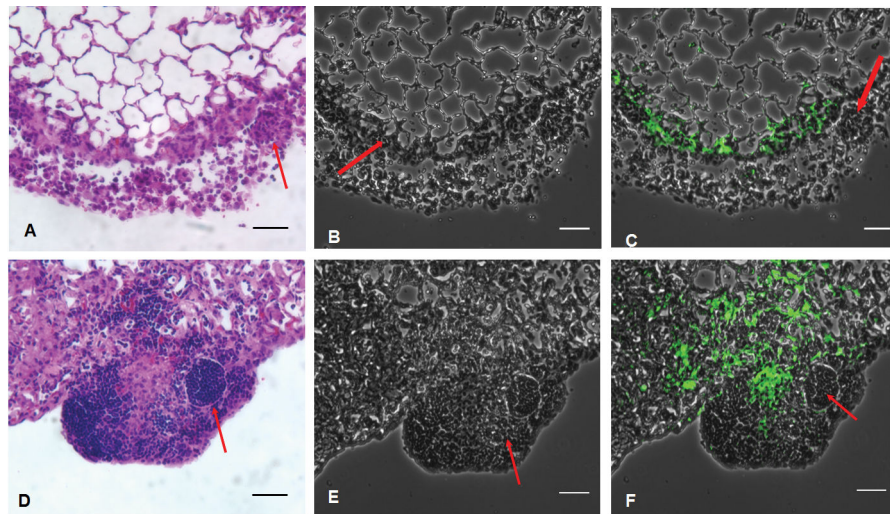


**Figure 3. Mouse lung tissue showing differential granuloma formation in HO-1<sup>+/+</sup> and HO-1<sup>-/-</sup> mice after 6 month *M. avium* infection or saline inoculation**  
 Mouse lungs were harvested following 6 month *M. avium* inoculation and visually inspected for presence of granuloma. Alternatively, lung tissue was fixed and stained with H&E to confirm granuloma formation. **(A)** Absence of granuloma in HO-1<sup>+/+</sup> control lung. **(B)** Presence of organized, mature granulomas in HO-1<sup>+/+</sup> lung inoculated with *M. avium* for 6 months. **(C)** Absence of granuloma in HO-1<sup>-/-</sup> control lung. **(D)** Diffuse infection and unorganized granuloma formation in HO-1<sup>-/-</sup> lung inoculated with *M. avium* for 6 months. **(E)** 200× H&E stain of normal tissue from HO-1<sup>+/+</sup> control lung. **(F)** 200× H&E stain showing organized granuloma formation in HO-1<sup>+/+</sup> lung inoculated with *M. avium* for 6 months. **(G)** 200× H&E stain of normal tissue from HO-1<sup>-/-</sup> control lung. **(H)** 200× H&E stain of diffuse infection and dysfunctional granuloma formation in HO-1<sup>-/-</sup> lung inoculated with *M. avium* for 6 months. Five mice were included in each treatment group (lungs are not shown to scale).



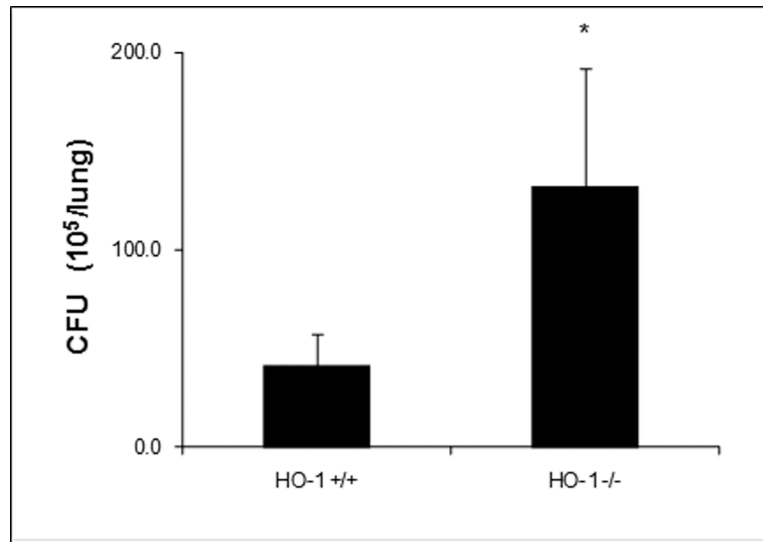


**Figure 4. HO-1<sup>-/-</sup> mice showed diffuse *M. avium* infection in lung tissue as opposed to HO-1<sup>+/+</sup> mice which contained the mycobacteria in mature, wellformed granulomas** Lung sections were stained with fluorescent auramine stain (BD Biosciences, Sparks, MD, USA) to show *M. avium* (green color) in lung tissue. **(A)** Auramine stain of HO-1<sup>+/+</sup> control lung. **(B)** Auramine stain of HO-1<sup>+/+</sup> lung infected with *M. avium* for 6 months. **(C)** Auramine stain of HO-1<sup>-/-</sup> lung after 6 months *M. avium* infection. White arrows show the presence of *M. avium*. **(D-F)** H&E stain corresponding to panels A-C.



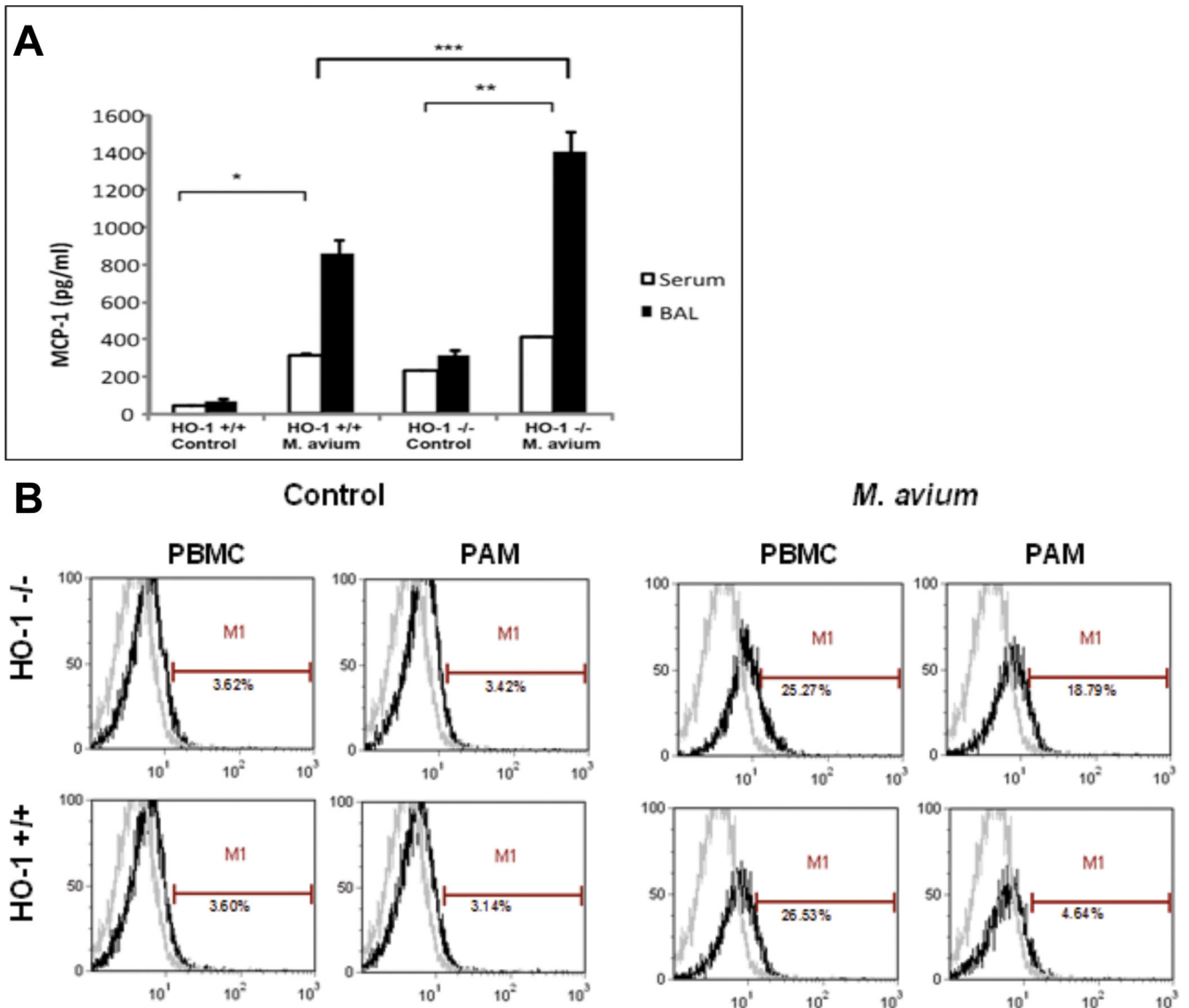
**Figure 5. HO-1 modulates development and recruitment of monocytes recruitment into granuloma**

HO-1<sup>-/-</sup> and HO-1<sup>+/+</sup> mice were infected with *M. avium* intratracheally as described in material and methods. At 6 weeks control and experimental mice were injected with PBMCs from GFP positive mice. The mice were euthanized 24 hrs post PBMC injection; the lungs were perfused with heparinised normal saline and fixed in 4% paraformaldehyde. The lungs were embedded in paraffin and 4 micron sections were prepared, stained with H&E (**A, D**). HO-1<sup>-/-</sup> mice demonstrated poorly organized granuloma formation (**A, B**). The HO-1<sup>+/+</sup> mice demonstrated highly organized granuloma (**D, E**). The red arrows points to the granuloma. No organized granuloma formation was present in the HO-1<sup>-/-</sup> mice (**A, B**). This is a single representative image of H&E stained images of three mice from each strain. (**C, F**) are the transmission microscopy images of pulmonary granulomas from HO-1<sup>-/-</sup> and HO-1<sup>+/+</sup> mice respectively. The HO-1 positive GFP labeled monocytes given intravenously tracked into the granuloma of HO-1<sup>+/+</sup> mice (**F**) but they remain dispersed in HO-1<sup>-/-</sup> mice (**C**). This is a single representative image of three similar images observed in six mice from each strain. The white arrows point to the granulomas. The green fluorescence indicated GFP positive monocytes. Magnification= 200×. Scale bar = 100 μm.



**Figure 6. *M. avium* CFU counts are higher in lung tissue of infected HO-1<sup>-/-</sup> mice as compared to lung tissue from infected HO-1<sup>+/+</sup> mice**

Mouse lung tissue was homogenized in 2ml of 7H9 broth medium. Then, 100  $\mu$ l of 10-fold serial dilutions were plated on 7H10 agar plates and incubated for 10-21 days in 37°C after which colony forming units were counted. CFU counts were significantly higher (\* P<0.05) in lung tissue of infected HO-1<sup>-/-</sup> mice as compared to HO-1<sup>+/+</sup> mouse tissue. Values represent the average CFU counts from four mice in either non-infected or *M. avium* infected groups.



**Figure 7. (A) MCP-1 levels are up-regulated in BALF and serum samples of *M. avium* infected HO-1<sup>+/+</sup> and HO-1<sup>-/-</sup> mice**

BALF and serum were collected and MCP-1 protein levels were measured by ELISA. MCP-1 levels were significantly increased in BALF of infected HO-1<sup>+/+</sup> mice as compared to HO-1<sup>+/+</sup> saline controls (\*P < 0.001) and HO-1<sup>-/-</sup> mice as compared to HO-1<sup>-/-</sup> saline controls (\*\* P < 0.001). MCP-1 levels are increased in mouse serum after 6 months *M. avium* infection and are significantly higher in infected HO-1<sup>+/+</sup> as compared with saline controls (\* P < 0.0001) and HO-1<sup>-/-</sup> mice (\*\* P < .0001) as compared with saline controls. MCP-1 levels were significantly higher in *M. avium* infected HO-1<sup>-/-</sup> mice then in *M. avium* infected HO-1<sup>+/+</sup> mice (\*\*\*) P < 0.05) Samples from five mice in each group were assayed in triplicate. **(B) CCR2 expression in PBMC and PAM of HO-1<sup>+/+</sup> and HO-1<sup>-/-</sup> mice post *M. avium* infection.** CCR2 receptor PAM in HO-1<sup>-/-</sup> mice with *M. avium* infection is significantly increased compared to *M. avium* infected HO-1<sup>+/+</sup> mice. CCR2 expression in PBMC and PAM in HO-1<sup>-/-</sup> mice with *M. avium* infection were comparable. This is a

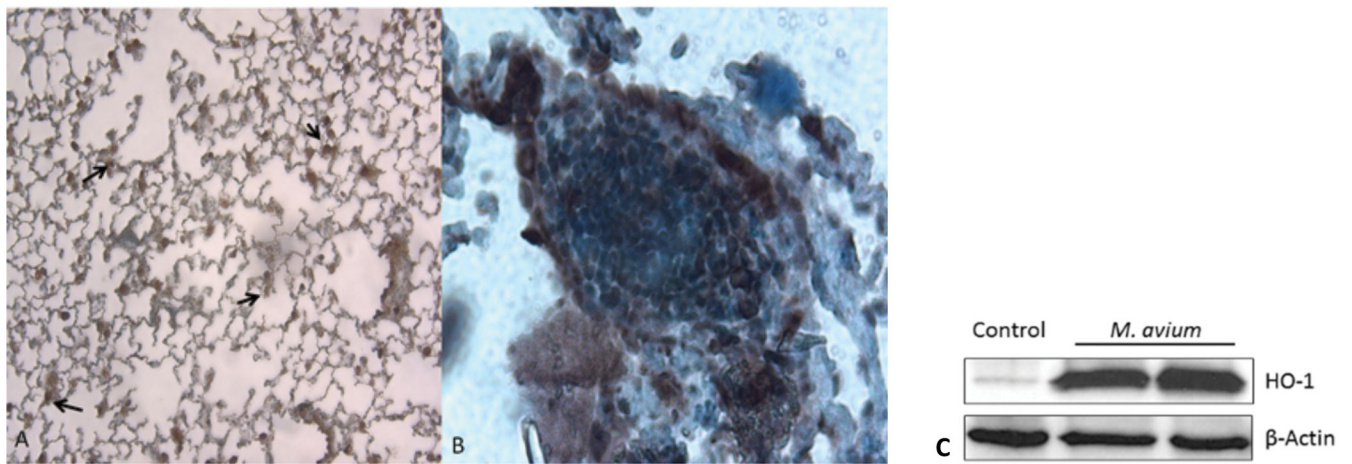
representative histogram of three experiments with similar observation. Control mice received saline.

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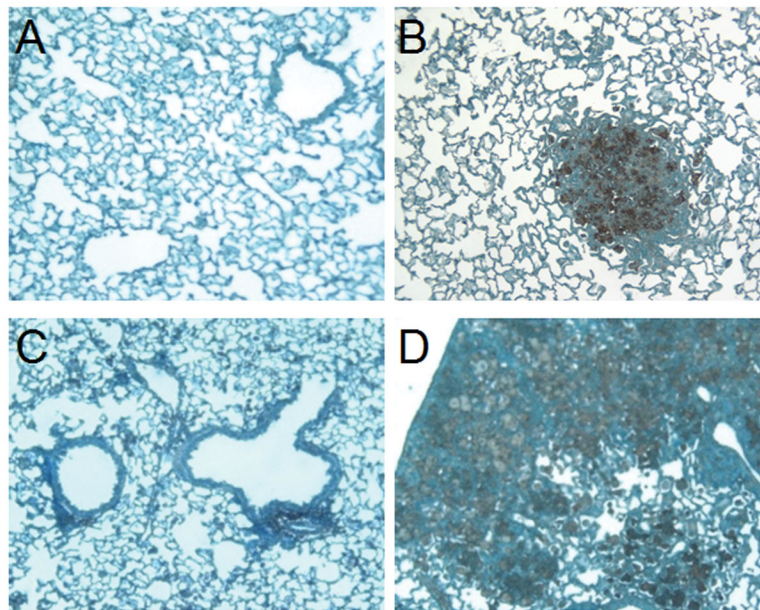
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**Figure 8. HO-1 protein expression is up-regulated and co-localizes to granuloma after *M. avium* infection in mouse lung tissue**

Paraffin fixed lung sections were stained using primary antibody against HO-1. **(A)** HO-1<sup>+/+</sup> mouse lung showing basal levels of HO-1 (brown) at 100× magnification. **(B)** HO-1<sup>+/+</sup> *M. avium* infected mouse lung showing increased expression of HO-1 (brown) in granuloma at 200× magnification. Nuclei were counterstained with hematoxylin (blue). Experiments were done in triplicate. **(C)** HO-1 protein levels were assessed in lung tissue of saline control mice (lane 1) and *M. avium* infected mice (lanes 2, 3) by western blot using primary antibody against HO-1. β-Actin was probed as an internal loading control. Blotted antibody was developed by horseradish peroxidase-conjugated secondary antibody and enhanced chemiluminescence detection system.



**Figure 9. CCR2 co-localizes to granuloma in HO-1<sup>+/+</sup> mouse lung tissue but shows a diffuse pattern of expression in HO-1<sup>-/-</sup> mouse lungs following *M. avium* infection**  
 Lung sections were stained for CCR2 receptor using anti-CCR2 antibody. (A) HO-1<sup>+/+</sup> lung tissue from saline inoculated mouse showing basal levels of CCR2 (brown). (B) HO-1<sup>+/+</sup> lung tissue from mouse inoculated with *M. avium* for 6 months. CCR2 (brown) expression was upregulated and co-localized to granuloma. (C) HO-1<sup>-/-</sup> saline inoculated mouse lung section showing basal levels of CCR2 expression (brown). (D) Increased and diffuse CCR2 expression in HO-1<sup>-/-</sup> mouse lung tissue following 6 months *M. avium* infection. Blue color shows counter-stain with hematoxylin. All pictures were taken at a magnification of 100×. Lungs were harvested from five mice in each treatment group.

**Table 1**  
**Granuloma formation and *M. avium* dissemination in HO-1<sup>-/-</sup> and HO-1<sup>+/+</sup> mice**

Test/Parameter	HO-1 <sup>-/-</sup> (n=8)	HO-1 <sup>+/+</sup> (n=8)
Granuloma size	29 ± 10 μm	1100 ± 102 μm
Granuloma number	3 ± 1	16 ± 5
Morphometric index	1 ± 0.5	10 ± 0.2
<i>M. avium</i> dissemination (to liver, spleen)	cultures positive	cultures negative
Mortality	50% by 6 weeks	None by 6 weeks

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