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**Recommended Citation**

Gitlin, Leonid; Benoit, Loralyn; Song, Christina; Cella, Marina; Gilfillan, Susan; Holtzman, Michael J.; and Colonna, Marco, "Melanoma differentiation-associated gene 5 (MDA5) is involved in the innate immune response to Paramyxoviridae infection in vivo." PLoS Pathogens. 6, 1. e1000734. (2010).  
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Melanoma Differentiation-Associated Gene 5 (MDA5) Is Involved in the Innate Immune Response to Paramyxoviridae Infection In Vivo

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Abstract

The early host response to pathogens is mediated by several distinct pattern recognition receptors. Cytoplasmic RNA helicases including RIG-I and MDA5 have been shown to respond to viral RNA by inducing interferon (IFN) production. Previous in vitro studies have demonstrated a direct role for MDA5 in the response to members of the Picornaviridae, Flaviviridae and Caliciviridae virus families (+) ssRNA viruses but not to Paramyxoviridae or Orthomyxoviridae (−) ssRNA viruses. Contrary to these findings, we now show that MDA5 responds critically to infections caused by Paramyxoviridae in vivo. Using an established model of natural Sendai virus (SeV) infection, we demonstrate that MDA5−/− mice exhibit increased morbidity and mortality as well as severe histopathological changes in the lower airways in response to SeV. Moreover, analysis of viral propagation in the lungs of MDA5−/− mice reveals enhanced replication and a distinct distribution involving the interstitium. Though the levels of antiviral cytokines were comparable early during SeV infection, type I, II, and III IFN mRNA expression profiles were significantly decreased in MDA5−/− mice by day 5 post infection. Taken together, these findings indicate that MDA5 is indispensable for sustained expression of IFN in response to paramyxovirus infection and provide the first evidence of MDA5-dependent containment of in vivo infections caused by (−) sense RNA viruses.

Introduction

Innate pathogen sensors detect viral products and respond by initiating a signaling cascade that leads to rapid anti-viral response involving secretion of type I IFNs (i.e. IFN-α and IFN-β) and inflammatory cytokines (i.e. IL-6 and TNF-α) [1]. In particular, type I IFNs restrict infection by inhibiting viral replication within cells and by stimulating the innate and adaptive immune responses. Once induced, secreted IFN-α and IFN-β bind to the IFNα receptor on the cell surface in an autocrine or paracrine manner. Activation of this receptor initiates the JAK/STAT signal transduction pathways [2,3] and the expression of IFN-inducible genes [4]. These gene products increase the cellular resistance to viral infection and sensitize virally-infected cells to apoptosis [5]. In addition, type I IFNs directly activate DC and NK cells, and promote effector functions of T and B cells, thus providing a link between the innate response to infection and the adaptive immune response [6,7].

Several viral sensors have been identified that belong to the Toll-like receptor (TLR) and RIG-I like receptor (RLR) families [8]. TLRs are expressed on the cell surface and/or in endosomal compartments [9]. TLR3 recognizes double stranded RNA (dsRNA), a molecular pattern associated with replication of single stranded RNA (ssRNA) viruses as well as the genomic RNA of dsRNA viruses [10]. TLR7 and TLR8 recognize ssRNA [9,11,12], whereas TLR9 recognizes unmethylated CpG-containing DNA [13]. RLRs are cytoplasmic proteins that recognize viral nucleic acids that have gained access to the cytosol [14–19]. The RLR family consists of three known members: retinoic acid-inducible gene I (RIG-I), melanoma differentiation-associated gene 5 (MDA5), and LGP2. RIG-I and MDA5 both contain a DExD/H box helicase domain that binds dsRNA, a C-terminal domain and two N-terminal caspase recruitment domains (CARDs) that are involved in signaling [8,17,20,21]. LGP2 contains a helicase domain but lacks the CARDs, and its precise contribution to antiviral signaling remains ambiguous [17,22].

Though RIG-I and MDA5 share common downstream signaling via activation of IPS-1 (also called MAVS, VISA or Cardif) and IRF3 [23–26], these helicases exhibit distinct substrate specificity. In this regard, RIG-I has been shown to preferentially...
Author Summary

The innate immune system possesses an array of sensory molecules which are purposed in detecting viral nucleic acids. Our understanding of how these molecular sensors detect viral nucleic acids continues to evolve. Herein, we demonstrate that MDAs, a member of the RIG-I-like receptor family, is involved in the detection of paramyxovirus infection in vivo. Specifically, MDAs appears to trigger antiviral cytokines that inhibit paramyxovirus replication. In this regard, mice that are deficient in MDAs are unable to express sustained levels of these cytokines and thus succumb to extensive viral propagation and disease. Our findings are largely discordant from previous in vitro studies using cultured cells, where it has been shown that RIG-I and not MDAs is involved in the innate response to negative sense RNA viruses. Thus, our data provides strong evidence of MDAs-based detection of negative sense RNA viruses, and furthermore underscore the importance of organism-based analysis of the innate system.

Results

Infection with SeV causes increased morbidity and mortality in MDA5−−/− mice

In order to assess an in vivo role for MDA5 in containment of paramyxovirus infection, we infected MDA5−−/− mice with Sendai virus (SeV). Mice on a C57BL/6 (B6) background were selected for these experiments as the 129 strain is lethally susceptible to SeV at extremely low inocula [46], thus prohibiting assessment of loss of MDA5 function on this background. A dose of 200,000 pfu was administered to mice by intranasal delivery, an infection method that typically results in acute, non-lethal bronchiolitis in B6 mice. As a gross determinant of virus-induced morbidity, % body weight for infected WT and MDA5−−/− mice was monitored for 2 weeks post infection (PI). Though essentially identical % weight loss values were observed up until day 8 PI; onwards, weight loss in MDA5−−/− mice was significantly more severe (p<0.05) (Figure 1A). Correspondingly, histological analysis of lung sections obtained from day 12 PI MDA5−−/− mice revealed consolidation of the lung parenchyma as well as notable PAS-positive airway cells, an indication of mucus hyper-secretion (Figure 1B). Severe histopathology was not observed in the lung sections obtained from control mice at this time point. In addition, we compared survival following increasing inocula of SeV (Figure 1C). Though MDA5−−/− mice were not susceptible to the 200K pfu SeV dose, MDA5−−/− mice fully succumbed to 400K and 600K pfu SeV, between 9–14 days PI. In contrast, control mice were fully resistant to the 400K pfu dose, though 40% mortality was observed for controls infected with the 600K dose. Thus MDA5−−/− mice exhibit enhanced morbidity and susceptibility to SeV infection relative to control mice.

To more fully assess SeV susceptibility, we extended our analysis of the histological changes seen in the lungs of SeV-infected MDA5−−/− mice. H&E stained sections obtained from day 2 PI (not shown) and day 5 PI (Figure 2A) lungs demonstrated similar patterns of bronchiolitis, though peribronchial lymphoid cufing that formed in the lungs of control mice appeared moderately thicker and more densely populated than those of MDA5−−/− mice (Figure 2A). FACS analysis of lung-derived leukocytes at d2, d5, and d8 PI revealed no significant differences in lymphoid and myeloid subpopulations (neutrophils, cDC, macrophage and alveolar macrophage; data not shown). Significantly, at d5 and d8 PI, FACS analysis revealed equal relative numbers of lymphoid subpopulations (CD3+, CD19+ and NK1.1+); CD69 expression profiles on these subsets were comparable between strains (data not shown). By d9-PI, significant pathology was observed in the lungs of SeV-infected MDA5−−/− mice (Figure 2A), despite the fact that comparable numbers of SeV-specific CTL were generated in both strains at this time point (Figure 2B). Grossly, lungs dissected from SeV-infected MDA5−−/− mice exhibited enhanced areas of hemorrhage relative to control lungs (data not shown). Microscopic analysis revealed epithelial cells that were notably hyperplastic with abundant micropapillary projections. Additionally, severe bronchointerstitial pneumonia was observed, with alveolar walls adjacent to affected airways thickened and congested with chronic inflammatory cell infiltrates and hyperplastic type II pneumocytes, a lung injury pattern consistent with SeV susceptibility [46,47]. In comparison, sections obtained from control mice at these later time points exhibited moderate changes to the airway epithelium and mild interstitial infiltration (Figure 2A).
MDA5<sup>−/−</sup> mice demonstrate increased susceptibility to SeV propagation

As susceptibility to SeV infection correlates with increased viral burden [48], we next assessed viral replication in wild type and MDA5<sup>−/−</sup> mice using a combined approach of real-time PCR analysis and specific staining for SeV antigens. Initially, at d2 PI, IF staining of viral antigens in lung sections appeared comparable between the two strains. By d5 PI, SeV antigens exhibited restrained expression in the airways of control mice (Figure 3A top of panel). In contrast, the bronchioles of MDA5<sup>−/−</sup> mice remained notably positive for SeV antigens at this time point (Figure 3A bottom of panel). More striking however, was the observation that parenchyma tissues proximal to infected airways stained conspicuously for SeV antigens in MDA5<sup>−/−</sup> mice at d5 PI. In SeV resistant strains of mice, SeV infection is typically restricted to the mucociliary epithelium of the conducting airways, including the trachea, bronchi and bronchioles [49,50]. Viral replication that extends to the alveolar spaces is a feature commonly seen in susceptible strains of mice [51]. Accordingly, this pattern of infection supports a role for MDA5 in controlling the replication of SeV during in vivo infection.

To confirm this finding, we measured viral RNA levels from WT and MDA5<sup>−/−</sup> mice infected with 200K pfu SeV using real-time PCR analysis. Assessment was made using primer/probe sets designed to amplify SeV genome (3′ untranslated region) and SeV N gene (genomic and transcript) (Figure 3B and C). Using this strategy, an approximate 5 fold increase in SeV genome copy number/Gapdh mRNA was detected on d5 PI, though significant differences were also observed on d2 and d8 PI. Analysis of N gene revealed ~2 fold increase in expression on days 2 and 5, though there were no significant differences by d8 PI. Thus it appears that MDA5 contributes in part to the containment of SeV replication in vivo.

Cytokine response to SeV infection is altered in MDA5<sup>−/−</sup> mice

Though SeV is a potent inducer of type I IFN in the mouse, functioning via several distinct pathways, it possesses several mechanisms by which it can counteract the IFN response. Despite this property, induction of IFN expression [14,52], particularly type I and II, is critical in the containment of SeV infection in vivo as underscored by the profound SeV susceptibility seen for mice deficient in STAT1<sup>−/−</sup> mice [53]. As MDA5 is known to induce expression of type I IFN in vitro in response to polyI:C stimulation and viral infection [17], we sought to directly assess the ability of MDA5<sup>−/−</sup> mice to express IFN in response to SeV infection. In this regard, WT and MDA5<sup>−/−</sup> mice were infected with 200K pfu SeV and subsequently assessed for cytokine expression by real-time PCR analysis over the acute period. While both strains demonstrated comparable mRNA levels at d2 PI, type I IFN expression was dramatically dampened in MDA5<sup>−/−</sup> mice at d5 PI (Figure 4A and B). Unexpectedly, significant decreases in expression of Ifn-γ, Il-28b (Ifn-l3) and Tnf-α mRNA were also observed in the lungs of MDA5<sup>−/−</sup> mice compared to the WT cohort, with the most dramatic difference observed for Il-28b mRNA expression (Figure 4C, D and E). In contrast, Il-1β, and Il-
mRNA levels were not significantly different across strains, though the levels of *Il-6* mRNA was markedly increased in MDA5/2/2 mice following infection (Figure 4F, G and H). Accordingly, MDA5 appears to control the expression of SeV-induced anti-viral cytokines, particularly type I, II and III IFNs, during the late acute period (d5 PI), but does not appear to be involved during the immediate early response. Importantly, the decrease in IFN expression coincides with expanded viral propagation in the MDA5/2/2 mice, suggesting that reduced IFN expression during this time point accounts for the corresponding increased viral burden.

Induction of IFN expression transactivates expression of a number of IFN response genes through a signal transduction cascade involving JAK/STAT activation. MDA5 and RIG-I are among the genes induced by IFN signaling in vitro [20]. To determine the expression profile of MDA5 and RIG-I in the airways of mice infected with SeV, mRNA was measured by real-time PCR analysis from whole lung homogenates obtained from WT mice infected with 200K pfu SeV. Expression of *Mda5* and *Rig-i* mRNA was significantly increased at d2 and d5 PI, though the levels began to decline by d8 PI (Figure 5A). Lastly, to determine the tissue distribution of MDA5 expression, lung sections from d5 PI mice were stained with anti-MDA5 polyclonal antibodies. Visualization of MDA5-specific staining was performed using tyramide-based amplification. IF microscopic analysis of affected airways revealed a pattern of MDA5 expression that was primarily restricted to the airway epithelium, though expression was also detected in cells of the proximal interstitium, in particular, in cells that appeared to resemble type II pneumocytes and alveolar macrophage (Figure 5B). Sections from MDA5/2/2 mice did not stain for MDA5, confirming the specificity of anti-MDA5 staining. Accordingly these findings indicate that MDA5 is induced following SeV infection and that the lack of expression in MDA5/2/2 mice accounts for the phenotype described at the later time point.

**Discussion**

Our understanding of innate immune factors that recognize and respond to pathogens has greatly expanded over the last decade. A major component of the RNA virus detection system in mammals involves members of the RLR family, including RIG-I, MDA5, and LGP2 [1]. Elucidating a role for the RLRs in virus-induced IFN production has been facilitated by the availability of
RIG-I\(^{-/-}\) and MDA5\(^{-/-}\) mice [32,33]. Initial observations using embryonic fibroblasts and bone marrow derived DCs generated from these mice revealed striking phenotypes including a failure to produce IFN in response to a wide cross-section of viruses and nucleic acids, and an inability to contain viral replication. Specifically, MDA5 was found to be the sole receptor for picornaviruses and caliciviruses (\((+)^{\text{ssRNA}}\) viruses) [32,33,35], whereas RIG-I was described as the receptor for \((-)^{\text{ssRNA}}\) viruses such as paramyxoviruses and orthomyxoviruses, as well as for \((+)^{\text{ssRNA}}\) viruses belonging to the Flaviviridae family [16,33]. However, our understanding of these virus recognition systems in vivo is limited, in part because RIG-I\(^{-/-}\) mice die perinatally.

The precise molecular patterns of virus replication recognized by RIG-I and MDA5 are still not fully clear. Initially, a mimic of viral dsRNA, polyIC, was found to bind and activate RIG-I. However, ensuing research identified 5’-tripholphosphorylated ssRNA as the major RIG-I inducer [27,28]. Furthermore, in vitro data obtained using knockout mice suggested in fact that MDA5, and not RIG-I, recognizes polyIC, thereby formulating a recognition model whereby RIG-I recognizes short 5’-tripholphosphorylated RNAs, while MDA5 recognizes dsRNA structures irrespective of the 5’ cap [8,30–32]. However, more thorough dissection of the helicase binding function and activation process has recently determined that the picture is indeed more complex than previously thought [29,40]. In this regard, both helicases have been shown to recognize dsRNA, in a manner that is likely dependent on its length, while RIG-I demonstrates the added ability to respond to 5’-tripholphosphorylated ssRNA products. To complicate these paradigms, there is increasing evidence that viruses have evolved various properties aimed at antagonizing or degrading viral sensors. Thus, our understanding of viral recognition by the RLR helicases is evolving.

With respect to molecular sensing of paramyxovirus infection, both 5’-tripholphosphorylated ssRNA and long dsRNA species are likely present in SeV-infected cells, thereby implicating both MDA5 and RIG-I in the antiviral sensing process. However, in vitro studies concur that cultured embryonic fibroblasts and bone marrow-derived DC cells detect SeV RNA chiefly through RIG-I, whereas MDA5 and TLR3 are dispensable [33,34,41,54]. TLR7 and TLR8 in myeloid cells have also been shown to recognize SeV RNA in vitro as well [55]. Regardless, it cannot be excluded that in vivo, other RNA sensors, including MDA5, may contribute, at least in part, to anti-SeV responses. Indeed, a recent study by Yount et al. has demonstrated that MDA5 can detect SeV DI particles in vitro [45]. The relevancy of this recognition system in vivo is uncertain; certainly in our hands, using SeV/52, which has a limited ability to form DI particles, as per PCR-based analysis (data not shown), MDA5 appears to exert a significant effect on

**Figure 3.** SeV replication is enhanced in MDA5\(^{-/-}\) mice. WT and MDA5\(^{-/-}\) mice infected with 200K pfu SeV were assessed for A) SeV replication by IF detection of SeV antigens and by real time PCR analysis of B) SeV genome and C) SeV N gene expression. N = 4, error bars refer to SEM, * P < 0.05; ** P < 0.005. doi:10.1371/journal.ppat.1000734.g003
viral containment. Most importantly however, SeV encodes a V protein that specifically binds to and blocks MDA5 signaling in vitro [14,44]. Thus, it is possible that MDA5 does, indeed, detect SeV in vitro, but that it is functionally curtailed by the V protein in this circumstance. Interestingly, in our hands, in vivo infections using SeV with V protein deletion resulted in no real effect on mortality or type I IFN induction across strains (data not shown), likely explained by the fact that deletion of V protein in SeV markedly attenuates virulence and pathogenicity in vivo [56].

While initial characterization of MDA5-deficient cells has not supported a role for MDA5 in containment of SeV, these studies have been limited to observations made in cultured embryonic fibroblasts and in vitro-derived dendritic cells; populations which are not primary targets for SeV replication in the course of the natural infection. Rather, SeV replication mostly occurs in the airway epithelium of the conducting airways [49,50]. For these reasons, we hypothesized that SeV propagation would be sensitive to the MDA5 status of the host in the context of an in vivo infection. Indeed, the epithelial cells constitutively express MDA5 at low levels and subsequently up-regulate expression in response to SeV (Figure 5B), a finding that supports the relevance of this RNA helicase to SeV and other airborne infections. Interestingly, MDA5 deficiency did not influence the composition of the inflammatory infiltrate (data not shown), implying that the immune defect is largely restricted to the airway epithelium, the site of viral replication. This is compatible with our earlier findings using STAT-1−/− chimeras, wherein we observed that loss of IFN response in the stromal compartment alone accounted for the immune deficiency to SeV [53]. We therefore sought to further assess the significance of MDA5 in the control of SeV infection in vivo. In this regard we have demonstrated that MDA5 controls SeV replication and spread through induction of type I IFNs, but that this effect appears late (d5 PI), as IFN gene transcription is not impaired on d2 PI (Figure 4). It is likely that the initial IFN response is sufficient to initiate a range of immune responses, such that the late reduction in IFN transcripts results only in a 2–3 fold change in LD50 (Figure 1). Whether this specific IFN pattern remains true for other viruses as well remains to be tested. This surprising collapse of the host type I IFN response at d5 PI is accompanied by parallel decreases in the level of Il-28b and Tnf-α expression (Figure 4), and, curiously, decreased Ifn-γ transcript levels. This later observation may reflect a selective role for MDA5 in the induction of IFN-γ expression by NK cells. Lastly, the MDA5 status does not appear to influence the levels of IL-1β, or IL-10 or the ability of the host to mount a virus-specific CTL response. However, the levels of Il-6 mRNA in whole lung

Figure 4. MDA5 is required for sustained expression of cytokines in response to SeV infection. Real time PCR analysis of whole lung homogenates obtained from WT and MDA5−/− mice infected with 200K pfu SeV for expression levels of A) Ifn-α2, B) Ifn-β, C) Ifn-γ, D) Il-28b, E) Tnf-α, F) Il-1β, G) Il-6 and H) Il-10 mRNA. N=4, error bars refer to SEM, * P<0.05, ** P<0.00001. doi:10.1371/journal.ppat.1000734.g004
homogenates derived from d2 and d5 PI MDA5−/− was markedly increased, suggesting the induction of compensatory mechanisms in the context of MDA5 deficiency that could potentially account for the enhanced morbidity and mortality seen in the MDA5−/− mice.

An additional concern raised by these data is the relative contribution of MDA5 and RIG-I in the response to virus. In light of the existing literature [33,40], it seems likely that RIG-I is responsible for the normal IFN response to SeV early in the infection. Indeed, as depicted in Figure 5A, RIG-I is strongly induced early on during infection. Why the later IFN response depends on MDA5 is not known. MDA5 is encoded by an IFN-upregulated transcript, and it remains possible that it is the accumulation of MDA5 that allows for the subsequent MDA5-dependent IFN response on d5 PI. Yet other IFN-induced genes, notably RIG-I, are also upregulated by IFN, which should provide additional antiviral protection in vivo. Interestingly, SeV encodes a nested set of C proteins that have been shown to impede IFN signaling through direct inhibition of STAT signaling [41,42] and which are also known to strongly antagonize RIG-I function [41]. Furthermore, SeV-V proteins have been shown to have direct inhibitory effects on both MDA5 and RIG-I signaling [41,44]. Thus it remains possible that the effects of SeV V and C proteins have an accumulative effect on RIG-I function that essentially overwhelms this sensor at d5 PI, and that in this context, MDA5 plays an essential role in containment of SeV. Since assessment of the relative contribution of RIG-I and MDA5 in containment of SeV infection in vivo is not possible, a possible next step in assessing the importance of MDA5 function would involve assessment in MDA5−/− and IPS-1−/− strains.

We envision several possibilities that could potentially explain this dramatic effect of MDA5. The first is that, in the absence of MDA5, the balance between virus replication and the IFN response is disrupted sufficiently, such that by d5 PI, virus replication has overwhelmed the response in a qualitative fashion—presumably through direct cytotoxic effects or via the overproduction of immunosuppressive C proteins. This possibility is supported by the fact that SeV is replicating to higher levels in the MDA5−/− lung already by d2 PI (Figure 3B). Indeed, in support of this hypothesis, we observe a striking increase in SeV replication that spreads extensively into the interstitium of MDA5−/− lungs compared to controls. Another possible explanation, which we have not assessed, is an apoptotic response potentially mediated by MDA5. In this scenario, MDA5 would instruct or sensitize infected cells to commit suicide so as to shut down viral replication in infected cells. Indeed, ectopic expression of MDA5 in a melanoma cell line has been shown to inhibit colony formation, presumably through induction of apoptosis [20], and IPS-1 overexpression induces cell death, as well [57]. In fact, SeV-dependent apoptotic signaling requires IRF3 [58]. In the case of MDA5 deficiency, loss of pro-apoptotic activity could lead to a robust increase in viral replication and enhanced IFN blockade through overexpression of SeV C proteins. This possibility is favored by the fact that, despite a normal IFN response on d2 PI (Figure 4), the virus is found to be replicating at higher titers (Figure 3B).

It seems likely that the inability of the MDA5−/− animals to sustain an IFN response leads to increased viral replication and dissemination on d5 PI, thus causing significantly higher morbidity and mortality in the knockout cohort (Figure 1). It is important to note, however, that the effects of MDA5 deficiency on SeV...
replication are much milder than expected if MDA5 were the sole target of the V protein. Indeed, SeV V mutants (SeVΔV) are severely attenuated; replication is demonstrably abrogated in the lungs by d2 PI [36]. IRF3 deficiency of the host restores SeVΔV pathogenicity, suggesting that the mutant virus acts by blocking IRF3 signaling [59]. Yet disease caused by SeV in MDA5Δ/Δ mice is milder than the disease seen in the IRF3−/− animals. Consequently, we believe that the V protein must have additional targets besides MDA5. In this regard, it has recently been demonstrated that Lgp2 encodes a helicase epitope that is akin to the MDA5 helicase, the portion of MDA5 that binds paramyxovirus V proteins [60], thereby suggesting that Lgp2 may be an additional V protein target. In this case, a MDA5-LGP2 double knockout mouse may potentially phenocopy the IRF3 mutation in its response to SeV infection.

Taken together, our findings demonstrate that MDA5 significantly contributes to the response to paramyxovirus and constitute the first in vivo demonstration of MDA5 activity against a negative-strand virus. As such, it appears likely that MDA5 has a wider specificity as a viral nucleic acid receptor than initially believed, and that the initial clear-cut cases of either MDA5 or RIG-I being the specificity as a viral nucleic acid receptor than initially believed, and that the first in vivo demonstration of MDA5 activity against a negative-strand virus.

Materials and Methods

Mouse generation, maintenance and infection

Control C57BL/6j (B6) mice used in these experiments were purchased from JAX. MDA5Δ/Δ mice [32] were backcrossed onto the B6 background to 99.9% congenicity. For in vivo SeV infection, Sendai/52 Fushimi strain was instilled intranasally into deeply anesthetized mice and at the indicated time points, mice were humanly sacrificed for harvest of lung tissue. Virus was purchased from the ATCC and subject to two rounds of in vitro plaque purification in Vero cells to eliminate the presence of DI particles. A clone thus identified was then subject to a single round of amplification in embryonated chicken eggs following inoculation of ~1000 PFU. 24–36 hr post inoculation, SeV was isolated from the allantoic fluids and diluted in phosphate-buffered solution to generate a viral stock that was subsequently characterized on the basis of in vivo infectious properties. Calculation of PFU was performed by standard plaque assay using either Vero E6 cells or LLC-MK2 cells. Importantly, propagation under these conditions does not favor the formation of DI particles, a process that occurs most frequently when virus is repeatedly passaged at high MOI. Indeed, PCR analysis of stock virus indicated the absence of DI genomes. The methods for mice use and care were approved by the Washington University Animal Studies Committee and are in accordance with NIH guidelines.

FACS

Single cell lung suspensions were made from minced lung tissue subjected to collagenase/hyaluronidase/DNase I digestion. Staining of surface markers was performed using FcR block and fluorochrome-conjugated mAbs. To immunophenotype the immune infiltrate, specific combinations of mAbs were chosen which discern granulocytes (Ly6G+), macrophages (F4/80+), pDC (Siglec-H+), mDC (CD11c+CD8α+), cDC (CD11c+), NK cells (NK1.1+CD11c-), T cells (CD3+CD4+CD8+), and B cells (CD19+). SeV-specific PE-labeled pentamer K8:FPAGYNPAL (NP 324-332) was purchased from Proimmune; cells were stained with CD6 and counterstained with propidium iodide, F4/80 and CD19 to eliminate background. Activation status was determined using specific mAbs for MHC-II, NK2D, and CD69. Samples were acquired on a FACScalibur (BD Biosciences) and analyzed using Cellquest software.

Analysis of mRNA and virus-specific RNA

RNA was purified from lung homogenate using Trizol Reagent (Invitrogen). RNA was treated with RNase-free DNase I (Ambion) to eliminate genomic DNA. RNA was converted to cDNA using the High-Capacity cDNA Archive kit (Applied Biosystems). Target mRNA and viral RNAs were quantified by real-time PCR using specific fluorogenic probes and primers and the Fast Universal PCR Master Mix system (Applied Biosystems). Primer sets and probes for mouse Ifn-α2 (Mm00833961_s1), Ifn-β (Mm00439532_s1), Ifn-γ (Mm00801778-m1), Il-20b (Mm00663660_g1), Tnf-α (Mm00443259_g1), Mda5 (Mm00459183_m1), Il-β (Mm004454227_g1), Il-6 (Mm00446190_m1), Il-10 (Mm00439616_m1) mRNA and SeV genome and Gapdh mRNA were purchased from Applied Biosystems. Samples were assayed on the 7500 Fast Real-Time PCR System and analyzed using the 7500 Fast System Software (Applied Biosystems). Levels of specific gene expression were standardized to Gapdh mRNA expression levels.

Histology

Lungs were perfused and fixed with 4% paraformaldehyde. Tissue was embedded in paraffin, cut into 5 um sections and adhered to charged slides. Sections were deparaffinized in Citrosolv (Fisherbrand), hydrated, and in the case of IF-microscopy, treated to heat-activated antigen unmasking solution (Vector Laboratories, Inc). H&E and PAS sections were visualized by brightfield microscopy. Expression analysis was performed by IF using chicken polyclonal anti-SeV (Jackson ImmunoResearch Laboratories, Inc) and rabbit polyclonal anti-mouse MDA5 (Axxora Life Sciences, Inc). Biotinylated secondary antibodies were purchased from Vector Laboratories, Inc). SeV and MDA5 expression was visualized using tyramide-based signal amplification with Alexa Fluor 488 or 594 fluorochromes (Invitrogen). Slides were counterstained with DAPI mounting media (Vector Laboratories, Inc). Microscopy was performed using an Olympus BX51 microscope.

Statistical analyses

Real-time PCR data was analyzed using an unpaired Student’s t-test. If variances were unequal, Welch’s correction was applied. Charted values represent mean ± SEM. Survival statistics were determined using Kaplan-Meier analysis of paired cohorts. P values below 0.05 were regarded as being significant for all analyses. Experiments were repeated a minimum of three times.

Author Contributions

Conceived and designed the experiments: M. Holtzman, M. Colonna. Performed the experiments: L. Benoit, C. Song, M. Cella. Analyzed the data: L. Gitlin, L. Benoit, M. Holtzman. Contributed reagents/materials/analysis tools: L. Gitlin, M. Cella, S. Gillillan. Wrote the paper: L. Gitlin, L. Benoit.

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