

Role of zinc-finger anti-viral protein in host defense against Sindbis virus

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Abstract

Accumulating evidence indicates that type I interferon (IFN) mediates the host protective response to RNA viruses. However, the anti-viral effector molecules involved in this response have not been fully identified. Here, we show that zinc-finger anti-viral protein (ZAP), an IFN-inducible gene, plays a critical role in the elimination of Sindbis virus (SINV) *in vitro* and *in vivo*. The loss of ZAP greatly enhances the replication of SINV but does not inhibit type I IFN production in primary mouse embryonic fibroblasts (MEFs). ZAP binds and destabilizes SINV RNA, thereby suppressing the replication of SINV. Type I IFN fails to suppress SINV replication in ZAP-deficient MEFs, whereas the ectopic expression of ZAP is sufficient to suppress the replication of SINV in MEFs lacking the expression of type I IFN and the IFN-inducible genes. ZAP-deficient mice are highly susceptible to SINV infection, although they produce sufficient amounts of type I IFN. Therefore, ZAP is an RNA-sensing anti-viral effector molecule that mediates the type-I-IFN-dependent host defense against SINV.

Keywords: host defense, IFN, innate immunity, pattern-recognition receptor, RNA degradation

Introduction

Pattern-recognition receptors (PRRs) detect viral nucleic acids and induce an anti-viral innate immune response (1, 2). Retinoic acid-inducible gene I (RIG-I), also called DEAD (Asp-Glu-Ala-Asp) box polypeptide 58 (DDX58), and melano-ma differentiation-associated gene 5 (MDA5), also called interferon (IFN)-induced with helicase C domain 1 (IFIH1), are RNA helicases that detect cytosolic viral double-stranded (ds) RNAs (3, 4), whereas cyclic GMP-AMP synthase (cGAS) detects cytosolic viral dsDNA (5, 6). After cytosolic viral nucleic acids are detected, RIG-I-like receptors (RLRs) and cGAS trigger the activation of IFN regulatory factor 3 (IRF3), a transcription factor that induces the expression of type I IFN and IFN-inducible genes. Endosomal Toll-like receptor 7 (TLR7) and TLR9 detect extracellular viral single-stranded (ss) RNA and ssDNA, respectively, and activate IRF7 to induce the expression of type I IFN and the IFN-inducible genes (7–10). Thus, nucleic-acid-sensing PRRs are essential to the establishment of an anti-viral state by mediating the expression of type I IFN and IFN-inducible genes.

The anti-viral effector molecules responsible for the type-I-IFN-mediated host defense against viruses are not fully understood. Among the IFN-inducible proteins, zinc-finger anti-viral protein (ZAP), also called zinc-finger CCCH type, anti-viral 1 (ZC3HAV1), is a well-characterized anti-viral effector molecule (11). ZAP was originally identified as the anti-viral effector molecule that inhibits murine leukemia virus (MLV) infection (12). ZAP binds to MLV transcripts via its CCCH-type zinc-finger (C3H ZF) domains and reduces the level of MLV transcripts (13). After it detects the MLV transcripts, ZAP associates with the exosome complex, an RNA degradation structure, and destabilizes the MLV transcripts (14). Endogenous ZAP is involved in the anti-MLV response in mouse embryonic fibroblasts (MEFs) (15). Recent studies have shown that Sindbis virus (SINV), Ebola virus, Marburg virus, MLV, HIV-1 and hepatitis B virus (HBV) are also targeted by ZAP (12, 16–19). However, the roles of ZAP in type-I-IFN-mediated virus elimination and in anti-viral responses *in vivo* are unclear. In this study, we examined these issues and

show the essential involvement of ZAP in the host defense against viruses.

Methods

Reagents

High-molecular-weight poly rI:rC, LPS, R848, oligodeoxynucleotide (ODN) 1585, ODN 1668, 5'ppp-dsRNA and IFN stimulatory DNA (ISD) were purchased from InvivoGen. Anti-VSV-G (A190-131A) antibody was purchased from Bethyl Laboratories. Anti-Myc antibody (2276S) and mouse IgG1 isotype control antibody (5415S) were purchased from Cell Signaling Technology. Horseradish-peroxidase-conjugated anti- β -actin antibody (sc-1615) was purchased from Santa Cruz Biotechnology. Recombinant mouse IFN- β was purchased from ProSpec. Recombinant mouse granulocyte macrophage colony-stimulating factor (GM-CSF) and human Fms-like tyrosine kinase 3 ligand (FLT3L) were purchased from Peprotech. Lipofectamine™ 2000 was purchased from Invitrogen. Anti-SINV E2 antibody was originally developed by D. E. Griffin (Johns Hopkins University School of Medicine, USA) and obtained from Y. Yoshinaka (Tokyo Medical and Dental University, Japan).

Plasmids

The retroviral expression vector pMRX-ires-puro was kindly donated by S. Yamaoka (Tokyo Medical and Dental University, Japan). The cytomegalovirus-promoter-driven expression vector pcDNA3.1(+) was purchased from Invitrogen. A synthesized cDNA fragment encoding the Myc-tag sequence was cloned into pMRX-ires-puro and pcDNA3.1(+), generating pMRX-Myc-ires-puro and pcDNA3-Myc, respectively. cDNA fragments encoding ZAP-L, ZAP-S, ZAP-N and ZAP-C were amplified from an MEF cDNA library with PCR and cloned into pMRX-Myc-ires-puro, generating pMRX-ZAP-L-Myc-ires-puro, pMRX-ZAP-S-Myc-ires-puro, pMRX-ZAP-N-Myc-ires-puro and pMRX-ZAP-C-Myc-ires-puro, respectively. A cDNA fragment encoding ZAP-L was cloned into pcDNA3-Myc, generating pcDNA-ZAP-L-Myc. pEGFP-N1 was purchased from Clontech. pdsTE12Q was kindly provided by B. Levine (University of Texas Southwestern Medical Center, Dallas, TX, USA).

Mice and cells

The *Zc3hav1*^{+/-} mice (C57BL/6 and 129 mixed background) have been described previously (15) and were backcrossed to C57BL/6 mice over eight generations. Age-matched C57BL/6 mice were used as the control *Zc3hav1*^{+/+} mice. The *Ddx58*^{+/-}/*Ilfih1*^{+/-} mice have been described previously (20). The *Irf3*^{+/-}/*Irf7*^{+/-} mice were kindly donated by T. Taniguchi (The University of Tokyo, Japan). All animal experiments were performed in accordance with the guidelines of Osaka University and were approved by the Ethics Committee of the Research Institute for Microbial Diseases of Osaka University.

The *Zc3hav1*^{+/-}, *Ddx58*^{+/-}/*Ilfih1*^{+/-} and *Irf3*^{+/-}/*Irf7*^{+/-} mice were crossed separately, and primary MEFs were prepared from pregnant female mice on embryonic day 13.5, as described previously (15). Spontaneously immortalized wild-type MEFs have been described previously (21). Primary MEFs were used in most *in vitro* experiments. Because the transfection efficiency of the plasmid is quite low in primary MEFs, immortalized

wild-type MEFs, which are competent for plasmid transfection, were used in the experiment shown in Fig. 5(A). Plate-E packaging cells were kindly donated by T. Kitamura (The University of Tokyo). Vero cells have been described previously (22). The MEFs, Plat-E cells and vero cells were maintained in Dulbecco's modified Eagle's medium supplemented with 10% fetal bovine serum, penicillin (100 U ml⁻¹) and streptomycin (100 U ml⁻¹). To prepare the primary dendritic cells, mouse bone-marrow cells were treated for 6 days with GM-CSF (10 ng ml⁻¹) or FLT3L (20 ng ml⁻¹) in RPMI 1640 medium supplemented with 10% fetal bovine serum, 50 mM 2-mercaptoethanol, penicillin (100 U ml⁻¹) and streptomycin (100 U ml⁻¹).

Viruses

SINV and VSV have been described previously (4, 23). SINV RNA was synthesized from pdsTE12Q using SP6 RNA polymerase (Takara Bio), according to the manufacturer's protocol. In a mouse model of SINV infection, 10-day-old pups were subcutaneously inoculated with 100 plaque-forming units (pfu) of SINV. The viral titers were determined with 50% tissue culture infectious dose (TCID₅₀) assays, as described previously (22). A retroviral vector was produced by Plat-E cells transfected with pMRX-ires-puro, as described previously (24).

Enzyme-linked immunosorbent assays

The ELISA kits for IFN- α and IFN- β were purchased from PBL Interferon Source. The ELISA kits for CXCL10 and IL-6 were purchased from R&D Systems. The levels of IFN- α , IFN- β , IL-6 and CXCL10 were measured with ELISAs, according to the manufacturers' instructions.

RNA immunoprecipitation assay

The interaction between SINV RNA and ZAP was detected with the Ribocluster Profiler™/RIP-Assay kit (MBL), according to the manufacturer's instructions.

Immunoblotting

Total cell lysates were separated by SDS-PAGE and immunoblotted with the indicated antibodies, as described previously (24).

Northern blot analysis

Total RNA was isolated from cells with an RNeasy Kit (Qiagen), according to the manufacturer's instructions. The RNA samples (8 μ g each) were separated by electrophoresis, transferred to a nylon membrane and hybridized for 12 h with the indicated ³²P-labeled probes, prepared with the MegaPrime™ DNA Labelling System (GE Healthcare). The nylon membrane was washed with 2 \times saline sodium citrate (SSC) and 0.2 \times SSC and exposed to x-ray film to quantify the relative RNA levels. The probe was designed to target SINV RNA nucleotides 4048–4259.

Quantitative reverse transcription-PCR

Total RNA was isolated from cultured cells with the ZR RNA MicroPrep™ kit (Zymo Research). Total RNA was isolated

from the brain with an RNeasy Lipid Tissue Mini Kit (Qiagen). Reverse transcription (RT) was performed with ReverTra Ace® reverse transcriptase (Toyobo). For the quantitative PCR, cDNA fragments were amplified from the RT products with Realtime PCR Master Mix (Toyobo). The fluorescence from the TaqMan probe was detected with the StepOnePlus™ Real-Time PCR System (Applied Biosystems). The expression of *Irfn1*, *Cxcl10*, *IL-1b* and *IL-6* mRNAs was normalized to that of β -actin mRNA. The experiments were performed in accordance with the manufacturer's instructions.

Statistical analysis

The Student's *t*-test was used to analyze all data. A *P* value of <0.05 was considered significant.

Results

ZAP suppresses SINV replication in primary MEFs

SINV is a positive-sense RNA virus belonging to the genus *Alphavirus* in the family *Togaviridae*. SINV causes fever, rash, arthralgia and encephalopathy in humans and animals (25). Type I IFN is critically involved in the host defense against SINV (26). Therefore, we used SINV as the model pathogen in the present study. We first examined the involvement of endogenous ZAP in the anti-viral response to SINV in primary MEFs. The replication of SINV was greatly enhanced in primary *Zc3hav1*^{-/-} MEFs (Fig. 1A and C), although the replication of vesicular stomatitis virus (VSV) was normal (Fig. 1B and D). These findings indicate that endogenous ZAP contributes to the suppression of SINV replication in primary MEFs.

The ZAP long isoform has zinc-finger domains at its N-terminus and a poly(ADP-ribose) polymerases (PARP) domain at its C-terminus, whereas the ZAP short isoform has zinc-finger domains at the N-terminus but lacks the PARP domain (Fig. 2A). Previous studies have shown that the zinc-finger domains of ZAP are sufficient to trigger an anti-viral response, but another study has shown that both

the zinc-finger and PARP domains of ZAP are involved in the anti-viral response in the 293 cell line (27, 28). Therefore, the role of the PARP domain of ZAP in the anti-viral response is still unclear. We assessed the involvement of the zinc-finger and PARP domains of ZAP in the anti-viral response to SINV. The complementation of the ZAP N-terminal portion, but not that of ZAP C-terminal portion restored the anti-viral activity against SINV in *Zc3hav1*^{-/-} MEFs (Fig. 2B), confirming the

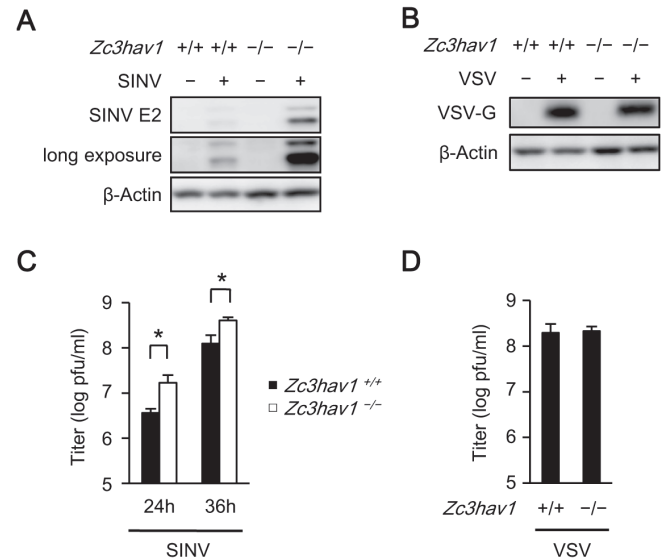


Fig. 1. Loss of ZAP enhances the replication of SINV in primary MEFs. (A and B) Primary *Zc3hav1*^{+/+} and *Zc3hav1*^{-/-} MEFs were infected with SINV [multiplicity of infection (MOI) = 0.1] (A) or VSV (MOI = 0.1) (B) for 24 h. The levels of viral proteins and β -actin in the cell lysates were measured with immunoblotting. (C and D) Primary *Zc3hav1*^{+/+} and *Zc3hav1*^{-/-} MEFs were infected with SINV (MOI = 0.1) (C) for the indicated periods or VSV (MOI = 0.1) (D) for 24 h. The viral titers in the culture supernatants were determined with TCID₅₀ assays. The experiments were performed three times and representative data are shown (means \pm SD of three independent samples). **P* < 0.05.

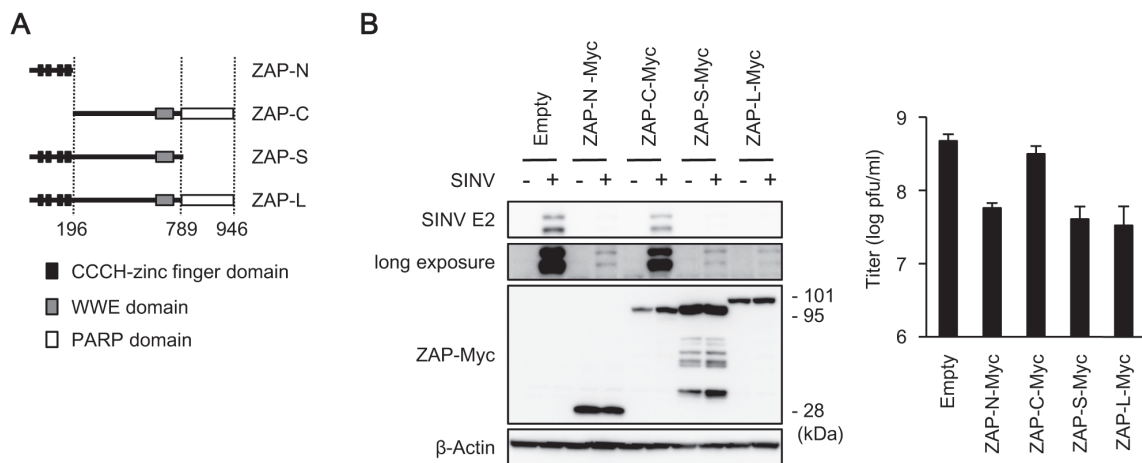


Fig. 2. N-terminal zinc-finger domains of ZAP mediate the anti-viral response to SINV. (A) Schematic of ZAP. (B) Primary *Zc3hav1*^{-/-} MEFs retrovirally transduced with ZAP-N-Myc, ZAP-C-Myc, ZAP-S-Myc and ZAP-L-Myc were infected with SINV (MOI = 0.05) for 24 h. The levels of viral proteins, β -actin and Myc-tagged proteins in the cell lysates were measured with immunoblotting. The viral titers in the culture supernatants were determined with TCID₅₀ assays. The experiments were performed at least three times and representative data are shown (means \pm SD of three independent samples).

importance of the zinc-finger domains of ZAP in the suppression of SINV replication. Complementation of both the ZAP long isoform and the ZAP short isoform in *Zc3hav1^{+/-}* MEFs restored the anti-viral activity against SINV (Fig. 2B), indicating that the PARP domain of ZAP is not essential for the suppression of SINV replication in primary MEFs.

ZAP reduces SINV RNA levels independently of type I IFN production

Type I IFN and chemokines are critically involved in anti-viral host defenses (29). ZAP is reported to be involved in the regulation of RIG-I signaling in the 293 cell line (30). Therefore, we assessed the involvement of ZAP in the production of these factors. The SINV-induced production of IFN- β , a type I IFN, and chemokine (C-X-C motif) ligand 10 (CXCL10), an IFN-inducible chemokine, was enhanced in *Zc3hav1^{+/-}* MEFs and *Zc3hav1^{+/-}* conventional dendritic cells (Fig. 3A and B). However, the VSV-induced production of IFN- β and CXCL10 was normal in *Zc3hav1^{+/-}* MEFs (Fig. 3A). Furthermore, the production of IFN- β and CXCL10 was normal in *Zc3hav1^{+/-}* MEFs stimulated with the RLR and cGAS ligands (Fig. 4A). These findings indicate that ZAP is not essential for the RLR- or cGAS-mediated expression of type I IFN or IFN-inducible genes in primary mouse cells and suggests that an increase in SINV causes the enhanced activation of RLR in primary *Zc3hav1^{+/-}* MEFs. *Zc3hav1^{+/-}* conventional dendritic cells normally produce CXCL10 after they are stimulated with the TLR4, TLR7 and TLR9 ligands (Fig. 4B). *Zc3hav1^{+/-}* plasmacytoid

dendritic cells normally produce IFN- α and IFN- β after they are stimulated with the TLR9 ligand (Fig. 4C). These findings also indicate that ZAP is not essential for the TLR-mediated expression of type I IFN or IFN-inducible genes.

We next examined whether ZAP directly affects SINV replication. Although a previous study showed that ZAP reduced the level of luciferase mRNA fused with fragmented SINV RNA (13), there is as yet no direct evidence to support the destabilization of SINV RNA by ZAP or to confirm that ZAP binds to SINV RNA. Ectopically expressed ZAP greatly reduced the level of transfected SINV RNA in wild-type MEFs (Fig. 5A). Stably expressed ZAP was also immunoprecipitated with transfected SINV RNA in wild-type MEFs (Fig. 5A and B). The interaction between ectopically expressed ZAP and transfected SINV RNA was barely detectable, probably because the level of SINV RNA was extremely low (data not shown). These findings indicate that ZAP detects SINV RNA to induce its degradation.

Type I IFN requires ZAP to suppress SINV replication in MEFs

Type I IFN suppresses SINV replication (26). Therefore, we examined the involvement of ZAP in the type-I-IFN-mediated suppression of SINV replication. Treatment with IFN- β induced the expression of ZAP mRNA and suppressed SINV replication in wild-type MEFs (Fig. 6A). However, treatment with IFN- β failed to suppress SINV replication in *Zc3hav1^{+/-}* MEFs (Fig. 6B). Although SINV replication was enhanced in *Ddx58^{+/-}Ilfih1^{+/-}* and *Irf3^{+/-}Irf7^{+/-}* MEFs, which are deficient in the expression of type I IFN and IFN-inducible genes, the ectopic expression of ZAP was sufficient to suppress SINV replication in these MEFs (Fig. 6C). These findings indicate that ZAP is an anti-viral effector molecule responsible for the type-I-IFN-mediated anti-viral response to SINV.

ZAP mediates host defense against SINV

A number of studies have shown that ZAP suppresses viral replication in immortalized and cancer cells (31). Therefore, we examined the involvement of ZAP in the anti-viral response *in vivo* using *Zc3hav1^{+/-}* mice. SINV replicated efficiently in the brain in infant *Zc3hav1^{+/-}* mice (Fig. 7A). The expression of *Ifnb1* and *Cxcl10* was elevated in infant *Zc3hav1^{+/-}* mice infected with SINV (Fig. 7B). The expression of *Ii1b* and *Ii6*, which are associated with the severity of SINV-induced encephalopathy (32, 33), were also elevated in infant *Zc3hav1^{+/-}* mice infected with SINV (Fig. 7C). These findings indicate that ZAP protects infant mice from SINV infection.

Discussion

Although we and others have demonstrated the importance of ZAP in anti-viral responses, the molecular mechanisms underlying the elimination of viruses by ZAP are still unclear. ZAP suppresses the replication of SINV, Ebola virus, Marburg virus, MLV, HIV-1 and HBV (12, 16–19), indicating that ZAP targets positive-sense RNA viruses, negative-sense RNA viruses, retroviruses and DNA viruses. However, ZAP does not suppress the replication of VSV, a negative-sense RNA virus. ZAP inhibits viral protein translation and viral RNA

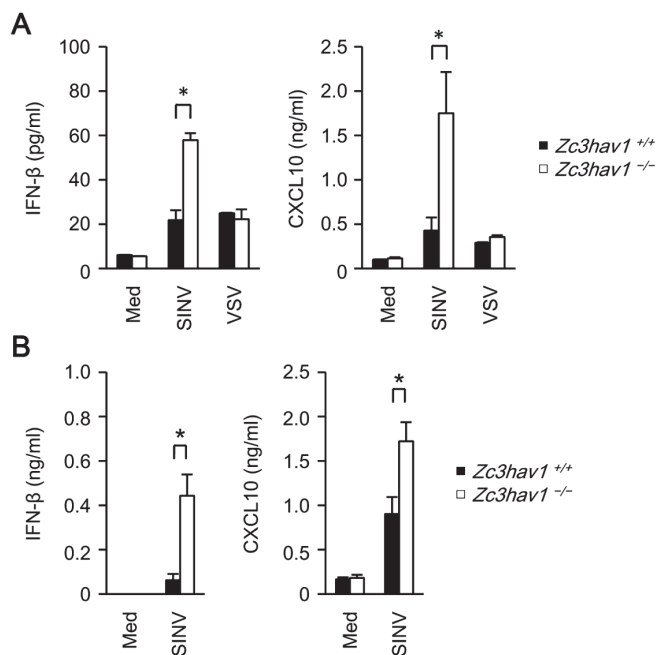


Fig. 3. ZAP is not essential for SINV-induced type I IFN production in primary MEFs or dendritic cells. (A) Primary *Zc3hav1^{+/-}* and *Zc3hav1^{+/+}* MEFs were infected with SINV (MOI = 1) or VSV (MOI = 1) for 24 h. (B) *Zc3hav1^{+/-}* and *Zc3hav1^{+/+}* GM-CSF-induced bone-marrow dendritic cells were infected with SINV (MOI = 5) for 24 h. The levels of IFN- β and CXCL10 in the culture supernatants were measured with ELISAs. The experiments were performed three times (means \pm SD of three independent samples). * $P < 0.05$.

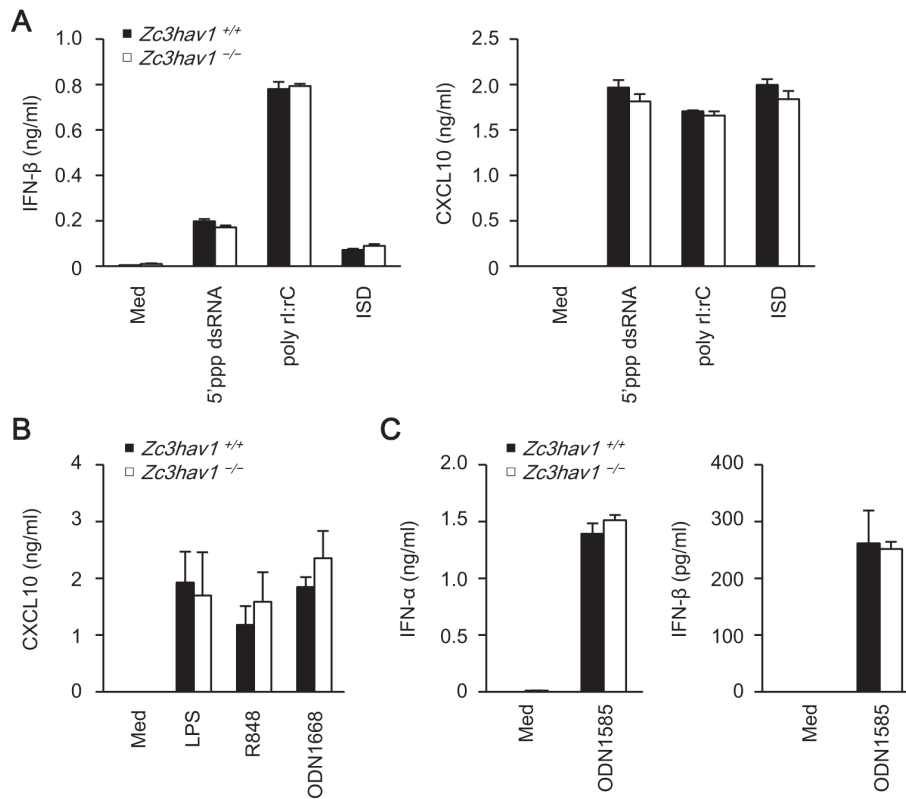


Fig. 4. ZAP is not essential for nucleic-acid-induced type I IFN production in primary MEFs or dendritic cells. (A) Primary *Zc3hav1*^{+/+} and *Zc3hav1*^{-/-} MEFs were stimulated with 5'ppp-dsRNA (1 $\mu\text{g ml}^{-1}$), poly rI:rC (1 $\mu\text{g ml}^{-1}$) and ISD (1 $\mu\text{g ml}^{-1}$), together with Lipofectamine™ 2000 for 24 h. The levels of IFN- β and CXCL10 in the culture supernatants were measured with ELISAs. (B) *Zc3hav1*^{+/+} and *Zc3hav1*^{-/-} GM-CSF-induced bone-marrow dendritic cells were stimulated with LPS (TLR4 ligand, 100 ng ml^{-1}), R848 (TLR7 ligand, 10 $\mu\text{g ml}^{-1}$) and ODN 1668 (TLR9 ligand, 1 μM) for 24 h. The levels of CXCL10 in the culture supernatants were measured by an ELISA. (C) *Zc3hav1*^{+/+} and *Zc3hav1*^{-/-} FLT3L-induced bone-marrow dendritic cells were stimulated with ODN 1585 (TLR9 ligand, 5 μM) for 24 h. The levels of IFN- α and IFN- β in the culture supernatants were measured with ELISAs. The experiments were performed three times (means \pm SD of three independent samples).

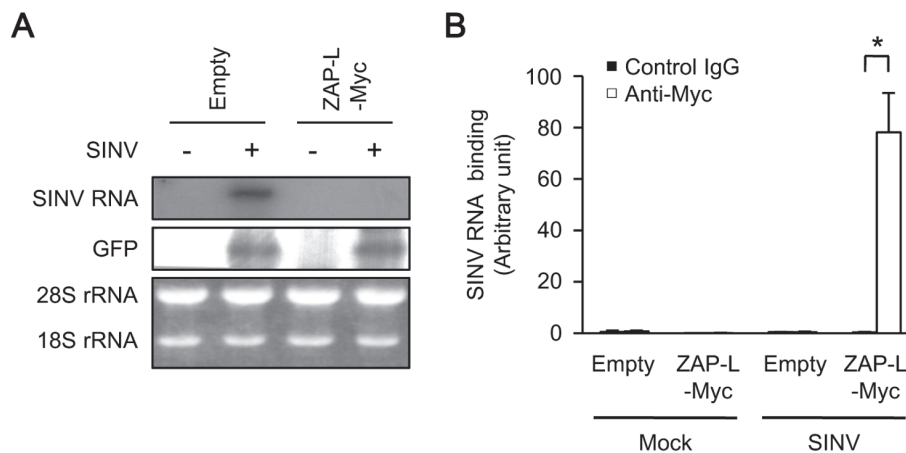


Fig. 5. ZAP binds to and reduces SINV RNA. (A) Immortalized wild-type MEFs were transiently transfected with SINV RNA and an enhanced green fluorescent protein (EGFP)-expressing plasmid with or without the ZAP-L-Myc-expressing plasmid for 24 h. The total cellular RNA was subjected to a northern blotting analysis for SINV RNA. (B) Immortalized wild-type MEFs retrovirally transduced with or without ZAP-L-Myc were transiently transfected with SINV RNA for 24 h. The level of SINV RNA binding to ZAP-L-Myc was measured with RNA immunoprecipitation coupled to quantitative RT-PCR. The level of immunoprecipitated SINV RNA was normalized to that of the input SINV RNA to determine the SINV RNA binding activity. The experiments were performed at least three times and representative data are shown (means \pm SD of three independent samples). * $P < 0.05$.

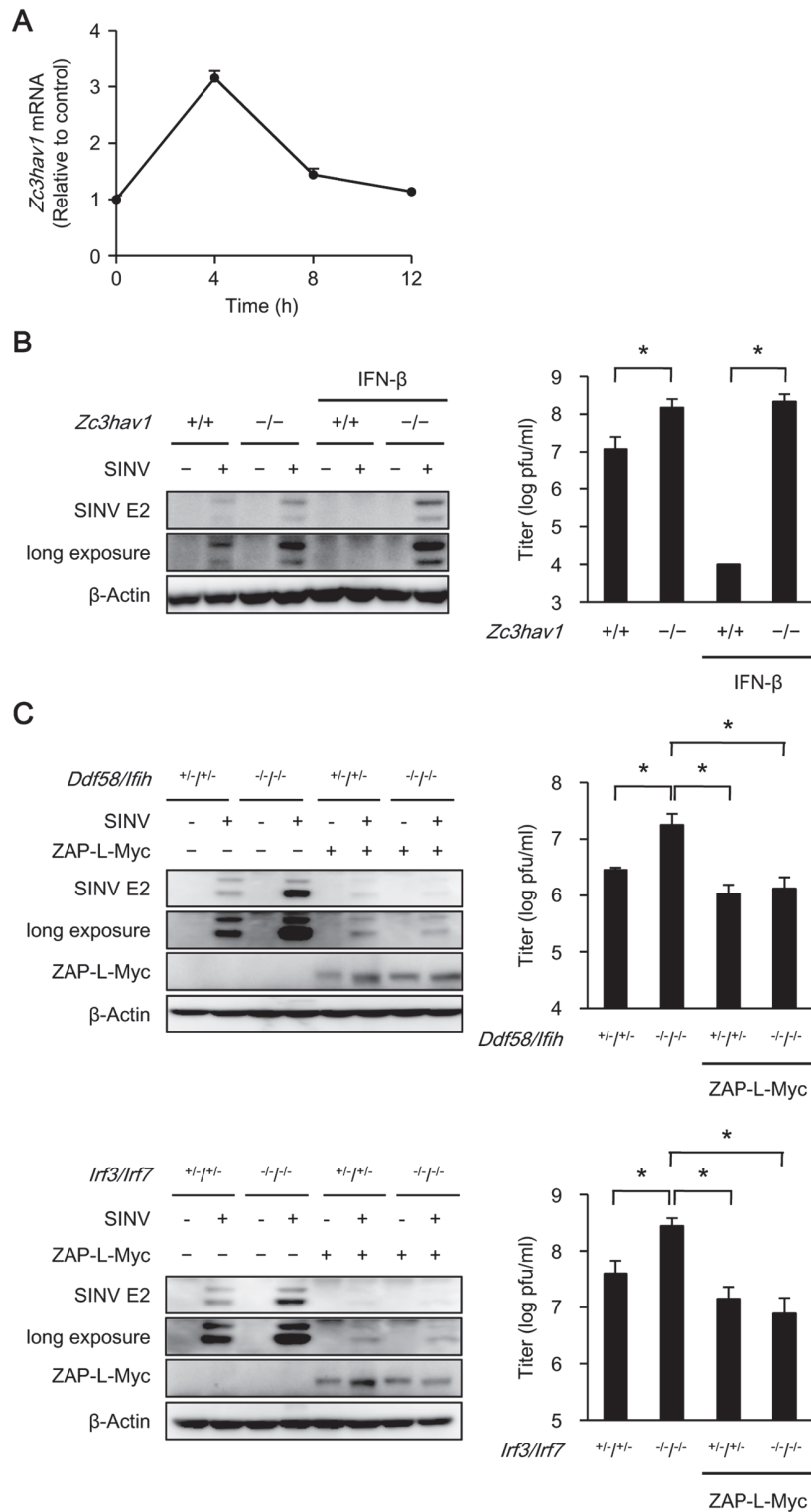


Fig. 6. ZAP is essential for the type-I-IFN-induced anti-viral response to SINV. (A) Primary wild-type MEFs were stimulated with IFN- β (50 pg ml⁻¹) for the indicated periods. The level of Zc3hav1 mRNA was measured with quantitative RT-PCR. (B) Primary *Zc3hav1*^{+/+} and *Zc3hav1*^{-/-} MEFs were treated with or without IFN- β (50 pg ml⁻¹) for 12h and then infected with SINV (MOI = 0.05) for 24h. The levels of SINV E2 and β -actin in the cell lysates were measured with immunoblotting. The viral titers were determined with TCID₅₀ assays after 24h. (C) Primary *Ddx58*^{+/+/+}*Ilfih*^{+/+}, *Ddx58*^{+/+/+}*Ilfih*^{-/-}, *Irf3*^{+/+/+}*Irf7*^{-/-} and *Irf3*^{+/+/+}*Irf7*^{+/+} MEFs retrovirally transduced with or without ZAP-L-Myc were infected with SINV (MOI = 0.05) for 24h. The levels of SINV E2, β -actin and Myc-tagged proteins in the cell lysates were measured with immunoblotting. The viral titers were determined with TCID₅₀ assays after 24h. The experiments were performed three times and representative data are shown (means \pm SD of three independent samples). **P* < 0.05.

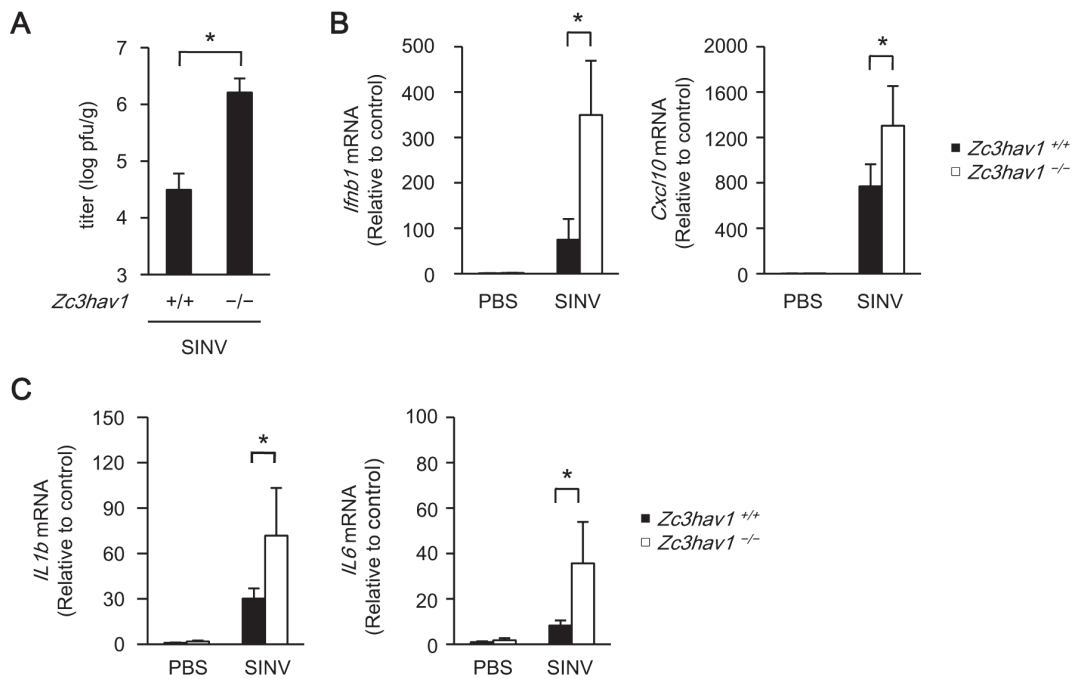


Fig. 7. Loss of ZAP renders mice susceptible to SINV infection. (A–C) Ten-day-old *Zc3hav1*^{+/+} and *Zc3hav1*^{-/-} mice ($n = 5$ each) were subcutaneously inoculated with SINV (100 pfu per mouse). The brains of the SINV-infected *Zc3hav1*^{+/+} and *Zc3hav1*^{-/-} mice were isolated on day 5. The viral titers (pfu per tissue weight) in the brains were determined with TCID₅₀ assays (A). The levels of *Ifnb1*, *Cxcl10*, *Il1b* and *Il6* mRNAs in the brains were measured with quantitative RT–PCR (B and C). The experiments were performed five times (means \pm SD of five independent samples). * $P < 0.05$.

splicing and destabilizes viral RNA (34). These findings raise important questions. How does ZAP sense a wide variety of viruses? How does ZAP specifically identify its target viruses? How does ZAP distinguish the different modes of anti-viral actions? In future studies, we will address these questions to better understand the anti-viral innate immune response.

Members of the C3H ZF protein family can bind to RNA and are conserved from yeasts to mammals. In mammals, >50 members of the C3H ZF protein family have been identified (35). This family is involved in a variety of cellular events, including RNA synthesis, mRNA splicing, mRNA degradation and protein translation, and thus regulates the innate and acquired immunity of the host (36–38). ZAP is a member of the C3H ZF protein family and acts as an RNA-sensing anti-viral effector molecule. In addition to ZAP, ZFP36 and regnase-1, other C3H ZF protein family members, sense viral RNA and are involved in the regulation of viral replication (39–41). Therefore, the C3H ZF protein family seems to play a major role in the direct elimination of viruses. However, the anti-viral activities of the majority of C3H ZF protein family members have not yet been investigated. It would be interesting to identify the C3H ZF protein family members involved in host defenses against viruses in a future study.

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