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Parul Kapil
Cleveland State University

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GLIA SPECIFIC INNATE RESPONSES AND THEIR INFLUENCE ON MURINE
CORONAVIRUS INDUCEDENCEPHALOMYELITIS

PARUL KAPIL

Bachelor of Science
Ch Charan Singh University, Haridwar, India
July 1999

Master in Microbiology
Gurukul Kangri Vishwavidalya
July 2001

Submitted in partial fulfillment of requirements for the degree
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This thesis/dissertation has been approved for
the Department of Biological, Geological,
and Environmental Sciences and for the
College of Graduate Studies of
Cleveland State University

By

Dr. Cornelia Bergmann, Chairperson     Date
Cleveland Clinic

Dr. Robert Silverman     Date
Cleveland Clinic

Dr. Crystal Weyman     Date
Biological, Geological, and Environmental Sciences

Dr. Vincent Tuohy     Date
Cleveland Clinic

Dr. William Baldwin     Date
Cleveland Clinic

Dr. Bruce Lamb     Date
Cleveland Clinic
DEDICATIONS

This thesis is dedicated to my parents, Rameshwar and Rekha Sharma, for their endless love support and encouragement.
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Writing this section is a bittersweet experience as one side it marks as an end of an era of my life as a graduate student while other side it indicates beginning of a new future. In my journey of past 5 years I have gained several experiences and I am indebted to the many people.

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GLIA SPECIFIC INNATE RESPONSES AND THEIR INFLUENCE ON MURINE CORONAVIRUS INDUCED ENCEPHALOMYELITIS
PARUL KAPIL

ABSTRACT

Control of viral infections of the central nervous system (CNS) is challenging, as the host has to balance anti-viral activity and immune mediated damage to non-renewable cells. The present study focuses on the role of glia and macrophages in regulating initial viral control and subsequent T cell activity in a murine model of neurotropic coronavirus induced encephalomyelitis. As induction of IFNα/β, pro- and anti-inflammatory factors depend on activation of pattern recognition receptors (PRR), macrophages, microglia and oligodendroglia were isolated from the CNS of infected mice to characterize their gene expression patterns. Oligodendroglia did not induce Ifnα/β expression following intracranial inoculation of both virus or the PRR agonist poly I:C, consistent with limited expression of PRRs and factors critical in the IFNα/β pathway. By contrast, induction of Ifnα/β by infected microglia and macrophages coincided with a broad and high basal expression pattern of IFNα/β signaling components and a protective antiviral state. Furthermore, an anti-inflammatory property of IFNα/β was suggested by abrogation of early Il1ra expression in both microglia and macrophage in the absence of IFNα/β signaling. Microglia and macrophages also induced a mixture of pro-inflammatory (Il12) and anti-inflammatory (Il1ra, Ym1/2) genes, which shape the adaptive immune response. IL12 and Ym1/2 stimulate T cells towards IFNγ and IL13 producing cells, respectively. The absence of IL12
decreased IFNγ and increased IL10 production, resulting in amelioration in clinical disease without altering virus control. Under the influence of distinct T cell mediated stimuli both microglia and macrophage retained a mixed pro- and anti-inflammatory phenotype defined by increased expression of iNos and Arg1, respectively. Sustained expression of Arg1 and increased phagocytic activity indicated an overall bias towards an anti-inflammatory phenotype with time. Finally, analysis of the role of PKR, an IFNα/β induced anti-viral gene, demonstrated only a modest direct anti-viral role. Surprisingly however, PKR deficiency reduced production of the anti-inflammatory cytokine IL10 by CD4 T cells, coincident with increased diffuse demyelination. These results demonstrate how a tightly linked network between innate and adaptive responses regulates viral control as well as tissue damage and disease in a viral encephalomyelitis model.
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CHAPTER I

INTRODUCTION

Various RNA and DNA viruses can infect the Central nervous system (CNS) and pose a serious threat as they can induce many different pathological outcomes such as encephalitis, meningitis, and demyelination. Viruses can affect CNS homeostasis in two ways; 1. They can directly infect and destroy CNS resident cells leading to neuronal damage, 2. The immune response to the virus can mediate damage to the CNS (Griffin, 2003). Neurotropic viruses generally invade the host through peripheral routes, but can enter the CNS by axonal spread or through the hematogenous route either as free virus or by transport in infected cells. Once viruses enter in the CNS, infected cells can mount a local innate immune response, which recruits peripheral immune responses to control the virus. However, due to the unique physiological and immune features of the CNS discussed below, viruses often evade the immune system and establish persistent or latent infection. Persistent infection in the CNS can be kept under control by the immune system in the absence of overt clinical symptoms. However, immune suppression or dysregulation may lead to loss of viral control resulting in fatal damage to the CNS. Viruses that can
infect the human nervous system include Herpes Simplex Virus-1 (HSV-1), Human Cytomegalovirus (HCMV), West Nile Virus (WNV), Poliovirus, Measles virus, Rabies virus, Human Immunodeficiency virus (HIV) etc (McGavern and Kang, 2011).

1.1 Host Barriers and Innate Immune Responses Associated With CNS Infection

Controlling virus infections in the CNS is a complicated task for the host because of the potentially damaging consequences of immune responses. Damage to non-renewable cells of the CNS is associated with increased permeability of vascular endothelium and influx of inflammatory cells releasing proteolytic enzymes and pro-inflammatory molecules, which can mediate cytotoxicity. Not surprisingly, therefore, the CNS is structurally protected by meningeal coverings and the blood brain barrier to minimize invasion of micro-organism as well as peripheral leukocytes. Furthermore, limited expression of major histocompatibility (MHC) molecules on CNS resident cells and absence of dedicated lymphatic drainage system (Galea et al., 2007) make it difficult to initiate adaptive immune responses. Overall, these features protect non-renewable CNS resident cells from infection as well as an overly active immune system. Furthermore, local production of the anti-inflammatory cytokine TGFβ and neurotrophins, as well as interactions between microglia and neurons through maintain an anti-inflammatory environment in the CNS (Hoek et al., 2000, Johnson et al., 1992, Neumann et al., 1998).
Nevertheless, following infection CNS resident cells quickly initiate expression of innate immune molecules. Glial cells can recognize an array of molecular patterns associated with micro-organisms (PAMPs) and endogenous danger signals (DAMPs) through receptors collectively known as Pathogen Recognition Receptors (PRRs) (Piccinini and Midwood, 2010, Hanke and Kielian, 2011). Signaling via PRRs leads to induction of a variety of innate inflammatory cytokines, type-1 Interferons (IFNα/β), chemokines. Rapid production of IFNα/β slows virus replication and constrains dissemination prior to emergence of virus specific adaptive immune responses. Furthermore, PRRs such as Toll like receptors-3 (TLR3) and IFNα/β induced protein kinase RNA dependent (PKR), can induce NFκB pathway, resulting in expression of a variety of pro-inflammatory genes (Alexopoulou et al., 2001). Although the pattern and magnitude varies with specific virus infection and the cell type infected, the most commonly induced factors are IL1, IL6, IL12, TNFα, CCL2, CCL5 and CXCL10 (Griffin, 2003). Furthermore, up-regulation of pro-inflammatory genes triggers the recruitment of leukocytes to the CNS (Engelhardt and Ransohoff, 2005, Griffin, 2003). Lymphocytes are activated in the periphery and restimulated by CNS resident cells presenting viral antigen. However, the pattern of pro-inflammatory molecules within the CNS can influence the pattern and magnitude of T cell cytokine expression. Anti-inflammatory molecules, such as IL10, TGFβ further limit the noxious effect of CNS inflammation to prevent tissue damage. IL10 primarily acts to suppress macrophages and DC function by inhibiting expression of MHC class II and co-stimulatory molecules such as CD80/CD86 and
production of proinflammatory cytokines and chemokines, including IL12 (Mosser and Zhang, 2008). The successful outcome of an infection thus relies on fine-tuning between pro- versus anti-inflammatory responses to control the infection while minimizing tissue damage.

1.2 Activation of CNS Resident Cells Following Viral Infection

Two types of PRR that recognize RNA viruses include endosomal receptors, Toll Like receptors (TLRs) and cytoplasmic receptors, Retinoic acid inducible gene-I (RIG-I) like receptors (RLRs, MDA5 and RIG-I). Microglia and astrocytes are the most prominent cell types in CNS in producing a large array of cytokines and chemokines. Microglia are the CNS resident macrophages and act as immune sentinels in the CNS. Murine microglia expresses all TLRs (TLR 1-9) at basal levels (Bsibsi et al., 2002, Olson and Miller, 2004). Activation of microglia by various exogenous and endogenous TLR ligands contributes to their either neuroprotective or neurotoxic phenotype. Primary murine astrocytes also express a wide variety of TLRs at basal levels, however at lower amounts than microglia. Specifically, robust expression of TLR3 in astrocytes under physiological conditions indicates their responsiveness against viral infections. (Bsibsi et al., 2006, Carpentier et al., 2005, Suh et al., 2009). Neurons and oligodendroglia can also express/induce a more limited array of TLRs; however, their role in induction of innate immune response following CNS infection is not well defined (Bsibsi et al., 2002, Suh et al., 2009). These studies indicate that activation of innate immune responses in the CNS is not homogeneous but rather tailored according to cell type and environmental cues (figure 1).
Figure 1. Role of TLR Signaling in Glial Cells Following CNS Infection (adapted from Carpentier et al., 2008). (A) Virus recognition, (B) Glial Cell Activation, (C) Recruitment and Activation of Peripheral Cells.

1.3 Experimental Models to Study Immune Responses in CNS

Due to limitations in obtaining CNS samples to study human viral CNS infection, several experimental viral models have been developed to understand mechanism regulating immune responses in the CNS. Among infections associated with demyelination are Theiler's murine encephalomyelitis virus (TMEV) infection and neurotropic coronavirus infection. TMEV causes persistent infection in microglia and macrophage; infected microglia induce production of various innate and co-stimulatory molecules and were able to efficiently process and present virus and myelin epitopes to infiltrating CD4 T cells (Olson et al.,
Furthermore, administration of IFNβ resulted in reduced infiltration of myelin specific CD4 T cells and reduced demyelination, which was associated with increased expression of the anti-inflammatory cytokine IL10 in the CNS (Olson and Miller, 2009). Similarly, Stimulation of TLRs on glial cells by Herpes Simplex Virus-1 (HSV-1), induces IFNα/β, IL15, TNFα and CCL2 expression (Conrady et al., 2010). IFNαβ mediates anti-viral effects by inducing proteins such as RNaseL and Protein Kinase R (PKR) (Samuel, 2001). In addition, the absence of IL10 prevents the severity of inflammatory responses caused by HSV-1 (Dakhama et al., 2009). Furthermore, treatment with both IFNβ and IL10 in mice infected with HSV-1 resulted in suppression in virus spread and prevented corneal inflammation (Minagawa et al., 1997).

These models indicate that the pattern of innate immune responses induced by various viruses is critical in dictating the outcome of an infection. To better understand the role of individual resident glia cells in mounting innate immune response and shaping the adaptive immune response in the CNS, we utilized murine gliatropic coronavirus CNS infection as a model of virus-induced encephalomyelitis.

1.4 Coronavirus (JHMV) Mouse Model to Study Encephalitis and Demyelination in CNS

Coronaviruses (CoV) are the largest positive sense single stranded RNA viruses (27-32kb genome). These belong to the group Nedovirales (nedo means nested in latin) which are characterized by their production of a nested set of mRNA with common 3’ end to express various viral gene products (Figure 2
MHV replication cycle). Virions are composed of four structural proteins namely spike protein (S), enveloped protein (E), membrane protein (M) and nucleocapsid protein (N) coated genome (Figure 2). The virus assembles in the Golgi Body and accumulates in cytoplasmic membrane-bound vesicles which subsequently fuse with the plasma membrane resulting in virus exocytosis (Masters, 2006).

Coronaviruses are the cause of 1/3 of common colds. While primarily infecting the respiratory tract, coronaviruses have also been reported to cause CNS infection (Bonavia et al., 1997, Yeh et al., 2004, Xu et al., 2005). The biggest outbreak of coronavirus (Severe Acute Respiratory Syndrome-coronavirus (SARS-CoV) ) infection was reported in 2002-2003 near Hongkong, which soon spread to 23 countries claiming 916 lives (WHO, retrieved 2008-10-13). With the exception of SARS CoV, human coronavirus infections have not received much attention due to their benign disease course. However, murine coronavirus infections of their natural rodent hosts are excellent models to study acute hepatitis, respiratory disease and encephalomyelitis associated with persistence. Specifically a virus variant of the neurotropic John Howard Muller strain of coronavirus (JHMV) has prominently been used to study control of virus replication in the CNS and immune mediated demyelination. The monoclonal antibody (mAb) selected variant of JHMV designated V2.2-v1 (characterized by point mutation and deletion in spike protein), used throughout in this study is gliatropic virus. Following intracranial inoculation virus initially replicates in ependymal cells and spreads to microglia, astrocytes, and oligodendroglia. Neurons are rarely infected. Infectious virus in the CNS peaks at day 5 post-
infection (p.i.), but is generally controlled by day 21 p.i. in a T cell dependent manner (Bergmann et al., 2006).

Figure 2. Mouse Hepatitis Virus (MHV) Replication (Bergmann et al., 2006).

Virus replication leads to activation of glial cells and expression of cytokines/chemokines that contribute to disruption of the blood brain barrier (BBB) integrity. The influx of T cells is associated with signs of encephalitis that can be observed ~7 days post infection (p.i.) (Fleming et al., 1986). Infiltrating T
cells control infectious virus, although viral antigen and RNA can be detected for the life of the rodent (Bergmann et al., 2006).

1.5 Role of Immune Response Following JHMV: Virus Clearance versus Pathogenesis

1.5.1 Innate immune response following JHMV and their interaction with adaptive immune response

Innate immune response in the CNS: JHMV infection initiates rapid induction of pro-inflammatory chemokines and cytokines. Early expression of CXCL1 and CCL2 is linked to CNS infiltration of neutrophils and macrophages, respectively. Macrophages constitute the major part of total early infiltrates at day 3, 5, and 7 p.i. Furthermore, abrogation of chemokines, CXCL1, CCL2, CCL5 and CXCL10 has been reported to prevent infiltration of T cells to the CNS (Liu et al., 2001, Glass et al., 2004, Hosking et al., 2009, Savarin et al., 2010). Similarly, JHMV infection induces pro-inflammatory cytokines such as IL1β, IL6, IL12, TNFα, iNOS (inducible form of nitric oxide synthase) (Parra et al., 1997, Rempel et al., 2005). Upregulation of these pro-inflammatory cytokines even in absence of cellular infiltrates indicates a direct response of CNS resident cells to JHMV. All these cytokines are involved in enhancing leukocyte infiltration to the CNS, but can also induce apoptosis. IL12 drives the differentiation of naïve CD4 T cells towards IFNγ producing Th1 cells. Treatment with recombinant IL12 has been shown to protect from lethal hepatotropic mouse hepatitis virus (MHV) in C57Bl/6 mice, but not in IFNγ R-/- mice indicating IFNγ dependent protective role of IL12 during hepatitis (Schijns et al., 1996). To evaluate the specific role of IL12
following JHMV infection in CNS, we utilized mice deficient in IL12 only (p35/-mice), mice deficient in IL23 only (p19/- mice), and mice deficient in both IL12 and IL23 (p40/- mice) as described in chapter 4.

Interferons (IFNs): Initially coronavirus were thought to not to induce IFNα/β production. However, the importance of IFNα/β signaling in controlling viral dissemination became apparent in vivo following peripheral as well as CNS infection of mice deficient in type-1 IFN receptor (IFNAR/-) (Ireland et al., 2008, Cervantes-Barragan et al., 2007). The absence of classical IFNα/β producing “plasmocytoid dendritic cells (pDCs)” in the CNS emphasize the importance of the ability of CNS resident cells to produce IFNα/β (Serafini et al., 2000). Very recently, in vitro and ex vivo studies indicate that microglia/macrophage, dendritic cells, and oligodendroglia (N20.1 cell line) can produce IFNα/β following CNS as well as peripheral MHV (coronavirus) infection (Rose and Weiss, 2009, Roth-Cross et al., 2008, Li et al., 2010). However the response of fully differentiated oligodendroglia compared to microglia in vivo following virus infection has not been studied. Thus, chapter 2 focuses on induction of IFNα/β and its response in microglia versus oligodendroglia in CNS following viral infection.

IFNs play a very important role following infection, not only by inducing antiviral genes (RNaseL, Protein Kinase RNA dependent (PKR)), but also by regulating immune response via the induction of antigen presentation molecules (MHC class I and classII), pro-inflammatory genes (IL12p70, IL6, iNOS) as well as anti-inflammatory genes (IL1 receptor antagonist, IL10). We previously reported a noncanonical role of the antiviral gene RNaseL following JHMV
infection. This study indicated that while RNaseL does not contribute to overall virus replication but rather affects virus control in macrophages and microglia following CNS infection. Furthermore, RNaseL-/- mice exhibited accelerated and severe demyelination and axonal damage compared to wt mice (Ireland et al., 2009). To further understand the downstream mechanism of IFNα/β mediated anti-viral response, we evaluated the role of another well-studied anti-viral gene; PKR in JHMV mediated CNS infection. Chapter 3 focuses to understand the role PKR in virus replication as well as in modulating pro- and anti-inflammatory cytokines.

1.5.2 Adaptive immune response

Both T cell mediated as well as humoral immune responses are pertinent to the control of JHMV infection at multiple levels.

Virus Control: The inability of irradiated mice to clear virus following JHMV infection emphasizes the importance of both T as well as B cells. Antigen specific CD8 T cells contribute a major part in controlling infectious virus from the CNS (Bergmann et al., 1999, Bergmann et al., 1993, Castro and Perlman, 1996, Stohlman et al., 1995). CD8 T cells control virus replication in different cell types using different effector mediators; virus is controlled in astrocytes and microglia by perforin dependent cytolysis, while IFNγ secretion is important in controlling virus replication in oligodendroglia (Bergmann et al., 2003, Bergmann et al., 2004, Gonzalez et al., 2006, Parra et al., 1999). CD4 T cells also contribute to virus control by supporting CD8 T cell function and directly affecting virus replication (Savarin et al., 2008, Stohlman et al., 2008). While T cells are indispensable in
controlling virus replication, however, virus specific antibodies provide protection from recurrence of infectious virus during persistence (Lin et al., 1999).

Pathology: One of the hallmarks of JHMV infection is demyelination. Demyelination was originally thought to be due to cytolytic infection of oligodendroglia. However, the absence or sparse demyelination in SCID or irradiated mice despite high oligodendroglia infection suggests a contribution of adaptive immunity (Wang et al., 1990, Houtman and Fleming, 1996). Restoration of demyelination after transfer of spleen as well as purified T cells supports a role of T cells in destruction of myelin (Houtman and Fleming, 1996, Fleming et al., 1993).

1.5.3 Anti-inflammatory response

Suppression of infectious virus progresses to persistent infection accompanied by chronic ongoing demyelination following JHMV infection (Weiner, 1973). The inability to completely eliminate viral antigen and RNA suggests an important role of anti-inflammatory responses designed to preserve CNS function at the expense of virus persistence. During acute JHMV encephalomyelitis anti-inflammatory cytokines IL10 and IL4 are upregulated with the same kinetics as pro-inflammatory cytokine IFNγ and TNFα (Parra et al., 1997). Lack of IL10, but not IL4 resulted in increased morbidity/mortality without altering virus clearance (Lin et al., 1998). Furthermore, a reengineered JHMV strain to produce IL10 confirmed a protective role of IL10. Increased IL10 production resulted in reduced microglial activation, expression of pro-
inflammatory cytokines and chemokines, infiltration of lymphocytes to the brain, and demyelination (Trandem et al., 2011).

The balance between pro- versus anti-inflammatory responses is critical to achieve effective viral control, but minimal tissue damage. Activation of microglia and macrophages play an important role following JHMV mediated CNS infections based on their role as primary innate responders. The plasticity of activated pro-inflammatory versus immunomodulatory microglia/macrophage thus raises an interest in their respective roles in downregulating the immune response as virus is controlled in the CNS. In chapter 5 we explored the existence of pro as well as anti-inflammatory phenotypes across CNS resident microglia and infiltrating macrophages following JHMV infection.
CHAPTER II

OLIGODENDROGLIA ARE LIMITED IN TYPE I INTERFERON INDUCTION
AND RESPONSIVENESS IN VIVO

2.1 Abstract

Type I interferons (IFNα/β) provide a primary defense against infection. Nevertheless, the dynamics of IFNα/β induction and responsiveness by central nervous system (CNS) resident cells in vivo in response to viral infections are poorly understood. Mice were infected with a neurotropic coronavirus (CoV) with tropism for oligodendroglia and microglia to probe innate Antiviral Responses during acute encephalomyelitis. Expression of genes associated with the IFNα/β pathways were monitored in microglia and oligodendroglia purified from naïve and infected mice by fluorescent activated cell sorting. Compared to microglia, oligodendroglia were characterized by low basal expression of mRNA encoding viral RNA sensing pattern recognition receptors (PRRs), IFNα/β receptor chains, interferon sensitive genes (ISG), as well as kinases and transcription factors critical in IFNα/β signaling. Although PRRs, including CoV RNA sensing Mda5, and ISGs were upregulated by
infection in both cell types, the repertoire and absolute mRNA levels were more limited in oligodendroglia. Furthermore, although oligodendroglia harbored higher levels of viral RNA compared to microglia, *Ifna/β* was only induced in microglia. Stimulation with the dsRNA analogue poly (I:C) also failed to induce *Ifna/β* in oligodendroglia, and resulted in reduced and delayed induction of ISGs compared to microglia. The limited Antiviral response by oligodendroglia was associated with a high threshold for upregulation of *Ikke* and *Irf7 transcripts*, both central to amplifying IFNα/β responses. Overall, these data reveal that oligodendroglia from the adult CNS are poor sensors of viral infection and suggest they require exogenous IFNα/β to establish an antiviral state.

2.2 Introduction

Type I interferons (IFNα/β) are critical innate cytokines that limit viral replication and dissemination prior to the emergence of adaptive immune responses. In addition to inducing an antiviral state, IFNα/β exert numerous other functions including induction of apoptosis, mobilizing innate cells, and regulating adaptive immune responses (Barton, 2008, Borden et al., 2007, Stetson and Medzhitov, 2006, Vilcek, 2006). As these pleiotropic effects can be detrimental if left unregulated, the innate Antiviral response has to be controlled to minimize cell injury and maintain cellular homeostatic functions. Although an early IFNα/β response contributes to viral control and disease outcome, its precise regulation is especially critical in the central nervous system (CNS), where the health of non renewable neurons as well as resident glia is vital for the maintenance of host physiological function. IFNα/β production and signaling is
thus tightly regulated by both the basal and inducible expression of numerous factors involved in the IFNα/β pathway.

During viral infections the ability to induce IFNα/β is determined by activation of pattern recognition receptors (PRR) following interaction with pathogen associated molecular patterns, which typically constitute viral RNA or DNA structures. PRRs comprise members of the Toll-like receptor (TLR) family and the cytosolic helicase sensors, retinoic acid-inducible gene 1 (RIG-I) and melanoma differentiation-associated antigen 5 (MDA5) (Kawai and Akira, 2009, Mogensen, 2009). PRR recognition of viral RNA triggers the activation and nuclear localization of interferon regulatory factors (IRF) IRF3 and IRF7 leading to IFNα/β induction. By binding to the IFNα/β receptor secreted IFNα/β initiates a signaling cascade leading to transcription of a variety of interferon sensitive genes (ISG). ISG encode both direct antiviral factors as well as PRR and signal transduction components, such as IRF7 and STAT1 (Borden et al., 2007). This amplification loop thus elevates the ability of cells to induce and respond to IFNα/β. In general IRF7 is considered the master switch in IFNα/β induction following various viral infections due to its role in amplifying IFNα/β production (Honda and Taniguchi, 2006). Since the pattern as well as levels of PRR, IRF and IFNα/β receptor expression and activation varies with each cell type, initial IFNα/β induction and amplification can be very distinct depending on the cell types infected (Colonna, 2007, Stewart et al., 2005). Furthermore, due to the potency of IFNα/β in inhibiting viral replication, many viruses have evolved mechanisms to antagonize the IFNα/β pathway, either at the induction or
signaling level (Levy and Garcia-Sastre, 2001, Sen, 2001). Microglia and astrocytes are primary sentinels within the CNS parenchyma responding to stimuli triggered by either microbial infection or degenerative events (Dong and Benveniste, 2001, Hanisch, 2002, Paul et al., 2007). This function is partially attributed to basal expression of a vast array of PRRs (Bsibsi et al., 2002, Carty and Bowie, 2011, Hanke and Kielian, 2011, Okun et al., 2009, van Noort and Bsibsi, 2009). *Irf3* mRNA and to a lesser extent *Irf7* mRNA, are constitutively expressed in the CNS. While *Irf7* expression is highly inducible, *Irf3* transcripts remain constant following infection (Ousman et al., 2005). However, their relative expression in different cell types has not been extensively explored. Consistent with PRR activation, microglia, astrocytes and neurons all induce IFNα/β (Paul et al., 2007). However, the vast majority of information on innate responses is derived from primary glia and neuronal cultures established from neonates and may not reflect responsiveness of fully differentiated cells. Distinct from *in vitro* studies, IFNα/β production *in vivo* is highly restricted as indicated by the small proportion (less than 5%) of infected neurons with detectable IFNα/β expression in mice infected with lymphocytic choriomeningitis virus (Delhaye et al., 2006). Furthermore, IFNα/β inducible *Irf7* transcripts mapped closely to sites of viral RNA (Ousman et al., 2005), supporting highly focal IFNα/β responses. Overall however, little is known about responsiveness of distinct CNS cell types to viral infections *in vivo*. Specifically oligodendroglia appears to express a limited TLR repertoire at basal levels compared to microglia (Hanke and Kielian, 2011,
Okun et al., 2009, Paul et al., 2007). They also do not appear to contribute to active proinflammatory cytokine secretion (Cannella and Raine, 2004).

Infection with the neurotropic mouse hepatitis virus (MHV) strain JHM (JHMV), belonging to the positive strand RNA coronavirus (CoV) family, was used to better characterize the ability of oligodendroglia to mount innate antiviral immune responses in vivo based on its prominent infection and persistence in oligodendroglia (Bergmann et al., 2006). In adult mice JHMV infection causes an acute encephalomyelitis, which resolves into a persistent infection of the spinal cord associated with demyelination. In addition to oligodendroglia, this virus also infects microglia and infiltrating macrophages during acute infection; however, neuronal and astrocyte infection is sparse (Ireland et al., 2008). CoV members are poor inducers of IFNα/β in numerous cell types due to their 5’RNA structures mimicking self RNA (Daffis et al., 2010, Zust et al., 2011). However, IFNα/β is induced in microglia, macrophages, and plasmacytoid dendritic cells (pDC) (Cervantes-Barragan et al., 2007, Roth-Cross et al., 2008). Importantly, a protective effect of IFNα/β in vivo was highlighted by uncontrolled MHV replication in mice deficient in IFNα/β signaling (Cervantes-Barragan et al., 2007, Ireland et al., 2008). Furthermore, protection following peripheral MHV infection is associated with TLR7 dependent IFNα/β production by pDC (Cervantes-Barragan et al., 2007). The naïve CNS is devoid of pDC (Serafini et al., 2000), implicating infected glia as primary candidates contributing to protective IFNα/β production following CNS infection. This is supported by MDA5 mediated IFNα/β production in infected microglia (Roth-Cross et al., 2008).
In the present study we compared the relative IFNα/β responsiveness of oligodendroglia and microglia following infection with JHMV or intracerebral inoculation of the double stranded (ds)RNA mimic poly (I:C). Both glia populations were isolated directly from the CNS to monitor changes in mRNA expression. The results are the first to imply an inherent block in IFNα/β transcription by oligodendroglia, coincided with limited expression of PRR and signaling molecules downstream of PRR activation. The dependence of oligodendroglia on IFNα/β from other cellular sources to induce an antiviral state in vivo highlight the potent monitoring functions of microglia and suggest a mechanism to preserve oligodendroglial homeostatic functions.

2.3 Results

2.3.1 Limited expression of PRR and signal transduction molecules in oligodendroglia

Induction of IFNα/β by murine CoV is mediated via TLR7 in pDC and via MDA5 in microglia/monocytes (Cervantes-Barragan et al., 2007, Roth-Cross et al., 2008). Furthermore, both MDA5 and RIG-1 act as PRR in an oligodendroglia cell line (Li et al., 2010). However, the expression of these viral sensors in oligodendroglia in the adult CNS is unknown. To analyze their capacity to initiate innate responses, oligodendroglia and microglia were isolated from spinal cords of uninfected PLP-GFP mice by FACS based on their respective CD45− GFP+ and CD45lo/int F4/80+ phenotypes (Malone et al., 2008). Spinal cords were used as donor tissue based on higher oligodendroglia yields compared to brains. The potential of mature oligodendroglia to recognize viral RNA in comparison to
microglia was assessed by comparing basal transcript levels of genes encoding the viral RNA sensors MDA5, RIG-I, TLR3, and TLR7 (Kawai and Akira, 2007). Whereas transcription of all four PRR was readily detected in microglia, *Mda5*, *Rig-I* and *Tlr3* transcripts were significantly reduced and *Tlr7* was below detection in oligodendroglia (Figure 3A). Basal levels of PRR mRNAs are low in the resting CNS relative to lymphoid tissue, with the exception of *Tlr3* (McKimmie et al., 2005). Nevertheless, PRR are rapidly upregulated during viral encephalitis initiated by Semliki Forest virus, Rabies virus, and Venezuelan Equine Encephalitis virus (McKimmie et al., 2005, Sharma and Maheshwari, 2009). However, the relative contribution of infiltrating leukocytes versus resident CNS cells was not directly assessed. Microglia and oligodendroglia were therefore isolated from JHMV infected mice to evaluate upregulation of PRRs during the course of infection. Microglia upregulated *Mda5*, *Rig-I*, *Tlr3* and *Tlr7* transcripts 2-5 fold over basal levels, with peak expression at days 3-5 post infection (p.i.) (Figure 3B). Relative to the modest fold increase in microglia, *Mda5* and *Rig-I* transcripts were vastly increased in oligodendroglia. *Rig-I* expression reached levels similar to microglia by day 3 p.i. mounting to an ~70 fold increase over naïve levels (Figure 3). Nevertheless, *Mda5* mRNA levels in oligodendroglia remained overall lower compared to microglia. Similarly, mRNAs encoding the endosomal PRR TLR3 and TLR7 were also induced in oligodendroglia; however, absolute levels remained low and the peak of expression was delayed relative to microglia. *Mda5* and *Rig-I* induction in oligodendroglia, as well as all RNA sensing PRR in microglia, correlated with induction of *Ifna* and *Ifnb* mRNA
throughout days 3-5 p.i. in total CNS RNA following JHMV infection (Ireland et al., 2008, Malone et al., 2008).

Figure 3. Oligodendroglia are limited in basal and inducible expression of viral RNA sensing PRRs. Microglia and oligodendroglia purified from spinal cords of naïve (A) or JHMV infected (B) PLP-GFP mice were assayed for Mda5, Rig-I, Tlr3, and Tlr7 transcripts at the indicated times p.i. Data represent the average of 3 separate experiments ± standard error of the mean (SEM). * denotes $p \leq 0.05$ comparing microglia to oligodendroglia at each timepoint; # indicates $p \leq 0.05$ compared to basal levels for each cell population.

To verify that the increase in PRR transcripts is directly attributed to IFNα/β signaling, mRNA expression was assessed in microglia and oligodendroglia from naïve and JHMV infected wt and IFNAR$^{−/−}$ mice. Although Mda5 and Rig-I mRNA levels were diminished by 8-9 fold in microglia from naïve
IFNAR<sup>-/-</sup> relative to wt mice, they were only slightly reduced in oligodendroglia (Figure 4). These results are reminiscent of previous observations that PRR expression in neurons is controlled by basal IFNα/β signaling (Shrestha et al., 2003). Importantly, the absence of IFNα/β signaling abrogated upregulation of Mda5 and Rig-I transcripts in both microglia and oligodendroglia following JHMV infection (Figure 4). Early induction of Mda5 and Rig-I, both classified as ISG, thus demonstrated that oligodendroglia respond to exogenous IFNα/β <i>in vivo</i>, similar to microglia. IFNα/β thus enhances the potential to recognize invading RNA viruses, thereby facilitating early innate responses.

Figure 4. MDA5 and RIG-I upregulation is IFNα/β dependent. Wt and IFNAR<sup>-/-</sup> mice infected with JHMV were used to compare induction of Mda5 and Rig-I mRNA in oligodendroglia and microglia purified from spinal cords. Data are average of 3 individual experiments ± SEM as described in Fig. 3. * denotes p<0.05 comparing wt to IFNAR<sup>-/-</sup> populations at each timepoint.
2.3.2 IFNα/β is not induced in oligodendroglia

IFNα/β signaling is crucial to prevent dissemination of gliatropic CoV within the CNS parenchyma (Ireland et al., 2008). Furthermore, IFNα/β induction in response to CoV infection has been demonstrated in microglia in vitro and in vivo (Roth-Cross et al., 2008) as well as in an oligodendrogial cell line (Li et al., 2010). Given the distinct basal and inducible levels of Mda5 transcripts in mature CNS derived glial populations, the contribution of infected oligodendroglia and microglia to IFNα/β induction was assessed in vivo by comparing Ifnβ and Ifnα4 mRNA relative to viral replication in each population. Viral loads were monitored by measuring viral mRNA encoding the N protein, the most abundant viral gene in infected cells (Skinner and Siddell, 1983). JHMV-N transcripts prevailed in microglia at day 3 p.i., but decreased by days 5 and 7 p.i. (Figure 5A). By contrast, although viral mRNA was near detection levels at day 3 p.i. in oligodendroglia, the levels increased ~20 fold relative to microglia by day 5 p.i. Despite subsiding by day 7 p.i., viral mRNA levels remained significantly elevated relative to microglia. These results not only assured that infected cells were recovered by the isolation procedure, but clearly indicated that virus predominated in microglia early, but subsequently prevailed in oligodendroglia. Concomitant with infection, sparse but detectable basal levels of Ifnβ and Ifnα4 transcripts increased in microglia (Figure 5 B). Ifnβ levels were sustained through day 5 p.i., while Ifnα4 levels were only transiently upregulated. The contribution of infiltrating infected monocyte derived macrophages (Ireland et al., 2009) to Ifnα/β induction was also assessed. Viral RNA loads in macrophages
were less than 50% of those in microglia and declined with similar kinetics; *Ifna/β* transcripts reflected the magnitude of viral RNA transcripts similar to microglia (data not shown). By contrast, neither *Ifnα4* nor *Ifnβ* mRNA were detected in oligodendroglia from naïve or infected mice (Figure 5B), despite higher viral RNA loads as well as elevated *Mda5* and *Rig-I* mRNA expression at day 5 p.i. (Figure 4).

The failure of oligodendroglia to induce *Ifna/β* following infection suggested that MDA5 and RIG-I are not activated, and/or that downstream signaling factors are limiting. PRR signaling requires several mediators including the inducible kinase IKKε (also known as Ikbke or Ikki) and the transcription factors IRF3 and IRF7. While IRF3 is constitutively expressed IRF7 is induced by IFNα/β and amplifies IFNα/β production (Honda and Taniguchi, 2006, Honda et al., 2005). Therefore to investigate the potential for PRR signaling,
Oligodendroglia were examined for expression of mRNAs encoding IKKε, IRF3 and IRF7. All three mRNAs were regulated in a cell type specific manner (Figure 6). Microglia from naïve mice expressed higher basal Ikkε transcripts, which were modestly, but progressively upregulated throughout day 7 p.i. (Figure 6). By contrast Ikkε induction was not detected in oligodendroglia until day 7 p.i., at which time levels reached those in microglia. Irf3 mRNA was readily detected at basal levels in both microglia and oligodendroglia from naïve mice, but remained constant or even declined following infection (Figure 6). Lastly, Irf7 transcripts were sparse in both microglia and oligodendroglia derived from naïve mice (Figure 6), consistent with lower basal CNS expression relative to Irf3 (Ousman et al., 2005). However, in contrast to the induction of Irf7 mRNA in microglia by day 3 p.i., induction was only modest and significantly delayed in oligodendroglia following infection (Figure 6). As the Irf7 gene is an ISG, its upregulation in microglia was consistent with IFNα/β upregulation. By contrast, the delayed response in oligodendroglia implied more selective ISG induction and/or distinct IFNα/β receptor signaling thresholds.

Figure 6. Oligodendroglia are limited in basal and inducible expression of PRR associated signaling factors during JHMV infection. Microglia and oligodendroglia from naïve or JHMV infected mice were assayed for Ikkε, Irf3 and Irf7 transcripts as indicated in Fig. 3. Data represent the average of 3 separate experiments ± SEM. * denotes p≤0.05 comparing microglia to oligodendroglia at each timepoint; # indicates p≤0.05 compared to basal levels for each cell population.
2.3.3 Oligodendroglia induce ISG delayed relative to microglia

Upregulation of the ISGs Rig-I, Mda5, Tlr3 and Irf7 support the notion that oligodendroglia respond to IFNα/β following infection. However, optimal induction was delayed and absolute levels were not as robust as in microglia. These results suggested that oligodendroglia are exposed to limiting amounts of IFNα/β in the microenvironment, or that they are intrinsically more limited than microglia in IFNα/β responsiveness. While the first possibility is difficult to address in vivo, the second may reflect reduced expression of the IFNα/β receptor (IFNAR) or downstream signaling components. We therefore assessed regulation of transcripts encoding the IFNAR, as well as STAT1 and IRF9, which act downstream of IFNα/β signaling. IFNα/β binds a common receptor complex consisting of two transmembrane proteins, IFNAR1 and IFNAR2. As IFNAR2 exists as a transmembrane and soluble isoform generated by alternative splicing (Domanski et al., 1995), we then examined basal expression patterns of IFNAR1, and both IFNAR2 isoforms in microglia and oligodendroglia. Basal mRNA levels of the Ifnar1 chain were ~2-fold higher in microglia compared to oligodendroglia. However, transcripts encoding both the soluble as well as the transmembrane form of Ifnar2 were more than 5-fold higher in microglia compared to oligodendroglia (Figure 7A), suggesting more limited responsiveness of oligodendroglia to IFNα/β. Stat1 and Irf9 transcript levels were also elevated in microglia compared to oligodendroglia derived from naïve mice (Figure 7B). STAT1 is inducible by both IFNα/β and IFNγ, while IRF9 is constitutively
expressed in most non CNS cell types (Borden et al., 2007). Consistent with a
response to IFNα/β Stat1 mRNA indeed increased ~10-fold by day 3 p.i. and
remained elevated until day 7 p.i. in microglia (Figure 7B). By contrast, a
reduced capacity of oligodendroglia to respond to IFNα/β was supported by the
modest upregulation of Stat1 transcripts at days 3 and 5 p.i. The prominent
increase of Stat1 transcripts prominently at day 7 p.i. suggested IFNγ mediated
upregulation. Irf9 transcripts were only upregulated 2-fold in microglia by 3 p.i.
and subsequently declined to or below basal levels (Figure 7B). Nevertheless,
despite lower basal levels and apparent constitutive expression, Irf9 transcripts in
oligodendroglia rapidly increased by ~15-fold at day 3 p.i., but did not increase
further at day 7 p.i. Although activation and nuclear localization of STAT1 and
IRF9 remains to be evaluated, the overall limited basal levels of Ifnar1 and Ifnar2,
as well as basal and inducible levels of Stat1 and Irf9 transcripts in
oligodendroglia during JHMV infection, support an overall paucity in ISG
transcriptional activation.

Figure 7. Oligodendroglia have limited IFNα/β receptor signaling capacity. Microglia and
oligodendroglia from naïve or JHMV infected mice were assayed for transcripts encoding the IFNAR1,
soluble IFNAR2 (IFNAR2s) and IFNAR2 transmembrane (IFNAR2TM) chains (A) or Stat1 and Irf9 transcripts
(B). Data in (A) represent the average of 2 experiments; data in (B) represent the average of 3 independent
experiments ± SEM as indicated in Fig. 3. * denotes p≤0.05 comparing microglia to oligodendroglia at each
timepoint; # indicates p≤0.05 compared to basal levels for each cell population.
The notion of limited ISG transcriptional activation in oligodendroglia was further supported by analysis of a subset of ISG encoding factors with anti-viral activity. Basal levels of transcripts encoding protein kinase R (Pkr), and the OAS/RNaseL pathway (Oas2), known to modulate translational activity, were all elevated in microglia derived from naïve mice relative to oligodendroglia (Figure 8). Transcripts encoding adenosine deaminase specific for RNA (Adar1), which limits viral replication via introduction of RNA mutations (Bass, 1997), were also more abundant in microglia (Figure 8). By contrast, Ifit2 RNA was expressed at similar levels in both cell types derived from naïve mice. Importantly, in microglia all transcripts, particularly Oas2 and Ifit2 RNA, were rapidly increased to peak levels by day 3 p.i. and declined thereafter. With the exception of Oas2, induction was also evident at day 3 p.i. in oligodendroglia; however, increases were most prominent by day 7 p.i., yet never reached the peak absolute values observed in microglia (Figure 8). These data demonstrate that oligodendroglia not only express lower steady state levels of ISGs, but also upregulate transcription with delayed kinetics relative to microglia in response to CoV infection. Furthermore, delayed ISG expression in oligodendroglia supported a more prominent responsiveness to IFNγ than IFNα/β. Overall the data clearly suggest that oligodendroglia require paracrine IFNα/β signals to induce ISG during gliatropic CoV infection.
Figure 8. Delayed induction of anti-viral ISG in oligodendroglia. Microglia and oligodendroglia from naïve or JHMV infected mice were assayed for Pkr, Oas2, Adar1, and Ifit2 transcripts as indicated in Fig. 3. Data represent the average of 3 separate experiments ± SEM. * denotes p ≤ 0.05 comparing microglia to oligodendroglia at each timepoint; # indicates p ≤ 0.05 compared to basal levels for each cell population.

2.3.4 Inability of oligodendroglia to induce Ifna/β genes is virus independent

To test whether oligodendroglia are intrinsically limited in IFNα/β production, or if limited IFNα/β production is specific to JHMV infection, we assessed IFNα/β and ISG induction following intracranial injection of the synthetic PRR ligand Poly (I:C). Poly (I:C) is a strong agonist of both TLR3 and the RIG-like receptors RIG-I/MDA5 (Alexopoulou et al., 2001, Kato et al., 2006). IFNα/β upregulation was initially tested in total tissue to characterize the time
course of Poly (I:C) induced IFNα/β responses in the CNS. *Iifen*β* expression was maximal at 2 hours and *Ifna*4 mRNA was elevated between 2 and 12 hours post poly (I:C) inoculation (Figure 9). By contrast, peak induction of the ISGs *Ifit*1 and *Ifit*2 was delayed until 12 hours (Figure 9). Cell type specific gene analysis of IFNα/β and ISG after intracranial poly (I:C) injection was thus performed in purified microglia and oligodendroglia at 4 hours as an intermediate timepoint for optimal *Ifna*/β expression and at 12 hours.

![Figure 9.Kinetics of CNS IFNα/β responses to Poly (I:C).](image)

Wt mice were injected intracranially with 200ug poly (I:C) to assess kinetics of *Iifen*β, *Ifna*4, *Ifit*1, and *Ifit*2 mRNA induction in spinal cords. Data represent the mean ± SEM for 3 mice per group; # indicates *p* ≤ 0.05 compared to basal levels.

In microglia *Ifna*4 and *Iifen*β transcripts were clearly induced by 4 hours and expression of *Iifen*β was sustained until 12 hours (Figure 10). Importantly, the levels were 30-40 fold higher than those induced by viral infection (see Figure 5), confirming the significantly increased magnitude of the response. Nevertheless, *Ifna*/β mRNA remained near detection thresholds in oligodendroglia (Figure 10). However, *Mda5* and *Rig-I* levels were increased in both cell populations after poly (I:C) injection relative to the modest 4-5 fold increases induced by virus.
infection (Figure 10). Furthermore, similar to the trend observed during infection, Rig-I transcripts were significantly increased over Mda5 transcripts in oligodendroglia. Nevertheless, Ikkε and Irf7 transcripts were elevated in both microglia and oligodendroglia at 4 and 12 hours after poly (I:C) (Figure 10) relative to viral injection (Figure 6). While limiting IFNα/β levels thus clearly contributed to suboptimal ISG induction in both glial populations following infection, the mechanism(s) underlying the paucity of IFNα/β induction in oligodendroglia remains elusive. Representative of antiviral ISG expression, Ifit1 and Ifit2 mRNA were also significantly increased in microglia at 4 hours, but subsided by 12 hours post poly (I:C) administration. By contrast, in oligodendroglia Ifit1 and Ifit2 levels induced at 4 hours were maintained at similar (Ifit1) or even increased levels (Ifit2) at 12 hours (Figure 10). Even under these conditions of optimal in vivo stimulation, oligodendroglia exhibited tentative and delayed ISG induction. These results support the notion that in vivo oligodendroglia are intrinsically impaired in inducing IFNα/β in response to dsRNA stimuli but also in responsiveness to IFNα/β suggesting an overall dampened capacity for innate responses.
Figure 10. Distinct in vivo responses of oligodendroglia and microglia to poly (I:C). Microglia and oligodendroglia purified from spinal cords at 4 or 12 hours post poly (I:C) injection were analyzed for expression of Ifna4, Ifnβ, Mda5, Rig-I, Ikkε, Irf7, Ifit1, and Ifit2 transcripts. Data are representative of two independent experiments each with 10 pooled mice per experiment.

2.4 Discussion

PRR initiated innate responses in the CNS are critical for the induction of IFNα/β, as well proinflammatory cytokines and chemokines. While PRR expression and activation in microglia and astrocytes has been extensively explored in vitro (Butchi et al., 2010, Carpentier et al., 2008, Jin et al., 2011, So and Kim, 2009), PRR expression and associated signaling components in glia in vivo are poorly defined, especially in oligodendroglia. In this study we compared the capacity of oligodendroglia relative to microglia to mount IFNα/β mediated
innate immune responses. Glial specific responses were monitored within the context of the CNS following intracerebral infection of adult mice with gliatropic JHMV or injection of poly (I:C). Microglia were prominent inducers of *Ifna/β* mRNA following infection, consistent with IFNα/β induction in primary microglia as well as bone marrow derived macrophages infected with a heterologous MHV (Roth-Cross et al., 2008). However, the inability of oligodendroglia to induce *Ifna/β* mRNA, despite their high load of viral RNA, contrasted with *in vitro* studies demonstrating MDA5 and RIG-I dependent IFNα/β induction in the infected N20.1 oligodendroglia cell line (Li et al., 2010). This discrepancy likely reflects differences in expression and/or induction of IFNα/β pathway components, rather than distinct viral loads, as even poly (I:C) did not induce *Ifna/β* in oligodendroglia *in vivo*.

The restricted ability of CoV to induce IFNα/β was recently attributed to the O-methylated cap structure of the viral RNA, making it difficult for the host to distinguish viral from self mRNA (Daffis et al., 2010, Zust et al., 2011). However, in contrast to cultured fibroblasts, bone marrow derived DC, astrocytes and neurons the ability of pDC, macrophages/microglia, and an oligodendroglia cell line to induce IFNα/β via TLR7, MDA5 and MDA5/RIG-I dependent pathways (Cervantes-Barragan et al., 2007, Li et al., 2010, Roth-Cross et al., 2008, Zhou and Perlman, 2007) suggests sufficient PRR activation by viral RNA structures to mediate protective responses in select cell types. Our data indicate that cell type specific basal expression, as well as inducible components in the innate signaling pathway constitute critical signaling thresholds determining IFNα/β induction. In
the adult CNS oligodendroglia differed significantly from microglia in reduced basal transcript levels of viral RNA sensing PRRs and Ikkε. However, basal Irf3 transcripts were only reduced by ~50%, and Irf7 transcripts were barely detectable in either cell population, consistent with higher basal Irf3 than Irf7 levels observed in whole naïve brains (Ousman et al., 2005). Reduced basal PRR levels in oligodendroglia were reminiscent of sparsely expressed PRRs in primary neurons (Carpentier et al., 2008), and supported the superior initiation of Ifnα/β responses of microglia. This concept is consistent with the inability of primary cultured neurons and astrocytes to induce IFNα/β expression following MHV infection (Roth-Cross et al., 2008).

Optimal IFNα/β induction requires amplification through IFNAR to elevate PRRs and IRF7 (Honda and Taniguchi, 2006, Honda et al., 2005, Malmgaard et al., 2002). However, in contrast to Ifnα/β induction, IFNAR signaling was intact in oligodendroglia. Although IFNα/β protein was below detection in cell free supernatants derived from the JHMV infected CNS by ELISA (data not shown), IFNα/β sufficed to up regulate the ISGs Mda5, Rig-I, Stat1, Irf7, Pkr, Ifit2, and Adar1 in oligodendroglia. The ability of IFNα/β to induce ISGs was consistent with basal expression of IFNAR chains, as well as basal and inducible levels of Stat1 and Irf9 transcripts. Although induction above basal levels was greater in oligodendroglia than microglia for some genes, peak absolute mRNA levels were in general lower compared to those reached in microglia at days 3 and 5 p.i., as exemplified by Stat1 and Irf7 mRNA. Furthermore, peak expression for many genes did not correlate with maximal Ifnα/β mRNA levels, but rather with peak
IFNγ secretion at day 7 p.i. (Parra et al., 2010, Phares et al., 2011, Puntambekar et al., 2011). Distinct pattern of ISG expression in oligodendroglia compared to microglia may further reside in low Ikke mRNA levels. In addition to mediating PRR signaling, IKKε influences ISG induction through participation in JAK/STAT mediated IFNα/β signaling (Pham and Tenoever, 2010, Tenoever et al., 2007). Although not as severe as the complete abrogation of ISG induction in STAT1−/− mice, IKKε−/− mice lack induction of ~30% of ISGs, including Adar1 (Durbin et al., 1996, Tenoever et al., 2007). The biological consequences are demonstrated by impaired clearance of influenza virus infection by IKKε−/− mice, despite similar expression of IL2, IL6, IL12, IFNγ and RANTES and induction of anti-viral antibody (Tenoever et al., 2007). The paucity of Ikke mRNA and modest, if any, increase of Irf7 mRNA in oligodendroglia, despite abundant viral and Mda5 mRNA upregulation, may partially explain the absence of Ifna/β expression and more selective and delayed ISG induction during infection. However, even the strong IFNα/β response elicited in the CNS by poly (I:C) did not overcome the inability of oligodendroglia to induce Ifna/β, despite over 400 fold increases in Ikke and 85-400 fold increases in Irf7 transcripts relative to those induced by infection. These results suggest additional restrictions intrinsic to oligodendroglia in the initiation of IFNα/β production.

The results thus indicate a reliance of oligodendroglia on external production of IFNα/β to induce an antiviral state. This concept is consistent with data indicating that microglia are a dominant source of FNa/β within the CNS during JHMV induced encephalomyelitis (Roth-Cross et al., 2008). The biological
relevance of suboptimal IFNα/β responses is clearly evident from differential viral control in both cell types. A direct response to infection by microglia is supported by the correlation between peak viral and Ifnα/β mRNA. Furthermore, the early decline in viral RNA, prior to infiltration of T cells, indicates autocrine IFNα/β contributes to the rapid control viral spread in this cell type. By contrast, although oligodendroglia were initially infected to a lesser extent than microglia, viral RNA increased over 2 logs between days 3 and 5 p.i., coincident with the absence of Ifnα/β and tentative induction of antiviral ISGs. These data indicate that oligodendroglia thus rely largely on T cell effector function, prominently IFNγ, for viral control (Parra et al., 2010). Oligodendroglia are indeed highly responsive to IFNγ as indicated by upregulation of class I MHC molecules and associated antigen processing components during peak IFNγ, but not IFNα/β induction during JHMV infection (Malone et al., 2008). Moreover, specific blockade of IFNγ receptor signaling on oligodendroglia prolongs JHMV infection in this cell type (Gonzalez et al., 2005, Parra et al., 2010). Whether early events favoring robust infection of oligodendroglia prior to emergence of T cells, contribute to ultimate persistence of JHMV in oligodendroglia, remains to be elucidated.

Overall our results demonstrate a limited role of oligodendroglia as both inducers of, and responders to, IFNα/β compared to microglia. Limited innate antiviral activity may predispose oligodendroglia to be potent responders to IFNγ (Gonzalez et al., 2005, Popko and Baerwald, 1999). While there is no evidence supporting toxicity of IFNα/β (Akwa et al., 1998), more restricted induction of antiviral mediators, especially those also affecting host cell translation, may
circumvent apoptosis and guarantee maintenance of critical myelin housekeeping functions and survival. Our findings also contradict the hypothesis suggesting that virus initiated IFNα/β production by oligodendroglia drives inflammatory responses during viral induced demyelinating disease (Lipton et al., 2007). The low abundance of PRR at basal levels rather provide a mechanism underlying the apparent paucity of oligodendroglia to express cytokines during multiple sclerosis or other CNS inflammatory conditions (Cannella and Raine, 2004, Zeis et al., 2008). Our data thus support the notion that oligodendroglia are defensive players under inflammatory conditions.

Figure 11. Model for differential innate glia responses to JHMV infection
CHAPTER III
PKR: UNCONVENTIONAL ROLE OF AN ANTI-VIRAL FACTOR FOLLOWING CNS INFECTION

3.1 Abstract

IFNα/β plays an eminent role in restricting viral tropism and limiting dissemination within the CNS following infection with the glia tropic coronavirus JHMV. However, which among the many IFNα/β stimulated innate antiviral factors are protective in the CNS is largely unknown. RNaseL and Protein Kinase RNA dependent (PKR) are the best-characterized innate anti-viral proteins. Although the absence of RNaseL resulted in elevated infection of microglia/macrophages, it was insufficient to alter overall JHMV control or CNS inflammation. Nevertheless, RNaseL deficiency resulted in mortality coincided with accelerated demyelination and axonal degeneration, suggesting RNaseL activation provides protective effects independent of viral control. To identify the role of other innate antiviral pathways in JHMV pathogenesis, we explored a potential contribution of PKR, an inhibitor of host cell translation and modulator of NFkB mediated gene transcription using PKR deficient mice (PKR-/-). Infected
PKR-/− mice exhibited no defect in expression of *Ifnα/β* or induced anti-viral genes *Ifit1* and *Ifit2* suggesting no over deficits in the IFNα/β pathway. Surprisingly, however, PKR deficiency significantly reduced expression of NFκB mediated pro-inflammatory cytokines and chemokines. Viral burden in the CNS was initially increased by ~10-fold in PKR-/− mice relative to wt mice, but histological examination revealed no discernable differences in viral antigen distribution or tropism. Furthermore, virus was subsequently controlled with similar kinetics indicating effective CD8 T cell recruitment and function, although PKR deficiency was associated with more extensive accumulation of infiltrating cells in the perivascular space, Lower *Il21* and *Il10* expression further suggested modulation of CD4 T cell activity in the absence of PKR. Lastly, the onset of axonal damage appeared accelerated and demyelination more diffuse in PKR-/− mice. Nevertheless, the overall pathology resembled that of wt rather than RNaseL-/− mice and the absence of PKR did not result in high mortality. A subtle effect of PKR on JHMV pathogenesis was supported by a similar phenotype of mice dually deficient in RNaseL and PKR as in single RNaseL-/− mice. In summary the results suggest that RNaseL and PKR only exert subtle direct antiviral activity during JHMV infection. Moreover, while PKR activation can profoundly influence the cytokine/chemokine milieu without overt consequences on disease, RNaseL activation mediates protective function by mitigating demyelination.
3.2 Introduction

Type-1 IFN signaling initiates transcription of numerous genes resulting in an antiviral state characterized by elevated expression of numerous anti-viral factors (Borden et al., 2007, Trinchieri, 2010). Many of the antiviral molecules directly block viral replication by interfering with translational activity, either by degrading viral and host RNA or directly inhibiting protein synthesis. Nevertheless, effective antiviral functions differ not only between viruses themselves, but also the cell types infected. Moreover, although distinct anti viral factors all minimize viral spread, altered cellular transcriptional and translational profiles can also significantly impact host cell function, for example resulting in apoptosis. The most characterized anti-viral mechanisms are the 2'-5' oligoadenylate synthetase (OAS)/RNaseL and protein kinase double stranded RNA (PKR) pathways (Silverman, 2007b, Silverman, 2007a, Garcia et al., 2007). Both depend on activation of latent enzymes to be functional, which serves to minimize altered cellular functions in non-infected, bystander cells. Cell type dependent antiviral function is illustrated by RNaseL dependent control of West Nile virus (WNV) in MEF, but RNaseL independent IFNβ mediated virus clearance in either peripheral or cortical neurons. Furthermore, increased mortality of WNV infected dual RNaseL/PKR−/− mice is attributed to increased peripheral infection, but not enhanced virus dissemination to the CNS (Samuel et al., 2006). Virus dependent protection is highlighted by impaired control of Sindbis virus in the absence of RNaseL and PKR in mouse embryonic fibroblasts (MEF) and bone marrow derived macrophages (Ryman et al., 2002), but IFNβ
mediated inhibition of dengue virus is independent of the PKR/RNaseL pathways (Diamond and Harris, 2001). PKR play a crucial role in antiviral defense following vesicular stomatitis virus infection (VSV) in vitro. Furthermore, 100% mortality and uncontrolled virus replication was observed in PKR -/- mice following intranasal VSV infection (Stojdl et al., 2000).

IFNα/β mediated innate responses are also critical for limiting tropism, controlling virus spread and preventing mortality following infection with the glia tropic, murine coronavirus, JHMV (Ireland et al., 2008). However, the type-1 IFN downstream anti-viral factors mediating protection are largely unknown. RNaseL plays a modest role in reducing infection of macrophage/microglia, but does not alter overall virus clearance in the CNS (Ireland et al., 2009). Surprisingly however, RNaseL mediated protection from accelerated axonal damage, demyelination and death. The pathological differences, independent of viral control, suggested dysregulation of as yet unknown RNaseL functions in the CNS during inflammatory conditions.

Based on uncontrolled JHMV spread in the CNS in absence of IFNα/β signaling, but limited effects of RNaseL, we further explored other antiviral mediators by determining the role of PKR. PKR is an intracellular sensor of stress. Viral dsRNA as well as other ligands such as PKR activating protein (PACT) and polyanionic molecules such as dextran sulfate, chondroitin sulfate, poly (L-glutamine) and heparin can directly activate PKR (Bergeron et al., 2000). In addition to viral RNA, some highly structured cellular RNA molecules, such as Tumor Necrosis factorα (TNFα) and Interferonγ (IFNγ) mRNA also has been
shown to activate PKR (Ben-Asouli et al., 2002, Osman et al., 1999). The ability of PKR to respond to multiple stimuli emphasizes its broader role as a signaling molecule against different types of physiological stress. Activated PKR depicts kinase activity and undergoes autophosphorylation. The main target of phosphorylation by activated PKR is the eukaryotic translation initiation factor, eIF2\(\alpha\). Phosphorylation of eIF2\(\alpha\) leads to inhibition of translation and induction of apoptosis in host cells, therefore inhibiting viral replication. PKR also induces transcriptional expression of pro-inflammatory genes and IFN\(\alpha/\beta\) by initiating the degradation of IkB, inducing IRF1 expression and indirectly mediating STAT1 phosphorylation. IkB degradation leads to activation of downstream transcription factor NFkB (Kumar et al., 1994, Bonnet et al., 2000). PKR mediated activation of NFkB is independent of its kinase activity. The N terminus (AA 1-265) of PKR, which includes the dsRNA binding domain (AA 1-181) and third basic domain (AA 181-265) has been shown to activate NFkB by directly binding and activating the IKK complex (Bonnet et al., 2006). PKR can also associate with TNF receptor associated factor (TRAF) to mediate activation of NFkB (Gil et al., 2004). Other than inducing pro-inflammatory cytokines, PKR is also demonstrated to be involved in inducing anti-inflammatory cytokine IL10 in monocyte/macrophages in response to polyI:C, LPS and distinct infections such as mycobacterium, and sendai virus (Cheung et al., 2005, Chakrabarti et al., 2008). Several studies also support a link between PKR and cellular immunity. For example, PKR limits HIV replication by reducing surface expression of CD4 T cells in vitro in jurkat cells (Nagai et al., 1997). PKR affects ability of CD8 T cells to control replication of
Lymphocytic Choriomeningitis Virus (LCMV) without altering CD8 T cell their effector function (Nakayama et al., 2010). Lastly, impaired proliferation of CD8 T cells in response to mitogen stimulation during systemic lupus erythematosus (SLE) is associated with increased PKR activation and therefore reduced protein synthesis (Grolleau et al., 2000).

PKR is also implicated to participate in neuro-degenerative processes. Following poliomyelitis infection PKR deficient mice show no difference in virus replication, but were protected from infection mediated extensive spinal cord damage (Scheuner et al., 2003). A separate study links PKR activation as an early biomarker of neuronal cell death in Alzheimer’s disease (Peel and Bredesen, 2003). Contribution of PKR in neurotoxicity was attributed to its ability to increase apoptosis by activating fas associated protein with a death domain (FADD) and subsequently activating caspase 3 and 8 in recombinant virus infected cells as well as in Aβ treated cells and in APPSLPS1 K1 mice (Transgenic mouse to study AD) (Gil and Esteban, 2000, Couturier et al., 2010). In the EAE autoimmune model of multiple sclerosis, PKR is mainly expressed by oligodendroglia, T cells and some macrophages (Chakrabarty et al., 2004). In summary, these findings implicate PKR can control numerous aspects of immunity ranging from direct antiviral activity via translational inhibition, to induction of pro- as well as anti-inflammatory molecules, and modulation of adaptive responses. Moreover, PKR is also demonstrated to contribute in neurotoxicity by inducing apoptotic pathway.
In the absence of PKR we observed higher virus titers early during infection at day 3, 5 and 7 p.i. following JHMV infection. However, virus in PKR-/− mice was subsequently controlled by day 10 p.i., with similar kinetics as in wt mice. This indicated an early antiviral effect of PKR, which was overcome by effective CD8 T cell function. We were therefore surprised that reduced expression of pro-inflammatory cytokines and chemokines such \textit{Il6}, \textit{Ccl5}, and \textit{Cxcl10} had no apparent effect on leukocyte recruitment. Nevertheless, a hallmark of infected PKR-/− mice was accumulation of cells in the perivascular space. Specifically the distribution of CD4 T cells was skewed towards retention in the perivascular space indicating impaired entry into the CNS parenchyma. Moreover, in the absence of PKR, CD4 T cells were impaired in IL10, but not \textit{IFNγ} production. While these results all indicated substantial immune modulation by PKR, ultimate virus clearance and disease outcome were surprisingly similar to wt mice.

3.3 Results

\textbf{3.3.1 PKR deficiency does not alter IFNα/β induction, reduces NFkB mediated gene expression and only affects virus replication modestly in the CNS following JHMV infection}

PKR is constitutively expressed at low levels and can signal to activate IFNα/β after binding to dsRNA. Activated PKR can also bind to and degrade the inhibitor of NFkB (IκB), resulting in NFkB activation (Kumar et al., 1994). Once activated NFkB translocates to the nucleus and regulates expression of pro-inflammatory genes. To initially identify the role of PKR in IFNα/β induction and
NFkB activation in the infected CNS, we determined the expression of IFNα/β and NFkB induced genes using quantitative real time PCR analysis (Figure 12). Induction of Ifnα4 and Ifnβ mRNA peaked in spinal cords at day 5 p.i. in both mice and declined thereafter. Although Ifnα4 mRNA levels were similar, Ifnβ mRNA was reduced at day 5 p.i. in PKR-/- mice. Nevertheless, no differences were observed in expression of the Ifit1 and Ifit2 genes, two ISG that are highly sensitive to IFNα/β induction. Both Ifit1 and Ifit2 transcripts were highly upregulated by day 5 p.i., remained elevated at day 7 p.i., and declined thereafter (Figure 12A). By contrast, expression of NFkB induced cytokines and chemokines was significantly impaired, albeit not universally (Figure 12B). While the proinflammatory genes Ccl2, Il1β, IL6, Tnfa, Ccl5 and Cxcl10 were all upregulated in spinal cords of both groups at day 5 p.i., Il6, Ccl5 and Cxcl10 were selectively reduced 2-fold in the absence of PKR. Ccl2 and Il1β mRNA levels both increased to similar relative levels at day 7 p.i., and subsequently declined, demonstrating no overt effect of PKR. Il6 mRNA remained at 50% the levels of wt mice and declined with similar kinetics at days 7 and 10 p.i. By contrast, Ccl5 and Cxcl10 chemokine mRNAs increased by day 7 in wt mice, but remained low throughout days 7 and 10, barely increasing over the day 5 levels in PKR-/- mice. By day 7 p.i. PKR-/- mice also exhibited decreased mRNA levels of Tnfa, which was sustained throughout day 10 p.i.
Figure 12. PKR deficiency resulted in reduced expression of NFκB induced pro-inflammatory genes, but does not alter IFNα/β dependent antiviral genes. Expression of NFκB and IFNα/β induced genes were determined using Real time PCR analysis. (A) Expression of Ifnα/β and dependent genes Ifit1 and Ifit2 from spinal cord of PKR-/- mice and wt mice. (B). Expression of selected NFκB dependent genes, Il1β, Il6, Tnfa, Ccl2, Ccl5, and Cxcl10 relative to Gapdh in spinal cords of PKR-/- and wt mice. Data is represented as average of n = 3 mice/ group/ time-point ± standard error mean (SEM).* p ≥ .05.

To determine an effect of PKR on virus replication viral loads in brains and spinal cords were determined by plaque assay and PCR, respectively. Infection is always initiated in the brain and rapidly spreads to spinal cords, where it
preferentially persists. In wt mice, infectious virus peaks at days 3-5 p.i., starts to be controlled at day 7 p.i., and is reduced to undetectable levels by plaque assay by day 14 p.i. We observed ∼5-10 fold higher virus titers early during the infection at days 3, 5 and 7 p.i. in PKR-/- mice compared to wt mice; however, at day 10 p.i. virus replication was controlled similarly as wt mice (Figure 13). Analysis of viral RNA in spinal cords revealed no evidence of an early antiviral effect of PKR. Analysis of viral antigen distribution by histology revealed no overt differences in either infected cell numbers or cellular distribution (data not shown). However it is plausible that infectious virus is higher in the absence of PKR if viral mRNA is translated more efficiently. These results indicates that PKR plays a modest antiviral role early during the infection, but virus can be cleared with similar efficiency as in wt mice once CD8 T cells infiltrate to the CNS (days 7-10 p.i.).

Figure 13. Virus replication in PKR-/- mice is enhanced during early phase of infection but clear with similar kinetics as wt mice. Virus replication in brain was determined by plaque assay on brain supernatants at indicated time-points post infection. Data is represented as average of n = 3 mice/ group/time-point ± standard error mean (SEM). (n = 9 mice/group/time-point). Viral RNA was determined in spinal cords relative to Gapdh. * p ≥ .05.
3.3.2 Increased CNS T cell infiltration, but accumulation of CD4 T cells in the perivascular space in the absence of PKR

Reduced levels of Il6 and T cell attracting chemokines Ccl5 and Cxcl10 suggested alteration in leukocyte infiltration into the CNS, yet efficient virus control contradicted this notion. Therefore, composition of infiltrating cells was determined in CNS of PKR -/- and wt mice using FACS analysis for the cell specific surface markers. We observed no difference in number of overall infiltrates (CD45^{hi} cells) to the CNS in the absence of PKR. Similarly, monocytes infiltrating the CNS, characterized by a CD45^{hi}/F4/80^{+} phenotype, were not statistically different between both mouse groups at any time p.i. consistent with similar monocyte recruiting Ccl2 expression. CXCL10 and CCL5 expression is reported to attract T cells to the CNS following JHMV infection (Glass et al., 2004, Liu et al., 2001). Therefore alterations in T cell accumulation were determined.

Both CD4 and CD8 T cells accumulate in the CNS to maximal levels between 7-10 days following JHMV infection and account for ~ 60% of total CD45^{hi} infiltrates. Compared to wt mice, CD8 T cells were significantly higher at day 7 p.i. and CD4 T cells at both day 7 and 10 p.i. within brain infiltrates in PKR-/- mice (Figure 14).

![Figure 14. Increased number of infiltrating T cells to the CNS in PKR-/- mice.](image)

Number of infiltrating, Total lymphocytes (CD45hi), macrophages (CD45hi F4/80+), CD8 T and CD4 T cells derived from the brain of infected wt and PKR-/- mice was determined using flow cytometric analysis. Data is average of three separate experiments ± standard error mean (SEM). (n = 9 mice/group/time-point). * p ≥ .05.
Potential differences in distribution of leukocytes within the CNS in both groups of mice were also analyzed by histochemical analysis of spinal cord sections. Hemotoxylin/eosin staining showed accumulation of infiltrating cells in the perivascular space at day 7 and day 10 p.i. (Figure 15A). To quantify the relative distribution of subtypes of infiltrating lymphocytes, laminin staining was used to visualize the endothelial cells and parenchymal basement membrane to define the outer and inner layers of perivascular space where leukocyte accumulate before trafficking to the parenchyma of the CNS (Savarin et al., 2010). Monocytes have been reported to enhance perivascular accumulation of lymphocytes (Savarin et al., 2010). However, we did not detect any difference in either recruitment of monocyte (Figure 14) or function of monocyte determined by phagocytic ability (data not shown). Furthermore, only a transient increase in CD8 T cell numbers was observed at day 7 p.i. and similar virus control at day 10 p.i. in PKR-/- mice compared to wt (Figure 14) suggested no defect in their functional ability. We specifically focused on CD4 T cells based on their increased numbers at both day 7 and 10 p.i. To distinguish CD4 T cells accumulated in perivascular space compared to total infiltrating CD4 T cells in the spinal cord, sections were stained for laminin and CD4 T cells. In wt mice, CD4 T cells preferentially accumulate in perivascular space at day 7 p.i., accounting to ~50% of total CD4 T cells in the perivascular space, eventually CD4 T cells migrate to the parenchyma at day 10 p.i. as observed only ~20% CD4 T cells retaining in the perivascular space while ~80% localize in parenchyma at this time p.i. (Figure 15B). In the absence of PKR, the frequency
Figure 15. Trafficking of lymphocytes from perivascular space to CNS parenchyma is influenced in PKR-/- mice compared to wt mice. (A) Cross-sections of spinal cords at day 10 p.i. were stained for Hematoxylin and Eosin (H & E). Enhanced perivascular inflammation is consistent with increased T cell accumulation by flow cytomteric analysis. Data is representative of three separate experiments. (B) Accumulation of CD4 T cells in the perivascular space was determined using staining spinal cord sections with CD4 and laminin in PKR-/- mice relative to wt mice. Staining of CD and laminin indicates the perivascular accumulation of CD4 T cells. Data is representative of n = 3 mice/group at day 10 p.i. Quantification of frequency perivascular CD4 T cells within total infiltrating Cd $ T cells is presented as graph form. Data is average of total 15 sections between n = 3 mice/group/time-point ± SEM. (C) Expression of activated PKR in infiltrating CD4 T cells to the CNS was determined by double staining spinal cord sections for PKR-p and CD4 in wt mice. Data is representative of n = 3 mice/group at day 7 p.i. * p ≥ .05.
of perivascular CD4 T cells was elevated to ~70% at day 7 p.i., which was
decreased to ~50% at day 10 p.i. but still remained significantly higher than wt
mice. (Figure 15B). Therefore first time we demonstrate that PKR can influence
CD4 T cells migration across the glia limitans following JHMV infection. To
identify direct role of PKR in regulation of CD4 T cell function, first we analyzed if
CD4 T cells express activated PKR (phosphorylated-PKR [PKR-p]) following
JHMV infection by utilizing fluorescent microscopy on spinal cord sections.
Positive staining for PKR-p in some CD4 positive T cells indicate that not all CD4
T cell can express activate PKR (PKR-p) under viral infection (Figure 15C).

3.3.3 Absence of PKR influences CD4 T cell function

PKR has been implicated in regulating IL10 production in monocyte and
CD8 T cell antiviral function (Nakayama et al., 2010, Cheung et al., 2005,
Chakrabarti et al., 2008), however its effect on CD4 T cell has not been explored.
Co-expression of activated PKR in CD4 T cells, and increased retention of CD4 T
cells in perivascular space in absence of PKR would indicate a defect in CD4 T
cell function in PKR/-/- mice. To investigate the effect of PKR on CD4 T cell
activity, expression of cytokines mainly produced by CD4 T cells during JHMV
infection was determined using real time PCR analysis. Following JHMV infection,
IFN\textsubscript{\gamma} is produced by both CD4 as well as CD8 T cells, while IL10 is produced at
high level by CD4 T cells and at lower extent by CD8 T cells (Puntambekar et al.,
2011) and IL21 is mainly secreted by CD4 T cells (Phares et al., 2011). The
absence of PKR did not affect production of IFN\textsubscript{\gamma} at either mRNA (Figure 16A) or
protein level in the CNS following virus infection. However, \textit{Il10} (~ 6-7 fold) as
well as *Il21* (∼ 2 fold) transcripts were reduced at day 7 and 10 p.i. in PKR-/- mice compared to wt mice (Figure 16A).

Figure 16. CD4 T cell function is specifically affected in PKR-/- mice relative to wt mice. (A) Effector function of CD4 T cells was assessed by measuring mRNA expression of cytokines mainly produced by CD4 T cells following JHMV infection (high IFNγ, high IL10 and IL21). Data is represented as average of n = 3 mice/group/time-point ± SEM. (B) Ability of CD8 and CD4 T cells to produce IFNγ and IL10 was determined using intracellular cytokine assay in splenocytes after stimulation with CD + and CD4 specific peptides, S510 and M133, respectively. Frequency of IL10 producing CD4 T cells is presented as representative flow graph and data is also quantified as average of n = 3/group ± SEM. Reduced effector functions associated primarily with CD4 T cells suggests reduced IL10 may contribute to enhanced pathology. * p ≥ .05.
Reduced expression of \textit{Il10} and \textit{Il21} in CNS following JHMV infection indicates a defect in CD4 T cell to produce these cytokines in the absence of PKR. To verify a role of PKR in modulating CD4 T cell function, CD4 T cells were activated by peripheral infection and analyzed for cytokine secretion. C57Bl/6 mice were intra peritoneal injected with DM strain of JHMV (plaque variant of JHMV), which produces excellent peripheral response (Stohlman et al., 2008). Splenocytes were isolated 7 days p.i. and stimulated with MHC class II M133 peptide for 5 days to bias composition of spleen cells towards CD4 T cells enriched phenotype. Following restimulation with M133 peptide for 6 hours and then analyzed for IL10 or IFN\textsubscript{\gamma} production by intracellular staining. We observed reduced ability of CD4 T cells to secrete IL10 in the absence of PKR, but no defect in their ability to produce IFN\textsubscript{\gamma} compared to wt mice (Figure 16B). PKR has been implicated in regulating IL10 expression in monocytes, however first time in this report we demonstrated PKR mediated regulation of IL10 production in CD4 T cells.

\subsection*{3.3.4 PKR deficiency results in more diffuse demyelination, but does not contribute to clinical disease or survival}

IL10 deficiency results in exacerbated demyelination following JHMV infection (Trandem et al., 2011). Due to the significant effects of PKR deficiency on IL10 transcription, spinal cords were assessed for myelin loss. Luxol fast blue (LFB) staining on cross sections of spinal cords from infected PKR-/- and wt mice revealed accelerated and more diffuse demyelination in absence of PKR. At day 7 p.i. demyelination was evident by very local vacuolization accompanied by
occasional swollen axons within the lesions in wt mice, but lesions were more pronounced in the absence of PKR, although axonal loss was not yet evident.

Demyelination in both groups was more extensive by day 10 p.i., but more widespread in PKR-/- mice (Figure 17). However, enhanced demyelination did not affect progression or severity of clinical disease or survival rate of PKR-/- compared to wt mice out to 21 days p.i. (Figure 18).

**Figure 17. Demyelination but not axonal damage increased in PKR-/- mice.** The extent of demyelination was determined by staining cross sections of spinal cord with Luxoal fast Blue (LFB) from wt and PKR-/- mice at day 10 p.i. Data is representative of three separate.

**Figure 18. Clinical disease and mortality is not altered in absence of PKR.** (Clinical disease and survival rate between wt and PKR-/- mice was determined following intracranial JHMV infection. Data is representative of three separate (n ≥ 15 mice/group/time-point ± SEM).
3.3.5 Dual deficient mice in PKR and RNaseL depicts overall similar phenotype as RNaseL-/- mice

Studies in mice deficient in either RNaseL or PKR indicated modest, but distinct roles in early antiviral function; nevertheless, RNaseL deficiency resulted in mortality, while PKR deficiency did not. To evaluate to what extent reduced elevated viral titers and reduced early proinflammatory responses by PKR deficiency may contribute to pathogenesis in RNaseL-/- mice, mice dually deficient in RNaseL and PKR (DRP-/-) were infected with JHMV. The additional absence of PKR did not influence mortality, severity or progression of clinical disease relative to RNaseL-/- mice (Figure 19A), and ∼80% succumbed to infection by day 10-12 p.i. We also observed higher viral burden similar to PKR-/- mice by ∼10-fold throughout acute infection, although it declined with kinetics similar to wt and RNaseL-/- mice (Figure 19B). Higher virus titer even at day 10 p.i. in DRP-/- mice (Figure 19B) compared to PKR-/- or wt mice (Figure 13) indicated an additive effect of the absence of both RNaseL and PKR. To assess a defect in T cells potentially contributing to elevated virus titers at day 10 p.i., CNS infiltrating cells were examined for number and composition using flow cytometry. However, no alterations in either total numbers or composition were noted comparing DRP-/- to wt mice (Figure 19C). To confirm a reduction in NFkB dependent pro- inflammatory cytokines/ chemokines (Figure 12B), mRNA expression of pro-inflammatory cytokines Il6, Tnfα and chemokines Ccl5 and Cxcl10 was examined in DRP-/- mice relative to wt mice.
Surprisingly, we did not observe any defect in expression of either cytokines or chemokines in DRP-/- mice compared to wt mice (Figure 20). These results indicate that absence of RNaseL rescued from impaired NFkB mediated gene expression observed in absence of PKR.
Figure 20. Dual deficiency of PKR and RNaseL reverses PKR mediated reduction in expression of NFκB induced pro-inflammatory genes. Expression of NFκB induced genes Il6, Tnfα, Ccl5, and Cxcl10 were determined relative to Gapdh in spinal cords of DRP-/- mice relative to wt mice, using real time PCR analysis. Data is represented as average of n = 3 mice/group/time-point ± SEM.

3.4 DISCUSSION

PKR and RNaseL are two well-characterized IFNα/β induced anti-viral molecules. The absence of RNaseL did not affect overall JHMV clearance from the CNS, although virus burden was elevated in CNS derived macrophage/microglia (Ireland et al., 2009). Nevertheless, RNaseL deficiency resulted in high mortality and accelerated demyelination and axonal damage. The limited antiviral effect of RNaseL suggested other anti-viral molecules play a more prominent role in preventing initial viral spread throughout the CNS following JHMV infection. PKR was chosen as a candidate based on its ability to block translation, but also to indirectly inducing IFNα/β production (O’Malley et al., 1986). A recent report further suggested that PKR can affect the ability of CD8 T cells to control virus replication, but without altering effector function in response to LCMV infection (Nakayama et al., 2010). Unlike the LCMV model, the absence of PKR did result in higher virus titers early during JHMV infection, but we could
not detect a cell type specific effect by histology. Furthermore, there were no
global defects in expression of IFNα/β or the ISGs, *Ifit1* and *Ifit2*. Similar
expansion of virus-specific CD8 T cells and kinetics of viral control also ruled out
alterations in the CD8 T cell response.

Reduced expression of cytokine/chemokine such as *Il6*, *Tnfα*, *Ccl5* and
*Cxcl10* in the absence of PKR is attributed to defective transcriptional activation
of NFκB mediated pro-inflammatory genes (Brain Williams, 1994, PNAS). All
these genes have been reported to play an important role in recruiting and
facilitating infiltration of leukocytes through the BBB into the CNS parenchyma.
Specifically expression of CXCL10 and CCL5 has been shown to facilitate T cell
recruitment, predominantly CD4 T cells to the CNS following JHMV infection
(Glass et al., 2004, Liu et al., 2001). It was therefore surprising that the lack of
PKR resulted in a transient increase in CNS infiltrating CD8 T cells at day 7 p.i.
and significantly higher number of CD4 T cells at day 7 and day 10 p.i.
Nevertheless, a large fraction of CD4 T cells accumulated within the perivascular
space, suggesting they were inhibited from entering the parenchyma, or were
continuously recruited from the periphery. Various studies have indicated the
important role of integrins as well as chemokines in migration of lymphocytes
across the blood vessels to the perivascular space (Ley et al., 2007). Reduced
monocyte infiltration also resulted in increased retention of T cells in the
perivascular space following JHMV infection (Savarin et al., 2010). However, we
did not observe any deficiency in the monocyte recruiting chemokine *Ccl2*,
consistent with similar monocyte infiltration in PKR-/- mice compared to wt mice.
The mechanism underlying enhanced perivascular T cell accumulation may reside in differential positioning of chemokines (McCandless et al., 2006, McCandless et al., 2008, Savarin et al., 2010), or overall reduced proinflammatory cytokine/chemokine expression.

Previous reports have implicated role of PKR in upregulation of the anti-inflammatory cytokine IL10 in a NFkB, ERK1/2 and JNK1 dependent manner as a response to mycobacterium, bacterial lipopolysaccharide, dsRNA and sendai virus infection (Chakrabarti et al., 2008, Cheung et al., 2005, McCandless et al., 2006, McCandless et al., 2008, Savarin et al., 2010). IL21 has been shown to induce IL10 in a mouse model of autoimmunity of systemic lupus erythematosus (SLE) (Spolski et al., 2009) (Roassne spoliski, 2009, Ji). In response to JHMV infection, both IL10 and IL21 are mainly produced by CD4 T cells (Phares et al., 2011, Puntambekar et al., 2011). Here we demonstrated for the first time, that PKR controls the expression of both IL10 and IL21 in CD4 T cells. The inability of CD4 T cells to produce IL10 following ex vivo stimulation with peptide (M133) demonstrated that the defect in IL10 was both at the transcriptional and protein level. However, IFNγ production by CD4 T cells was not affected and CD8 T cells were capable of effective viral clearance in PKR−/− mice indicating that PKR specifically affects the ability of CD4 T cells to produce IL21 and IL10.

PKR also exacerbates pathology in various neurological diseases. Strong induction of activated PKR (phosphorylated PKR) in hippocampal neurons has been associated with their loss in pathologies associated with Alzheimer’s disease, Huntington disease, and Parkinson’s disease (Bando et al., 2005, Peel...
and Bredesen, 2003). Similarly, the absence of PKR provides protection from extensive spinal cord damage during poliomyelitis infection (Scheuner et al., 2003). Contrary to these reports, we observed accelerated and more diffuse demyelination in absence of PKR following JHMV CNS infection. The enhanced pathology observed in PKR-/- mice may be attributed to reduced expression of the anti-inflammatory cytokine IL10 (Trandem et al., 2011). IL10 reduces the expression of pro-inflammatory cytokines such as IL6, TNFα and IFNγ expression in the CNS following JHMV infection (Lin et al., 1998). Lastly, although an imbalance between pro- versus anti-inflammatory responses in the CNS has been linked to increased severity of clinical symptoms, progression of clinical disease and mortality was not altered in PKR-/- compared to wt mice. These data supported the notion that IFNγ, which was not altered, drives clinical symptoms following JHMV infection. Moreover, double deficient mice in PKR and RNaseL demonstrated overall similar phenotype as RNaseL-/- mice. This finding would indicate that reduced expression of pro vs anti-inflammatory response observed in PKR-/- mice was mitigated in absence of RNaseL.

In summary, the absence of PKR contributed to increased viral replication, albeit with little consequences on T cell mediated viral control. Surprisingly however, PKR deficiency resulted in significantly reduced expression of both pro- as well as anti-inflammatory responses, without affecting overall pathogenesis. These results are the first to demonstrate numerous effects of PKR at multiple levels in CNS resident cells as well as CD4 T cells. While the role of PKR in
regulating pro- versus anti-inflammatory responses during CNS infection had no overt overall phenotype, its effect in other inflammatory diseases remains to be determined.
CHAPTER IV

IL12, BUT NOT IL23, DEFICIENCY AMELIORATES VIRAL ENCEPHALITIS WITHOUT AFFECTING VIRAL CONTROL¹

Kapil, P., Atkinson, R., Ramakrishna, C., Cua, D.J., Bergmann, C.C. and Stohlman, S.A.

4.1 Abstract

The relative contribution of IL12 and IL23 to viral pathogenesis has not been extensively studied. IL12p40 mRNA rapidly increases following neurotropic coronavirus infection. Infection of mice defective in both IL12 and IL23 (p40⁻/⁻), or IL12 (p35⁻/⁻) and IL23 (p19⁻/⁻) alone, revealed that the symptoms of coronavirus induced encephalitis are regulated by IL12. IL17 producing cells never exceeded background levels, supporting a redundant role of IL23 in pathogenesis. Viral control, tropism, and demyelination were all similar in p35⁻/⁻, p19⁻/⁻ and wild type (wt) mice. Reduced morbidity in infected IL12 deficient mice was also not associated with altered recruitment or composition of inflammatory cells. However, IFNγ levels and virus specific IFNγ secreting CD4 and CD8 T cells

¹ This work has been published in the form of a research article (Parul Kapil, Roscoe Atkinson, Chadran Ramakrishna, Daniel J. Cua, Cornelia C. Bergmann and Stephen A. Stohlman. 2009 . JVI. 83. 5978-5986).
were all reduced in the CNS of infected p35\(^{-/-}\) mice. Transcription of the pro-inflammatory cytokines \(Il1\beta\) and \(Il6\), but not \(Tnf\) were initially reduced in infected p35\(^{-/-}\) mice, but increased to wild type levels during peak inflammation. Furthermore, although \(Tgf\beta\) mRNA was not affected, IL10 was increased in the CNS in the absence of IL12. These data suggest that IL12 does not contribute to antiviral function within the CNS, but enhances morbidity associated with viral encephalitis by increasing the ratio of IFN\(\gamma\) to protective IL10.

4.2 Introduction

Resistance to infection, as well as the extent of clinical symptoms, is directly influenced by the pattern and magnitude of cytokine induction. Specifically, interferon gamma (IFN\(\gamma\)) is closely associated with control of many viruses and other intracellular pathogens (Novelli and Casanova, 2004, Scott and Kaufmann, 1991). IFN\(\gamma\) secretion in turn is enhanced by interleukin-12 (IL12), which in concert with the related cytokine IL23, participates in regulating both innate and adaptive immune responses (Hunter, 2005, Langrish et al., 2004, Novelli and Casanova, 2004, Jouanguy et al., 1999, Trinchieri, 2003). IL12 is a heterodimeric cytokine composed of two subunits (IL12p35 and IL12p40) secreted primarily by macrophages and dendritic cells. It is a potent inducer of IFN\(\gamma\) secretion by CD4 T cells, even in the presence of T regulatory cells (King and Segal, 2005). It also increases cytokine secretion as well as cytolytic potential of NK cells and CD8 T cells (Trinchieri, 2003). IL12 is rapidly induced in the periphery and the central nervous system (CNS) following a variety of viral infections (Coutelier et al., 1995, Parra et al., 1997, Trinchieri, 2003, Novelli and
Casanova, 2004); however, its contribution to virus resistance varies depending upon tissue tropism and predominant effector mechanisms employed to control virus replication. For example, IL12 is critical to control murine cytomegalovirus, while it is dispensable for immune responses to lymphocytic choriomeningitis virus; IFNγ is nevertheless required for resistance to both these infections (Orange and Biron, 1996b, Oxenius et al., 1999, Novelli and Casanova, 2004). Although IFNγ is a critical component in the protective host immune response to many viruses, not all viral infections are associated with IL12 mediated enhancement of the IFNγ response (Novelli and Casanova, 2004, Trinchieri, 2003).

IL23 shares both the IL12p40 subunit and the IL12 receptor (IL12R) β1 subunit with IL12 (Hunter, 2005, Trinchieri, 2003, Langrish et al., 2004) and similar to IL12, IL23 is primarily secreted by macrophages and dendritic cells. The role(s) of IL23 in innate and adaptive immunity have only recently begun to come to light (Hunter, 2005, Langrish et al., 2004). A primary consequence is induction of CD4 T cells secreting IL17, which play a prominent role in neutrophil recruitment to sites of inflammation (Aggarwal et al., 2003, Kolls and Linden, 2004, Langrish et al., 2004, McGeachy and Cua, 2008). Unlike the IL12R, IL23R is not expressed by naïve T cells but is present on memory T cells (McGeachy and Cua, 2008). Differential receptor expression is consistent with IL23 independent induction of IL17 secreting CD4 T cells (Langrish et al., 2004, McGeachy and Cua, 2008); however, it is required for sustained expression of autoimmune T cell effector function associated with autoimmune demyelination of the CNS (Cua et al., 2003, McGeachy and Cua, 2008). During chronic
mycobacterial and HIV infections IL23 functions as a regulator of the protective IFNγ response (Khader et al., 2005, Novelli and Casanova, 2004) but also dampens immune pathology during herpes simplex infection of the eye (Kim et al., 2008). However, IL23 has a limited capacity to influence resistance in the absence of IL12 (Khader et al., 2005, Lieberman et al., 2004).

Infection of the CNS with the neurotropic JHM strain of mouse hepatitis virus (JHMV) induces an acute encephalomyelitis accompanied by myelin loss. Virus replicates in microglia, astrocytes and oligodendroglia, but only rarely in neurons and is controlled by a vigorous inflammatory response localized to the CNS (Bergmann et al., 2006). CD8 T cells exert crucial antiviral functions via perforin and IFNγ mediated mechanisms (Bergmann et al., 2006, Bergmann et al., 2003, Parra et al., 1999); however, recent data indicates that CD4 T cells alone can also mediate virus clearance (Savarin et al., 2008, Stohlman et al., 2008). The critical requirement for IFNγ in combating JHMV infection is demonstrated by prolonged virus replication, and increased clinical disease and mortality in IFNγ deficient mice (Parra et al., 1999). In the absence of IFNγ, virus preferentially resides in oligodendroglia demonstrating a cell specific action of IFNγ within the CNS (Gonzalez et al., 2006). Furthermore, morbidity and mortality are dramatically increased in JHMV infected immunodeficient recipients of CD4 T cells unable to secrete IFNγ (Pewe et al., 2002, Savarin et al., 2008). Although demyelination, a hallmark of JHMV infection, is an immune mediated manifestation of tissue destruction rather than a direct viral cytopathic effect (Bergmann et al., 2006, Matthews et al., 2002), a contribution of IFNγ to myelin
loss is controversial. Similar myelin loss in infected IFNγ⁻ mice compared to controls (Parra et al., 1999), as well as demyelination induced by IFNγ deficient CD4 or gamma delta T cells in the context of immunodeficient hosts implicate an immune component other than IFNγ in contributing to demyelination (Bergmann et al., 2003, Dandekar and Perlman, 2002). By contrast, T cell transferred into infected immunodeficient recipients suggest that IFNγ secretion by CD4 T cells limits demyelination (Pewe et al., 2002), while IFNγ secretion by CD8 T cells contributes to demyelination (Bergmann et al., 2003, Bergmann et al., 2004, Pewe and Perlman, 2002).

IFNγ thus plays a key role in regulating clinical disease, mortality, control of virus replication and myelin loss following JHMV infection of the CNS. To determine the contribution of IL12 to JHMV pathogenesis, as well as a possible indirect contribution of IL23 to myelin loss, antiviral immune responses and tissue damage were examined in infected IL12/IL23 (p40⁻⁻), IL12 (p35⁻⁻), and IL23 (p19⁻⁻) mice. The data demonstrate that IL12 enhances the magnitude of the IFNγ response in the CNS following infection, albeit without affecting viral control. However, decreased clinical disease in IL12 deficient mice indicates that IL12 induced IFNγ and a concomitant reduction in IL10 contributes to the disability associated with JHMV induced encephalomyelitis. By contrast, no differences in demyelination were detected in JHMV infected mice deficient in either IL12 or IL23, suggesting that neither of these cytokines influences virus induced demyelination.
4.3 Results

4.3.1 IL12 enhances JHMV induced encephalitis

Increased transcription of IL12p40 occurs within the CNS rapidly following JHMV infection (Parra et al., 1997) implicating a possible role for IL12 and/or IL23 in the pathogenesis of JHMV encephalitis. Recent data further suggested that IL12p35 mRNA is rapidly induced and that IL23p19 mRNA transcription increases subsequent to virus clearance from the CNS (Held et al., 2008). However, the latter results were not correlated with IL12p40 mRNA expression. As previous reports relied upon semi-quantitative measures of mRNA, we sought to confirm the activation of IL12 and/or IL23 during JHMV infection of the CNS by real time PCR. Expression of p40 mRNA was below detection levels in the CNS of naïve animals, but was rapidly induced following infection (Figure 21), consistent with previous data (Parra et al., 1997). Although p40 mRNA declined rapidly after day 5 post infection (p.i.), it remained above basal levels to day 14 p.i. In contrast to the mRNA encoding the IL12 and IL23 common p40 chain, both the IL12 specific p35 subunit and the IL23 specific p19 subunit were detected at relatively low levels in the CNS of naïve animals (Figure 21). However, expression of both p35 and p19 mRNA remained at or near basal levels following JHMV infection (Figure 21). Despite the inability to detect upregulation of these mRNAs, the increase of IL12/23 p40 mRNA suggested that either IL12 or IL23 may play important roles in the pathogenesis of JHMV.
The potential role(s) of IL12 and IL23 on JHMV induced encephalomyelitis were thus investigated in IL12/IL-23 deficient (p40\(^{-/-}\)) mice. Following infection of wt mice clinical symptoms of encephalitis were initially detected at day 7 p.i., progressed to a peak between days 10-12 p.i. and then began to decline (Figure 22). By contrast, initial onset of symptoms was delayed in p40\(^{-/-}\) mice (Figure 22). Furthermore, although clinical symptoms in infected p40\(^{-/-}\) mice peaked at days 10-12 p.i. similar to wt mice, severity was substantially reduced (Figure 22). To determine a dominant contribution of either IL12 or IL23 to morbidity, progression and severity of clinical disease was examined following infection of mice specifically deficient in either IL12 (p35\(^{-/-}\)) or IL23 (p19\(^{-/-}\)). Delayed onset, as well as a significant reduction in peak symptoms, was only observed following infection of IL12 deficient mice (Figure 22). IL23 deficient mice exhibited no difference in onset or maximum clinical symptoms compared to wt mice (Figure 22). Mortality was also similar (~20%) in infected wt and p19\(^{-/-}\) mice. By contrast,
mortality in infected p35\textsuperscript{-/-} and p40\textsuperscript{-/-} mice was reduced (~5\%) compared to wt mice. These data indicate that IL2, but not IL23, plays a critical role in regulating the morbidity and mortality associated with acute viral encephalitis. Subsequent studies thus focused on analysis of p35\textsuperscript{-/-} mice.

Figure 22. IL12 deficiency decreases the symptoms of virus-induced encephalitis. JHV-infected p40\textsuperscript{-/-}, p35\textsuperscript{-/-}, and p19\textsuperscript{-/-} mice were examined for disease severity at various times p.i. The data are representative of two separate experiments for each strain with n>20 per experiment ± SEM. * p ≤ 0.05 (compared to wt). (Kapil et al., 2009)
Clearance of infectious virus from the CNS of p35\(^{-/-}\) and control mice was compared to assess whether reduced morbidity reflected enhanced control of virus replication. Infectious virus peaked in the CNS to similar levels at day 5 p.i. It was subsequently controlled with equal efficiency in both groups and by day 14 p.i. infectious virus was at the limit of detection in both groups of mice (Figure 23). The apparent redundancy of IL12 in clearance of infectious virus is consistent with the absence of altered CNS virus replication in mice treated with anti-IL12 antibody (Held et al., 2008). Similarly, no alteration in virus replication within the CNS was detected in mice treated with anti-IL23 antibody (Held et al., 2008) and no differences were detected in virus replication or clearance comparing JHMV infected p19\(^{-/-}\) and wt mice (data not shown). These data indicate that while IL12 plays an important role in the onset and severity of morbidity, it does not influence virus clearance.

![Figure 23. IL12 does not influence control of infectious virus.](image)

*Figure 23. IL12 does not influence control of infectious virus.* Virus replication in the CNS of p35\(^{-/-}\) and wt mice following JHMV infection. The data are representative of two independent experiments and show the mean of four individual mice per time point ± SEM. (Kapil et al., 2009)
4.3.2 IL12 does not alter CNS inflammation

Reduced morbidity during acute infection suggested that leukocytes infiltration into the CNS might be altered in the absence of IL12. However, analysis of CD45^hi bone marrow derived inflammatory leukocytes showed no difference in overall recruitment into the CNS of JHMV infected p35^{−/−} mice compared to wt mice throughout infection (Figure 24A). Recruitment of inflammatory cells was also similar following infection of p19^{−/−} mice (data not shown). There were also no differences in recruitment of Ly6G^{+} neutrophils or F4/80^{+} monocytes within the CD45^hi inflammatory populations at any time point p.i. in either p35^{−/−} or p19^{−/−} mice (data not shown). Lastly, to determine if reduced clinical disease correlated with altered recruitment of T cell subsets, CD8 and CD4 T cells within the infiltrating lymphocyte populations were examined. A slight decrease in total CNS infiltrating CD8 T cells was detected in infected p35^{−/−} relative to wt mice at days 7 and 10 p.i., which only reached statistical significance at day 7 p.i. (Figure 24B). Virus specific tetramer^{+} cells specific for the dominant S510 epitope encoded within the viral spike glycoprotein comprised ~ 40% of CD8 T cells at day 7 and ~60% at day 10 p.i. in both groups (data not shown). Similar recruitment of virus specific CD8 T cells was consistent with the ability of the p35^{−/−} mice to control infectious virus (Figure 23). The absence of IL12 was also associated with a slight decrease in CD4 T cell CNS infiltration throughout infection (Figure 24C), reaching statistical significance at day 10 p.i. In summary, although IL12 did not affect recruitment of innate immune cells, total CD8 and CD4 T cells were reduced at 7 and 10 days p.i., respectively.
4.3.3 IFNγ is compromised in the absence of IL12

In addition to its antiviral function (Bergmann et al., 2006, Gonzalez et al., 2006, Parra et al., 1997), IFNγ has been suggested to regulate clinical symptoms during JHMV induced encephalitis (Pewe et al., 2002). IFNγ is primarily secreted by activated T cells during JHMV infection, with no contribution by NK cells (Zuo et al., 2006, Savarin et al., 2008, Bergmann et al., 2006, Gonzalez et al., 2006). Although IL12 is a powerful inducer and enhancer of IFNγ secretion by T cells,
IL12 independent activation of IFNγ producing T cells has been described following a number of viral infections (Novelli and Casanova, 2004, Trinchieri, 2003). Thus, the frequency of CNS derived virus specific T cells producing IFNγ was examined. In the absence of IL12, frequencies of virus specific CD8 and CD4 T cells secreting IFNγ were consistently reduced at day 7 p.i. and. Although slightly variable between independent assays, on average the frequencies of CD4 and CD8 T cells were reduced by 30% - 40% compared to T cells from the CNS of wt mice. Overall, the impact of IL12 deficiency was greater on the CD4 T cell population at day 7 p.i. (Figure 25B). The enhancing affect of IL12 on IFNγ secretion by CD4 T cells becomes especially evident when the data reflect total numbers of virus specific T cells per brain (Figure 25C). At day 10 p.i. the proportion of IFNγ secreting CD8 T cells was comparable in wt and p35−/− mice (~36%); however, the fraction of CD4 T cells secreting IFNγ in the CNS of the p35−/− mice remained at ~60% of wt levels (data not shown). By contrast, no difference in the frequency of IFNγ producing CD4 T cells was detected in CLN (Figure 25D) or spleen (data not shown), excluding a peripheral impairment of T cell activation and expansion. Furthermore, reduced frequencies of virus specific IFNγ producing cells were not attributable to an increase in IL17 secreting cells, as these remained below 0.5% in both the CD4 and CD8 T cell compartments in wt as well as p35−/− mice (data not shown).
Figure 25. IL12 enhances IFNγ production and Class II expression. Frequencies of CNS derived IFNγ secreting T cells at day 7 p.i. (A) CD8 T cells following stimulation with S510 peptide and (B) CD4 T cells with M133 peptide. Data are representative of two separate experiments analyzing cells combined from n = 4-6 individuals (C) Number of CNS derived CD4 T cells producing IFNγ after ex vivo peptide stimulation in infected wt and p35-/- mice at days 7 and 10 p.i. Data represent the mean of two independent experiments ± SEM. (D) Frequencies of CLN derived IFNγ secreting CD4 T cells at day 7 p.i. following stimulation in presence or absence of M133 peptide. Data are representative of two separate experiments analyzing cells combined from n = 4-6 individuals (E) IFNγ dependent MHC class II expression on CD45+ microglia (boxed cells) at days 7 and 10 p.i. (F) Cell free homogenates of the infected CNS were examined for IFNγ by ELISA. Data show the mean of n= 4 per time point ± SEM and are representative of two separate experiments. Asterisks indicate statistical significance p≤ 0.05, compared to wt. (Kapil et al., 2009)

During JHMV infection MHC class II expression on microglia correlates with IFNγ levels within the CNS (Bergmann et al., 2003). To determine if the absence of IL12 also compromised IFNγ secretion in the CNS in vivo, microglia were examined for upregulation of major histocompatibility complex class II molecules. Microglia from infected p35−/− mice was indeed characterized by an
initial delay in class II expression relative to Wt mice (Figure 25E). However, by
day 10 p.i. class II expression on microglia was identical in both groups. IFNγ
within the CNS was measured by ELISA to assess to what extent the delay in
class II upregulation correlates with IFNγ protein levels. In infected Wt mice,
IFNγ was detectable at day 5 p.i., increased significantly by day 7 p.i., and then
declined substantially by day 10 p.i. By contrast, in absence of IL12, IFNγ was
undetectable at day 5 p.i., and only increased modestly by day 7 p.i. (Figure 25F).
IFNγ levels within the CNS were similarly low in both groups by day 10 p.i.
(Figure 25F), when infectious virus was already reduced. These data suggest
that even under optimal re-stimulation conditions, the capacity of T cells derived
from an IL-12 deficient CNS environment to produce IFNγ is reduced (Figure
25C). This deficit was more strongly manifested in vivo, where MHC/antigen
density, and thus T cell stimulation, is more limited. Significantly reduced IFNγ
levels in the CNS of p35−/− mice correlated with delayed class II expression on
microglia and reduced morbidity. Nevertheless, similar and sustained class II
expression at day 10 p.i. suggested that the reduced IFNγ in the absence of IL12
were still sufficient to drive optimal class II expression late during infection and
sustain viral control.

4.3.4 Viral induced demyelination is IL12 and IL23 independent

IFNγ is critical for the control of JHMV infection in oligodendroglia
(Gonzalez et al., 2006, Parra et al., 1999); however, demyelination in the host
with an otherwise intact immune system is independent of IFNγ (Gonzalez et al.,
2006, Parra et al., 1999). Although demyelination associated with experimental
autoimmune encephalitis is dependent upon IL23 expression (Cua et al., 2003, Langrish et al., 2004, McGeachy and Cua, 2008) the mechanism of myelin loss following JHMV infection is unclear (Matthews et al., 2002). Therefore, potential alterations in inflammation, demyelination and viral tropism in p35\(^{-/-}\) and p19\(^{-/-}\) mice were compared with Wt mice. Despite the delayed onset of clinical symptoms and reduced severity in p35\(^{-/-}\) mice (Figure 22) no difference in inflammation or cells expressing viral antigen were detected compared to wt (Figure 26). Although a slight increase in myelin loss was apparent in infected p35\(^{-/-}\) mice (Figure 26) it did not reach statistical significance due to variability between individuals. Similarly, the inability to secrete IL23 did not alter viral induced inflammation or demyelination (Figure 26). These data are consistent with an IFN\(\gamma\) independent mechanism of demyelination (Gonzalez et al., 2006).

Figure 26. IL12 and IL23 do not influence viral pathogenesis. Longitudinal sections of spinal cords from JHMV infected wt, p35\(^{-/-}\) and p19\(^{-/-}\) at day 10 p.i. The extent of demyelination as shown by luxol fast blue (LFB). Viral antigen (arrows) detected by immunohistochemical analysis for viral nucleocapsid using mAb J.3.3. Bar = 200 microns. (Kapil et al., 2009)
4.3.5 IL12 regulation of innate cytokine, chemokine and anti-inflammatory cytokines

Ameliorated clinical symptoms in the absence of IL12 suggested three possible mechanisms: 1) diminished activation of pro-inflammatory genes; 2) reduced chemokine expression; or 3) increased activation of anti-inflammatory genes. Expression of genes associated with the acute JHMV response or with induction or severity of clinical disease were compared in p35−/− and wt mice. Based on the crosstalk between IL12 and type I interferons in regulating IFNγ responses (Cousens et al., 1997, Cousens et al., 1999, Orange and Biron, 1996b, Novelli and Casanova, 2004, Trinchieri, 2003), IFNα4 and IFNβ mRNA levels were initially compared. Transcription of both mRNAs in the CNS was reduced by ~40%-50% in the absence of IL12 (data not shown). Nevertheless, the reduction was insufficient to alter infectious virus load or increase the tropism for neurons (Figure 26), previously shown to correlate with the inability to respond to IFNα/β (Ireland et al., 2008). IL12 enhances secretion of IFNγ indirectly via enhanced IL1β and TNF secretion, thereby initiating production of inducible nitric oxide synthase (iNOS) in addition to other IFNγ inducible genes (Novelli and Casanova, 2004). Decreased IFNγ in the absence of IL12 was indeed reflected in decreased CNS expression of the mRNA encoding iNos (data not shown). Furthermore, consistent with their role in regulating clinical symptoms Il6 and Il1β mRNA were reduced in infected p35−/− mice at day 5 p.i. (Figure 27A). However, no difference in transcription of these genes was detected at day 7 p.i. (Figure 27A), when the p35−/− group exhibited a significant decrease in morbidity (Figure
Although expression of *Tnf* mRNA was also slightly reduced in the CNS of infected IL-12−/− mice at day 5p.i., the difference did not reach statistical significance (Figure 27A). These data suggest that the initial decrease in pro-inflammatory cytokines contributes to the delayed onset of clinical symptoms.

Reduced recruitment of T cells into the CNS of infected p35−/− mice suggested a possible defect in chemokine induction following infection in absence of IL12. However, *Cxcl10* mRNA expression was only transiently reduced ~2 fold at day 5p.i. in the absence of IL12. *Cxcl10* mRNA increased to wt levels by day 7 p.i. and no differences were observed in expression of *Cxcl9* or *Ccl5* mRNA (Figure 27B). Although expression levels were equivalent by day 7 p.i., early differences presumably shape the responsiveness to chemokines and cytokines later during infection, thus maintaining overall decreased morbidity.

Based on their association with diminished clinical symptoms of EAE (Prud’homme, 2000), expression of *Tgfβ* and IL10 were examined as down regulatory candidates in JHMV infected p35−/− mice. In contrast to the pro-inflammatory cytokines, no significant alterations in *Tgfβ* mRNA were detected relative to wt controls (Figure 27C). However, IL10 was increased in the CNS of p35−/− mice at days 5, 7 and 10 p.i. (Figure 27D). Decreased morbidity was thus associated with decreased expression of pro-inflammatory cytokines as well as increased expression of the anti-inflammatory cytokine IL10.
IL12 alters the expression of pro-inflammatory and anti-inflammatory cytokines. Brains from infected wt and p35/- mice were compared for mRNA expression of pro-inflammatory cytokines (A) Il1β, Il6 and Tnfa; chemokines (B) Cxcl10, Cxcl9 and Ccl5 and the anti-inflammatory cytokines (C) Tgfβ1 and Tgfβ3 by real time PCR. Expression levels were normalized to Gapdh using the following formula: (2^{-\Delta\text{CT}} (\text{GAPDH} - \text{CT (Target)}) x 1000, where Ct is the threshold cycle. Data represent the mean of three mice per time point ± SEM. (D) IL10 levels in brains of infected wt and p35-/- mice determined by the ELISA. Data are representative of two experiments (n = 4 per time point) ± SEM. BD = below detection. Asterisks indicate statistical significance p ≤ 0.05, compared to wt. (Kapil et al., 2009)

4.4 Discussion

IL12 regulates both innate and adaptive immune responses and plays a major role in controlling bacterial and intracellular protozoa infections predominantly via stimulation of IFNγ (Jouanguy et al., 1999, Langrish et al., 2004, Trinchieri, 2003). Similarly, IL23 contributes to host defense against bacterial infections(Jouanguy et al., 1999, Trinchieri, 2003, Langrish et al., 2004). However, many viral infections resolve without IL12 participation (Trinchieri, 2003) and the contribution(s) of IL23 have not been extensively explored (Held et
Importantly, both cytokines also have the potential to influence the severity of pathological lesions and/or clinical disease (Hunter, 2005, Langrish et al., 2004, Trinchieri, 2003, Novelli and Casanova, 2004). Rapid CNS induction of IL12p40 mRNA in a neurotropic coronavirus induced demyelination model (Parra et al., 1997) thus prompted analysis of the relative contributions of IL12 and IL23 to viral clearance and disease severity. Although quantitative PCR analysis confirmed upregulation of Il12p40 mRNA (Parra et al., 1997), we were unable to detect statistically significant increases in either Il12p35 or Il23p19 mRNA following infection. Discrepancies with the previously reported early Il12p35 mRNA induction and delayed Il23p19 mRNA induction following clearance of infectious virus (Held et al., 2008) may reside in distinct viral inoculum doses or our quantitative measurements of PCR replicons. Nevertheless, similar kinetics of virus control in p35−/−, and p40−/− mice relative to wt mice, are consistent with reports demonstrating no enhancing effects of IL12 on mouse hepatitis virus clearance from the liver and CNS (Held et al., 2008, Schijns et al., 1998). Similarly, infected p19−/− mice provided no evidence for IL23 mediated enhancement of host defense (data not shown) consistent with our inability detect upregulation of p19 transcripts in JHMV infected Wt mice. A transient increase in virus replication observed by treatment with anti-IL23 (Held et al., 2008) could thus not be confirmed in p19−/− mice infected with slightly lower virus doses.
A primary role of IL12 is enhancement of cytolytic activity and IFNγ secretion by NK cells as well as T cells (Trinchieri, 2003). Nevertheless, IL12 independent T cell activation and IFNγ secretion has been noted following a number of viral infections including systemic lymphocytic choriomeningitis virus infection (Orange and Biron, 1996a) and Influenza virus infection of the respiratory tract (Monteiro et al., 1998). This may be attributed to type I IFN, which can also promote IFNγ secretion by lymphocytes (Cousens et al., 1999). Type I IFN and IFNγ are both essential to control acute JHMV infection as shown by the inability of either virus induced IFNα/β or IFNγ alone to achieve protection (Ireland et al., 2008, Parra et al., 1999). The absence of an IL12 affect on JHMV clearance from the CNS may thus be attributed to the rapid induction of IFNα (Ireland et al., 2008), which may act as both an inhibitor of IL12 and enhancer of IFNγ production (Cousens et al., 1999). Nevertheless, IFNγ levels were significantly decreased in the CNS of p35−/− mice, suggesting that type I IFN was insufficient to compensate for the absence of IL12 in augmenting IFNγ responses. Indeed, the modest decrease in IFNα/β mRNA levels in p35−/− mice supports a role of IL12 in promoting IFNγ responses locally within the CNS. Impaired NK cell function in this context was ruled out, as NK cells do not contribute to either IFNγ dependent MHC class II expression on microglia or virus control in otherwise immune competent mice (Zuo et al., 2006). Reduced IFNγ correlated with decreased frequencies as well as total numbers of virus specific CD8 and CD4 T cells derived from the CNS of infected p35−/− mice compared to wt mice. However, differences were not observed in frequencies of virus specific IFN-γ producing
CD4 T cells (Figure 25D) or CD8 T cells (Data not shown) in CLN or IFNγ producing CD4 or CD8 T cells in splenocytes (data not shown), consistent with a role of IL12 in directly promoting local T cell mediated IFNγ secretion within the CNS. Although control of JHMV in oligodendrocytes, the predominant cell type infected, is IFNγ dependent (Gonzalez et al., 2006, Parra et al., 1999), no differences in viral tropism, viral antigen distribution or viral clearance were noted comparing p35−/− and wt mice. This suggested that the CD8 T cell response in wt mice, the dominant adaptive effector response controlling viral replication (Bergmann et al., 2006), exceeds the minimum requirement to effectively inhibit viral replication. This is supported by similar results in IL15 and B cell deficient mice, which also show reduced numbers of virus specific CD8 T cells in the CNS following JHMV infection (Ramakrishna et al., 2002, Zuo et al., 2006).

A striking finding was the early effect of IL12 on clinical disease, without alterations in either virus replication or demyelination. A contribution of IL12 to clinical symptoms of viral induced encephalitis, independent of virus load, has not been previously reported to our knowledge and provides a novel insight into viral pathogenesis of the CNS. The absence of a direct influence of IL12 on tissue destruction is consistent with similar findings in infected mice treated with anti-IL12 antibody (Held et al., 2008) and other models of virus induced demyelination. IL12 contributes to limiting Semliki Forest virus infection of neurons, yet has no affect on subsequent demyelination (Keogh et al., 2002). Furthermore IL12 does not influence either the acute phase or demyelination induced by Theiler’s murine encephalitis virus (Inoue et al., 1998). The contribution of IFNγ to the
pathogenesis of CNS disease is complex, as it can act as a prominent mediator of pathogenesis during some experimental viral infections, but can also exert anti-inflammatory functions and decrease clinical symptoms (Anghelina et al., 2006, Kim et al., 2008, Savarin et al., 2008), similar to its role in autoimmunity (Kelchteermans et al., 2008). For example, inhibition of IFNγ enhances both morbidity and mortality following Semliki Forest virus, Theilers murine encephalitis virus, and JHMV induced CNS disease (Tomkins et al., 1988, Pullen et al., 1994, Parra et al., 1999).

IFNγ plays a critical role in shaping the inflammatory cells recruited into the CNS (Tran et al., 2000). It is thus difficult to assess how increased virus replication and altered inflammation, e.g. neutrophil accumulation within the CNS, both a result of IFNγ deficiency, contribute to the symptoms of acute encephalitis. This dilemma is evidenced by increased clinical symptoms and mortality of JHMV infected immunodeficient recipients of immune CD4 donor T cells deficient in IFNγ compared to wt CD4 T cells, independent of viral load (Savarin et al., 2008). Both neutrophil and monocyte recruitment in to the CNS, as well as pro-inflammatory cytokine levels were similar in the absence of IL12 at day 7 p.i., correlating reduced clinical disease most prominently with reduced IFNγ levels. The observation that clinical disease in not reduced in infected IL15−−, despite decreased virus specific CD8 T cells (Zuo et al., 2006), implicates the paucity of total and virus specific CD4 T cells in disease amelioration. Although CD4 T cells may contribute to viral clearance (Stohlman et al., 2008), they are more prominently implicated in the clinical symptoms of encephalitis (Anghelina et al.,
Deletion of the immunodominant CD4 T cell epitope dramatically reduced both clinical symptoms and mortality following a lethal JHMV challenge (Anghelina et al., 2006). Similarly, mutations of the Theiler’s murine encephalitis virus immunodominant CD4 T cell epitope ameliorate disease (Palma et al., 2002). Although IL23 dependent IL17 secreting CD4 T cells have recently been implicated in enhancing clinical symptoms during CNS autoimmune disease (Cua et al., 2003, Langrish et al., 2004) the frequencies of IL17 secreting T cells in the JHMV infected CNS remained near background levels in both wt and p35⁻/⁻ mice. A negligible contribution of IL17 secreting T cells during JHMV pathogenesis is supported by a similar disease course and demyelination in both p19⁻/⁻ mice and anti-IL23 antibody treated mice (Held et al., 2008) compared to wt mice. In contrast to experimental autoimmune encephalitis (Cua et al., 2003, Langrish et al., 2004), IL23 does not act as either a positive or negative regulator of viral induced CNS inflammation. These data support the notion that the IL12 dependent increase in virus specific IFNγ secreting CD4 T cells contributes to the expression of clinical symptoms. Furthermore, although no evidence for induction of anti-inflammatory TGFβ was detected, IL10 was modestly increased in the absence of IL12. JHMV infection of IL10⁻/⁻ mice resulted in increased encephalitis suggesting that IL12 dependent limitations on IL10 may also contribute to morbidity by increasing CD4 Th1 cell activation.

In summary, the study of JHMV induced encephalitis in IL12 and IL23 deficient mice defined IL12, but not IL23, as a controlling element in the enhanced expression of clinical symptoms during viral encephalitis. Importantly,
the absence of IL12 decreased IFNγ levels within the CNS; however, neither the absence of IL12 or IL23 impeded viral clearance or affected demyelination. These data support the notion that IL12 contributes to morbidity by enhancing IFNγ secreting CD4 T cells and reducing IL10 mediated protection.
CHAPTER V

KINETIC ANALYSIS OF PRO VERSUS ANTI-INFLAMMATORY GENES
EXPRESSED BY MICROGLIA/MACROPHAGES DURING CORONAVIRUS
MEDIATED CNS INFECTION

5.1 Abstract

The functional state of tissue macrophage, including CNS resident microglia, can have a significant impact on inflammation and pathogenic outcome in various diseases. Depending on external stimuli macrophages can exist in three major different functional states: the pro-inflammatory (M1) phenotype, anti-inflammatory (M2) phenotype, and deactivated phenotype. Microglia activation and recruitment of monocytes are among the first host responses to JHMV infection of the CNS. In this study we monitored alterations in gene expression by microglia and infiltrated macrophages to assess if these populations are 1) regulated similarly, and 2) acquire an anti-inflammatory phenotype as virus is cleared. Initial analysis of IFNγ, Il13 and Il10, stimuli regulating macrophage differentiation towards a pro vs anti-inflammatory phenotype, respectively, revealed similar upregulation with peak expression of all three cytokines at day 7 post infection (p.i.) in total CNS tissue. RNA analysis of
purified microglia and infiltrating macrophages showed microglia initially expressed the pro-inflammatory *Il12p40*, but subsequently rapidly upregulated anti-inflammatory *Il1ra* and *Ym1/2* (Chitinase like lectins). By contrast, in macrophages expression of *Il1ra* and *Ym1/2* peaked at day 3 (p.i.) and rapidly declined by day 7 p.i. Nevertheless, a strong effect of IFNγ was reflected by ~31 fold increase in expression of the M1 associated *iNos* gene in both microglia and macrophages during peak IFNγ expression at day 7 p.i. Surprisingly, however, the M2 specific *Arg-1* was also increased by ~72 fold at day 7 p.i. in both cell populations. Furthermore, phagocytic activity was similar in both microglia and macrophages and increased between day 7 and 10 p.i. suggesting increased activation. These results indicate that microglia and infiltrating monocytes exhibit overall similar regulation of pro and anti inflammatory factors at the population level. Moreover, the data suggest that both M1 and M2 populations coexist and are driven by early IFNαβ as well as T cell mediated IFNγ and IL10 production.

### 5.2 Introduction

Macrophages are innate immune cells that play a crucial role in regulating both innate and adaptive immunity in response to pathogen as well as tissue damage. The functional heterogeneity of macrophages, defined by differential activation states, was recently highlighted for various disease settings (Colton, 2009, Gordon, 2007); an overview of distinct stimuli leading to prominent gene expression characteristics is provided in Table 5.1.
Table 5.1.

Functional Heterogeneity of Macrophages

<table>
<thead>
<tr>
<th>Activation state</th>
<th>Stimulating agent</th>
<th>General function</th>
</tr>
</thead>
<tbody>
<tr>
<td>Innate activation</td>
<td>LPS, other TLR ligands</td>
<td>Phagocytic activity, Adaptive immunity</td>
</tr>
<tr>
<td>Classical activated (M1)</td>
<td>IFN(<em>\gamma), TNF(</em>\alpha)</td>
<td>Phagocytic activity, Pro-inflammatory cytokine production, Nitric Oxide (NO) production</td>
</tr>
<tr>
<td>Alternative activated (M2a; M2)</td>
<td>IL4, IL13</td>
<td>Increased phagocytic activity, Anti-inflammatory cytokine production, Arginase-1 production, tissue repair, reconstruction of extra cellular matrix</td>
</tr>
<tr>
<td>Acquired deactivation</td>
<td>TGF(_\beta), IL10</td>
<td>Immunosuppression</td>
</tr>
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</table>

Macrophages are widely distributed in the body and can be categorized into circulating monocytes and tissue resident macrophages. In the CNS microglia are the resident macrophages, which are originally derived from myeloid progenitors in the yolk sacs (Ginhoux et al., 2010). In the adult CNS,
microglia are on constant surveillance for any perturbation mediated by injury or infection.

As a consequence of CNS injury or infection microglia activation is phenotypically evident by thickening of highly ramified processes. Similar to and in concert with infiltrating macrophages they play an important role in promoting inflammation during microbial infection, but also provide protective roles by limiting ongoing tissue damage and by clearing cellular debris. These dual roles are reflected by the existence of both pro- and anti-inflammatory microglia and macrophages in various CNS inflammatory conditions. Multiple sclerosis (MS) is associated with both intracerebral M1 and M2 cells (Oleszak et al., 1998, Chiang et al., 1996). M2 macrophages were described as foamy cells that can ingest myelin-derived lipids and produce anti-inflammatory cytokines (Boven et al., 2006). The role of pro- and anti-inflammatory microglia and macrophages in MS pathology was further assessed using an experimental demyelinating rodent model termed Experimental Autoimmune Encephalitis (EAE). While, a balance between M1/M2 phenotype of microglia and macrophage resulted in mild EAE, an imbalance towards a prominent M1 phenotype promoted relapsing disease (Mikita et al., 2011). Similarly, perivascular macrophages and microglia exist in a mixed M1/M2 polarized state following HIV infection in the CNS (Benoit et al., 2008). M1 macrophages were relatively short lived, reduced the level of HIV replication but caused neurotoxicity, while an M2 activation state resulted in persistent viral infection but maintained neuronal homeostasis (Cassol et al., 2009). Overall, distinct regulation of microglia relative to infiltrating macrophages,
the co-existence of M1/M2 macrophages and microglia, and the relative contribution in propagating disease or maintaining recovery during CNS infection has not been well studied.

Microglia and macrophage express a wide array of pathogen recognition receptors (PRRs) which can be activated following recognition of pathogen associated molecular patterns (PAMPs), such as DNA and RNA structures (Takeda et al., 2003). Engagement of PRRs leads to induction of IFNα/β, as well as numerous pro-inflammatory cytokines and chemokines (Takeda and Akira, 2004). IFNα/β also induces transcription of the anti-inflammatory factor Il1ra in macrophages (Jungo et al., 2001, Sciacca et al., 2000). IFNα/β can further polarize macrophages towards an M2 phenotype following infection with Tropheryma whippelii (Al Moussawi et al., 2010). These data highlight the plasticity of pro- and anti-inflammatory gene expression by macrophages as a result of numerous autocrine as well as paracrine stimuli. They further suggest the integrative outcome of complex signaling pathways is difficult to predict in vivo, especially in situations where infiltration by T cells provides prominent changes in environmental cues, such as release of IFNγ and IL10.

Following intracerebral inoculation, JHMV initially targets ependymal cells and spreads to microglia, astrocytes and oligodendroglia, with oligodenrocytes being the most prominently infected cell type (Ramakrishna et al., 2004); (see Chapter 2). Infection elicits numerous proinflammatory factors resulting in recruitment of leukocytes. Driven by CCL2, monocytes constitute the largest portion of early CNS infiltrating cells (Savarin et al., 2010), and provide further
targets of infection (Ireland et al., 2009). Early infection is characterized mainly by pro-inflammatory cytokines, such as TNFα, IL1β, IL1α, IL6 and IL12 (Bergmann et al., 2006, Parra et al., 1997), with IFNγ only prominent as T cells infiltrate the CNS (Parra et al., 1999, Stohlman et al., 1995). Nevertheless, IL4, an anti-inflammatory signal, is also induced in response to JHMV infection (Parra et al., 1997). In vitro, infection induces IL1β in both primary microglia and macrophage and IL1α, IL6 and IL12 are all induced by microglia (Pearce et al., 1994, Rempel et al., 2004, Rempel et al., 2005, Stohlman et al., 1995). More recent data indicate IL10 is prominently produced by T cells (Puntambekar et al., 2011). These findings suggested JHMV infection induces complex cues with the potential to favor both pro- and anti-inflammatory microglia and macrophages.

The studies in this chapter thus set out to address three interrelated questions:

1. Are microglia and macrophage skewed from a prominently proinflammatory to an anti-inflammatory phenotype as virus is controlled or do they reveal a mixed phenotype throughout?
2. Do infiltrating macrophages display overall similar gene expression profiles as microglia over time, indicating exposure to similar environmental cues?
3. How does IFNαβ influence the microglia/macrophage gene expression with respect to M1/M2 associated genes?

To characterize distinct functional populations, microglia and macrophages were isolated from brains of C57Bl/6 mice at various times post JHMV infection and
mRNA prepared for real time PCR analysis. We detected induction of both pro- and anti-inflammatory factors at the transcript level, suggesting the simultaneous existence of both M1 and M2 phenotypes in both microglia and macrophages. Moreover, the expression profiles were overall similar between resident microglia and the infiltrating macrophages. One exception was high expression of *Il1ra* in early infiltrating monocytes, which was IFNα/β signaling dependent. With the exception of early *Il1ra*, anti-inflammatory markers already increased between days 5 and 7 p.i., Phagocytic activity of microglia as well macrophages was also increased between days 7 and 10, supporting an increasingly anti-inflammatory phenotype with increased potential to clear dying cells.

5.3 Results

5.3.1 The innate phase of JHMV infection is associated with abundant monocyte recruitment

Many approaches study the activation states of macrophages by *in vitro* stimulation with microbial agonist and cytokines (Murray and Wynn, 2011). However, *in vivo* these cell types are subjected to many different signals dependent on their localization. Resident tissue macrophages exist in an anti-inflammatory state by default to minimize continuous inflammation mediated by homeostatic cell turnover (Murray and Wynn, 2011). Initial warning signals at the site of infection or injury lead to activation of resident macrophages and trigger the recruitment of circulating monocytes. During JHMV infection, CCL2 is the major chemokine recruiting monocytes to the CNS (Savarin et al., 2010). (Figure 28) depicts the kinetics on monocyte recruitment and the relative numbers of
microglia and monocyte derived macrophages in the CNS during the first week of infection. Microglia are characterized as CD45lo F4/80+ cells (Figure 28). Infiltrating macrophages exhibit a CD45hiF4/80+ phenotype and comprise ~75% at day 3, and 5 p.i. and ~69% at day 7 p.i. of total infiltrating populations (Figure 28). The frequency of macrophages decline subsequently to ~36% at day 10 p.i. (Figure 28). These data reveals macrophages as a major infiltrating population early following infection, which declines by 2 weeks p.i.

**Figure 28.** Kinetics of CD45hi F4/80+ macrophage, and CD45lo F4/80+ microglia in the CNS following JHMV infection. Representative density plots of CNS derived cells stained with CD45 and F4/80 at day 3, 5, 7, and 10 p.i. CNS-infiltrating cells contain a prominent population of CD45hi F4/80+ macrophages early during the infection, while microglia are represented as CD45lo F4/80+ cells. The data represents three separate experiments at each time point, combining cells from 3-4 mice/time point/experiment.

### 5.3.2 JHMV replicates in microglia and infiltrating macrophages early during the infection

Following intracerebral infection JHMV spreads to a variety of resident cells. Microglia are one of the initial targets for infection consistent with their expression of the MHV receptor, carcinoembryonic antigen cell adhesion molecule 1 (CECAM-1 or MHVR) (Nakagaki and Taguchi, 2006, Ramakrishna et al., 2004). Similarly a subset of infiltrating macrophages is also infected (Ireland
et al., 2009). As virus replication drives initial PRR dependent innate responses, we compared the relative kinetics of viral replication in microglia and infiltrating macrophages by quantifying the level of viral RNA using primers specific for RNA encoding the nucleocapsid (N) protein, which are found on both genomic and the nested subgenomic RNAs (Baric and Yount, 2000, Schelle et al., 2005).

Microglia and infiltrating macrophages were purified by FACS and RNA viral N RNA expression real time PCR analysis Gapdh as a housekeeping gene. Viral RNA expression was maximal at day 3 p.i. and declined by day 5 and 7 p.i. in both microglia and macrophages (Figure 29), indicating innate stimulation decreases coincident with decreasing virus. This is indeed confirmed by kinetics of Ifnα/β levels in chapter 2.

![Figure 29. Kinetics of viral RNA in macrophage and microglia](image)

**Figure 29. Kinetics of viral RNA in macrophage and microglia.** Viral mRNA comprising N-protein encoding sequences was measured in microglia and macrophage populations isolated from brains of JHMV infected C57Bl/6 mice using real time PCR. Numbers depict mRNA levels relative to the house keeping gene GAPDH using the following formula \(2^{(C_T - C_T(\text{target}))} \times 1000\), where \(C_T\) is the threshold cycle. The data represent the average values of pooled cells from 5-6 mice of three individual experiments ± SEM.
5.3.3 Activation of microglia/macrophage following JHMV infection

To further characterize the activation state of microglia and macrophages within the CNS early following JHMV infection, we determined the initial induction of representative pro- and anti-inflammatory factors. The pro-inflammatory cytokine \( \text{Il12 p40} \) was barely detectable in microglia from naïve mice. However, we observed \( \sim 200 \) fold induction in \( \text{Il12 p40} \) as early as day 3 p.i. in microglia which was subsequently downregulated, but still remained \( \sim 55 \) fold and \( \sim 5 \) fold higher at day 5 and 7 p.i. respectively, compared to naïve levels (Figure 30). Similarly, expression of \( \text{Il12 p40} \) peaked at day 3 p.i. in infiltrating macrophages and was subsequently reduced. Overall, \( \text{Il12 p40} \) expression reflected the kinetics and magnitude of virus replication in both populations (Figure 29). These data supported a direct correlation between virus infection and induction of \( \text{Il12 p40} \).

We also assessed transcription of two genes associated with anti-inflammatory functions. While IL12 promotes IFN\(_\gamma\) secretion, the anti-inflammatory factor Ym1/2 has been reported to induce expression of the Th2 cytokines IL13 (yeping cai, 2009, JI). Ym1/2 has been described as secretory protein that shares a close homology with chitinases. (Jin et al., 1998). Ym1 and Ym2 are isomers sharing \( \sim 91\% \) similarity and therefore many studies do not differentiate between these isomers and collectively referred as Ym1/2 proteins. In the CNS, Ym1/2 are involved in inducing regeneration of olfactory receptor neurons (Giannetti et al., 2004). A second prominent anti-inflammatory mediator, which counteracts the innate pro-inflammatory cytokines IL1\(\alpha\) and IL1\(\beta\), is IL1Ra.
This antagonist competitively binds to IL1 receptor 1 without inducing signal transduction, thereby downregulating IL1α and β activity (Eisenberg et al., 1990, Seckinger et al., 1987). IL1Ra is also protective during brain injury (Hutchinson et al., 2007). Similar to Il12 p40, microglia from naïve mice exhibited very low expression of Il1ra, but expression of Ym1/2 was readily detectable, confirming a prominently anti-inflammatory state prior to infection. However, while Il-1ra increased ~74 fold, Ym1/2 expression stayed at basal level levels in microglia at day 3 p.i. (Figure 30). Expression of Il1ra was further induced to ~134 fold at day
5 p.i. and still remained ~52 fold higher over naïve levels at day 7 p.i. (Figure 30). Ym1/2 was also increased at day 5 p.i., and down-regulated by day 7 p.i., but the maximal increases was only ~6 fold relative to its basal expression (Figure 30). Contrasting delayed induction of anti-inflammatory response genes in microglia, Il1ra and Ym1/2 mRNA levels were overall much higher in infiltrating macrophages at day 3 p.i. Furthermore, although mRNA levels encoding these factors declined progressively in macrophages throughout days 5 and 7, they remained at elevated levels compared to microglia (Figure 30). Overall these data suggest that early infiltrating macrophages contribute substantially to an anti-inflammatory environment.

5.3.4 Expression of cytokines polarizing pro- and anti-inflammatory macrophages in the CNS following JHMV infection

Virus mediated activation of microglia and macrophages lead to induction of various cytokines, which in-turn mediates CNS recruitment of peripherally activated T cells. During intracerebral CNS infections initial activation, differentiation and expansion of T cells takes place in the draining cervical lymph nodes. T cells start accumulating in the CNS as early as day 5 p.i. and increase to maximal numbers between days 7 and 10 (Ireland et al., 2009, Kapil et al., 2009); chapter 4). During this time T cell effector function is further influenced by the CNS environment and responsible for clearance of the vast majority of infectious virus (Bergmann et al., 2003, Bergmann et al., 2004, Gonzalez et al., 2006, Parra et al., 1999, Stohlman et al., 1995). The key antiviral mediator is IFNγ (Kapil et al., 2009, Parra et al., 1999), while IL10 production by T cells
dampens T cell pathogenicity (Lin et al., 1998, Trandem et al., 2011). While IFN$\gamma$ is considered an activating cytokine for macrophages, IL10 mediated signaling to macrophages is associated with deactivation. Expression of these cytokines is thus likely to play a dominant role in influencing macrophage activity. Figure 31 shows the relative levels of IFN$\gamma$, $\text{Il10}$ and $\text{Il13}$, (a cytokine associated with alternative macrophage activation) in the CNS during JHMV infection. IFN$\gamma$ protein is detectable in the brain as early as day 5 p.i. peaks at day 7 p.i., and is significantly reduced, yet still detectable by day 10 (Figure 31). Similarly, the deactivation factor $\text{Il10}$, predominantly expressed by CD4 T cells (Puntambekar et al., 2011) also peaks at day 7, but stays elevated at day 10 p.i. (Figure 31). Finally, the increase in $\text{Il13}$ in the brain followed similar kinetics as IFN$\gamma$ and $\text{Il10}$, peaking at day 7 p.i., albeit at considerable lower overall levels. Overall the increases and decreases relative to naïve levels were also less pronounced (Figure 31). The overall expression profiles of IFN$\gamma$, $\text{Il13}$ and $\text{Il10}$ followed similar kinetics as previously described for the anti-inflammatory cytokines $\text{Tgf}\beta$ (Kapil et al., 2009) and cytokine $\text{Il4}$ (Parra et al., 1997). These data clearly demonstrate expression of cytokines known to polarize macrophages into a both pro- and anti-inflammatory phenotypes.
Figure 31. Kinetics of cytokine expression known to polarize distinct microglia/monocyte activation states. Protein levels of the pro-inflammatory cytokine IFN-γ was measured using ELISA and are represented as ng/brain (n=9 mice/time-point). Anti-inflammatory cytokines IL-13 and IL-10 were measured at the transcriptional level using real time PCR (n=6 mice/time-point). Data are represented as relative to GAPDH as depicted in Fig. 29. Values represent the average of three separate experiments ± SEM.

5.3.5 T cell mediated influence on M1/M2 profiles of microglia and macrophages following JHMV infection

IFNγ induces macrophages to produce iNOS, resulting in a classically activated M1 phenotype (Nathan et al., 1983). iNOS metabolizes amino acids to produce nitric oxide (NO) (Bronte and Zerviello, 2005, nat rev immunol). NO provides protection against intracellular pathogens, however, if released in to the extracellular milieu, it can cause damage to the neurons. In contrast, Th2 cytokines, IL4, and IL13 induce production of arginase1 and mediate polarization of macrophage towards an alternatively activated M2 phenotype (Gordon, 2007). Furthermore, overexpression of IL10 in macrophages shown to induce gene expression and activity of arginase1 and reduce production of reactive nitrogen intermediates (Schreiber et al., 2009). Arginase1, expression initiates shift in metabolism of L-arginine from NO production by iNOS towards generation of L-
ornithine, a precursor for polyamines and collagen, which is important for wound healing (Bansal and Ochoa, 2003, Durante et al., 2007). Furthermore, the cytokines IL10 and TGFβ induce a deactivated state in macrophages (Mantovani et al., 2004). Based on the presence of both IFNγ and anti-inflammatory cytokines (IL4, IL13, IL10 and Tgfβ) in the brain following JHMV infection, we monitored microglia and macrophages for expression of the respective signature M1/M2 molecules iNos and Arg-1.

Expression of iNos was upregulated ~3 fold at day 5 p.i. and ~31 fold at day 7 p.i. in microglia, compared to naïve levels (Figure 32). Expression of iNos was reduced at day 10 p.i. but still remained ~11fold higher than basal level (Figure 32). Expression of iNos in infiltrating macrophages follow the similar kinetics as microglia (Figure 32). Increasing iNos levels in both microglia and macrophages between days 5 and 7 p.i. is consistent with increasing IFNγ mediated signaling. To determine the existence of alternative state macrophages, we monitored expression of arginase1. In microglia expression of Arg-1 was upregulated ~8 fold at day 5 p.i. and further increased ~70 fold at day 7 p.i. and remained higher at day 10 p.i. relative to basal levels in microglia (Figure 32). Similarly, a major increase in Arg-1 was apparent in macrophages between days 5 and 7 p.i. (Figure 32). Although, the expression level of Arg-1 was reduced at day 10 p.i. but still remained elevated compared to day 5 (Figure 32). Overall, these results demonstrate the co-existence of both M1 and M2 specific gene patterns in both microglia and macrophages following JHMV infection. However sustained expression of arginase1 in microglia at day 10 p.i.
indicates a prevalence of anti-inflammatory phenotype in these cells. The data further demonstrate rapid adaptation of macrophages to environmental influences exerted by T cells.

Figure 32. Upregulation of mRNA characteristic of M1/M2. Microglia and macrophages were isolated from infected mice by FACS and RNA analyzed for IFNγ mediated upregulation of iNos as a representative M1 specific factors and Arg1 as a typical M2 specific factor. Expression relative to Gapdh was determined by real time PCR. The data represent the average values of pooled cells from 5-6 mice of three individual experiments ± SEM.

5.3.6 Increasing phagocytic activity of microglia and macrophage following infection

Phagocytosis of myelin debris by activated microglia and macrophages was reported to be an essential response to promote regeneration following MS (Boven et al., 2006). Furthermore, clearance of dead or dying cells by myeloid cells is critical to maintain a healthy microenvironment. To determine the
phagocytic activity of microglia and macrophage following JHMV infection we incubated brain-derived cells with fluorochrome labeled Ecoli particles (K12 strain) ex vivo and monitored the uptake in a phagocytic assay. Both microglia and macrophages showed high phagocytic ability at day 7 p.i. which was further increased at day 10 p.i. (Figure 33). The uptake was ∼40% at day 7 and ∼80% at day 10 p.i. in both microglia and macrophages (Figure 33). Increased phagocytic activity in microglia and macrophages as T cells accumulate and virus is controlled suggest an overall M2 phenotype. Whether the macrophage populations expressing M1 and M2 associated genes segregate into distinct populations, or represents a mixed phenotype remains to be determined by in situ hybridization. Future studies may also utilize bone marrow derived macrophages and primary microglial cultures exposed to external M1 or M2 specific stimuli to further assess gene expression and phagocytic activity in vitro.

Figure 33. Phagocytic activity of macrophages and microglia is increased as virus is controlled. Phagocytic ability of microglia and macrophages was determined by incubating percol gradient purified brain cells with fluorescein (494/518) labeled E coli particles (E2861) at 5 MOI (5 particles per one cell) at day 7 and day 10 p.i. The data is representative of two separate experiments. Cells were pooled together from 2-3 mice/time-point/experiment.
5.3.7 IFNα/β signaling plays an important role in induction of anti-inflammatory cytokine IL1Ra

IFNα/β induction in microglia initiates induction of various interferon-stimulated genes (ISGs), not only in the same cells, but also in neighboring cells such as oligodenroglia (Chapter 2). These ISGs include pro-inflammatory as well as anti-viral genes. However, IFNα/β also display anti-inflammatory properties as indicated by their ability to induce IL1Ra (Aman et al., 1994, Palmer et al., 2004). The importance of antiviral activity of IFNα/β during JHMV mediated CNS infection was established in mice deficient in IFN receptor chain 1 (IFNAR1-/-). IFNAR mice exhibited uncontrolled virus replication and pro-inflammatory responses and succumbed to infection within 8 days (Ireland et al., 2008). To determine the role of IFNα/β in upregulation of Il1ra, we initially monitored IFNα/β expression level in microglia and macrophages derived from brain of infected wt mice using real time PCR analysis. The levels of IFNα/β transcripts in microglia and macrophage correlated closely with viral RNA loads in both cell types (Figure 34A). Expression of both Ifna and Ifnβ transcripts peaked at day 3 p.i. in macrophages and was subsequently reduced at day 5 and 7 p.i. (Figure 34A). In microglia both Ifna and Ifnβ were upregulated at day 3 p.i.; while Ifna transcripts followed the same trend as in macrophages, expression of Ifnβ peaked at day 5 p.i. and was reduced at day 7 p.i. (Figure 34A). These data confirmed macrophages and microglia as source of IFNα/β in the CNS (Roth-Cross et al., 2008); Chapter2).
We subsequently isolated mononuclear cells from brains of IFNAR-/- mice and congenic control C57/Bl6 mice at different time post infection following JHMV infection to monitor $Il1ra$ levels. Induction of $Il1ra$ was significantly blocked in both microglia and macrophages in absence of IFNα/β signaling (Figure 34B). These results suggested that significantly increased expression of pro-inflammatory molecules in the CNS of JHMV infected IFNAR-/- mice may not only be attributed to uncontrolled viral spread, but also to significantly impaired anti-inflammatory $Il1ra$ expression.

Figure 34. Upregulation of IL1R antagonist mRNA in microglia and macrophage is IFNαβ dependent. (A) Ifnα and Ifnβ transcripts in FACS purified microglia and macrophages were determined using real time PCR. Data represent the average of three separate experiments ± SEM. (B). $Il1ra$ transcript levels were determined in both microglia and macrophage isolated by FACS from IFNAR-/- and wt mice following JHMV infection. The data is from a single experiment. All data is presented as relative to Gapdh. Cells were pooled together from 5-6 mice/time-point/experiment.
5.4 Discussion

Microglia and macrophages are dynamic cells, which continuously survey their surroundings for pathogenic agents and dangerous signals. In response to extracellular signals they maintain cellular homeostasis by phagocytosis of cellular debris and toxic substances. At the site of infection or injury, activation of microglia and infiltrating macrophages mediates protective effects, but also contributes to tissue damage (Yong and Rivest, 2009). This conflicting behavior can be attributed to the plasticity of these cell types, as both microglia and macrophage can adopt pro- and anti-inflammatory states, thereby exhibiting harmful or protective properties (Mantovani et al., 2004, Michelucci et al., 2009, Mosser, 2003). Polarization of microglia and macrophage towards an M1 phenotype is linked to increased production of pro-inflammatory cytokines and pathophysiology during HIV induced dementia (HAD) and Alzheimer’s disease (AD) (Minagar et al., 2002). Treatment with non-steroid anti-inflammatory drugs (NSAIDs) resulted in reduced amyloid plaque burden in mice and reduced risk for AD in humans (Stewart et al., 1997, Szekely et al., 2004, Zandi et al., 2002). Furthermore, M2 macrophages promote virus survival and persistence (Cassol et al., 2009).

During JHMV infection, microglia/macrophages are prominent cell types inducing pro-inflammatory responses, and their uptake of myelin is characteristic of ongoing demyelination during viral persistence. To determine the plasticity of activation states during JHMV mediated CNS infection, microglia and macrophages purified at different times post-infection from infected brains were
characterized for pro- and anti-inflammatory phenotypes. Comparative analysis of transcriptional regulation following CNS infection by JHMV revealed the existence of overlapping pro and anti-inflammatory (M1/M2) polarization in both microglia and macrophages. While an early increase in \( \text{Il12p40} \) and delayed upregulation of IFN\( \gamma \) stimulated \( \text{iNos} \) readily depicted an M1 phenotype, an early transient increase in \( \text{Ym1/2} \), followed by delayed upregulation of \( \text{Arg-1} \) characterized an M2 phenotype. A prevalent M2 phenotype coincident with viral control was supported by increased phagocytic potential of both microglia and macrophages between 7 and 10 days p.i. By contrast, transcription of \( \text{Il1ra} \) was most prominent in infiltrating macrophages during the initial 5 days of infection, suggesting a critical role of early accumulating monocytes in dampening exacerbated pro-inflammatory responses.

The coexistence of M1/M2 or presence of mixed phenotype macrophages is supported by similar observations in distinct model systems. A transcriptome analysis of monocytes during human cytomegalovirus infection reveals an atypical mixed M1/M2 phenotype biased towards an M1 phenotype (Chan et al., 2008). In contrast, patrolling monocytes initially display an M1 phenotype following Listeria monocytogens infection, which subsequently changed to an M2 phenotype during tissue remodeling (Auffray et al., 2007). Furthermore, an imbalance in equilibrium of M1/M2 expression towards M1 phenotype in microglia and macrophage in brain and blood monocytes has been linked to favor relapsing EAE (Mikita et al., 2011). These studies emphasize that distinct functional states of microglia/macrophage play a critical role in final outcome of
disease. An imbalance between various polarization states can result in early termination of protective microbial responses or aggravated progression of autoimmune responses.

IL1Ra is a soluble factor, which competes with IL1 (α and β) for the IL1 receptor (Carter et al., 1990, Seckinger et al., 1987), thus antagonizing IL1 mediated inflammation. IL1 is an innate cytokine, which plays a crucial role in leukocyte extravasation to site of infection or injury (Tamaru et al., 1998). Dysregulation in IL1 is linked with autoimmune and abnormal immune responses (Horai et al., 2000, Nicklin et al., 2000). Exogenous administration of IL1 exacerbates the course of EAE, while administration of IL1Ra suppresses disease (Jacobs et al., 1991, Martin and Near, 1995). IL1Ra production can be induced by various different signals including GMCSF, IL3, IL4 and IFNα/β (Jenkins and Arend, 1993, Sciacca et al., 2000). Both IFNα and IFNβ can induce IL1Ra production in different cell types in vitro, including monocytes (Jungo et al., 2001, Wan et al., 2008). Importantly, IFNα/β treatment in Hepatitis C viral infection as well as in autoimmune disease such as Rheumatoid Arthritis (RA) induces IL1Ra production in patients (Cotler et al., 2002, Palmer et al., 2004). Consistent with these findings, early expression of Il1ra in microglia and macrophages following JHMV infection correlates with peak of induction of Ifnα/β in these cell types. Moreover, the significant reduction in Il1ra expression in both macrophage and microglia in the absence of IFNα/β signaling (IFNAR−/− mice) supports IFNα/β as critical inducers of IL1Ra in the CNS. Elevated proinflammatory cytokines in the CNS of infected IFNAR mice further support an
anti-inflammatory property of IFNα/β, which may be mediated via IL1Ra following JHMV infection.

In summary, our findings demonstrate the existence of both pro- and anti-inflammatory microglia and macrophages in the CNS following JHMV infection. Moreover, the regulation of characteristic M1 and M2 associated genes in microglia follows similar kinetics as in infiltrating macrophages. While monocyte recruitment is associated with detrimental responses in EAE, their recruitment during JHMV infection does not have adverse consequences (Savarin et al., 2010), perhaps due to their overall more abundant expression of M2 genes. Furthermore, similar disease course when monocyte recruitment is abrogated suggests the function of microglia is sufficient to control inflammation. Finally, IFNα/β is not only crucial in controlling viral dissemination with in the CNS (Ireland et al., 2008), but is also required for expression of the anti-inflammatory factor Il1ra. Overall, the data indicate a delicate interaction between pro and anti-inflammatory responses in the CNS.
CHAPTER VI
DISCUSSION

In this thesis we have focused on innate responses following gliatropic coronavirus induced encephalomyelitis and how they impact adaptive responses and the pathogenesis. Recognition of viruses by various PRRs provides the first line of defense by activating the transcription factors, NFkB and Interferon Regulatory Factors (IRFs). These early responses limit viral spread by initiating transcription and activation of Interferon (IFN) induced antiviral mediators, such as RNaseL and double stranded RNA-dependent protein kinase (PKR). However they also regulate pro and anti-inflammatory responses, thereby contributing to the tissue damage if not appropriately balanced. There is a growing appreciation of distinct abilities of resident CNS cells to recognize and respond to microorganism, how responses are coordinated following infection in vivo is poorly understood.

Studies in mice defective in type1 IFN signaling (IFNAR-/- mice) demonstrated a critical protective role of type1 IFN signaling in preventing virus replication and dissemination following peripheral as well as CNS infection (Cervantes-Barragan et al., 2007, Ireland et al., 2008). Although plasmacytoid
dendritic cells (pDCs) were identified as primary type1 IFN inducers in the periphery, the contribution of CNS resident cells to IFNα/β production in vivo was not dissected. Our analysis of the relative contribution of microglia/macrophages and oligodendroglia in vivo initiating and propagating IFNα/β responses following coronavirus CNS infection revealed that oligodendroglia are incapable to produce IFNα/β. By contrast, despite being infected at lower levels, microglia initiated IFNα/β expression. Compared to microglia oligodendroglia also mounted a restricted response towards external IFNα/β to induce anti-viral state. Importantly, low basal and delayed inducible levels of mRNA encoding components of the IFNα/β pathway in oligodendroglia was reflected in increasing viral mRNA levels in oligodendroglia at a time when virus was already controlled in microglia. The ineffective innate control in oligodendroglia therefore appears to contribute to preferential virus spread in this cell types prior to accumulation of T cells with anti-viral activity. Nevertheless, although T cells reduce virus load in oligodendroglia, virus still persists preferentially cell in this cell type. Whether a stronger innate response in oligodendroglia could slow virus replication, thereby enhancing the chances of T cells to clear the virus remains to be determined. Oligodendroglia have important function other than myelin formation, including neuronal support by secreting neurotropic factors as well as in neurotransmission. Neurons are cells with little or no regenerative capacity and would be especially vulnerable to inflammation-mediated damage. Therefore, limited and narrowly focused innate responses to infection by oligodendroglia provides a mechanism to avoid attraction of infiltrating cells and to protect their CNS functions.
The role of IFN induced anti-viral genes is largely unknown in the CNS. We determined role of two well-described IFN induced antiviral genes, RNaseL and Protein Kinase RNA Dependent (PKR) following JHMV infection in the CNS. These latent enzymes are constitutively expressed at lower levels and further upregulated by IFNαβ. When activated by dsRNA these antiviral molecules can directly block viral replication by degrading viral as well as host RNA and by inhibiting protein synthesis. Absence of RNaseL resulted in modest increase in viral burden specifically in macrophage/microglia following JHMV infection in the CNS, without altering overall virus clearance (Ireland et al., 2009). However, unexpectedly RNaseL deficiency resulted in increased axonal damage, demyelination and mortality (Ireland et al., 2009). As RNaseL played a minimum role as an anti-viral factor, we focused on potential role of PKR. PKR activation results in phosphorylation of eIF2α, an eukaryotic translation initiation factor and Inhibitor of NFkB (IκB). Where, Phosphorylation of eIF2 leads to inhibition of translation in the host as well virus, phosphorylation of IκB leads to activation of potent transcription factor NF-κB and transcriptional induction of downstream pro- and anti-inflammatory genes. Our studies demonstrated a modest direct antiviral role of PKR, but a strong affect on transcriptional regulation of both and pro-inflammatory genes such as Il6, Tnfα, Ccl5, Cxcl10, and anti-inflammatory genes namely Il10. Surprisingly, reduced Il10 mRNA in the CNS, correlated with a direct defect in the ability of CD4 T cells to produce IL10. Furthermore, the absence of PKR resulted in increased accumulation of CD4 T cells in the perivascular space. Diminished Il10 expression observed in PKR/-/- mice was
consistent with enhanced and more diffused demyelination. PKR mediated regulation of IL10 in CD4 T cells suggested a virus independent effect and was consistent with PKR mediated IL10 regulation in macrophages. These results confirmed the notion that antiviral factor also play a significant role in immune regulation and cellular homeostatic functions during CNS inflammation. It may therefore be informative if RNaseL and PKR play a role in non-viral immune mediated demyelinating disease.

During microbial control or injury induced inflammation the host induces countermeasures to limit the tissue damage after virus infection. Macrophage and microglia have been identified to play an ambivalent role in inflammation based on their plasticity to acquire pro to anti-inflammatory phenotype following various CNS infection as well as autoimmune model such as EAE. We therefore explored role of infiltrating macrophage and microglia in regulation of inflammation following JHMV infection. Microglia and infiltrating macrophages exhibited mRNA expression characteristic of pro- as well as anti-inflammatory responses, indicating co-existence of several activation phenotypes. Overall similar expression patterns but different magnitude of gene expression between macrophage and microglia, suggests a cell type specific contribution in regulating pro and anti-inflammatory responses at different stages of infection. Microglia induced IL12p40 expression at ~200 fold as soon as at day 3 p.i. IL12p40 is a common subunit shared by both IL12 and IL23. While IL12 initiates CD4T cells towards IFNγ producing Th1 phenotype; IL23 initiates their activation towards IL17 producing Th17 phenotype. Our results indicate that IL12 but not IL23
deficiency resulted in ameliorated clinical disease without altering virus replication or demyelination. Reduced clinical disease in IL12 deficient mice was attributed to diminished pro-inflammatory response ($\text{Il6}$, $\text{Il1}\beta$ and IFN$\gamma$) and increased anti-inflammatory response (IL10). However, reduced level of IFN$\gamma$ was sufficient to control virus replication in the CNS. These results indicate that balance between pro and anti-inflammatory cytokines dictate a complex interplay between nature of the CNS insult and timing of the cytokine expression during disease.

Therefore, our findings emphasize the importance of cell specific innate immune responses following virus mediated CNS infection using coronavirus model. These studies further underscores the growing understanding of innate immune responses and its collaboration with adaptive immune response to combat acute and persistent infection and their role in successful or pathological outcome in general and in the CNS.
CHAPTER VII
MATERIALS AND METHODS

7.1 Mice

C57BL/6 (H-2D^B) -IL23-deficient (p19^-/-) mice were purchased from Schering-Plough, Biopharma (Palo Alto, CA). C57BL/6-IL12- deficient (p35^-/-) mice as well as -IL12 and -IL23 (p40^-/-) double deficient animals were purchased from Jackson Laboratory (Bar Harbor, ME). Mice expressing green fluorescent protein (GFP) under control of the proteolipid protein (PLP) promoter (PLP-GFP) (Fuss et al., 2000) were backcrossed 6 times with C57BL/6 mice prior to use. Congenic mice were used in subsequent experiments for deficient in type-1 IFN receptor-1 (IFNAR^-/-) (Ireland et al., 2008), PKR^-/- (Yang et al., 1995) and dual deficient in RNaseL and PKR (DRP^-/-) (Zhou et al., 1999). Mice were housed and bred in an accredited animal facility at the Cleveland Clinic, Cleveland, OH in accordance with IACUC guidelines. Age-matched C57BL/6 wild-type mice purchased from the National Cancer Institute (Frederick, MD) were used as controls in all experiments.
7.2 Virus Infection and Clinical Disease

6-7 week old mice were restrained and given an intracerebral injection of 250 plaque forming units (PFU) of the neutralizing monoclonal antibody (mAb) - derived neurotropic John Howard Mueller strain of mouse hepatitis virus (JHMV), variant 2.2v-1, in 30μl of endotoxin free Dulbecco’s modified phosphate-buffered saline (PBS). In PKR studies, virus was used at 1000 PFU. Severity of clinical disease was based on post-infection scores graded as 0 = healthy, 1 = ruffled fur and hunch back, 2 = reduced mobility & inability to upright, 3 = paralysis and wasting, and 4 = moribund or dead (Fleming et al., 1986).

7.3 Intracranial Inoculation of Poly (I:C)

TLR3 ligand Poly (I:C) (high molecular weight) was purchased from Invivogen, suspended in endotoxin-free phosphate buffered saline (PBS) to a final concentration of 200 µg/30 µl, aliquoted and stored at -20° C until use. Six week old C57BL/6 mice were injected intracerebrally with 200 µg poly (I:C) in 30 µl sterile PBS, similar to the JHMV infections as described. Brains and spinal cords were removed at 2, 6, 12, and 24, post inoculation, snap frozen in liquid nitrogen, processed and analyzed for mRNA expression.

7.4 Isolation of the Mononuclear Cells and Determination of the Virus Titer

Mononuclear cells were isolated from brain, spleen and cervical lymph nodes (CLN) as previously described (Ireland et al., 2008). Briefly, spleen and CLN tissues were dissociated, washed, and resuspended in RPMI supplemented with 25mM HEPES at 7.2 pH. Additionally, splenocytes were treated with Gey’s solution to lyse red blood cells. Brain mononuclear cells were isolated from
individual tissues homogenized in 4 ml of Dulbecco’s PBS using chilled Ten-Broeck glass homogenizers (7 ml). Following centrifugation at 400 x g for 7 minutes, clarified supernatants were collected and stored at -80°C. The brain cell pellet was resuspended in a final concentration of 30% Percoll (Pharmacia, Uppsalala, Sweden), underlaid with 1 ml of 70% Percoll, and the gradient was centrifuged for 30 min at 800 x g at 4°C. Mononuclear cells were collected from the interface between the 30 and 70% Percoll layers, washed twice with RPMI-HEPES and then viable cells were counted using a hemocytometer-based trypan blue exclusion assay. Virus titer in clarified brain supernatants was determined by plaque assay on a monolayer of DBT cells (a continuous murine astrocytoma cell line), as previously described (Fleming et al., 1986). Briefly, DBT cells were grown in RPMI supplemented with 7% new born calf serum (NBC) and 10% TPB in 20mm petri dish. After 72 hours, confluent monolayer of DBT cells were washed once with RPMI supplemented with TPB and infected with 200 µl of brain supernatant, either undiluted or at 10, 100 or 1000 fold dilutions at various time p.i. After 90 minutes incubation, cells were overlaid with 0.6% agarose in RPMI supplemented with 10% TPB. Plaques were counted after 48 hour incubation at 37°C.

7.5 Flow Cytometry

Cells isolated from CNS, Spleen and CLN were incubated with the 10% mouse serum and anti-mouse CD16/32 (clone 2.4G2; BD PharMingen, San Diego, CA) for 15 minutes on ice to block non-specific binding (Kapil et al., 2009). Cells were stained with flurochrome conjugated mAb as described in Table 7.1.
To detect JHMV-specific CD8 T cells, cells were stained with PerCP conjugated anti-CD8 and PE conjugated tetramer H-2D<sup>b</sup>-S510. To study the expression of MHC on infiltrating macrophages and on resident brain microglia, cells were stained with the anti-CD45 and F4/80 (Serotec, Raleigh, NJ). CD45<sup>high</sup>-F4/80 positive cells were distinguished as infiltrating macrophages, while CD45<sup>low</sup>-F4/80 positive cells were distinguished as microglia.

Table 7.1

Antibodies Utilized in Flow Cytometry

<table>
<thead>
<tr>
<th>Antibody</th>
<th>Clone</th>
<th>Fluorochrome</th>
<th>Vendor</th>
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<tbody>
<tr>
<td>CD4</td>
<td>RM4-5</td>
<td>FITC</td>
<td>BD Pharmingen</td>
</tr>
<tr>
<td>CD8</td>
<td>53-6.7</td>
<td>PE</td>
<td>BD Pharmingen</td>
</tr>
<tr>
<td>CD45</td>
<td>30-F11</td>
<td>APC, PerCP</td>
<td>BD Pharmingen</td>
</tr>
<tr>
<td>MHC Class II</td>
<td>2G9</td>
<td>PE</td>
<td>BD Pharmingen</td>
</tr>
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<td>1A8</td>
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</tr>
<tr>
<td>F4/80</td>
<td>C1:A3-1</td>
<td>APC, PE</td>
<td>Serotec</td>
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<tr>
<td>O4</td>
<td>O4</td>
<td>Na&lt;sup&gt;+&lt;/sup&gt;</td>
<td>In house</td>
</tr>
<tr>
<td>H-2D&lt;sup&gt;b&lt;/sup&gt;-S510 tetramer</td>
<td>-</td>
<td>PE</td>
<td>Beckman-Coulter</td>
</tr>
</tbody>
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7.6 Purification of Cells by Flow Activated Cell Sorting (FACS)

To identify infiltrating macrophages and CNS resident microglia, cells were stained with as described previously in section 6.5 with mAb for the surface receptor for macrophage, microglia and oligodendroglia. Infiltrating macrophages were characterized by their CD45<sup>hi</sup>/F4/80<sup>+</sup> phenotype, while CD45<sup>low</sup>/F4/80<sup>+</sup> cells
were distinguished as microglia. Mature oligodendroglia, were isolated utilizing transgenic mice (PLP-GFP/B6) that express green fluorescence protein (GFP) under control of an oligodendroglia specific promoter for proteolipid protein, PLP, to (Fuss et al., 2000) in conjunction with CD45 surface staining. Oligodendroglia from IFNAR-/- and Wt control mice were isolated utilizing CD45 and O4 staining. O4 antibody from hybridoma supernatant was obtained from Dr. Stephen Stohlman or Dr. Bruce Trapp's lab. It was titrated for use against cells isolated from PLP-GFP mice and detected with secondary anti-mus IgM (PE). Total protein concentration ranged from 0.1-0.5mg/mL. Staining strategy for specific cell type is described in Table 7.2. Following staining cells were maintained in 1X FACS buffer on ice for sorting. Purified cells were pelleted by centrifugation 400g for 7 minutes at 4°C to remove sheath fluid. Cells were resuspended in trizol and stored at -80°C. Typical yields for 6-7 naïve spinal cords were 40,000-80,000 oligodendroglia and 400,000-600,000 microglia.

Table 7.2

Classification of Cell Types for Identification by Flow

<table>
<thead>
<tr>
<th>Cell type</th>
<th>Phenotypic classification</th>
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</thead>
<tbody>
<tr>
<td>Mature oligodendroglia</td>
<td>CD45-ve GFP+ve (using PLP-GFP mice)</td>
</tr>
<tr>
<td>Oligodendroglia</td>
<td>CD45-ve O4+ve</td>
</tr>
<tr>
<td>Microglia</td>
<td>CD45lo</td>
</tr>
<tr>
<td>Macrophage</td>
<td>CD45hi CD11b+ve F4/80+ve</td>
</tr>
</tbody>
</table>

* O4 antibody from hybridoma supernatant was obtained from Dr. Stephen A. Stohlman.
7.7 Correlation between Cell Number and Ct Value

The CNS cell isolation procedure routinely yielded higher numbers of microglia and macrophages compared to oligodendroglia (see section 6.6). To validate that cell numbers were not a factor limiting detection of low abundance mRNA, we evaluated the minimum number of cells needed to achieve reliable PCR products by sorting limiting numbers of brain microglia (CD45lo CD11b+) from mice inoculated with the synthetic dsRNA mimic poly (I:C) 4h prior to isolation. Naïve brains were excluded as *Ifnα/β* transcripts were too close to detection thresholds. RNA was isolated from as few as 7500 cells up to 240,000 cells and analyzed for relative expression of *Gapdh, Ifnα4, and Ifnβ. Ifit1 and Ifit2* transcripts served as representative ISG. Titration curves show that the Ct values for all the genes were in the linear range to *Gapdh* Ct values (Suppl. Figure 3). *Ifnα4* transcripts that are expressed at low copy number were still in the linear range and reliably detected even below 30,000 cells. In the present study, a minimum of 30,000-40,000 cells obtained from cell preparations of 6-8 mice were thus used for qPCR analysis.

7.8 Ex Vivo Stimulation Of Splenocytes

Spleens were isolated 7 day p.i. from mice injected intraperitoneally with MHV DMN strain at 500 PFU. Splenocytes were isolated as described previously. To bias composition of spleen cells towards CD4 T cells, splenocytes were cultured in RPMI medium 1640 supplemented with 10% FCS, 1% L glutamine, 1% nonessential amino acids, 1% sodium pyruvate, .01% gentamycin and β-mercaptoethanol with CD4 specific peptide M133 for 5 days. Stimulated cells
were utilized to determine ability of these cells to produce intracellular cytokines such as IFNγ and IL10.

7.9 Intracellular Detection of Cytokines

Virus-specific production of IFNγ, TNFα and IL17 was evaluated in CNS-, spleen- and CLN-derived CD8 and CD4 T cells. 2x10^6 splenocytes and CLN-derived cells, and 5x10^5 CNS-derived cells were incubated with 3x10^5 El-4 or CHB3 feeder cells with or without JHMV-specific peptide for the stimulation of CD8 or CD4 T cells. Cells were stimulated either with or without 1μM S510 (CD8-epitope) or 5 μM M133 (CD4-epitope) peptide/ml in a total volume of 200μl of RPMI supplemented with 20% fetal calf serum (FCS) for 5 hr at 37°C with 1μl/ml Golgi Stop (BD PharMingen, San Diego, CA). Cytokine production by stimulated cells was detected by intracellular flow cytometry, as previously described (Kapil et al., 2009, Zuo et al., 2006). Briefly, cells were stained for surface expression of CD8, CD4 and CD45, fixed and then permeablized using the Cytofix/Cytoperm kit (BD PharMingen) according to the manufacturer’s protocol. Expression of intracellular cytokine production was detected using FITC conjugated mAb specific for IFNγ, PE conjugated IL17 mAb and APC conjugated IL10 Ab. Cells were analyzed with FACS Calibur flow cytometer (Becton Dickinson, Mountain View, CA) using FlowJo software (TreeStar, Inc., Ashland, OR).

7.10 Phagocytic Assay For Macrophage And Microglia

*Escherichia coli* (K-12 strain) BioParticles®, Alexa Fluor® 488 (invitrogen) conjugate were used as particle tracers to analyze phagocytosis by infiltrating macrophage and resident microglia cells from CNS of wt mice. Briefly,
mononuclear cells from brain were incubated along with bioparticles. Before using, particles were incubated with opsonizing reagent (invitrogen) in 1xPBS for 1 hour at 37°C and then washed twice with 1XPBS. Cells were incubated with beads for 1 hour at 37°C in 5% CO2 incubator. Cells were then washed 3X to remove particles from surface and than blocked and stained for CD45 and F4/80 as described previously. To avoid nonspecific signal from beads bound to the surface of microglia/macrophages, cells were quenched with trypan blue for 5 minutes at ice and washed 3X to remove extra dye. Cells were analyzed with FACS Calibur flow cytometer (Becton Dickinson, Mountain View, CA) using FlowJo software (TreeStar, Inc., Ashland, OR).

7.11 ELISA

A sandwich enzyme-linked immunosorbent assay (ELISA)-based system previously described (Kapil et al., 2009) was used to measure concentrations of IFNy and IL10 protein in brain supernatants. Briefly, ELISA plates were coated with an anti-mouse cytokine capture antibody specific for either IFNy or IL10 (BD PharMingen, San Diego, CA) at 1μg/ml and incubated overnight at 4°C. The plates were washed (0.5% Tween-20 in PBS) and blocked with 10% FCS in PBS for 1 hr at room temperature and washed again. Brain supernatants collected at different time points post-infection were incubated in capture antibody-coated ELISA plates for 2 hr at room temperature. Bound cytokine was detected with a secondary biotinylated detection antibody specific for either IFNy or IL10 (BD PharMingen) combined with Avidin-Horseradish Peroxidase (Av-HRP) and incubated for 1 hr at room temperature. TMB (BD PharMingen) was used as
colorometric substrate and the reaction was stopped using sulfuric acid (H₂SO₄). Absorbance levels were measured at 450 nm using a microplate reader (Bio-Rad) and SoftMax Pro software (Molecular Devices, Sunnyvale, CA). Cytokine levels were calculated using a standard curve of known concentrations of commercially available recombinant IFNγ or IL10 protein. Triplicate samples were averaged and standard error was calculated. Data are presented as ng of protein/brain ± SE.

7.12 Histology and Fluorescent Staining

Spinal cords were fixed in 10% formalin and embedded in paraffin. 10 μm sections were stained with hematoxylin and eosin or luxol fast blue to evaluate inflammation and myelin loss, respectively. The distribution of viral antigen was determined by immunoperoxidase staining (Vectastain-ABC kit; Vector Laboratory, Burlingame, CA), using the anti-JHMV mAb J.3.3 specific for the carboxyl terminus of the viral nucleocapsid protein as the primary Ab and horse anti-mouse as secondary Ab, and 3, 3'-diamionbenzidine substrate (Vector Laboratory). Sections were scored in a blinded manner for inflammation, demyelination, and viral antigen.

To identify accumulation of CD4 T cells within perivascular space, spinal cord tissues were isolated from perfused wt and PKR-/- mice as described previously (Ireland et al., 2009, Savarin et al., 2010). Tissues were snap frozen in liquid N2 in OCT and were stored at -80°C. Spinal cord sections were cut at 12μm thickness using microtome and stored at -80°C. Tissue sections were thawed and fixed with 4% PFA for 20 minutes each at room temperature.
Following permeabilization with 1% tritonX-100 at room temperature, nonspecific antibody binding was blocked using 1% bovine serum albumin and 10% goat serum. For perivascular accumulation of CD4 T cells, spinal cord sections were incubated at 4°C with mouse anti rat CD4 Ab and rabbit anti mouse laminin (Cedarline Laboratories, Ontario, Canada) overnight. Next day, tissue sections were washed and incubated with AF 594 conjugated anti rat and AF488 conjugated anti rabbit (Invitrogen) secondary antibodies at room temperature for 60 minutes. Following wash, sections were mounted with ProLong Gold antifade mounting media containing 4’,6-diamidino-2-phenylindole (Invitrogen, Carlsbad, CA). To further determine expression of PKR-p in CD4 T cells, spinal cord sections were incubated at 4°C with mouse anti rat CD4 Ab and rabbit anti mouse PKR-p (Cedarline Laboratories, Ontario, Canada) overnight and processed similarly as described previously.

7.13 Real Time PCR

Tissues were immediately frozen in liquid nitrogen and stored at -80°C. RNA was extracted from CNS tissues dissociated in TRIzol Reagent (Invitrogen, Carlsbad, CA) using sterile Tenbroeck glass homogenizers according to the manufacturer’s instructions. Briefly, after chloroform extraction, RNA was precipitated in isopropyl alcohol, washed with 75% ethanol and resuspended in RNase-free water (Gibco, Invitrogen, Grand Island, NY). To remove DNA contamination, RNA was treated with the DNase-treatment Kit (Ambion, Austin, TX), according to the manufacturer’s instructions. RNA integrity and concentration were evaluated by electrophoresis on a 1.2% formaldehyde
agarose gel. cDNA was synthesized using 2 ug of total RNA, Moloney Murine Leukemia Virus Reverse Transcriptase (Invitrogen), 10mM dNTP mix, and 20 uM random hexamer primers (Promega). Quantitative real-time PCR was performed for expression levels of cytokines and chemokines, anti-viral genes and anti-inflammatory genes was determined using either or SYBR Green chemistry or taqman chemistry (Applied Biosystems, Foster city, CA). List of genes is described in the table form. Expression was compared relative to the endogenous control glyceraldehyde-3-phosphate dehydrogenase (GAPDH) as previously described (Ireland et al., 2008). Reaction conditions for the 7500 Fast Real-Time PCR system (Applied Bio-system) for SYBR green were as follows: 95°C for 10 min and 40 cycles of denaturation at 94°C for 10s, annealing at 60°C for 30s and elongation at 72°C for 30s and for taqman were 95°C for 20 sec as holding temperature and then 40 cycles of denaturation at 95°C for .03s, annealing and extension at 60°C for 30s. All PCR were performed in 10 ul final reaction volumes consisted of specific master mix, 1mM of each primer, and 4 ul cDNA using the ABI 7500 fast PCR machine and 7500 software. Data presented reflect the induction level (n fold) relative to GAPDH, using the following formula,

\[(2^{\text{CT(GAPDH)} - \text{CT(Target)}}) \times 1000\]  

where Ct is the threshold cycle.
Table 7.3
List of Primer Sequences/Catalog Number for Genes Utilized in the Study

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<tr>
<th>Gene Name</th>
<th>Primer sequence/ Catalog #</th>
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<tr>
<td><strong>House keeping gene</strong></td>
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<tr>
<td>1. Glyceraldehyde 3-phosphate dehydrogenase (Gapdh)</td>
<td>F, 5’-TGACCACCAACTGCTTAG-3’</td>
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<tr>
<td></td>
<td>R, 5’-GGATGCAGGGATGATGTTC-3’</td>
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<tr>
<td><strong>JHMV primers</strong></td>
<td></td>
</tr>
<tr>
<td>1. Nucleocapsid gene of John Howard Muller strain of coronavirus (JhmvN)</td>
<td>F, 5’-GAGTGCCTATTGCCAATGGAA-3’</td>
</tr>
<tr>
<td></td>
<td>R, 5’-TTAAAGGAACGTCGGTTGTGTCT-3’</td>
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<tr>
<td><strong>Cytokines</strong></td>
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<tr>
<td>1. Interleukin-1β (Il1β)</td>
<td>Mm01336189_m1</td>
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<tr>
<td>2. Interleukin-1 receptor antagonist (Il1Ra)</td>
<td>F, 5’-AGATAGACATGGTGCCTATTGACCTT-3’</td>
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<td></td>
<td>R, 5’-CATCTCCAGACTTGGCACAAGA-3’</td>
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<tr>
<td>3. Interleukin-6 (Il6)</td>
<td>F, 5’-ACACATGTTCCTGGGAAATCGT-3’</td>
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<td>R, 5’-AAGTGCATCATTGGTTGCTATAC-3’</td>
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<tr>
<td>4. Interleukin-10 (Il10)</td>
<td>F, 5’-TTTGAATTCCCTGGGTTGAGAA-3’</td>
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<tr>
<td></td>
<td>R, 5’-GCTCCACTGCTTGCTTTATT-3’</td>
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| 5. | Interleukin-12p35 (Il12p35) | F, 5’-CACCCTTGCCCTCCTAAACC-3’  
R, 5’-CACCTGGCAGGTCCAGAGA-3’ |
| 6. | Interleukin-12p40 (Il12p40) | F, 5’-ACAGCACCAGCTTTCATCAG-3’  
R, 5’-CTTCAAAGGGCTTCATCTGCAA-3’ |
| 7. | Interleukin-13 (Il13) | F, 5’-TGACCAACATCTCCAATTGCA-3’  
R, 5’-TGTGTTAAAGTGTCGCTACTTGGGATTT-3’ |
| 8. | Interleukin-21 (Il21) | F, 5’-GGCACAGTATAGACGCTACGAATG-3’  
R, 5’-CCTTCACTTCTCCACTTGCA-3’ |
| 9. | Interleukin-23p19 (Il23p19) | F, 5’-CAGCAGCTCTCTCGGAAT-3’  
R, 5’-ACAACCATCTTACACTGGATACG-3’ |
| 10. | Interferon-α4 (Ifnα4) | Mm00833969_s1 |
| 11. | Interferon-β (Ifnβ) | Mm00439546_s1 |
| 12. | Interferon-γ (Ifnγ) | Mm01168134_m1 |
| 13. | Tumor necrosis factor-α (Tnfα) | F, 5’-GCCACCACGCCTTTCGTTCT-3’  
R, 5’-GGTCTGGGGCCATAGAATCGATG-3’ |
| 14. | Transforming growth-factor-β1 (Tgfβ1) | F, 5’-CCCGAAGCGGACTACTATGC-3’  
R, 5’-CGAATGTCTGACGTATTGAAGAACA-3’ |
| 15. | Transforming growth-factor- β3 | F, 5’-CAATTACTGCTTCCGCAACCT-3’  
R, 5’-AGCACCCTGCTATGGGTGTTG-3’ |
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<th>(Tgfβ3)</th>
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<td><strong>Chemo-kines</strong></td>
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<tr>
<td>1. Chemokine (C-C motif) ligand 2 (Ccl2)</td>
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<tr>
<td>F, 5’-AGCAGGTGTCCTCAAGAA-3’</td>
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<tr>
<td>R, 5’-TATGTCTGGACCATTCCCT-3’</td>
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<tr>
<td>2. Chemokine (C-C motif) ligand 5 (Ccl5/Rantes)</td>
</tr>
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<td>F, 5’-GCAAGTGCTCAATCTTGCA-3’</td>
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<tr>
<td>R, 5’-CTTCTCTGGGTGTGCACACA-3’</td>
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<td>3. Chemokine (C-X-C motif) ligand 9 (Cxcl9/Mig)</td>
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<td>F, 5’-GACGGTCCGCTGCAACTG-3’</td>
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<tr>
<td>R, 5’-GCTTCCCTATGGCCCTCATT-3’</td>
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<tr>
<td>4. Chemokine (C-X-C motif) ligand 10 (Cxcl10/Ip10)</td>
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<td>F, 5’-TGCACGATGCTCCTGCA-3’</td>
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<tr>
<td>R, 5’-AGGTCTTTGAGGGATTTGATG-3’</td>
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<tr>
<td><strong>Cell signaling enzymes</strong></td>
</tr>
<tr>
<td>1. Nitric oxide synthetase 2 (Nos2)</td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td>2. Arginase-1 (Arg1)</td>
</tr>
<tr>
<td>F, 5’ TGGGTGGATGCTCACAGA-3’</td>
</tr>
<tr>
<td>R, 5’-CAGGTTGCCCATGCAGATT-3’</td>
</tr>
<tr>
<td>3. Chitinase like protein (Ym1/2)</td>
</tr>
<tr>
<td>F, 5’-AATATCTGACGTTCTGAGGATAGA G-3’</td>
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<tr>
<td>R, 5’-GGCCAAACAGAAGAAATG-3’</td>
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<tr>
<td><strong>Anti-viral factors</strong></td>
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<td>1. Protein Kinase RNA dependent (Pkr)</td>
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<td>Mm00440966_m1</td>
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<td></td>
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<tr>
<td>---</td>
</tr>
<tr>
<td>1.</td>
</tr>
<tr>
<td>F, 5'-GACACCAGAATTCAAGGGAC-3'</td>
</tr>
<tr>
<td>2.</td>
</tr>
<tr>
<td>F, 5'-GTCAGCACAACCACAACC-3'</td>
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<tr>
<td>3.</td>
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<tr>
<td>Mm00442346_m1</td>
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<tr>
<td>2.</td>
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<tr>
<td>F, 5'-AAAAATGTCTGCTTCTTGAATTCTGA-3'</td>
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<td>3.</td>
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<tr>
<td>Mm00515153_m1</td>
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<td>4.</td>
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<tr>
<td>Mm00492606_m1</td>
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</table>
| 1. | Interferon regulatory factor-3  
    
    \( (Irf3) \) | Mm00516779_m1 |
| 2. | Interferon regulatory factor-7  
    
    \( (Irf7) \) | Mm00516788_m1 |
| 3. | Signal Transducers and Activators of Transcription  
    
    \( (Stat1) \) | Mm00439531_m1 |
| 4. | I-kappa-B kinase epsilon  
    
    \( (Ikk\varepsilon) \) | Mm00444862_m1 |

|   | Toll like receptor-3  
    
    \( (Tlr3) \) | Mm01207404_m1 |
|---|--------------------------------------------------------------------------------------|--|
| 5. | Toll like receptor-7  
    
    \( (Tlr7) \) | Mm00446590_m1 |
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subunit of the interferon alpha beta receptor that is required for signaling’, 


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alternative activation in macrophages and suppresses antituberculosis effector mechanisms without compromising T cell immunity', *Journal of Immunology*, vol. 183, pp. 1301-12.


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APPENDICES
<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Description</th>
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<tbody>
<tr>
<td>wt</td>
<td>Wild-type</td>
</tr>
<tr>
<td>IL</td>
<td>Interleukin</td>
</tr>
<tr>
<td>IL1Ra</td>
<td>Interleukin receptor antagonist</td>
</tr>
<tr>
<td>IFN</td>
<td>Interferon</td>
</tr>
<tr>
<td>Ko</td>
<td>Knock-out</td>
</tr>
<tr>
<td>IP</td>
<td>Intra peritoneal</td>
</tr>
<tr>
<td>IC</td>
<td>Intracerebral</td>
</tr>
<tr>
<td>p.i.</td>
<td>Post-infection</td>
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<tr>
<td>NFkB</td>
<td>Nuclear factor kappa beta</td>
</tr>
<tr>
<td>JHMV</td>
<td>John Howard Muller strain of coronavirus</td>
</tr>
<tr>
<td>PKR</td>
<td>Protein Kinase RNA dependent</td>
</tr>
<tr>
<td>OAS-2</td>
<td>2'5' Oligoadenylate synthetase</td>
</tr>
<tr>
<td>STAT</td>
<td>Signal transducer and activator of transcription</td>
</tr>
<tr>
<td>IFIT</td>
<td>Interferon-induced protein with tetratricopeptide repeats</td>
</tr>
<tr>
<td>SC</td>
<td>Spinal cord</td>
</tr>
<tr>
<td>CD</td>
<td>Cluster of differentiation</td>
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<tr>
<td>mAb</td>
<td>Monoclonal Antibody</td>
</tr>
<tr>
<td>MHC</td>
<td>Major Histocompatibility Complex</td>
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<td>TNF</td>
<td>Tumor necrosis factor</td>
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<tr>
<td>PRR</td>
<td>Pathogen recognition receptor</td>
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<tr>
<td>PAMP</td>
<td>Pathogen associated molecular pattern</td>
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<td>Abbreviation</td>
<td>Full Name</td>
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<td>MDA-5</td>
<td>Melanoma Differentiation-Associated Gene 5</td>
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<td>Retinoic acid inducible gene-I</td>
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<td>TLR</td>
<td>Toll like receptor</td>
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<td>Interferon regulatory factor</td>
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<td>Interferon stimulated genes</td>
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<td>CoV</td>
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<td>West nile virus</td>
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<td>John cunningham virus</td>
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<td>EAE</td>
<td>Experimental autoimmune encephalitis</td>
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<tr>
<td>MS</td>
<td>Multiple sclerosis</td>
</tr>
<tr>
<td>Poly I:C</td>
<td>Polyinosinic-polycytidylic acid</td>
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