Research Article

TRIM25 Identification in the Chinese Goose: Gene Structure, Tissue Expression Profiles, and Antiviral Immune Responses In Vivo and In Vitro

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The retinoic acid-inducible gene I (RIG-I) and the RIG-I-like receptor (RLR) protein play a critical role in the interferon (IFN) response during RNA virus infection. The tripartite motif containing 25 proteins (TRIM25) was reported to modify caspase activation and RIG-I recruitment domains (CARDs) via ubiquitin. These modifications allow TRIM25 to interact with mitochondrial antiviral signaling molecules (MAVs) and form CARD-CARD tetramers. Goose TRIM25 was cloned from gosling lungs, which possess a 1662 bp open reading frame (ORF). This ORF encodes a predicted 554 amino acid protein consisting of a B-box domain, a coiled-coil domain, and a PRY/SPRY domain. The protein sequence has 89.25% sequence identity with Anas platyrhynchos TRIM25, 78.57% with Gallus gallus TRIM25, and 46.92% with Homo sapiens TRIM25. TRIM25 is expressed in all gosling and adult goose tissues examined. QRT-PCR revealed that goose TRIM25 transcription could be induced by goose IFN-α, goose IFN-γ, and goose IFN-λ, as well as a35 s polyinosinic-polycytidylic acid (poly(I:C)), oligodeoxynucleotides 2006 (ODN 2006), and resiquimod (R848) in vitro; however, it is inhibited in H9N2 infected goslings for unknown reasons. These data suggest that goose TRIM25 might play a positive role in the regulation of the antiviral immune response.

1. Introduction

The tripartite motif family (TRIM) is a large family of proteins that share a common N-terminal RBCC motif (including a RING finger, one or two B-box, and a coiled-coil domain) and are involved in many biological processes, such as cell differentiation, apoptosis, transcription regulation, and cell signaling [1]. TRIM proteins play a critical role in a number of innate immunity responses and are critical for antiviral responses [2, 3].

The innate immune system is the first line of defense against detrimental pathogens, such as bacteria and viruses. Critical innate immune responses against viruses include constitutively expressed proteins with intrinsic antimicrobial properties and the inducible type I interferon (I-IFN) system. In general, pattern recognition receptors (PRRs), such as Toll-like receptors (TLRs), retinoic acid-inducible gene I-(RIG-I-) like receptors (RLRs), and NOD-like receptors (NLRs), take the lead in recognizing the pathogen-associated molecular pattern (PAMP) when pathogens invade organisms [4, 5]. Inducible innate immune responses are subsequently activated to increase I-IFN production by triggering downstream signaling pathways with a series of adaptor proteins that transmit downstream signals. The RLR protein plays a critical role in interferon responses during RNA virus infection. Over the past years, more and more TRIM proteins have been found to initiate the innate immune response via their capacity to act as an E3-ligase. In fact, there are seven lysine residues
(K6, K11, K27, K29, K33, K48, and K63) present in ubiquitin, on which polyubiquitin chains could form. Different polyubiquitin chains linked through different lysine have diverse functions [6]. K48 linked polyubiquitin chains are a signal for degradation, while the K63 linked polyubiquitin can be used to activate antiviral signaling pathways [7, 8]. Upon recognizing PAMP RNA, RIG-I hydrolyzes adenosine triphosphate (ATP) and undergoes a conformational change in which the RNA binding domain unfolds for subsequent interaction with PAMP RNA, while further releasing the CARD domain for MAVs interaction triggering the IFN signaling pathway. Meanwhile, E3-ligase TRIM25 modifies the CARD domain of RIG-I via ubiquitin, which allows it to interact with MAVs to form the CARD-CARD tetramer [9, 10]. In addition, a linear ubiquitin assembly complex was found to suppress RIG-I-mediated antiviral activity by binding human TRIM25 and eventually induces its degradation [11]. Recently, a number of studies revealed that TRIM family proteins play several diverse roles in antiviral innate immune responses. The TRIM proteins could inhibit virus replication at different stages, such as viral entry, transcription of viral genes, and viral release from the cells [1, 4, 11–14]. Another study reported that TRIMs adapt to regulate the innate immune response by attaching the K(lys)-27-linked polyubiquitin chains to the TRIM25 that can inform future studies investigating the interplay between bird TRIM25 and RIG-I. This study also elucidates the mechanisms by which TRIM25 activates the RIG-I signaling pathway to induce the production of interferon.

2. Materials and Methods

2.1. Virus and Animal Ethics Statement. All geese and goose embryos used in this study were purchased from the Sichuan Agricultural University Farm. The H9N2 (7.14 × 10<sup>12.64</sup> copies/mL) was kindly provided by the Shanghai Veterinary Research Institute, Chinese Academy of Agricultural Sciences. The GPV (10<sup>-6.6</sup> EID<sub>50</sub>/0.2 mL) strain was provided by the Avian Disease Research Center of Sichuan Agriculture University. The animal studies were approved by the Institutional Animal Care and Use Committee of the Sichuan Agricultural University and followed the National Institutes of Health Guide for the Care and Use of Laboratory Animals.

2.2. RNA Isolation and cDNA Synthesis. First, total RNA was isolated from tissues and cells using TRIZOL (Invitrogen, USA) following the manufacturer’s guidelines. Subsequently, the cDNA used for the amplification of goose TRIM25 and qRT-PCR was synthesized using the 5-All-in-one Kit (Abm, Canada) according to the manufacturer’s guidelines. All steps described above were performed under RNase-free conditions.

2.3. Cloning of Full Length Goose TRIM25. First, the goose TRIM25 sequence was amplified using the gene-specific primers G-TRIM25-F and G-TRIM25-R, which were designed based on the predicted goose sequences obtained from GenBank. Then, the full length TRIM25 cDNA, including the short 5'-untranslated regions (UTRs), was obtained by PCR using the specific primers designed in the previous step. The amplified PCR fragments were purified using the universal DNA Purification Kit (Tiangen, China) and sub-cloned into pMD19-T (Sigma–Aldrich, USA), which was then
transformed into high-efficiency DH5α competent cells. The positive clones were selected and then sent for sequencing. The full length cDNA sequence was assembled. Its continuity was confirmed by sequencing the cloned PCR product amplified with a pair of terminal primers. These primers were designed for quantitative reverse transcription polymerase chain reaction (qRT-PCR) based on the acquired sequences. Table 1 lists all primers used in the procedures described above.

2.4. Cell Culture and Treatment. The GEFs were isolated from 9-day-old goose embryos and seeded into individual wells of a cell culture plate. The PBMCs were isolated from goose peripheral blood using the goose peripheral blood mononuclear cell separation medium kit (TBD, China) according to the manufacturer’s guidelines. All isolated cells were cultivated in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% fetal calf serum at 37°C in a humidified 5% CO2 atmosphere for 24 hours. The media were supplemented with penicillin (100 U/mL) and streptomycin (100 μg/mL). The PBMCs were isolated from several healthy adult geese. The PBMCs obtained from the blood of each goose were seeded into several wells of a cell culture plate. Each well contained 200 μL of cell suspension with a density of 5 × 10^5 cells/mL. Then, the GEFs were divided into four groups and stimulated with either poly(I:C) (Sigma–Aldrich, USA) (100 μg/well), goose IFNα (60 μL/well), goose IFNγ (60 μL/well), or goose IFNβ (60 μL/well) and poly(I:C) (100 μg/well). PBS (60 μL/well) was chosen as the control. The PBMCs were divided into 7 groups with stimulation of H9N2 (10^5 TCID₅₀/50 μL), GPV (10^5 TCID₅₀/50 μL), poly(I:C) (100 μg/well), IFNα (60 μL/well), ODN2006 (100 μg/well), R848 (100 μg/well), or PBS (60 μL/well) as the control. Stimulation with poly(I:C), ODN2006, and R848 lasted 6 hours before the cells were collected into RNase-free centrifuge tubes with 900 μL of TRIZOL, while the cells treated with goose IFNα, IFNγ, and IFNβ were collected 24 hours after stimulation. The recombinant goose IFNs used in this study were expressed by pcDNA3.1(+)–goIFNα, pcDNA3.1(+)–goIFNγ, and pcDNA3.1(+)–goIFNλ in baby hamster kidney 21 cells (BHK 21).

2.5. Animal Experiment. The 1-day-old goslings were fed in the animal rooms for 3 days before challenge. Then, 18 healthy goslings were randomly chosen and divided into 2 groups. The first group was inoculated with 500 μL of H9N2 via intramuscular injection. The control group was inoculated with the same volume of PBS. On day 1, day 3, and day 7 after infection, three goslings from each group were randomly selected for tissue sampling (lung, small intestine, blood, bursa of Fabricius, trachea, and brain). All tissues were ground in liquid nitrogen and stored in TRIZOL. The total RNA was extracted from each tissue using the TRIZOL reagent according to the manufacturer’s instructions.

In addition, the tissue distribution profile of TRIM25 was performed in both healthy adult geese and goslings. The cDNA was reverse transcribed using the 5-All-in-one Kit (Abm, Canada) according to the manufacturer’s guidelines. All of the synthesized cDNA were stored at −80°C for later use.

2.6. The Detection of Goose TRIM25 mRNA Transcription Levels. First, the cDNA used to clone the full length TRIM25 and the mRNA used to assess the transcription levels were synthesized using the same procedures described above. Then, the qRT-PCR analysis was conducted to determine the TRIM25 transcription levels in the different organs of the experimental geese and the treated GEFs and PBMCs. Two gene-specific primers (sense primer G-TRIM25-RT-F and reverse primer G-TRIM25-RT-R (Table 1)) that amplified a 127 bp TRIM25 fragment were designed to detect the transcription levels of TRIM25. Two other primers amplifying a 172 bp β-actin fragment served as an internal control. The qRT-PCR reaction mixture contained 5 μL of EvaGreen qPCR MasterMix (Abm, Canada), 0.3 μL of each primer (10 pmol/μL), 0.4 μL of cDNA, and 4 μL of sterile water. The qRT-PCR was performed using the Bio-Rad CFX96 Real-Time Detection System (Bio-Rad, USA). The goose TRIM25 qRT-PCR procedure was as follows: 95°C for 3 min, followed by 39 cycles of 95°C for 10 s, 57°C for 30 s (61°C for β-actin). Finally, the qRT-PCR data were analyzed using the 2–ΔΔCT method with Bio-Rad CFX Manager Software.

2.7. Bioinformatic Analysis. The goose TRIM25 sequence was compared with the predicted TRIM25 sequence obtained from GenBank using the BLASTX and BLASTP search programs (http://blast.ncbi.nlm.nih.gov/Blast.cgi). The ORF and the protein domain were determined using the ORF Finder (http://www.ncbi.nlm.nih.gov/orffinder/). The protein structure motif was predicted using SMART (simple modular architecture research tool, http://smart.embl-heidelberg.de/). The phylogenetic tree was created using the neighbor-joining (NJ) method in the MEGA version 5.0 software package. Bootstrap values were calculated with 1000 iterations to estimate the robustness of the internal branches.

3. Results

3.1. Clone and Sequence Analysis of Goose TRIM25. The 1892 bp cDNA of TRIM25 (KX_364385) with a 1662 bp conserved ORF was cloned (Figure 1(a)). This clone coded a predicted 554 amino acid protein. Sequence analysis by SMART program revealed that goose TRIM25 ORF contains an N-terminal BCC motif which consist of a B-box domain and a coiled-coil domain; however for unknown reason no RING finger domain exists in goose TRIM25 (Figure 1(b)). The protein sequence of goose TRIM25 had 89.25% identity

<table>
<thead>
<tr>
<th>Name</th>
<th>Primer sequence 5′-3′</th>
</tr>
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<tbody>
<tr>
<td>G-TRIM25-F</td>
<td>GGTGCAGACGTGCGCTGAC</td>
</tr>
<tr>
<td>G-TRIM25-R</td>
<td>CAGACCAAGAATCATCAG</td>
</tr>
<tr>
<td>G-TRIM25-RT-F</td>
<td>CCACCACCTACGCTTTC</td>
</tr>
<tr>
<td>G-TRIM25-RT-R</td>
<td>GCCATAGCAATGCCAAT</td>
</tr>
<tr>
<td>β-Actin-F</td>
<td>CGGTGCATCAACAGGAGAA</td>
</tr>
<tr>
<td>β-Actin-R</td>
<td>GAAGGATGGCTGGAAGAG</td>
</tr>
</tbody>
</table>

Table 1: The list of primers.
Figure 1: Continued.
with *Anas platyrhynchos* TRIM25, 78.57% with *Gallus gallus* TRIM25, and 46.92% with *Homo sapiens* TRIM25 (Figure 2). The C-terminus of goose TRIM25 contains a PRY domain followed by a SPRY domain. In addition, the phylogenetic tree analysis suggested that goose TRIM25 is closer to *Anas platyrhynchos* TRIM25 and *Gallus gallus* TRIM25 (Figure 3).

3.3. The Effects of Agonist and Goose IFNs on Goose TRIM25 Transcription Levels in Treated PBMCs. It was reported that viral RNA and DNA could immediately trigger antiviral responses in human and murine cells. To investigate if goose TRIM25 is involved in the antiviral response, we determined the mRNA transcription levels of goose TRIM25 in PBMCs following treatment with H9N2, GPV, poly(I:C), ODN2006, IFNα, and R848. We found that H9N2 (P < 0.01) and GPV (P < 0.001) infection significantly upregulated TRIM25 mRNA levels in the treated PBMCs (Figure 6). TRIM25 mRNA levels were also upregulated by IFNα and poly(I:C), ODN2006, and R848 treatment, which served as positive controls.

3.5. The Effects of H9N2 on Goose TRIM25 Transcription Levels In Vivo. Three goslings were randomly sampled on day 1, day 3, and day 7 after infection with H9N2. QRT-PCR analysis found that goose TRIM25 expression levels were significantly downregulated in H9N2-infected goslings. On day 1 after infection, the transcription levels were downregulated in the blood and brain. On the day 3 after infection, the mRNA levels of goose TRIM25 in the lungs were significantly upregulated on day 7 after infection (Figure 7).

4. Discussion

The innate immune signaling pathways, such as RIG-I, are a critical component of the antiviral response. TRIM25 has a significant impact on the RIG-I signaling pathway. First, after RIG-I is activated following PAMP RNA recognition, TRIM25 modifies RIG-I with ubiquitin. Then, CARD domains of RIG-I and MAVs conjugate to form a tetramer. Finally, the interactions between the CARD domains of RIG-I and MAVs activate TBK1 and IKKe to initiate downstream signaling [24–27]. It was reported that TRIM25 exists in fish, humans, mice, chickens, and ducks. Domingo Miranzo-Navarro indicated that duck TRIM25 is an important component of the antiviral response by demonstrating that TRIM25 modifies RIG-I through ubiquitination, which was reported recently not to be related to ubiquitin anchoring [10]. Aquatic birds, particularly geese, contribute significantly to the transmission of many major pathogens, including AIV [28]. Though the complete genome of geese has been sequenced, many antiviral genes of geese are still unknown, including TRIM25.
Figure 2: Continued.
(b) Figure 2: Continued.
As with many other TRIM family proteins, TRIM25 has a conserved N-terminal region RBCC motif that consists of a RING finger domain, one or two B-box domains and a coiled-coil domain [3, 29]. The RING finger domain is defined by the following consensus sequence CX2CX(9–39)CX1–3HX(2–3)C/HX2CX(4–48)CX2C. The Cys and His residues form a cloud bond with the two zinc atoms and function as an E3-ubiquitin ligase (an enzyme involved in ubiquitination) [30]. This interaction between the E3-ubiquitin ligase and its target proteins is required for the antiviral response of most functional TRIM proteins. The B-box is also a zinc-binding motif, which is found exclusively in TRIM proteins.

**Figure 2:** Multiple comparisons of TRIM25 from several species. Multiple comparisons of goose (Anas cygnoides), Gallus gallus, Anas platyrhynchos, Homo sapiens, and Mus musculus TRIM25 nucleic acid sequences. Highlighted regions indicate the homology of TRIM25 between species.
**Figure 3**: Phylogenetic tree of goose (*Anser cygnoides*) TRIM25. The tree was constructed using MEGA 6.0 with the following TRIM protein sequences: *Homo sapiens* (NM_005082.4), *Mus musculus* (NM_009546.2), *Felis catus* (NM_001290251.1), *Gallus gallus* (NM_001318548.1), *Anas platyrhynchos* (XM_013092756.1), *Danio rerio* (NM_200175.1), *Pelodiscus sinensis* (XM_00125153.2), *Alligator sinensis* (XM_006017407.2), and the *Anser cygnoides* TRIM25 cloned.

**Figure 4**: Tissue distribution profiles of goose TRIM25 in goslings and adult geese. β-Actin was used as the internal control gene. The expression data are represented as the mean ± SEM (*n* = 3). BL (blood), Pr (proventriculus), HG (harderian gland), SI (small intestine), Cu (caecum), Ct (caecum tonsil), K (kidney), Bf (bursa of Fabricius), T (thymus), Lu (lung), Li (liver), Sp (spleen), P (pancreas), B (brain), Gi (gizzard), Tr (trachea), H (heart), and M (muscle).

[31, 32] and reportedly also functions as an E3 ligase [33]. The coiled-coil domain, the last domain of the RBCC, mediates the homodimerization/oligomerization in a large number of TRIM proteins, which may contribute to the diversity of their biological functions [34–36]. Mutation and deletion experiments have revealed that the coiled-coil domain may also be important for the formation of subcellular structures, including cytoplasmic or nuclear bodies (NBs) [35]. In general, the RING domain functions as an E3 ligase, while the C-terminal PRY/SPRY domain determines the specific target proteins for TRIM ubiquitination [2].

The 1892 bp full length goose TRIM25 cDNA cloned and sequenced in this study contained a 1662 bp ORF. It is longer than the predicted TRIM25 cDNA of *Anser cygnoides* listed in the GenBank database (accession numbers XM_013178947, XM_013178948). The predicted *Anser cygnoides* TRIM25 transcript variant X1 had a 1557 bp (transcript variant X2: 1638-bp) ORF with a short 5’UTR and a long 3’UTR, encoding 519-amino acid (transcript variant X2: 545-amino acid). The goose TRIM25 was cloned from gosling lung tissue, which had an RBCC motif in the N-terminal domain and a PRY/SPRY in C-terminal domain. Interestingly, the RING finger domain present in *Anas platyrhynchos* (XM_013092756.1) and *G. gallus* (XM_415653.4) is the most critical domain for E3-ubiquitin ligase activity [30]. However, according to our data (repeat 5RACE-PCR many times), the RING finger domain was absent in goose TRIM25 cDNA, which was consistent with the predicted goose TRIM25 sequence. Although the TRIM25 RING finger domain was suggested by previous studies to play an important role in the innate antiviral immune response [11, 20], one study suggested that B-Boxes could provide an E3 binding site, similar to the RING domain, thereby conferring E3 ligase activity to RINGless TRIMs [23]. It was revealed that the same E3 ligase activity was detected in the RING finger domain-deleted TRIM16 in vitro [37]. These data suggest that goose
**Figure 5:** Effects of goose IFNα, IFNγ, IFNλ, and poly(I:C) on goose TRIM25 in GEFs. The GEFs were treated with goose IFNα, IFNγ, and IFNλ for 24 hours and poly(I:C) for 6 hours. The expression data are represented as the mean ± SEM (n = 4). Differences in mRNA cytokine production in challenged cells were analyzed using the unpaired, two-tailed t-test (∗∗P < 0.01; ∗∗∗P < 0.001).

**Figure 6:** Effects of H9N2, GPV, poly(I:C), R848, ODN2006, and goose IFNs on goose TRIMM25 transcript levels in PBMCs. β-Actin was used as the internal control gene to detect the transcription levels of goose TRIM25. PBMCs were infected with either H9N2 or GPV for 24 hours. PBMCs were treated with poly(I:C), ODN2006, R848 for 6 hours, and IFNα for 24 hours. The expression data represent the mean ± SEM (n = 3). The differences in mRNA cytokine production in virus-challenged goslings were analyzed using the unpaired, two-tailed t-test (∗∗∗P < 0.001).

TRIM25, although lacking a RING domain, had the capacity to act as an E3 ligase and may function as an antiviral gene; however, this remains to be determined.

One could conclude from the tissue distribution profiles of goose TRIM25 that immune-associated tissues, such as blood, bursa of Fabricius, and harderian gland, had relatively high TRIM25 expression levels. In addition, it was more highly expressed in the tissues obtained from goslings when compared to those from adult geese. We found that TRIM25 expression levels were the highest in the blood of both geese and goslings; however, high TRIM25 transcription levels were also observed in gosling proventriculus and adult goose liver. It differed from the expression pattern of *G. gallus* TRIM25 where the highest levels were found in the lungs [20]. In contrast to the results of the current study, a previous study found that the highest TRIM transcription levels in felines were found in the heart [18].

ODN2006 is a synthesized oligonucleotide, determined in our previous study, is recognized by goose TLR21, and subsequently upregulates the expression of IFN[38]. Poly(I:C) is a synthesized oligonucleotide that not only functions as a TLR3 ligand but also activates the RIG-I and MDA5 signaling pathways, inducing the expression of IFNα [39, 40]. R848 is a TLR agonist, which previously has been used as a vaccine adjuvant. We found that there was significant upregulation of TRIM25 expression in ODN2006, poly(I:C), R848, and PBMCs treated with IFNα for 6 hours. It was reported that the TRIM family proteins could be induced by IFNs in humans [41]. This study investigated if goose IFNs, including IFNα, IFNγ, and IFNλ which were determined to be expressed in transfected BHK 21 in our previous study, can trigger goose TRIM25 expression. As expected, the expression of goose TRIM25 was significantly upregulated 24 hours after treatment with goose IFNα, IFNγ, and IFNλ protein in GEFs, which is consistent with findings reported in mammals.

Anna Figuiredo recently reported that KAP1, one of the TRIM family proteins, inhibits HIV-1 integration [14]. An
increasing number of TRIM proteins were determined to contribute to the antiviral response by mediating the antiviral signaling pathways in mammals [12, 42]. TRIM family proteins play a critical role in the innate immune antiviral response. Duck TRIM25 was found to play an important role in RIG-I-dependent antiviral signaling pathways [9]. QRT-PCR revealed that gosling TRIM25 expression levels were significantly downregulated in the trachea of H9N2-infected birds 1 day after infection and in the blood, brain, and small intestines 3 days after infection. As reported, the low pathogenic avian influenza virus is sensed by host TLRs, which in turn reduce the expression of IFNs to defend the host against the infection [43, 44]. Our in vitro study showed that the goose TRIM25 transcription levels were significantly upregulated by H9N2 and GPV treatment, which is consistent with previous studies [14, 19, 20, 45].

Interestingly, the differential expression of goose TRIM25 was detected in the H9N2-infected geese and PBMCs, which coincides with our previous findings. We designed an experiment in which the transcription levels of IFNα and IFNγ were measured in goslings and PBMCs infected with H9N2. The results revealed that IFNα expression was inhibited in the immune-related tissues of H9N2-infected goslings, such as the harderian gland and the bursa of Fabricius, while in PBMCs it was upregulated [45]. And the goMx transcription levels were upregulated in the trachea at 3 d.p.i. but downregulated in small intestine [46]. We speculate H9N2 utilized some unknown mechanism to evade the host immune response and decrease the production of IFNα and eventually decrease the production of TRIM25. Some reports indicated that the NS of the low pathogenic avian influenza virus can inhibit the host antiviral response and reduce IFNs production [47, 48]. These reports are consistent with our speculation.

5. Conclusion

In this study, we cloned goose TRIM25, determined its tissue distribution profiles, described its nucleotide sequence, and conducted both structural and phylogenetic analyses. The cloned full length goose TRIM25 cDNA contained a 1662-bp ORF but lacked the E3 ligase RING domain. QRT-PCR results revealed that goose TRIM25 is highly expressed in immune-associated tissues. TRIM25 transcription was induced by goose IFNs, poly(I:C), and ODN 2006, R848, GPV in H9N2-treated GEFs or PBMCs; however, it was inhibited in H9N2-treated goslings. A tentative conclusion could be drawn that goose TRIM25 transcription levels could be induced by goose IFNα, goose IFNγ, and goose IFNλ, as well as poly(I:C), ODN 2006 and R848 in vitro, but it is inhibited in H9N2-infected gosling through some unknown mechanism.

Abbreviations

RIG-I: Retinoic acid-inducible gene I
RLRs: Retinoic acid-inducible gene I- (RIG-I-) like receptors
IFNs: Interferon
TRIM25: The tripartite motif containing 25 proteins

CARDs: Caspase activation and recruitment domains
MAVs: Mitochondria antiviral signal molecules
ORF: Open reading frame
Poly(I:C): Polyinosinic-polycytidylic acid
ODN: Oligodeoxynucleotides
R848: Resiquimod
PRRs: Pattern recognition receptors
TLRs: Toll-like receptors
NLRs: NOD-like receptors
PAMP: Pathogen-associated molecular pattern
NEMO: NF-κB essential modulator
TRAF3: Tumor necrosis factor receptor associated factor 3
IRF3: Interferon regulatory factor 3
qRT-PCR: Quantitative reverse transcription polymerase chain reaction
ISG 15: Interferon stimulated gene 15
PBMCs: Peripheral blood mononuclear cell
GEFs: Goose embryo fibroblasts
UTRs: Untranslated regions
DMEM: Dulbecco’s modified Eagle’s medium
BHK 21: Baby hamster kidney 21 cells
ATP: Adenosine triphosphate
GPV: Gosling plague virus.

Competing Interests

The authors declare that there are no competing interests regarding the publication of this paper.

Authors’ Contributions

Yunan Wei and Shun Chen contributed equally as co-first authors of this work.

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