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# Mouse circulating extracellular vesicles contain virus-derived siRNAs active in antiviral immunity

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## Abstract

Induction and suppression of antiviral RNA interference (RNAi) has been observed in mammals during infection with at least seven distinct RNA viruses, including some that are pathogenic in humans. However, while the cell-autonomous immune response mediated by antiviral RNAi is gradually being recognized, little is known about systemic antiviral RNAi in mammals. Furthermore, extracellular vesicles (EVs) also function in viral signal spreading and host immunity. Here, we show that upon antiviral RNAi activation, virus-derived small-interfering RNAs (vsRNAs) from Nodamura virus (NoV), Sindbis virus (SINV), and Zika virus (ZIKV) enter the murine bloodstream via EVs for systemic circulation. vsRNAs in the EVs are biologically active, since they confer RNA–RNA homology-dependent antiviral activity in both cultured cells and infant mice. Moreover, we demonstrate that vaccination with a live-attenuated virus, rendered deficient in RNAi suppression, induces production of stably maintained vsRNAs and confers protective immunity against virus infection in mice. This suggests that vaccination with live-attenuated VSR (viral suppressor of RNAi)-deficient mutant viruses could be a new strategy to induce immunity.

**Keywords** antiviral RNAi; extracellular vesicles; live-attenuated virus vaccine; systemic antiviral RNAi; Zika virus

**Subject Categories** Immunology; RNA Biology

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## Introduction

Extracellular vesicles (EVs) are small membrane structures containing a complex cargo of proteins, lipids, and nucleic acids (Théry *et al*, 2018). EVs are produced by different types of cells and

can be detected in most body fluids including blood, saliva, urine, and cerebrospinal fluid (Srinivasan *et al*, 2019). EVs are classified as apoptotic bodies, microvesicles, or exosomes. Cells undergoing apoptosis release apoptotic bodies 1–5 µm in diameter. Microvesicles are generated via shedding/budding from the plasma membrane and are between 150 nm and 1 µm in diameter. Exosomes originate from multivesicular bodies (MVBs) with the cellular plasma membrane and small dimensions (30–150 nm) (Théry *et al*, 2018). EVs have essential roles in intercellular communication, both locally and systemically. Recent studies have demonstrated that EVs with viral components play dichotomous roles in viral infections and pathology (Nolte *et al*, 2016; Raab-Traub & Dittmer, 2017; Caobi *et al*, 2020; McNamara & Dittmer, 2020). Viruses often exploit the EV biogenesis pathway to promote viral infection and propagation. On the other hand, EVs can help the host cell to suppress virus infection by triggering antiviral responses and cytokine secretion. Sometimes, this enigmatic dual roles of EVs are observed in infections by the same virus (Caobi *et al*, 2020; Martins & Alves, 2020). Therefore, the role of EVs during viral infections needs to be further investigated.

Recent studies including ours have demonstrated production of abundant virus-derived small-interfering RNAs (vsRNAs) to target several distinct RNA viruses, such as influenza A virus (IAV), Zika virus (ZIKV), and Dengue virus (DENV), in mammalian cells (Li *et al*, 2013, 2017; Maillard *et al*, 2013; Qiu *et al*, 2017, 2020; Xu *et al*, 2019; Zhang *et al*, 2020). Infection of mouse embryonic stem cells (mESCs) and human neural progenitor cells (hNPCs) respectively by wild-type encephalomyocarditis virus (EMCV) and ZIKV triggers Dicer-mediated biogenesis of abundant vsRNAs only before the cells are induced to differentiate (Maillard *et al*, 2013; Xu *et al*, 2019). Reis e Sousa and colleagues have recently demonstrated a critical role for a Dicer isoform in vsRNA production to initiate antiviral RNAi in mouse and human stem cells (Poirier *et al*, 2021). In contrast, vsRNAs accumulate to high levels in both mESCs and mature baby hamster cells as well as infant mice in response to the infection with Nodamura virus (NoV) only after its VSR protein B2 is rendered either non-expressing or non-functional (Li *et al*, 2013;

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Maillard *et al*, 2013). Similarly, abundant vsiRNAs are produced by Dicer from IAV mutants not expressing its VSR non-structural protein 1 (NS1) in mature human and monkey cells after infection (Li *et al*, 2017). Zhou and colleagues also showed that the 3A protein of human enterovirus 71 (HEV71) is a dsRNA-binding VSR that suppresses Dicer-mediated production of vsiRNAs like B2 and NS1 despite no sequence similarity among the three mammalian VSRS (Qiu *et al*, 2017; Fang *et al*, 2021).

The role of antiviral RNAi in the innate immunity and its relationship to the interferon (IFN) response remain poorly understood (Berkhout, 2018; Ding *et al*, 2018; Guo *et al*, 2019; Maillard *et al*, 2019). It is well known that the IFN response directs a potent innate antiviral pathway. Independent studies have shown that the RNAi response is antiviral in undifferentiated cells, in which the canonical IFN system is not active (Maillard *et al*, 2013, 2019; Wu *et al*, 2019; Xu *et al*, 2019). In differentiated mammalian cells, antiviral RNAi is active against only mutant viruses defective in RNAi suppression (Li *et al*, 2013, 2017; Qiu *et al*, 2017, 2020; Han *et al*, 2020). Moreover, Dicer-mediated siRNA production and sequence-specific RNAi can be triggered by artificial long dsRNAs only when the IFN pathway is defective (Kennedy *et al*, 2015; Maillard *et al*, 2016). Notably, activation of siRNA production in cells from artificially introduced long dsRNA confers specific antiviral resistance to subsequent infections, indicating that mammalian cells can be “vaccinated” by RNA in a virus-specific manner (Maillard *et al*, 2016). Subsequently, Reis e Sousa and colleagues further showed that binding of LGP2 to Dicer inhibited cleavage of dsRNA into siRNAs *in vitro* and in cells but did not inhibit Dicer-dependent pre-miRNA processing (Van der Veen *et al*, 2018). Interestingly, a recent study found abundant accumulation of vsiRNAs from Dicer processing of viral dsRNA replicative intermediates in adult mice either wild-type or defective in the signaling of IFN, suggesting that vsiRNA biogenesis is distinct from Dicer processing of artificial long dsRNA (Han *et al*, 2020). Our recent work also revealed that viral dsRNA replicative intermediates made during authentic infection of mature somatic cells are efficiently processed by Dicer into vsiRNAs to direct antiviral RNAi (Zhang *et al*, 2021).

The control of viral infections in mammals requires signaling molecules to elicit an effective response and to establish systemic immunity at the organismal level. These signals must be amplified and disseminated throughout the organism to inhibit pathogen propagation and establishment of the infection (Hu & Shu, 2018; Hur, 2019). In mammals, cells contain pattern-recognition receptors (PRRs) to detect molecules associated with infection. These PRRs, once stimulated by their appropriate ligands, activate intracellular signaling cascades leading to transcription of IFN genes. Once secreted, IFN binds to the IFN receptor on neighboring uninfected cells and activates an intracellular signaling cascade leading to upregulation of several hundred IFN-stimulated genes (ISGs), many of which have direct or indirect antiviral activities (Schneider *et al*, 2014). Cell-autonomous antiviral RNAi has been shown to be functional in mammals, but whether there is a systemic immune response mediated by RNAi needs to be investigated (Berkhout, 2018; Ding *et al*, 2018; Maillard *et al*, 2019).

We have shown that infection with the VSR-deficient mutant of NoV, NoVΔB2, induces production of abundant vsiRNAs and is rapidly cleared in suckling mice in contrast to lethal infection by wild-type NoV (Li *et al*, 2013). Intriguingly, we subsequently found that

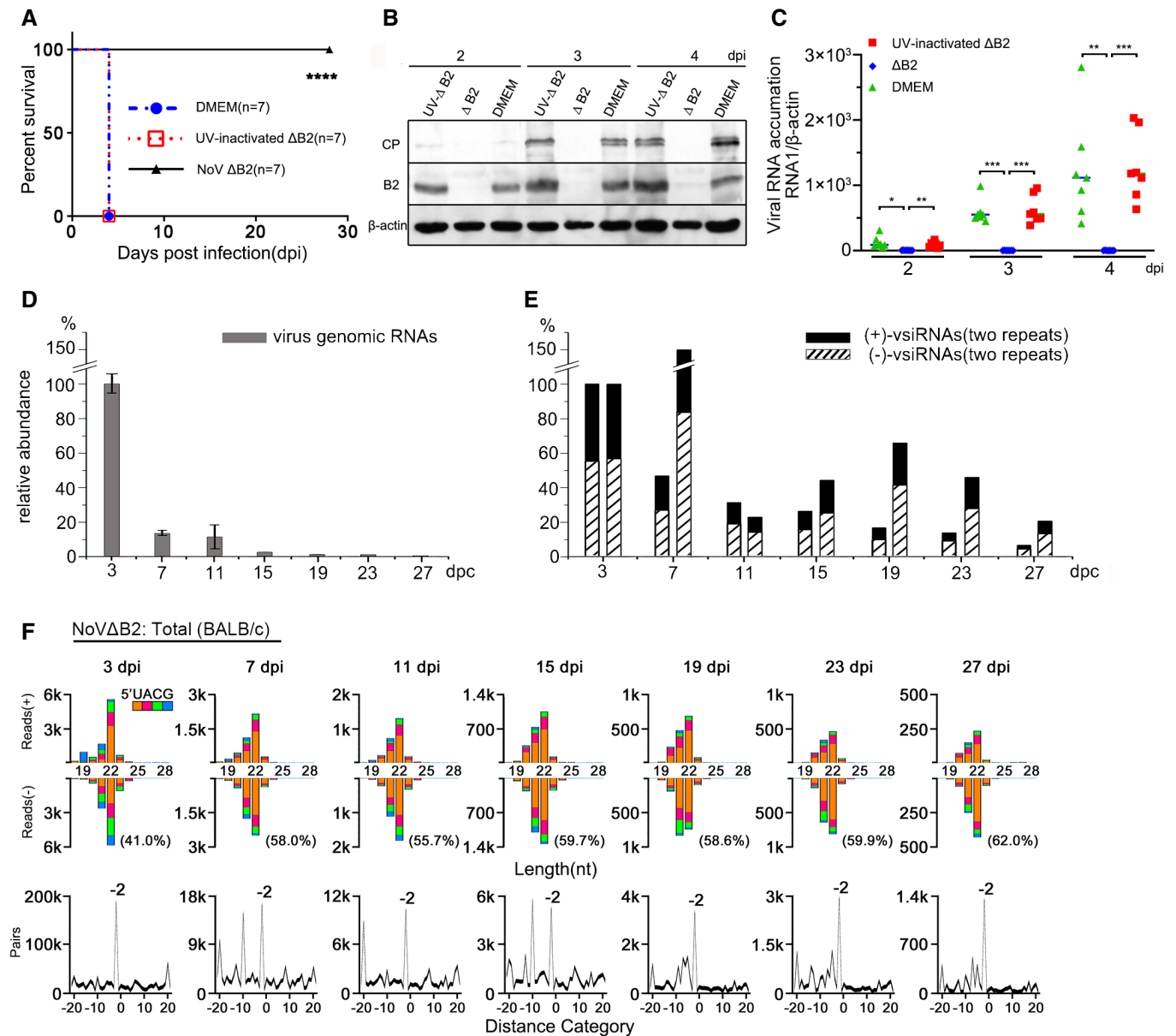
infant mice became fully protected against lethal NoV challenge after immunization with NoVΔB2 for only two days, suggesting rapid induction of protective immunity at whole organism level. A recent study has shown that infection of fruit flies with Sindbis virus (SINV) induces secretion of vsiRNAs-containing exosome-like vesicles circulating in the hemolymph sufficient to confer passive protection against SINV challenge in naive flies (Tassetto *et al*, 2017). This prompted us to investigate a potential role of EVs in NoVΔB2-triggered protection in mice. We demonstrate that immunization of suckling mice with NoVΔB2 induced accumulation of EV-associated vsiRNAs in the blood stream capable of conferring resistance against NoV challenge in naive mice. Notably, our findings indicate that murine EV-associated vsiRNAs mediate Ago2-dependent RNAi in cultured cells. We propose that the cell-autonomous and systemic antiviral RNAi are both active in mammals and that mutant viruses rendered deficient to suppress vsiRNA biogenesis can be explored as a new class of live-attenuated vaccines against animal and human RNA viruses.

## Results

### Vaccination with a live-attenuated, VSR-deficient NoV (NoVΔB2) confers complete protective immunity in mice against lethal infection

Our previous study have shown that unlike the 100% mortality observed 5 days post-NoV infection, suckling mice challenged by NoVΔB2 remained healthy for the duration of the experiment, up to 4 weeks post-inoculation (Li *et al*, 2013). If the viral siRNAs produced *de novo* in NoVΔB2-infected mice directed the clearance of NoVΔB2 by RNAi, they might confer specific protective immunity against secondary infection by NoV since NoVΔB2 and NoV differ only by three nucleotides. To test this hypothesis, we inoculated 6-day-old mice with buffer (mock), NoVΔB2, or NoVΔB2 inactivated by ultraviolet light (UV) irradiation, and two days later, challenged the mice with a lethal dose of NoV. The timing of NoV challenge was chosen according to the accumulation profile of viral siRNAs in the NoVΔB2-immunized neonatal mice. We found that all of the suckling mice pre-inoculated with either buffer or UV-inactivated NoVΔB2 developed hind limb paralysis 4 days after NoV challenge before death by 5 dpi. By contrast, none of the suckling mice immunized with NoVΔB2 exhibited any signs of disease up to 4 weeks after the secondary infection, indicating complete protection of the NoVΔB2-immunized mice (Fig 1A). Western blotting detected abundant expression of both the B2 and coat proteins, encoded by the NoV genomic RNAs 1 and 2, respectively, after NoV infection in the control mice pre-inoculated with either buffer or UV-inactivated NoVΔB2 (Fig 1B). However, neither viral protein was detectable in the NoVΔB2-immunized mice after NoV challenge (Fig 1B). RT-qPCR further verified successful infection by NoV in the two control groups of mice, but not in the mice immunized with NoVΔB2 (Fig 1C).

We showed recently that NoVΔB2-immunized suckling mice support significantly reduced replication of Sindbis virus (SINV) when engineered to contain a NoV fragment targeted by high densities of vsiRNAs in the immunized mice (Zhang *et al*, 2021), suggesting a role of vsiRNAs in the induced protection. Thus, we further



**Figure 1. Vaccination with a VSR-deficient mutant virus induces protection against secondary infection in neonatal mice.**

- A** At 2 days post vaccination with NoVΔB2, UV-inactivated NoVΔB2 (UV-ΔB2) or culture medium DMEM, BALB/c mice were challenged with WT NoV. The survival data were analyzed by the log rank test, \*\*\*\* indicates  $P < 0.0001$ ,  $n = 7$  per group.
- B, C** NoV levels in the vaccinated mice hindlimb muscle tissue at 2, 3, or 4 days post infection (dpi) with WT NoV were determined by Western blotting to detect the viral coat protein (CP) and B2 (B) or quantitative RT-qPCR to detect the viral genomic RNA1,  $n = 7$  per group (C). Staining of β-actin was used as a loading control (B). The viral RNA1 level was normalized by β-actin mRNA levels (C).  $n = 7$  per group. Horizontal bars represent for means. \* indicates  $P < 0.05$ , \*\* indicates  $P < 0.01$ , and \*\*\* indicates  $P < 0.001$  by Student's *t*-test.
- D, E** Time course analysis of the accumulation of NoVΔB2 RNA1 determined by RT-qPCR (D) and 21- to 23-nt vsiRNAs determined by deep sequencing (E, two independent sets) from BALB/c suckling mice infected with NoVΔB2. Mice,  $n = 3$ , in each independent analysis. Data are shown as mean  $\pm$  SD. Normalization was done by β-actin mRNA levels for NoVΔB2 RNA1 (D) or by mature miRNAs for vsiRNAs (E). Accumulation levels of NoVΔB2 RNA1 or vsiRNAs at 3 dpi are set as 100%, respectively.
- F** Properties of viral siRNAs (per million mature miRNAs) cloned and sequenced from the time course series of NoVΔB2-infected BALB/c suckling mice. Size distribution, 5' terminal nucleotide, and duplexes by 22-nt vsiRNAs are indicated. The percentage of 1 U vsiRNAs (21- to 23-nt) in each library is shown in parentheses.

investigated the *in vivo* stability of the vsiRNAs in suckling mice following the clearance of NoVΔB2. As described previously (Li *et al*, 2013), NoVΔB2 was rapidly cleared from the infected mice

following peak replication at 3 dpi (Fig 1D). Cloning and sequencing of the small RNAs from NoVΔB2-infected suckling mice at 3, 7, 11, 15, 19, 23, and 27 dpi in two independent repeats demonstrated

accumulation of vsiRNAs throughout the time course (Figs 1E and F, and EV1). The vsiRNAs cloned after virus clearance were similar to those cloned at earlier time points in the size distribution and the strand ratios. We found that the vsiRNAs sequenced from mice at 7 dpi and afterward were as highly enriched for 1 U vsiRNAs (Figs 1F and EV1) as found for those co-immunoprecipitated with AGOs (Li *et al.*, 2017; Zhang *et al.*, 2020). Consistently, we observed de-enrichment of the canonical vsiRNA duplexes among the total vsiRNAs cloned from the mice from later time points (Figs 1F and EV1). These findings indicate that the vsiRNAs remain stable *in vivo* up to two weeks following the clearance of the virus, which is likely a result of their association with Argonaute complexes.

### The vsiRNA profiles of serum EVs upon NoV $\Delta$ B2 infection

We next investigated whether murine vsiRNAs enter the blood stream for systemic circulation by sequencing the total small RNAs extracted from the whole blood upon NoV $\Delta$ B2 infection. The results showed that vsiRNAs were detectable by small RNA sequencing (Table EV1). To clarify the origin of the vsiRNAs, exosome-enriched EVs were precipitated from mouse serum through exosome isolation kit (Fig 2A). These EVs were physically homogenous, with a size distribution peaking at 110 nm in diameter as determined by nanoparticle tracking analysis (NTA) and electron microscopy (Fig 2B and C). To further confirm that the isolated particles were exosome-enriched EVs, we examined the accumulation of five proteins known as exosomal biomarkers, CD9, CD63, CD81, HSP70, and tsg101, by Western blotting. The results showed that these proteins are readily detectable in the isolated preparations, indicating that EVs isolated from the serum were enriched for exosomes (Fig 2D).

The small RNA sequencing was performed to characterize the RNA species in the EVs from mock and NoV $\Delta$ B2 infection. The results indicated that the length of these small RNAs formed two peaks at approximately 21–23 nt and 29–31 nt (Figs 2E and EV2A). We found that miRNAs and transfer RNA (tRNA) contents (tRNA-derived fragments, tRFs and tRNA-halves, tiRNAs) were also the two main types of small RNA species in the EVs as previously reported (El-Mogy *et al.*, 2018; Srinivasan *et al.*, 2019; Zhao *et al.*, 2020). However, tRNA contents were highly enriched in the EVs to levels higher than those of miRNAs (Fig 2E). In contrast, miRNAs were the most abundant small RNAs in the whole blood and blood clot samples. Notably, we detected the accumulation of the vsiRNAs not only in the whole blood, but also in the purified EVs from NoV $\Delta$ B2-infected mice (Figs 2F and G, and EV2A). The EV-associated vsiRNAs were similar in the size distribution, strand ratio, 1 U preference, and abundance to those sequenced from mouse hindlimb muscle tissues (Figs 2F and EV3). By comparison with the respective total miRNAs, the vsiRNAs were approximately 60-fold more abundant in the EVs (2.8% of the total) than those in the whole blood (0.05% of the total) (Fig 2F and H). Moreover, the distribution patterns of vsiRNA hot spots on the two positive-strand viral genomic RNAs were similar between the whole blood and EV libraries, indicating enrichment of the vsiRNAs in the EVs (Fig 2G).

To track the source of the EVs, we further compared the frequency of known miRNAs among the hindlimb, EVs, whole blood, and blood clot samples. In hindlimb muscle tissue, miR-1a-3p, which has been reported as one of the most highly expressed miRNAs in differentiated muscle tissues, accounted for 48.5% of the

total miRNAs (Fig 2H). In contrast, the accumulation of miR-1a-3p in whole blood and blood clot was relatively low. Instead, the miR-451a was the most abundant miRNA in these two blood samples. The most abundant EVs miRNA was miR-381-3p (15.4% of the total), whose expression level was low in the two blood samples, but ranked third (5.6% of the total) in the muscle tissue (Fig 2H). The accumulation patterns of specific cellular miRNAs and vsiRNAs indicate that although additional studies are necessary to conclude an origin from the virus-infected muscle tissues, EVs-associated vsiRNAs are unlikely from blood cells.

### The induction of EV-vsiRNAs in distinct mouse strains by NoV $\Delta$ B2, SINV, and ZIKV

To determine whether the accumulation of EV-associated vsiRNAs is mouse strain-specific, we profiled the vsiRNAs in the blood and the EVs in C57BL/6 infant mice infected with NoV $\Delta$ B2. Similar to BALB/c mice, a typical population of vsiRNAs was detected in both the whole blood and the purified EVs in C57BL/6 mice after infection with NoV $\Delta$ B2 at 3 and 9 dpi (Figs 3A and B, and EV2B and C, Table EV1). Interestingly, we noted that the abundance of EV-associated vsiRNAs from NoV $\Delta$ B2-infected C57BL/6 mice (1.1% of the total miRNAs at 3 dpi and 1.0% of the total miRNAs at 9 dpi) was similar to that from NoV $\Delta$ B2-infected BALB/c mice.

We have shown that the vsiRNAs are induced with SINV or ZIKV infections *in vivo* (Zhang *et al.*, 2020, 2021). Thus, we next purified the EVs from the mouse serum after infection with SINV or ZIKV using the exosome isolation kit. We detected the accumulation of abundant SINV vsiRNAs predominantly within 21 to 23-nt size range in the EVs of C57BL/6 mice infected with SINV at 2 dpi (Fig 3C and Table EV1). The 22-nt vsiRNAs of SINV found in the EVs were highly enriched for canonical siRNA duplexes with 2-nt 3' overhangs, and its abundance equivalent to 4.9% of the total EV-associated miRNAs (Fig 3C and Table EV1). We also isolated the EVs from the serum of A6 (type 1 IFN receptor knockout C57BL/6) mice infected with ZIKV at 4 dpi. We also detected the accumulation of abundant vsiRNAs predominantly in 21 to 23-nt size range in EVs from ZIKV-infected A6 mice (Fig 3D and Table EV1). The vsiRNAs of ZIKV (21–23-nt) found in the EVs were highly enriched for 22-nt canonical siRNA, and its abundance equivalent to 14.9% of the total EVs miRNAs (Fig 3D and Table EV1). These results indicate the circulation of abundant vsiRNAs via the EVs in mice after infection with NoV $\Delta$ B2, SINV, or ZIKV.

### The EV-associated vsiRNAs confer homology-dependent antiviral activity *in vitro* and *in vivo*

We next explored whether the systemic spread of the vsiRNAs in NoV $\Delta$ B2-infected mice provides protection against secondary virus infection. To this end, we investigated whether the vsiRNAs in the circulating EVs of NoV $\Delta$ B2-immunized mice play a role in the activation of specific virus resistance at the organismal level. In the first set of *in vitro* experiments, we examined NoV infection of baby hamster kidney fibroblasts (BHK-21 cells) after prior incubation with the EVs purified from NoV $\Delta$ B2-infected infant mice (Fig 4A and B). As controls, we purified EVs from infant mice either mock-inoculated or infected with SINV that also induced systemic spread of SINV vsiRNAs via EVs (Fig 3C). We found that NoV replicated to

significantly reduced levels in BHK-21 cells treated by two different concentrations of the EVs purified from NoV $\Delta$ B2-infected infant mice compared to the EVs from mock-inoculated mice (Fig 4B). In

contrast, we detected no significant differences in the accumulation of NoV RNAs in BHK-21 cells treated with EVs from either mock-inoculated or SINV-infected mice (Fig 4C). These results

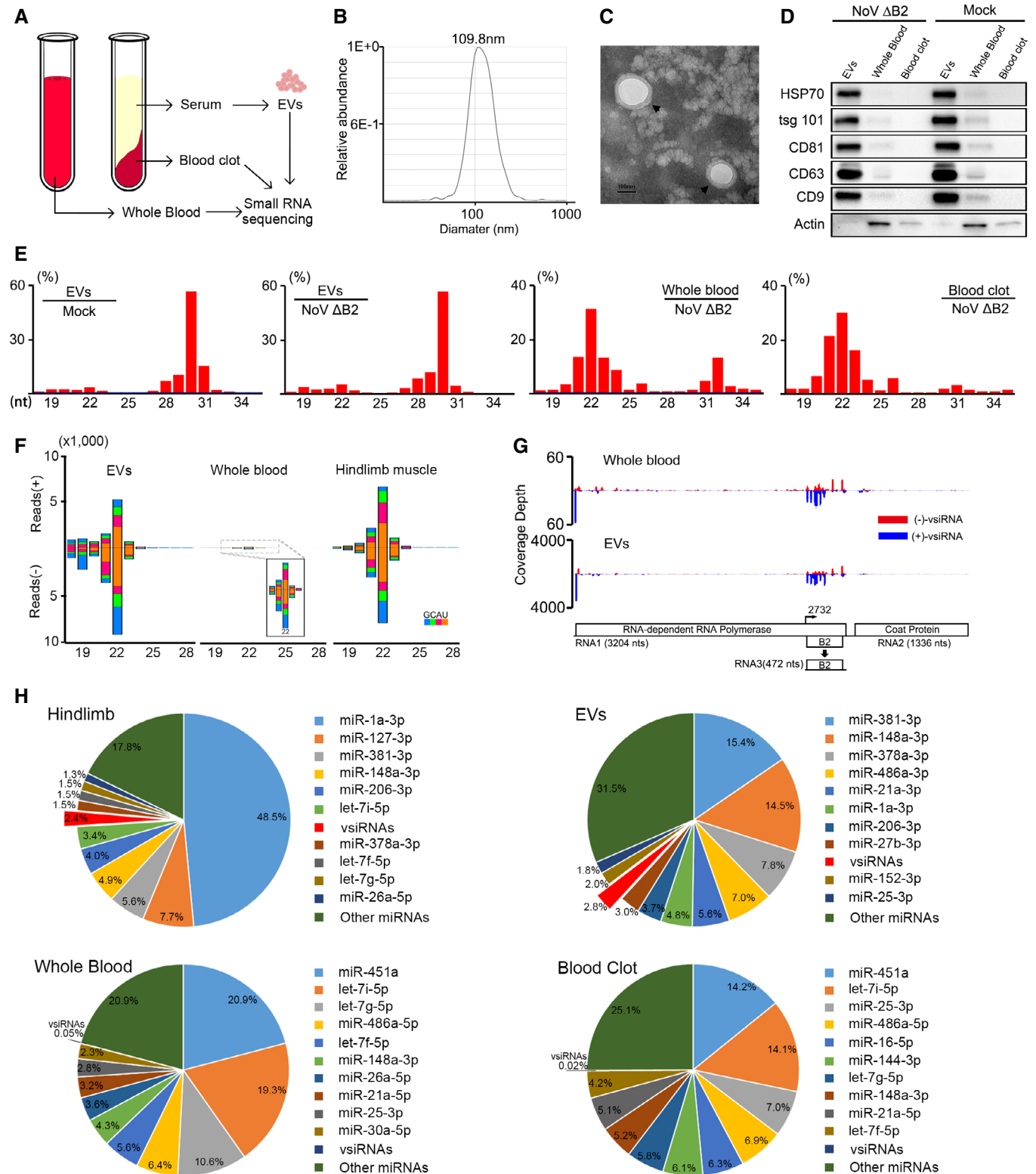


Figure 2.

**Figure 2. Characterization of viral siRNAs in murine EVs.**

- A Model of sample preparation for small RNA deep sequencing.
- B Size distribution of the purified EVs analyzed by NTA.
- C TEM images of purified EVs. Scale bar: 100 nm.
- D Western blotting of HSP70, tsg 101, CD81, CD63, and CD9 in EVs, whole blood, and blood clot derived from suckling mice infected with DMEM or NoVΔB2. Staining of β-actin was used as a loading control.
- E Size distribution of total sequenced small RNAs in the libraries constructed from EVs of DMEM challenged BALB/c suckling mice, and EVs, whole blood, blood clot of BALB/c suckling mice infected with NoVΔB2 at 3 dpi.
- F, G Size distribution of 18- to 28-nt vsRNAs (F) and genomic coverage depth of each nucleotide position by 21- to 23-nt vsRNAs (G) were analyzed from BALB/c suckling mice libraries infected by NoVΔB2 at 3 dpi.
- H Pie charts of the top ten most abundant miRNAs and vsRNAs in hindlimb, EVs, whole blood, blood clot from BALB/c suckling mice infected by NoVΔB2 at 3 dpi. Red charts indicate 18- to 28-nt vsRNAs abundance. The percentage of miRNA reads was shown above. The abundance of total miRNAs and 18- to 28-nt vsRNAs reads was set as 1.

demonstrate that pre-incubation with the EVs circulating in mice immunized with NoVΔB2 but not SINV inhibited NoV infection of BHK-21 cells, suggesting a specific antiviral activity of the EV-associated vsRNAs.

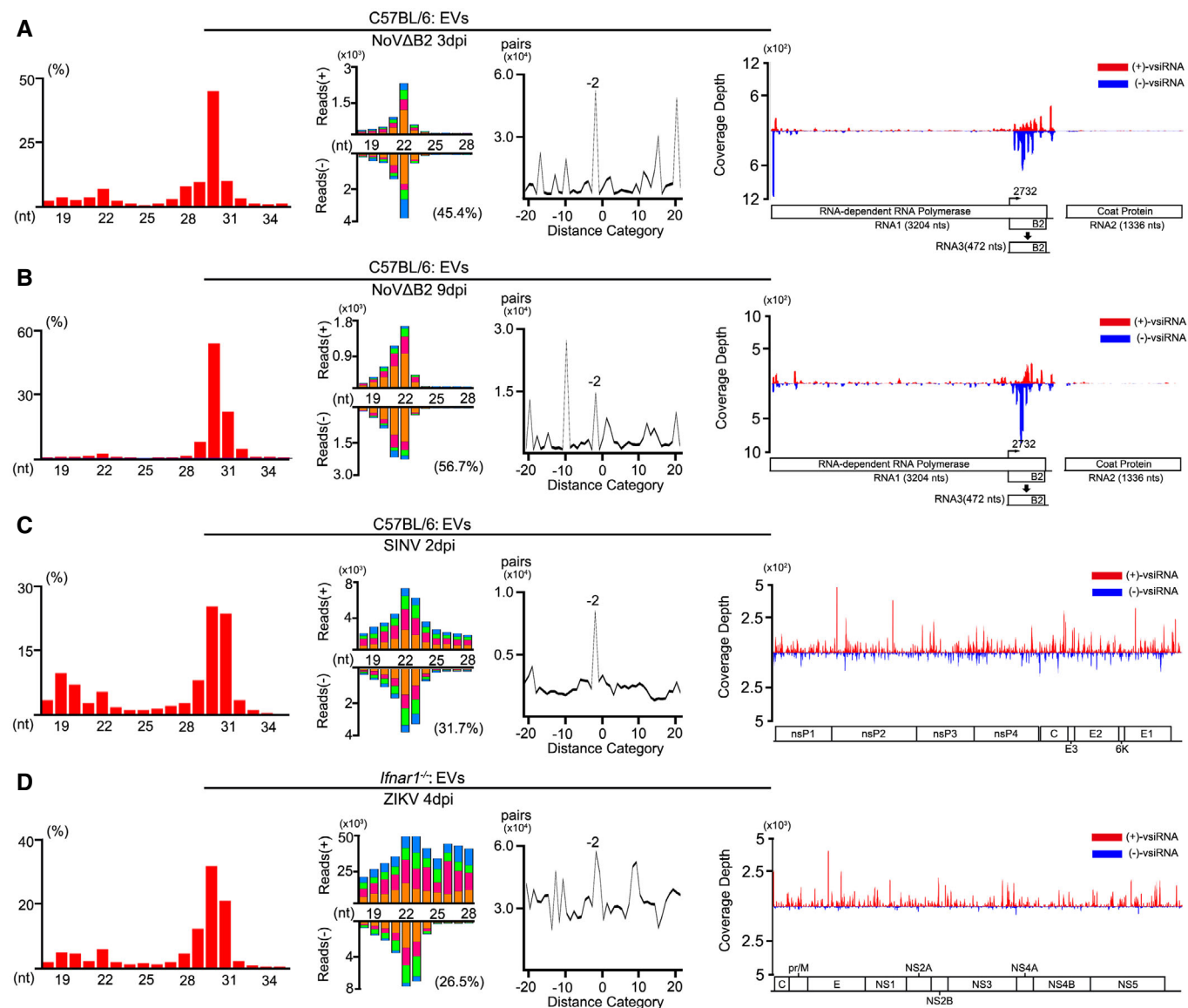
We performed another set of experiments to examine the RNAi activity of EV-associated vsRNAs in human 293T cells using a dual luciferase reporter system. A fragment of NoV RNA1 in sense orientation known to be targeted by high densities of EV-associated vsRNAs from NoVΔB2-infected infant mice was inserted into the 3' UTR of a dual luciferase reporter plasmid and the same length of GFP mRNA sequence was used in a control construct (Fig 4D). Four hours after transfection with one of the reporter constructs, the human 293T cells were treated by the EVs purified from NoVΔB2-infected infant mice or mock-inoculated mice. We observed significant suppression of the luciferase reporter containing the NoV fragment treated by the EVs purified from NoVΔB2-infected infant mice but not from mock-inoculated mice (Fig 4E). In contrast, treatment with the two different pools of EVs did not cause significant difference in the expression of the luciferase reporter containing the GFP fragment (Fig 4E). We further generated an Argonaute-2 knockout (Ago2 KO) line of 293T cells (Fig EV4) and performed the same luciferase reporter experiment using these cells. We found that the specific suppression of the luciferase reporter containing the NoV fragment by the EVs purified from NoVΔB2-infected infant mice was abolished in 293T-Ago2 KO cells (Fig 4F). These findings suggest that the specific antiviral effect of EVs from NoVΔB2-immunized infant mice is mediated by RNAi.

The next set of experiments examined the antiviral activity of the EV against NoV infection *in vivo* (Fig 4G). We challenged the infant mice with a lower dose of NoV (1/10 dose of vaccination experiment as described above) one day after i.p. injection with the EVs purified from NoVΔB2- or mock-inoculated mice, measured NoV titers at 1 and 3 days post challenge (dpc) by quantitative RT-PCR (RT-qPCR) detection of the NoV RNA1, and monitored survival of the challenged mice. As a control, infant mice immunized with NoVΔB2 for 1 day were also challenged by the same dose of NoV. At 1 dpc with NoV, accumulation of the viral RNA1 in both groups of mice injected with EVs was significantly lower than that in NoVΔB2-immunized mice (Fig 4H), possibly resulting from the on-going replication of NoVΔB2 rather than NoV. By 3 dpc, however, whereas NoV replicated to high levels in mice pre-injected with the EVs from mock-inoculated mice, NoV titers in mice pre-injected with the EVs from NoVΔB2-infected mice were as low as those in NoVΔB2-immunized mice

(Fig 4I). Among the three groups of mice challenged with NoV, viral B2 and coat proteins were detectable by Western blotting only in mice pre-injected with EVs from mock-infected mice (Fig 4J), indicating suppression of NoV infection by prior injection with either NoVΔB2 or the EVs from NoVΔB2-immunized mice. Notably, after NoV challenge, none of the seven infant mice immunized with NoVΔB2 for 1 day exhibited any signs of disease, and the mice pre-injected with the EVs purified from NoVΔB2-infected mice also survived significantly longer compared to those pre-injected with EVs from mock-infected mice (Fig 4K). We found that mice at 21 days post immunization with either live NoVΔB2 or EVs from NoVΔB2-immunized mice remained highly resistant to NoV challenge compared to those immunized with DMEM, UV-inactivated ΔB2, or EVs from mock-immunized mice (Fig EV5). These results show that the EVs circulating in mice during the course of antiviral RNAi activation and NoVΔB2 clearance conferred passive protection against NoV infection in naïve mice, suggesting an *in vivo* antiviral role of the EV in NoVΔB2 immunization-induced antiviral protection.

Extracellular vesicles can mediate communication between immune cells and other cell types upon virus infection (McNamara & Dittmer, 2020). EVs carry cytokines and cytokine-related RNAs that can affect recipient cells in the antiviral response (Nolte *et al*, 2016; Raab-Traub & Dittmer, 2017; Caobi *et al*, 2020). It is therefore possible that EVs from NoVΔB2-infected mouse serum may carry ISGs or dsRNAs and induce an IFN response to suppress reinfection. To investigate this possibility, we performed the EV immunization experiments in infant AG6 mice, which are deficient of both the type I and II IFN receptors in C57BL/6 background (Fig 5A). In control experiments, we detected only extremely low levels of NoVΔB2 RNA1 in mice injected with the EVs from either naïve or NoVΔB2-infected BALB/c suckling mice compared to those injected directly with NoVΔB2 (Fig 5B), suggesting presence of very few NoVΔB2 virions in the EVs purified from NoVΔB2-infected BALB/c suckling mice. We immunized infant AG6 mice for 12 h with the EVs purified from NoVΔB2- or mock-inoculated BALB/c mice before challenge with WT NoV, and compared NoV accumulation by RT-qPCR detection of the NoV RNA1 between the two groups of mice at 1 dpc (Fig 5A). We found that the accumulation of the viral RNA1 in mice immunized with the EVs from NoVΔB2-infected mice was significantly lower than those immunized with EVs from mock-inoculated mice (Fig 5C).

Furthermore, we determined whether vsRNAs produced from NoVΔB2-infected mice trigger the homology-dependent viral RNA



**Figure 3. Characterization of EV-vsIRNAs with distinct virus infections.**

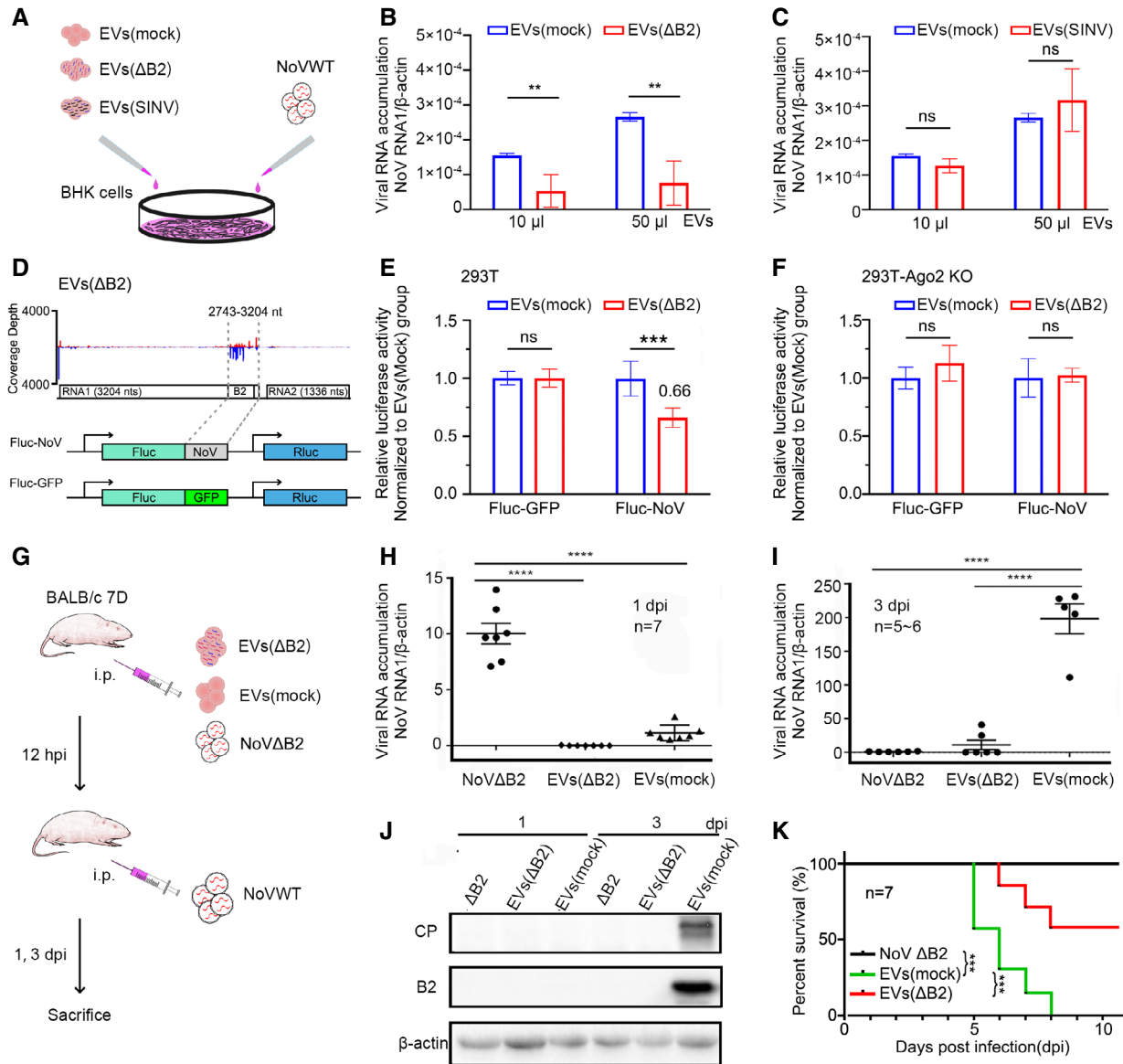
A–D Properties of total small RNA reads and vsIRNAs sequenced from EVs in C57BL/6 suckling mice infected with NoVΔB2 at 3 dpi (A) and 9 dpi (B), SINV at 3 dpi (C) and *Ifnar1*<sup>-/-</sup> mice infected with ZIKV at 4 dpi (D). Length distribution of total 18- to 35-nt reads in each library, size distribution of 18- to 28-nt virus-derived siRNAs, and viral genomic coverage depth of each nucleotide position by 21- to 23-nt vsIRNAs are presented as described in previous figures. Reads are shown as per million total mature miRNAs. 5' terminal nucleotide and the percentage of 1 U vsIRNAs and duplex pattern of the 22-nt vsIRNAs are indicated.

degradation guided by the vsIRNAs using Sindbis virus recombinants described recently (Zhang *et al*, 2021). SINV<sub>NoV</sub> contains an insert corresponding to a region of NoV genomic RNA1 targeted by high densities of vsIRNAs in NoVΔB2-infected mice whereas SINV<sub>GFP</sub> contains the sequence of GFP (Fig 5D). SINV<sub>NoV</sub> replicated to significantly lower levels than SINV<sub>GFP</sub> in the AG6 suckling mice immunized with the EVs from NoVΔB2-infected infant mice, but not in mice pre-inoculated with the control EVs, suggesting the EV-containing vsIRNAs were able to guide the homology-dependent antiviral response in the absence of IFN signaling (Fig 5E).

## Discussion

It is known that the EV composition changes drastically during virus infection (Nolte *et al*, 2016; Raab-Traub & Dittmer, 2017). EVs carry cytokines and cytokine-related RNAs that can affect viral pathogenicity, restrict viral propagation in recipient cells, and play an important role in the antiviral response. Different types of nucleic acids have been detected in EVs, such as miRNAs, lncRNA, mRNA, and various viral RNAs (Nolte *et al*, 2016; Raab-Traub & Dittmer, 2017). EV-associated host miRNAs produced by virus-resistant cells can confer resistance against various viruses (Delorme-Axford *et al*,





**Figure 4. Passive protection *in vitro* and *in vivo* conferred by isolated EV.**

A Scheme for protocol to determine the *in vitro* antiviral function of EVs from NoVΔB2-infected BALB/c suckling mice.

B, C 10  $\mu$ l or 50  $\mu$ l EVs stock from naïve [EVs (mock)] or NoVΔB2-infected [EVs ( $\Delta$ B2), B] or SINV-infected [EVs (SINV), C] mice were added to BHK cell culture medium along with NoV-WT, and then BHK cells were harvested at 24 hpi. Accumulation of NoV RNA1 in BHK cells was measured by RT-qPCR. Normalization was done by  $\beta$ -actin mRNA levels. Data from at least three independent experiments were combined. Data are shown as mean  $\pm$  SD. \*\* indicates  $P < 0.01$ , ns indicates no significant difference by Student's *t*-test.

D Diagram showing the 462-nt sequence in length from NoVΔB2 RNA1 was inserted into the 3' untranslated region (UTR) of firefly luciferase reporter mRNA, to be targeted by antisense vsRNAs with EVs from NoVΔB2-immunized BALB/c suckling mice. A translation initiation codon-deleted GFP sequence of the same length was inserted into the same site as control.

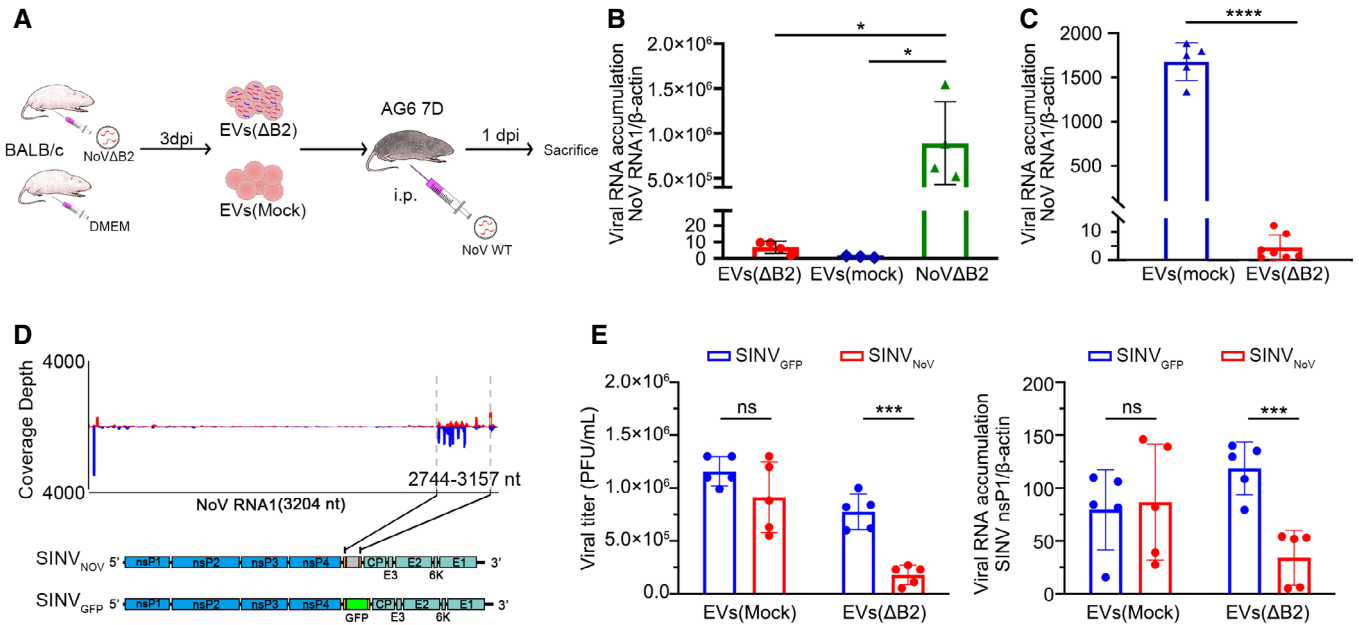
E, F Relative luciferase activity of the two reporter constructs after transfection of 293T cells (E) and 293T-Ago2 KO cells (F) and treatment by different pools of EVs, EVs ( $\Delta$ B2), or EVs (Mock). Error bars indicate standard deviation of three independent replicates. \*\*\* indicates  $P < 0.001$ , ns indicates no significant difference by Student's *t*-test.

G Experimental design for the *in vivo* functional analysis of EVs. In brief, mice were first immunized with live NoVΔB2, EVs (mock), or EVs ( $\Delta$ B2) and challenged with WT NoV at 12 h post immunization. The NoV-challenged mice were sacrificed at 1 and 3 days after challenge infection for determining the viral load or monitored daily for survival.

H, I Accumulation of NoV RNA1 was determined by RT-qPCR in NoVWT-infected BALB/c suckling mice at 1 (H) or 3 dpi (I).  $n = 5-7$  per group. Data are shown as mean  $\pm$  SD. \*\*\*\* indicates  $P < 0.0001$  by Student's *t*-test.

J Total proteins were extracted from the hindlimb muscle tissue of NoVWT-infected BALB/c suckling mice to measure the viral coat protein (CP) and B2 protein by Western blotting. Staining of  $\beta$ -actin was used as a loading control.

K Survival curve of WT NoV-challenged suckling mice immunized with live NoVΔB2, EVs ( $\Delta$ B2), or EVs (mock).  $n = 7$  per group. \*\*\* indicates  $P < 0.001$ . All data were analyzed by log rank test.



**Figure 5. The EV vsirRNAs of NoV induce homology-dependent virus resistance in AG6 mice.**

- A Schematic overview of antiviral function of EVs. Briefly, EVs from naïve or NoVΔB2-infected BALB/c suckling mice were extracted 3 days post infection, and then injected into AG6 suckling mice. After 12 h post immunization, AG6 suckling mice were challenged by WT NoV and then sacrificed at 1 dpi.
- B NoV replication was determined by RT–qPCR from hindlimb muscle of AG6 suckling mice challenged with NoVΔB2 as control or EVs from naïve or NoVΔB2-infected BALB/c suckling mice at 3 dpi. Normalization was done by β-actin mRNA levels. \* indicates  $P < 0.05$  by Student’s *t*-test.  $n = 3–4$  per group. Data are shown as mean ± SD.
- C Relative viral RNA1 accumulation levels at 1 dpi in NoV WT-infected AG6 mice 12 h after injection with EVs from NoVΔB2 or mock-inoculated BALB/c mice were measured by RT–qPCR. Normalization was done by β-actin mRNA levels. \*\*\*\* indicates  $P < 0.0001$  by Student’s *t*-test.  $n = 5–6$  per group. Data are shown as mean ± SD.
- D Diagram of recombinant SINV<sub>NoV</sub> (top) and SINV<sub>GFP</sub> (bottom). The gene of B2 protein was inserted at downstream of the duplicated subgenomic promoter sequence as shown.
- E AG6 suckling mice were challenged with EVs from naïve or NoVΔB2-infected BALB/c suckling mice and then inoculated by SINV<sub>NoV</sub> and SINV<sub>GFP</sub> respectively. Viral titer (PFU/ml) in the hindlimb muscle of infected mice was measured by a standard plaque assay and normalized by tissue mass, and the viral genomic RNA accumulation was determined by RT–qPCR amplification of the viral nsP1 coding region. Normalization was done by β-actin mRNA levels. \*\*\* indicates  $P < 0.001$ , ns indicates no significant difference by Student’s *t*-test,  $n = 5$  per group. Data are shown as mean ± SD.

2013). Notably, a previous study demonstrated EV-mediated delivery of herpesvirus-encoded miRNAs from infected cells into non-infected recipient cells (Kalamvoki *et al*, 2014). In this work, we show for the first time that abundant vsirRNAs accumulate in serum EVs upon infections by three distinct RNA viruses. We further show that both BHK cells and mice appear to be partially protected from reinfection with WT NoV when immunized by the EVs originated from the NoVΔB2-infected mice. The induced protection comes at least in part from nucleotide sequence homology-dependent antiviral immunity, suggesting a role for the systemic accumulation of vsirRNAs induced by NoVΔB2. However, other proteins or nucleic acids from the EVs with NoVΔB2 infection may also simultaneously vaccinate the naïve mice, allowing the mice to establish an immune protective effect.

In our current work, EVs were isolated from the serum by an exosome-enriched protocol. Several established methods used for EV isolation, such as differential ultracentrifugation, immunomagnetic-bead separation, density gradient centrifugation, chromatography, precipitation-based separation, and ultrafiltration, can also be used to isolate virions (Théry *et al*, 2018; Caobi *et al*, 2020). Thus, it is

often difficult to fully separate high-purity EVs from virions since they are of similar size, density, and biochemical content (Caobi *et al*, 2020). However, our isolated EVs from the serum of NoVΔB2-immunized suckling mice contained little infection-competent virions as verified in AG6 infant mice (Fig 5B). The small non-enveloped particles (28 nm in diameter) of NoV (Newman *et al*, 1978) and the severe *in vivo* defects of NoVΔB2 in replication (Li *et al*, 2013) may contribute to our successful separation of the EVs from the virions. In contrast, our use of the same isolation protocol in this study did not allow the purification of the EVs from the larger enveloped virions (65 nm in diameter) of SINV (Griffin, 2013), suggesting that optimization is necessary to obtain virion-free EVs from SINV-infected cells.

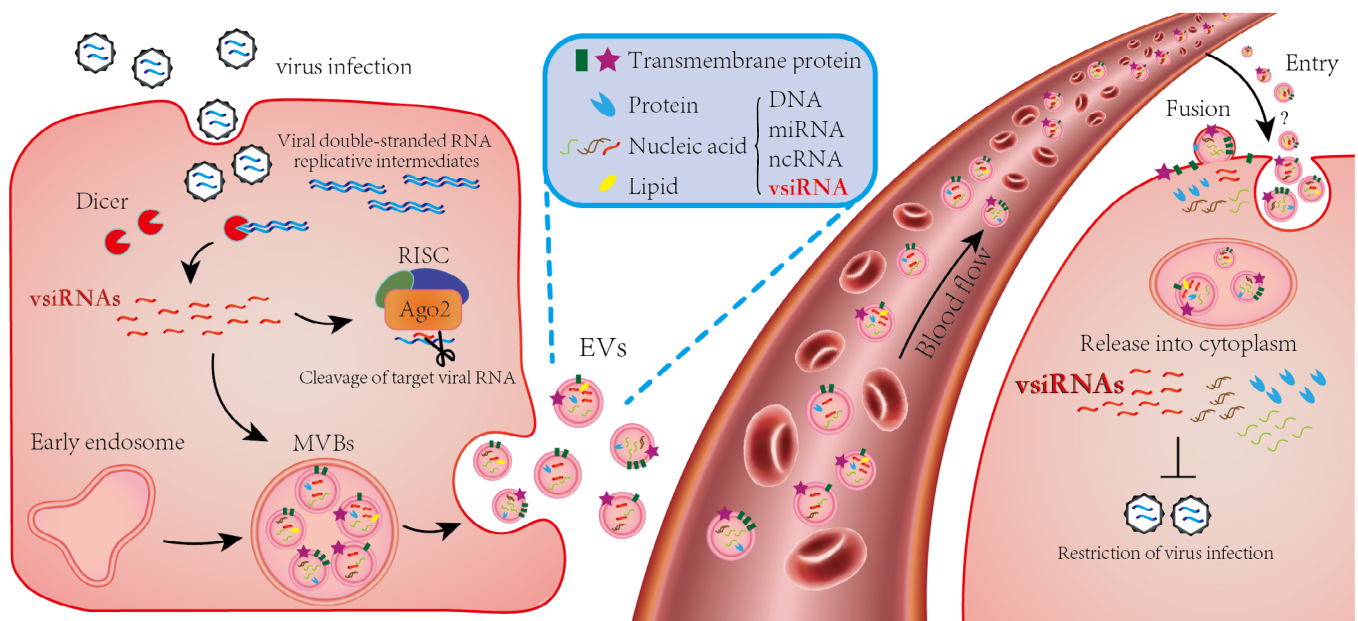
In addition to the ZIKV-induced antiviral RNAi response characterized in human NPCs (Xu *et al*, 2019), our recent study demonstrates that abundant vsirRNAs targeting ZIKV were also produced in the central neuron system (CNS) and muscle tissues of *Ifnar1*<sup>-/-</sup> mice upon virus infection (Zhang *et al*, 2020). A growing body of evidence indicates that EVs mediate the restriction of ZIKV infection (Caobi *et al*, 2020). For instance, Avraham *et al* showed that

exosomes carrying the chromosome 19 microRNA cluster (C19MC) of miRNAs attenuate ZIKV infection of U2OS cells in an IFN-independent manner (Bayer *et al*, 2018). Janis *et al* showed that semen prevents ZIKV attachment to the target cells by the anti-ZIKV activity of EVs; however, which components of semen EVs confer this inhibition remains unknown (Müller *et al*, 2018). Here, we detected abundant 21- to 23-nt vsiRNAs (equivalent to 14.9% of the total EVs miRNAs) in the serum EVs from ZIKV-infected mice, indicating that these EVs-associated vsiRNAs may have potential antiviral functions. Recently, Wu and colleagues provide evidence that EVs engineered to contain a host restriction factor, IFN-induced transmembrane protein 3 (IFITM3), can effectively move across the placental barrier to suppress ZIKV infection of fetuses (Zou *et al*, 2021). Thus, knowledge of the phenotypes and functions of EVs generated in response to natural viral infection may be applied to improve anti-virus treatment in the future.

Fang *et al* (2021) have recently achieved antiviral protection in mice using synthetic peptides designed to specifically inactivate the VSR of HEV71, demonstrating the functional importance of antiviral RNAi in mammals and a new therapeutic strategy of antiviral treatment. Maillard *et al* (2016) have shown that transfection with artificial long dsRNA specifically vaccinates cultured IFN-deficient cells against infection with viruses bearing a homologous sequence by RNAi. However, whether viral dsRNA replicative intermediates made during the authentic virus infections can trigger similar immune protection in an IFN-competent animal is not known. Results from this study show that infant mice were fully protected against lethal NoV challenge only one or two days after immunization with NoVΔB2, a live-attenuated mutant of NoV rendered defective to suppress vsiRNA biogenesis. Importantly, the vsiRNAs

accumulated in the hindlimb muscle tissues and the serum EVs of NoVΔB2-immunized mice were similar in the size distribution, strand ratio, 1 U preference, and abundance and these vsiRNAs were stably maintained in the immunized mice up to 27 days post vaccination and at least more than one week after the mutant virus is cleared. These findings suggest a role for the EVs in the spread of the protective immunity induced by NoVΔB2 immunization to uninfected cells and a new strategy for the development of animal and human RNA virus vaccines using mutant viruses attenuated by inactivating their VSRs.

In summary, we show for the first time that mammalian vsiRNAs of three positive-strand RNA viruses entered the blood stream for systemic circulation via EVs. The population of EV-associated vsiRNAs is indistinguishable from those found in the limb muscle tissues of the same mice immunized with NoVΔB2. Notably, immunization of infant mice with either NoVΔB2 or vsiRNA-containing EVs induced specific protective immunity against challenge infection, suggesting a critical role for the systemic spread of vsiRNAs. Similar to antiviral RNAi in fruit flies (Li *et al*, 2002; Galiana-Arnoux *et al*, 2006; Van Rij *et al*, 2006; Wang *et al*, 2006; Tassetto *et al*, 2017), antiviral RNAi is not only active in the infected cell but also sends immune signals to prevent viral infection in distant non-infected cells in mammals (Fig 6). In addition to the biogenesis of vsiRNAs from Dicer processing of the viral dsRNA replicative intermediates, insect vsiRNAs are further amplified using viral DNA templates reverse-transcribed from the invading RNA viruses (Goic *et al*, 2013; Tassetto *et al*, 2017; Poirier *et al*, 2018). There are reports of viral DNA synthesis from non-retroviral RNA viruses by endogenous reverse transcriptases in mammalian cells (Klenerman *et al*, 1997; Geuking *et al*, 2009; Shimizu *et al*, 2014). Thus, it will be



**Figure 6. The EV-vsiRNAs confer antiviral activity upon antiviral RNAi activation.**

VsiRNAs are induced in virus-infected cells by dicing the viral double-stranded RNA replicative intermediates and loaded into RISC for subsequent cleavage of viral RNA genomes. Part of the vsiRNAs enter the EVs together with miRNAs, ncRNAs, proteins, and lipids. EVs in MVBs are secreted to the extracellular system. When attached to recipient cells, vsiRNAs in EVs are released into the cytoplasm to restrict the virus replication.

interesting to investigate a possible contribution of an insect-like vsRNA amplification mechanism to the prolonged accumulation of vsRNAs and the spread of the induced protection in mice immunized by VSR-deficient mutant of NoV.

## Materials and Methods

### Viruses and cell culture

Wild-type NoV and mutant NoV $\Delta$ B2 strains used in this study were as described previously (Li *et al*, 2013). SINV was rescued from the plasmid of pSVN1, a gift from Dr. C.M. Rice. ZIKV strain (SZ01) was provided from Shanghai Public Health Clinical Center, Fudan University. ZIKV stocks were propagated in Vero cells after inoculating at a multiplicity of infection (MOI) of 0.01 and harvesting supernatants after 5 days post infection. Viruses were harvested and titered as previously described (Zhang *et al*, 2020). Human embryonic kidney (293T) cell was gift from Dr. B. Cullen. Baby hamster kidney cells (BHK) and African green monkey kidney epithelial cells (Vero) were purchased from the American Type Culture Collection (ATCC). All cells were cultured in Dulbecco's modified Eagle's medium (DMEM, Gibco) containing 10% fetal bovine serum (Gibco).

### Animals

BALB/c and C57BL/6 mice were purchased from Shanghai SLAC Laboratory Animal Co., Ltd. and Jackson labs (Bar Harbor, ME). *Ifnar1*<sup>-/-</sup> mice (A6) were bought from Cyagen Biosciences (Suzhou, China). *Ifnar1*<sup>2-/-</sup> mice (AG6) were a gift from Dr. Y. Xu. All the animal experiments in China were carried out under the guidelines of the Institutional Animal Care and Use Committee, Fudan University of China. Animals in USA were housed in the Animal Resources Facility according to the guidelines described under the federal Animal Welfare Regulations Act and the procedures were approved by the Institutional Animal Care and Use Committee at the University of California, Riverside.

### NoV infection

WT NoV or NoV $\Delta$ B2 preparations shown to contain  $7 \times 10^6$  copies of genomic RNA1 from the titrated set of stocks was inoculated to each of suckling mice of 6 to 8 days old after birth by intraperitoneal injection (i.p.) as described previously (Li *et al*, 2013). Total RNAs were extracted from the hindlimb muscle tissues of mice after infection with NoV $\Delta$ B2, which were used to determine the accumulation of NoV $\Delta$ B2 RNA1 by RT-qPCR and the construction of small RNA libraries as described. Whole blood samples were obtained from NoV $\Delta$ B2 infected suckling mice at 3 dpi and 9 dpi, and then used to isolate EVs.

### Vaccination

For the suckling mouse immunization and challenge study, 6-day-old BALB/c mice were inoculated by i.p. with NoV $\Delta$ B2 (shown to contain  $7 \times 10^6$  copies of genomic RNA1) either with or without UV-inactivation, or with the same volume of DMEM (mock). To inactivate NoV $\Delta$ B2 by UV, virus stocks were placed on ice in a

XL-1000 UV crosslinker (Spectronics corporation, Rolling Meadows, IL), and irradiated twice at  $1 \text{ J/cm}^2$  for 30 min. Virus inactivation was verified by *in vivo* infection. Two days after immunization, mice were challenged by i.p. inoculation with WT NoV (shown to contain  $7 \times 10^6$  copies of genomic RNA1 for BALB/c suckling mice). One suckling mouse immunized with DMEM, UV-inactivated NoV $\Delta$ B2, or NoV $\Delta$ B2 was euthanized 2, 3, or 4 days after WT NoV challenge to determine virus titers in the hind limb tissues by RT-qPCR and Western blotting. All suckling mice immunized with DMEM or UV-inactivated NoV $\Delta$ B2 were euthanized 4 days after WT NoV challenge since they began to show hind limb paralysis. The vaccination experiments were repeated two additional times.

### Construction of SINV<sub>NoV</sub>

The pSVN1 and pTE/5'2J/GFP (SINV expression EGFP) and pTE/5'2J were gifts from Dr C.M. Rice (Rockefeller University, New York, NY, USA). In pTE/5'2J, the gene of interest can be inserted at a multiple cloning site (MCS) downstream of the duplicated subgenomic promoter sequence (Pierro *et al*, 2003). pTE/5'2J/NoV were constructed by ligating PCR products of NoV RNA1 flanked by *Xba*I sites at 5' and *Apa*I sites at 3' into the MCS of pTE/5'2J.

### SINV Viruses production

SINV, SINV<sub>GFP</sub>, and SINV<sub>NoV</sub> viruses were rescued separately from the plasmids of pSVN1, pTE/5'2J/GFP, and pTE/5'2J/NoV. Briefly, plasmids containing SINV genomic RNAs were linearized with *Xho*I and transcribed *in vitro* using an SP6 mMESSAGE mMACHINE kit (Ambion). Purified SINV genomic RNAs were transfected into BHK cells by TransIT<sup>®</sup>-mRNA Transfection Kit (Mirus Bio, WI). Viruses were harvested and titered as previously described (Zhang *et al*, 2021).

### SINV and ZIKV infection

50 PFU of SINV was inoculated to C57BL/6 suckling mice and  $10^4$  PFU of ZIKV was inoculated to *Ifnar1*<sup>-/-</sup> suckling mice by i.p. Whole blood samples were obtained from SINV-infected suckling mice at 2 dpi and ZIKV-infected suckling mice at 4 dpi, and then used to isolate EVs.

### EVs isolation

Whole blood samples were obtained from infected or mock-infected suckling mice at respective dpi for each virus. The whole blood samples were placed in 37°C, thermostated for 15 min, then spun at 3,000 g, 30 min to collect serum. Centrifugation was performed twice to remove cells and cell debris, and serum were used to isolate EVs according to manufacturer's protocol of Exosome isolation kits: ExoQuick Exosome Isolation Reagent (System Biosciences, Palo Alto, CA, USA). Serum was placed into 1.5 ml tube with ExoQuick Exosome Precipitation Solution at a ratio of 4:1. Mixture were thermostated at 4°C for an hour then centrifuged at 1,500 g for 30 min and supernatant were aspirated. EVs were resuspended in 1× PBS. The isolated EVs were prepared for Transmission Electron Microscope (TEM) (FEI, Holland) and Nanoparticle Tracking Analysis (NTA) (Particle Metrix-PMX, Germany). The presence of five

exosome marker proteins, HSP70, tsg101, CD81, CD63, and CD9, were confirmed by Western blotting. EVs obtained by precipitation were quantified for particle number by NTA. The concentration of EVs stock was suspended in PBS to  $8 \times 10^{11}$  particles/ml. EVs, whole blood, and blood clot derived from different virus-infected suckling mice were used for RNA extraction by TRIzol (Invitrogen, Carlsbad, CA). Total RNAs obtained were used for the construction of small RNA libraries as described, and proteins samples were used for Western blot analysis (Li *et al*, 2017).

### Function assay of EV *in vitro* and *in vivo*

For *in vitro* study, BHK cells were seeded in a 12-well plate at a density of  $5 \times 10^5$ /well one day before infection. Cell culture medium was removed to opti-MEM and then the cells was inoculated with 10  $\mu$ l or 50  $\mu$ l EVs stock (stock concentration,  $8 \times 10^{11}$  particles/ml) and then infected with NoV WT (shown to contain  $7 \times 10^5$  copies of genomic RNA1). The cells were cultured at 30°C for next 24 h, and were harvested for the extraction of total RNAs using TRIzol according to the manufacturer's protocol. For *in vivo* study, 7-day-old BALB/c and AG6 suckling mice were inoculated with NoV $\Delta$ B2 (shown to contain  $7 \times 10^6$  copies of genomic RNA1),  $2.4 \times 10^{10}$  EVs (from NoV $\Delta$ B2-infected mice), or  $2.4 \times 10^{10}$  EVs (from mock-infected mice). 12 h after inoculation, the mice were infected with WT NoV (shown to contain  $7 \times 10^5$  copies of genomic RNA1; note, it is the 1/10 dose of vaccination experiment as described previously) by i.p. The hindlimb muscle tissues were isolated to obtain total RNAs and proteins for detecting RNA1 accumulation, Capsid, and B2 protein expression at 1 and 3 dpi. Survival was monitored daily. To exclude the effect of NoV particle existing in EVs, NoV replication was determined by RT-qPCR from hindlimb muscle of AG6 suckling mice challenged with NoV $\Delta$ B2 as control or EVs from naïve or NoV $\Delta$ B2-infected BALB/c suckling mice at 3 dpi.

### 293T Ago2-KO cell generation

293T cells were seeded in a 60 mm dish the day before transfection with 6  $\mu$ g eIF2c (Ago2)-targeting CRISPR plasmids (Santa Cruz sc-400813) by lipofectamine 2000 according to the manufacturer's instructions. Two days after transfection, culture medium was replaced with DMEM containing puromycin at a dose of 2  $\mu$ g/ml. Single clones were separated at 2 weeks post treatment. Positive Ago2-KO clones were identified and confirmed by siRNA knockdown analysis. In brief, siRNAs targeting GAPDH mRNA or control siRNAs were transfected into the Ago2-KO cells and the loss of knockdown ability was determined by comparing the expression levels of GAPDH mRNA in 293T and Ago2-KO cells after transfection with the specific and control siRNAs.

### Luciferase reporter assay

The pmirGLO-NoV/GFP reporters were constructed by cloning a fragment (nt 2743–3204) of NoV $\Delta$ B2 genomic RNA1 in sense orientation (to be targeted by the antisense vsRNAs) or GFP (nt 4-465 of the coding sequence) into pmirGLO plasmid at XbaI site, by ClonExpress II One Step Cloning Kit (Vazyme Biotech Co., Ltd). Luciferase reporter assay was performed as described by the manufacturer's instructions. Briefly, 293T cells in 12-well plates were transfected with 1  $\mu$ g

of pmirGLO-NoV/GFP plasmid DNA by lipofectamine 2000. At 4 h post-transfection, culture medium was replaced with fresh medium containing pre-diluted EVs at a density of  $5 \times 10^9$  particles/well. Luciferase activity was analyzed using the Dual Luciferase Reporter Assay Kit (Vazyme Biotech Co., Ltd) and measured on Fluoreskan Ascent FL (Thermo Scientific) after 24 h post-transfection. The normalized relative luciferase activity was calculated by dividing the firefly luciferase value with Renilla luciferase value.

### Western and Northern blot analyses

Western blotting analysis was performed as described previously (Li *et al*, 2017). Antibodies to NoV B2 and coat protein were described previously (Li *et al*, 2013). Antibodies to exosomal marker proteins, CD9, CD63, CD81, HSP 70, and tsg 101 (sc-13118, sc-5275, sc-166029, sc-24, sc-7964), were purchased from Santa Cruz Biotechnology, Santa Cruz, CA.

### *In vivo* recombinant SINV reporter experiments

For AG6 suckling mice, 7-day-old mice were inoculated by i.p. with  $2.4 \times 10^{10}$  EVs (from NoV $\Delta$ B2-infected mice) or  $2.4 \times 10^{10}$  EVs (from mock-infected mice). 12 h after inoculation, the mice were infected by i.p. with SINV<sub>GFP</sub> or SINV<sub>NoV</sub> viruses of 500 PFU. Each group of four suckling mice were euthanized one day after SINV infection to determine virus titers in the hind limb tissue by viral plaque assays as described (Zhang *et al*, 2021). In brief, hindlimb muscles were isolated from infected or mock-infected suckling mice. After homogenization, 1 ml DMEM were added to per 50 mg tissue. The tissue homogenates were used for plaque assay. BHK cells were plated at a density of  $5.0 \times 10^5$  cells/ well in 3 ml DMEM, 10% FBS, on 6-well plates and incubated at 37°C. On the next day, 1 ml of 10-fold gradient dilutions of supernatants was added to each well. After incubation of 1 h, supernatants were discarded and the cells were overlaid with 5 ml 1 $\times$ MEM containing 0.5% agarose in each well. One day after incubation, the agarose gels were removed, and cell layer was stained with Coomassie Brilliant Blue staining solution. Virus content of the supernatants was calculated as plaque forming units (PFU)/ml.

### Long-term protection

Six-day-old BALB/c suckling mice were vaccinated with NoV $\Delta$ B2, UV-inactivated  $\Delta$ B2, EVs ( $\Delta$ B2), or EVs (Mock) as the same dose of NoV vaccination or *in vivo* function assay of EVs. At 21 days post treatment, the mice were infected with WT NoV by intraperitoneal injection (shown to contain  $2.8 \times 10^6$  copies of NoV genomic RNA1, grown mice were challenged four times dose of virus as much as suckling mice based on body weight). Infected mice were sacrificed at 3 dpi, and relative accumulations of NoV RNA1 were determined by RT-qPCR.

### RNA extraction, RT-qPCR, and data analysis

For virus infection and exosomal function assay *in vitro* and *in vivo*, total RNAs were extracted using Trizol reagent (Thermo) from hindlimb muscle of infected mice according to the manufacturer's instructions as described (Zhang *et al*, 2020) and reverse-

transcription were done by 1<sup>st</sup> strand synthesis kit (Vazyme Biotech Co., Ltd). Virus accumulation of mice tissues was determined by RT-qPCR using ChamQ Universal SYBR qPCR Master Mix (Vazyme Biotech Co., Ltd). Primers were shown in Table EV2. GraphPad Prism was used for all statistical analyses, which was carried out by unpaired *t*-test or the log rank test for the survival data. All experiments were repeated at least three times.

### Construction of small RNA libraries

RNA preparations in this study were used for the construction of small RNA libraries by the method that depends on the 5' monophosphate of small RNAs as described previously with the TruSeq Small RNA Sample Preparation Kit of Illumina (San Diego, CA).

### Deep sequencing and bioinformatic analysis of small RNAs

Libraries of small RNAs were cloned from the RNA samples (mice  $n = 3$ , in each independent analysis) and sequenced by Illumina HiSeq 2000/2500. 26 libraries in total were sequenced from this work (Table EV1). Adapters were removed and reads containing any N base were removed. Trimmer reads were then mapped to virus genome references or compared to mature miRNAs of mouse (mmu21-). Alignment was done by Bowtie 1.2.2 with perfect match. Reads mapped to positive and negative strands were used to define positive and negative vsRNAs separately. For scheme of vsRNAs size distribution and genome mapping results, reads were normalized to per million miRNA. All the references used were downloaded from web sources. Subsequent bioinformatic analysis of virus-derived small RNAs was carried out using in-house Perl scripts as described previously. 22-nt vsiRNAs in each library were used to calculate overhang length of complemented duplexes by a previously described algorithm (Li *et al*, 2017). Detailed information of vsRNAs in each library are shown in Table EV1. The reference sequences used in this study are either identical with those described previously or as listed below:

- 1 NoV RNAs 1 and 2: AF174533.1 and AF174534.1
- 2 NoVΔB2 RNAs 1 and 2: the same as NoV except for 3 substitutions in RNA1: U2745C, U2754C, C2757G.
- 3 SINV: J02363.1
- 4 ZIKV: KX253996.1
- 5 Mature miRNAs and miRNA precursors: miRBase 21 (<http://www.mirbase.org/>).

### Data availability

The RNA sequencing data have been deposited to the database under the accession number PRJNA716839. All SRR data can be achieved through SRA-toolkit. Go to: <https://www.ncbi.nlm.nih.gov/Traces/study/?acc=PRJNA716839>

**Expanded View** for this article is available online.

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### Author contributions

**Yang Li:** Conceptualization; Resources; Data curation; Formal analysis; Supervision; Funding acquisition; Validation; Investigation; Methodology; Writing—original draft; Project administration; Writing—review & editing. **Yuqiang Zhang:** Data curation; Formal analysis; Validation; Methodology; Writing—original draft. **Yunpeng Dai:** Data curation; Software; Formal analysis; Validation; Visualization; Methodology. **Jiaxin Wang:** Formal analysis; Validation; Methodology. **Yan Xu:** Software; Formal analysis. **Zhe Li:** Formal analysis; Methodology. **Jinfeng Lu:** Resources; Software. **Yongfen Xu:** Resources. **Jin Zhong:** Resources. **Shou-Wei Ding:** Conceptualization; Supervision; Funding acquisition; Writing—original draft; Writing—review & editing.

In addition to the CRediT author contributions listed above, the contributions in detail are:

YZ, YD, JW, ZL, and YL performed all infection experiments. YD and YX performed all bioinformatics analyses of small RNA libraries. YFX and JZ assisted with infection experiments in mice. JL assisted with writing the manuscript. YL designed experiments and interpreted results. YL and S-WD conceived the study and wrote the final manuscript.

### Disclosure and competing interests statement

SWD and YL declare to be named as inventors on an issued patent (US 10,034,929B2), claiming subject matter related to the use of an attenuated virus defective in RNAi suppression as a vaccine to induce antiviral protection described in this paper. The other authors declare that they have no conflict of interest.

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