Rift Valley fever virus inhibits a pro-inflammatory response in experimentally infected human monocyte derived macrophages and a pro-inflammatory cytokine response may be associated with patient survival during natural infection

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A R T I C L E   I N F O

Rift Valley fever virus (RVFV) causes significant morbidity and mortality in humans and livestock throughout Africa and the Middle East. The clinical disease ranges from mild febrile illness, to hepatitis, retinitis, encephalitis and fatal hemorrhagic fever. RVFV NSs protein has previously been shown to interfere in vitro with the interferon response, and RVFV lacking the NSs protein is attenuated in several animal models. Monocytes and macrophages are key players in the innate immune response via expression of various cytokines and chemokines. Here we demonstrate that wild-type RVFV infection of human monocyte-derived macrophages leads to a productive infection and inhibition of the innate immune response via decreased expression of IFN-α, IFN-β and TNF-α. Using a recombinant virus lacking the NSs protein, we show that this effect is mediated by the viral NSs protein. Finally, analysis of RVF patient samples demonstrated an association between a pro-inflammatory cytokine response and patient survival.

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Background

Rift Valley fever virus (RVFV) is a mosquito-borne hemorrhagic fever virus that causes high morbidity and mortality in humans and livestock. It was first identified in 1931 in Kenya after isolation from a sheep in the Rift Valley (Daubney et al., 1931). The virus has caused disease throughout continental Africa, Madagascar, Yemen and Saudi Arabia (Bird et al., 2009). Recent reports of mosquito vector capacity in North America make this virus not only a scourge on the developing world, but also a potential threat to the US (Turell et al., 2010).

RVFV is a veterinary pathogen that infects cattle, goats, and sheep. Up to 90% mortality has been reported in newborn animals and as high as 30% in adult animals (Swanepoel and Coetzer, 1994). Consistent with its high degree of pathogenicity in juvenile animals, RVFV is also abortigenic; 40–100% of pregnant animals will abort during an outbreak leading to “abortion storms” (Daubney et al., 1931; Swanepoel and Coetzer, 1994). Furthermore, livestock caretakers are exposed to virus in the process of caring for sick and dying animals; both blood and amniotic fluid contain high quantities of virus.

The virus can be transmitted to humans by contact with infected livestock or by the bite of an infected mosquito. Infected individuals typically have a mild disease consisting of fever, malaise, and myalgia. A small percentage of individuals will develop severe disease manifested as hepatitis, encephalitis, retinitis or hemorrhagic fever, which are the hallmarks of fulminant RVFV clinical disease. The overall case fatality is estimated at 0.5–1%. However, in patients whose clinical illness is sufficiently severe to bring them to the attention of medical personnel, case fatality has been reported to be as high as 29%, as was seen in the Kenya 2006–2007 outbreak (Centers for Disease Control and P., 2007). Laboratory findings that are frequently present in RVFV infected patients include leukopenia, thrombocytopenia, and elevated liver transaminases, indicative of the hepatitis that is often associated with infection.

RVFV is a member of the family Bunyaviridae. It is an enveloped virus that has a negative stranded RNA genome consisting of three fragments, aptly named S (small), M (medium), and L (large). The S segment encodes two proteins, a nucleocapsid protein that coats the viral genome in the virion, and a non-structural protein (NSs). The M segment encodes two viral glycoproteins that are expressed on
the surface of the virion, and a nonstructural protein (NSm). The L segment encodes the viral RNA polymerase that is responsible for both transcription and replication of the virus (Fields et al., 2007).

The NSs protein is especially interesting in that it is a filamentous nuclear protein expressed by a virus that replicates and assembles in the cytoplasm of infected cells (Yadani et al., 1999). Several investigators have evaluated the role of the NSs protein in altering the host immune response. Initial studies utilized a naturally occurring variant that has a deletion in the S segment such that the NSs protein is truncated, cytoplasmic and rapidly degraded (Muller et al., 1995; Vialat et al., 2000). This variant, known as clone 13, was attenuated in wild type (WT) mice but lethal in IFN α/β receptor deficient mice, and was a potent inducer of Type 1 interferons, unlike the WT virus (Billecocq et al., 2004; Bouloy et al., 2001; Vialat et al., 2000). Clone 13 has also been shown to be immunogenic and protective in sheep (Dungu et al., 2010). More recently, a reverse genetics system has become available for RVFV, thereby facilitating studies of viral pathogenesis (Gerrard et al., 2007; Habjan et al., 2008b; Ikekami et al., 2006). This system has been used to generate viruses with full gene deletions in NSs or mutations of specific regions of the gene. Viruses with whole gene deletions have become live attenuated vaccine candidates since they provided protection in the Wistar–Furth rat model (Bird et al., 2008). This reverse genetics system has proven to be a powerful tool in the study of NSs-mediated pathogenesis.

At the molecular level, the NSs protein interacts with components of the general transcription factor, TFIIF, leading to a generalized down-regulation of host-cell transcription in infected cells (Le May et al., 2004). In addition, specific interactions of NSs with transcription factors YY1 and SAP30 lead to silencing of the IFN-α promoter in mouse fibroblasts (Le May et al., 2008). Via its SAP30 interacting domain, NSs also interacts with pericentromeric chromosomal sequences and causes chromosomal segregation defects in mouse fibroblasts and fetal sheep kidney cells (Mansurogulli et al., 2010). NSs facilitates proteasomal-mediated degradation of PKR, a protein that is important in sensing the presence of dsRNA, shutting down protein synthesis and signaling apoptosis in infected cells (Habjan et al., 2009; Ikekami et al., 2009). The NSs protein has multiple functions in alteration of the innate immune response: generalization and specific transcriptional down-regulation of genes active in innate immunity and targeted degradation of factors involved in the innate immune response. Given the varied spectrum of clinical illness resulting from infection there is clearly a wide array of factors involved in the innate immune response. It is noteworthy that although all experiments were performed with an moi of 5, for the experiments done with donors A and B, there appeared to be more viral RNA present at the 1 h time point for the ΔNSs infected cells. Despite the fact that there was slightly more input RNA, the WT virus still replicated to higher levels than the ΔNSs virus by 24 hpi (Fig. 3). These data demonstrate that primary human MDM are permissive for RVFV infection. Macrophages could modification of the initiation and/or functionality of the innate immune response in macrophages. To test this hypothesis, we infected human monocyte derived macrophages (MDM) with WT RVFV or recombinant RVFV lacking the NSs gene (ΔNSs RVFV) and assessed replication, virus production, cytopathic effects, and the expression of cytokines under these conditions. Furthermore, data from these experiments, and those done by others with different viral hemorrhagic fever viruses, led us to hypothesize that a pattern of cytokine secretion in infected individuals might be predictive of survival. This hypothesis was supported by analysis of the cytokine expression patterns in human serum samples from the Saudi Arabian 2000–2001 RVFV outbreak.

**Results**

**Infection of macrophages with RVFV**

CD14 positive MDM from 4 separate donors were used for these experiments to control for donor-to-donor variability. Cells were infected with either the WT or ΔNSs RVFV and supernatants were analyzed at various times post-infection by plaque assay to quantitate viral production. As indicated in Fig. 1, there was slight donor-to-donor variability in the maximal titers at 24 h post infection (hpi); however, WT virus grew to 0.5 to 1 log higher titers than the ΔNSs virus for each donor. Virus was detected as early as 12 hpi for both WT and ΔNSs viruses, indicating that the ΔNSs virus kinetics were not delayed compared to the WT virus (data not shown). In contrast, infection of Vero cells with WT or ΔNSs RVFV produces equivalent titers (1.5 × 10^7 PFU/mL and 4.6 × 10^7 PFU/mL respectively). Interestingly, by 48 hpi, there was 80–90% CPE with cell death in the MDM infected with WT virus and little to none in the MDM infected with the ΔNSs virus, and this effect could be seen as early as 12 hpi (Fig. 2). It is also of note in this figure that ΔNSs infected cells have an activated phenotype, being larger and having greater variability in morphology than the mock infected cells. WT infected cells are small and rounded up. By 72 hpi all WT infected cells were dead.

The RNA from WT or ΔNSs RVFV infected MDM from 3 of the donors was analyzed by real time RT-PCR to assess viral replication. Absolute Ct values were corrected by normalization to 18S RNA levels for each sample. Both WT and ΔNSs viruses replicated with similar kinetics; an increase in viral RNA was detected routinely by 12 hpi (Fig. 3). The 1 hpi time point represents the amount of input virus. It is noteworthy that although all experiments were performed with an moi of 5, for the experiments done with donors A and B, there appeared to be more viral RNA present at the 1 h time point for the ΔNSs infected cells. Since the liver and the CNS are main sites for RVFV mediated disease, it is possible that infection of macrophages could represent an important early target and a mechanism for viral spread. Given the known function of the NSs protein in alteration of the innate immune response, we hypothesize that a key component of RVFV pathogenesis is the initiation and/or functionality of the innate immune response in macrophages.
be an early and important in vivo target of infection. Given the known role of the NSs protein in immune modulation, we hypothesized that macrophages infected with the ΔNSs virus would exhibit a different pattern of cytokine secretion than those infected with the WT virus.

**Cytokine secretion in RVFV infected MDM**

MDM were mock-infected or infected with WT, ΔNSs, γ-WT or γ-ΔNSs virus. Gamma-irradiated (designated by the Greek letter “γ”) control viruses were used to distinguish non-specific cytokine secretion related to supernatant components, including inactivated virions, from those that were a result of active viral infection. Supernatants were collected at 0, 6, 12, 24 and 48 hpi and were analyzed for a selected panel of cytokines. A 12-plex panel of analytes including RANTES, MIP-1α, MIP-1β, IL-1RA, MCP-1, IP-10, IL-8, IFN-α2, TNF-α, IL-12, IL-1β, and IL-6 were examined on the Luminex platform and IFN-β levels were measured by ELISA. The most striking results were obtained for IFN-α2 (A), IFN-β (B), and TNF-α (C) (Fig. 4). IFN-α2 is an interferon alpha subtype that has potent antiviral activity in many different responder cell types (Hilkens et al., 2003; Hiscott et al., 1984). IFN-β is another Type 1 interferon that is known to inhibit viral replication and induce apoptosis of virally infected cells (Paul, 2008). TNF-α is a pro-inflammatory cytokine that plays a role in the activation of endothelial gene expression, activation of neutrophils and is a mediator of shock, sepsis, and vascular leakage (Paul, 2008). TNF-α, IFN-α2 and IFN-β were secreted by human MDM that were infected with ΔNSs virus but not by MDM that were infected with WT virus and only minimally by γ-irradiated viruses. In the ΔNSs virus infected MDM, expression of TNF-α was detectable as early as 6 hpi and IFN-α2 was detectable as early as 12 hpi; only the 24 h data is shown. IFN-β levels were only assessed at the 24 hpi time point. Unlike the data presented earlier for TNF-α, IFN-α2 and IFN-β, these cytokine patterns were not specific to cells infected with replicating virus. IL-8 and IL-1RA were not significantly elevated under any experimental condition except for donor D and donor B respectively, highlighting the importance of using multiple donors in experiments with primary cells (data not shown).

**Fig. 2.** WT virus causes marked CPE in infected cells while ΔNSs virus does not. MDM were mock-infected, or infected with WT, or ΔNSs RVFV. At 12 hpi cells were photographed under white light using the 20× objective to demonstrate the CPE caused by WT virus.

**Fig. 3.** ΔNSs RVFV replicates to lower levels than wild-type RVFV in MDM. MDM were infected with WT or ΔNSs RVFV. RNA was purified from cells at various times post infection and analyzed by real time PCR. WT virus (black squares with solid lines) replicated to higher levels than the ΔNSs virus (white squares with dotted lines) on cells from the same donor. Data are presented as inverse Ct value at various times post infection. 3 different donors are represented in the figure. RNA from the 4th donor was not available for testing.
for IFN-α and TNF-α were measured. 4 different donors are represented in the
lane. Aminotransferase, PT=prothrombin time, PTT=partial thromboplastin time, CI=con
noted laboratory value was not necessarily always available for each patient so the number of samples that were used in calculating the mean for the fatal and non-fatal cases re-
shown). Finally, there was no IL-12, IL-1β or IL-6 expression detected from any donor regardless of experimental condition.

Human serum cytokine analysis

After demonstrating that the macrophage is a susceptible cell type and that the virus is able to alter the innate immune response in macrophages, it followed that we might expect a perturbation of the cytokine response in severely infected individuals. We hypothesized that a suppression of the pro-inflammatory innate immune response by WT RVFV could play a role in viral pathogenesis. In order to test this hypothesis, we utilized human serum samples from the RVFV outbreak that occurred in Saudi Arabia in 2000–2001. The clinical and epi-
demiological data from his outbreak have been published (Madani et al., 2003). We were able to identify 26 samples from 26 different patients for which there was sufficient sample and for which the clinical outcome was known. Of the 26 cases, 6 were fatal and 20 were non-
fatal. All patients were hospitalized and exhibited fever and gastrointestinal symptoms (nausea, vomiting or diarrhea), 7 had jaundice, 3 had bleeding manifestations, 7 had CNS disturbances and none had vision changes. The samples were collected at the time of presentation and were from 1 to 14 days post onset of symptoms. There were 20 male and 6 female patients ranging in age from 17 to 90 years with an average age of 52 years. Key mean laboratory values in these patients are presented in Table 1. The patients all demonstrated clinical and labora-
tory findings typical for severe RVFV disease. The fatal cases had signif-
icantly more thrombocytopenia, coagulopathy and transaminase elevation, as has been previously reported (Madani et al., 2003).

Patient serum samples were analyzed in duplicate using a large multiplex assay to determine the concentration of 39 different cyto-
kines: EGF, Eotaxin, EGF-2, Flt-3 ligand, Fractalkine, G-CSF, GM-CSF, GRO, IFN-α2, IFN-γ, IL-10, IL-12 (p40), IL-12 (p70), IL-13, IL-15, IL-
17, IL-1RA, IL-1α, IL-1β, IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, IL-8, IL-9, IP-
10, MCP-1, MIP-3, MDC (CCL22), MIP-1α, MIP-1β, TGF-α, TNF-α, TNF-β, VEGF, sCD40L, and sIL-2R. There were no detectable levels of TNF-α or IFN-α2 in the samples from fatal or non-fatal cases. Howev-
er there were 5 cytokines that demonstrated a statistically significant difference (p<0.05) between fatal and non-fatal cases by a two-sample t-test (Fig. 5). Two pro-inflammatory cytokines, sCD40L (a mediator of B cell activation) and GRO (a mediator of neutrophil activation), were elevated in non-fatal cases as compared to fatal cases.

IL-1RA is the receptor antagonist for IL-1, a potent pro-inflammatory pyrogen, so therefore IL-1RA has immunosuppressive properties. IL-
1RA binds to the IL-1 receptor with high affinity and prevents receptor dimerization and downstream signaling (Paul, 2008). IL-1RA levels in fatal cases were a log higher than IL-1α levels. Additionally, IL-1RA levels were significantly higher in fatal vs non-fatal cases, leading to an overall immunosuppressive effect in fatal cases.

Finally, IL-10, a cytokine that is well known to be suppressive to the cell-mediated immune response, was elevated in fatal vs. non-fatal cases. In summary, a pro-inflammatory cytokine response was associat-
ed with increased survival while actively or passively suppressed cytokine response was associated with increased risk of fatality.

Discussion

Our studies have demonstrated that MDM are permissive for RVFV infection and that infection with WT virus leads to CPE and cell death.
Furthermore, we have studied the role of the NSs protein and determined that NSs deficient viruses do not replicate as well as WT RVFV in MDM. Since these two viruses replicate to equivalent levels in Vero cells, which are unable to produce interferon (Desmyter et al., 1968; Emeny and Morgan, 1979), it may be the case that the IFN response that is stimulated in ΔNSs RVFV infected MDM’s is responsible for the decrease in viral titers and the lack of CPE during infection with this virus.

Macrophages may play a role in the pathogenesis of WT RVFV. An infected macrophage would be unable to signal a pro-inflammatory response secondary to the inhibitory effects of the NSs protein. In addition, the intracellular anti-viral mechanisms would be rendered inactive because of NSs mediated inhibition of expression of type I IFN’s and virally mediated degradation of PKR. These many effects of the NSs protein could usurp a sentinel cell and convert it into a virus factory. The macrophage might also act as vehicle to transmit the virus to its target organs, the liver and the CNS. There is clearly precedence in the literature for viruses using the monocyte/macrophage to gain entry to the CNS in the case of Hepatitis C virus, Junin virus, Dengue virus, and HIV (Grap and Kaul, 2010; Koenig et al., 1986; Medeot et al., 1995; Miagostovich et al., 1997; Wilkinson et al., 2009). Further studies will need to be done to fully define the role of the macrophage in RVFV in vivo pathogenesis.

Our studies have demonstrated NSs-mediated inhibition of TNF-α, IFN-α2, and IFN-β expression in RVFV infected MDM’s. It was noted that several cytokines were activated by infection with gamma irradiated viruses. This non-specific activation (i.e., did not require viral gene expression or replication) could be secondary to the presence of viral RNA and/or protein in these inoculates or could represent activation by factors that were carried over in the supernatants during virus preparation. It is well known that surrogates for viral RNA such as poly I-C can activate a cytokine response in exposed cells, so these results were not surprising. However, it was quite striking that all of the non-specific activations were significantly diminished by the presence of the NSs protein in cells infected with WT virus. The NSs protein led to a striking, generalized down-regulation of all of the studied cytokines.

IFN-α2, IFN-β and TNF-α were elevated only in cells that were productively infected with ΔNSs virus. Expression of these cytokines required active viral transcription and/or replication. The RIG-I-like RNA helicases, RIG-I and MDA-5, are cytoplasmic viral RNA detector molecules that recognize ssRNA containing a 5′ triphosphate and dsRNA respectively (Paul, 2008). In one study, RIG-I recognized the 5′ triphosphate of a transfected RVFV genome and this led to downstream activation of the IFN-β promoter (Habjan et al., 2008a). This intracellular molecular sensor would be activated during ΔNSs infection of macrophages and initiate the signaling cascade that leads to IFN and NF-κB activation and Type 1 IFN and TNF-α transcription respectively. Our findings are consistent with previously published results that demonstrate the importance of the NSs protein in inhibition of Type 1 IFN’s in RVFV infected cells and animals. However, the finding of TNF-α inhibition by NSs is novel, and would be expected given our knowledge of the signaling mechanisms involved during viral infections.

Previous studies of other hemorrhagic fever viruses such as Ebola virus and CHF virus have demonstrated release of pro-inflammatory cytokines in vitro cell culture and in animal models and have reported an association between a pro-inflammatory response and increased fatality, when examining clinical specimens (Connolly-Andersen et al., 2009; Ergonul et al., 2006; Gupta et al., 2001; Hutchinson and Rollin, 2007; Papa et al., 2006; Stroher et al., 2001; Villinger et al., 1999). In contrast, Lassa virus appears to down-regulate the immune response; macrophages and dendritic cells are not activated by infection nor do they produce inflammatory cytokines when infected (Baize et al., 2004; Lukashevich et al., 1999). Additionally, activation of dendritic cells or macrophages by poly I-C, LPS or IFN-α prior to infection led to down-regulation of Lassa virus replication (Baize et al., 2006) and we have observed similar results with RVFV (data not shown). In animal studies done in cynomolgus macaques, survival from Lassa virus infection was associated with lower viral loads, faster antibody response, activation of an early type I IFN response, high activated monocyte counts and circulating activated T cells (Baize et al., 2009). In clinical samples from Lassa virus infected patients it has been reported that there are lower levels of the pro-inflammatory cytokines IL-8 and IP-10 in fatal cases (Mahanty et al., 2001). Taken together these data support the idea that a critical part of the pathogenesis of Lassa virus is preventing the activation of the immune response. It would follow that survival rates are higher in individuals who can activate this response despite virally mediated inhibition. The data that we have presented here using samples from human RVFV cases demonstrated a similar phenomenon,
where survival is associated with a robust pro-inflammatory cytokine response. Our data demonstrating an association between pro-inflammatory cytokines and human survival during RVFV infection is limited by the fact that we only have data for 6 fatal cases. Unfortunately, these types of samples are very difficult to obtain. However, it is striking that we found statistical significance given that our samples were obtained from 1 to 14 days post onset of symptoms. There was no significant difference between the time of presentation of the fatal vs nonfatal cases, (mean of 4.8 days vs 3.75 days; p = 0.41) lending even more credence to our data. All of our cases clearly represented severe disease since they came to the attention of medical personnel and exhibited derangement in their laboratory parameters. We suspect that if we were able to obtain data from mild cases and compare them to severe cases, the cytokine effects that we have seen would be even more pronounced.

Elucidating the factors that determine why some patients are able to mount a pro-inflammatory response and survive while others do not remains an area for future study. Genetic heterogeneity as the basis of differential susceptibility to RVFV infection has been well established in the rat and mouse model (Anderson et al., 1987; Anderson et al., 1991; do Valle et al., 2010; Peters and S lone, 1982; Ritter et al., 2000). Variable expression of interferon regulated genes were demonstrated recently in mouse embryo fibroblasts (MEFs) from BALB/cByJ mice versus the more susceptible MBT/Pas mouse (do Valle et al., 2010). In this study, the authors also report increased expression of Ifnβ1 and Ifnα4 transcripts upon infection of MEFs with a ΔNSs virus as compared to a WT virus. They were examining RNA at very early time points in infection (~9 h) and this might explain why they were able to see some expression of Ifn transcripts in cells infected with WT virus. The known heterogeneity of response to infection with RVFV in humans and animals is consistent with the heterogeneity of response that we saw amongst our four donors. One might predict that donor B, the donor with the highest cytokine levels in our study, would have a better outcome upon infection with RVFV than the other donors.

In reality, it is most likely that a combination of genetic and environmental factors are responsible for disease outcome. While we cannot rule out a specific genetic predisposition to fatal disease in a small proportion of the population, it is more likely that a person’s immune status at the time of infection (e.g. concurrent infections, nutritional status, stress level, etc.) is responsible for the lack of response that leads to a fatal outcome. We would predict that early and vigorous medical intervention, possibly targeting specific virulence factors, such as NSs, could significantly improve disease outcomes by maximizing the response potential of any given human genotype to viral infection.

**Materials and methods**

**Virus and cells**

All work with live virus was performed under BSL-4 conditions in a positive pressure suit. RVFV ZH501 (Bird et al., 2007b) or RVFV ΔNSs (Bird et al., 2008) were propagated in Vero E6 cells by infecting at an MOI of 0.1. Supernatants were collected 3 or 4 days post infection, clarified by centrifugation, aliquoted, and stored at −80 °C. Peripheral blood mononuclear cell pheresis products were obtained from healthy human donors at Emory Hospital. Pheresis products were diluted 1:1 with PBS (without calcium or magnesium), layered onto Histopaque (Sigma) or Ficoll-Paque (GE Healthcare), and mononuclear cells were purified per the manufacturer’s instructions. After purification, cells were washed several times in PBS and resuspended in MACS buffer (Miltenyl Biotech). Magnetically coupled CD14 antibodies (Miltenyl Biotech) were used to selectively purify the CD14 positive cells per the manufacturer’s instructions. CD14 positive cells were stored at −80 °C in freezing medium (90% FBS, 10% DMSO) until use.

**Infections**

CD14 positive cells were seeded onto 24 well plates in complete media (RPMI with 5% FBS, 100 U/mL of penicillin, 100 μg/mL of streptomycin, and 2 mM l-glutamine), and allowed to mature to macrophages by 5 days of adherence in culture. Cells were re-fed with fresh media every 2 days. Cells were then infected at an MOI of 5 with either ZH501, ΔNSs, gamma irradiated viruses (γ-ZH501 or γ-ΔNSs) that were inactivated by irradiation with 5×10⁶ rads, or mock infected with conditioned media. After allowing 1 h for adsorption, the inoculum was removed, cells were washed 3× with PBS, and then re-fed with complete media. At defined times post infection, supernatants were collected, centrifuged to pellet any debris, and stored at −80 °C for future analysis. Cells were lysed in NA lysis buffer (ABI) for RNA purification and stored at −80 °C until purification was performed.

**Plaque assays**

Vero E6 cells were plated onto 6 well plates at a density of 70%. The following day, supernatants were diluted serially in complete media and 200 μl of each dilution was placed per well in duplicate. Inocula were allowed to adsorb for 1 h with rocking every 15 min to prevent drying. Each well was then overlaid with 3 mL of overlay media (0.6% Seakem ME agarose, 1× EMEM, 10% FBS, 100 U/mL of penicillin, 100 μg/mL of streptomycin, and 2 mM l-glutamine) and incubated at 37 °C. After 3 days, cells were fixed in 10% formalin, agarose was removed and monolayers were stained with crystal violet and washed in PBS. Plaques were counted on a white light trans-illuminator.

**Real time PCR**

RNA was purified from cells that had been lysed in NA lysis buffer (ABI) according to the manufacturer’s instructions. Ten microliters of total RNA was used for an 18S assay (ABI) that allowed for normalization between samples. Twenty microliters of RNA was used in a RVFV assay that has been previously described (Bird et al., 2007a). Reactions were performed on an ABI 7500 real time PCR machine.

**Cytokine assays**

MDM-culture supernatants were gamma irradiated (5×10⁶ rads) to inactivate infectious materials prior to cytokine analysis. Cytokine assays were performed in duplicate according to the manufacturer’s instructions (Millipore-Milliplex MAP Kit) and analyzed on a Luminex 200 IS platform.

**ELISA**

The IFN-β ELISA was performed on the same supernatants that were used in the Luminex assays and according to manufacturer’s instructions (Invitrogen).

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