Immunogenicity and Efficacy of Coxsackievirus A16 Vaccine Candidate Formulations in a Mouse Model

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Abstract

Besides enterovirus 71 (EV71), Coxsackie virus A16 (CVA16) is a major etiologic agent of hand, foot and mouth disease (HFMD), causing infections in millions of children under 5 years of age each year. The progress made in the development of inactivated EV71 vaccines encourages research aiming at developing a CVA16 vaccine for better prevention of HFMD and control of spreading of a rapidly evolving virus. The immunogenicity and efficacy of a CVA16 vaccine candidate were examined in a mouse model. Several vaccine formulations using different adjuvants and formalin-inactivated or non-inactivated full and empty virus particles were compared. It was observed that the CVA16-P4-L731 vaccine induced strong B and T cell immune responses in mice. The mouse antisera contained the highest titers of neutralizing antibodies reported so far and prevented infection of cells in vitro by prototype A and subgenotype B2b. Pups born to immunized mice were protected from disease and death following challenge by more than 10,000 LD50 of homologous and heterologous viruses. Taken together, the data suggest that the CVA17-P4-L731 is a promising vaccine candidate, either independently or as a valuable component for a combined [EV71+CVA16] bivalent vaccine.

Keywords: CVA16; HFMD; Immunogenicity; Vaccine efficacy; Vaccine formulations; Mouse pup model

Introduction

Coxsackievirus A16 (CVA16) is a serotype of the type species Human enterovirus A, genus Enterovirus in the family Picornaviridae [1,2]. The virus contains a single-stranded, positive-sense RNA genome of approximately 7,400 nucleotides in size packaged inside a non-enveloped icosahedral capsid composed of 60 copies of protomers of each of the proteins VP1 to VP4.

The genome can act as mRNA and is translated in a cap-independent manner into a single polyprotein which is subsequently processed by virus-encoded into the structural and nonstructural proteins. The virion becomes infectious following the maturation cleavage of the interim product VP0 into the VP2 and VP4 [3] by an autocatalytic mechanism [4,5].

CVA16 and EV71 are the major pathogens of hand, foot and mouth disease (HFMD) [6] among a dozen of other enteroviruses [7], such as CVA4, 5, 6, 9, 10, 12 and CVB3 [8-15], affecting mainly children under 5 years of age [16]. CVA16 is associated less with severe neurological complications than EV71, although this has been reported [17-19].

Data from the Chinese CDC shows that since the outbreak of HFMD in 2008, the reported cases are increasing and peaked in 2012 at more than two millions in China [19]. EV71 and CVA16 as well as other serotypes of non-polio enteroviruses co-circulated [20] and caused indistinguishable clinical symptoms.

The viruses have evolved rapidly and genome recombination has occurred [21]. The phase 1, 2 and 3 clinical trials of EV71 vaccines are promising [22-29] and EV71 vaccine is available in China. Still, a bivalent or even a multivalent vaccine is required for prevention of the spread of these viruses and complete control of HFMD [16,30-32], since the EV71 vaccine does not provide cross-protection against Coxsackievirus A16 and circulating lineages of other HFMD-associated enteroviruses [29,33].

Like other RNA viruses, enteroviruses are known to have a high mutation rate due to the low fidelity of the virus-encoded RNA-dependent RNA polymerase (RdRp) and due to frequently occurring recombination events [21,34]. CVA16 has evolved quickly since it was first associated with HFMD in 1959 [9,35].

The genotype B and its subgenotypes and clusters have emerged [20]. Additional antigen variants might exist within several serotypes of enteroviruses on the basis of reduced or nonreciprocal cross-neutralization between variants [13,14]. Therefore, a vaccine candidate against HFMD should stimulate immune responses against the prevailing subgenotypes as well as the homologous vaccine strain [3]. In addition, for an inactivated vaccine high yields in a cell culture system and formulation with an optimal adjuvant are also very important [36].

The aim of this study was to screen a CVA16 vaccine candidate against different isolates and their variants in Vero cells, aiming at fulfillment of the criteria mentioned above. The CVA16-P4-L731 was selected from plaque-purified clones after serial passaging of clinical isolates in Vero cells.
Inactivated and non-inactivated full particles (FP) and empty particles (EP) were administered to Balb/C mice at different dosages and with different adjuvants. The vaccine candidate induced high titers of neutralizing antibodies against itself, genotype A and subgenotypes of B. The vaccine protected one-day-old mouse pups born to immunized mothers passively from disease and death after challenge with CVA16 viruses at extremely high doses of LD_{50}.

Materials and Methods

Ethics statement

Animal experiments were performed in accordance with the guidelines of Chinese Council on Animal Care. The research protocol was approved by Animal Care and Use Committee of WIBP.

Cells and viruses

African green monkey kidney (Vero) and human rhabdomyosarcoma (RD) cells were grown in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% of fetal calf serum (FCS) (Gibco, Sera), 100 U/ml penicillin and 0.1 mg/ml streptomycin. Virus stocks were produced in flasks at an MOI of 0.01 in DMEM without FCS.

The 2- and 10-stack cell factories (Corning Inc) were used for a large-scale propagation of viruses at an MOI of 0.001. The supernatant of infected cells was harvested when a cytopathic effect (CPE) of 80% was reached. Viruses were titrated by the method of 50% tissue culture infective dose (TCID_{50})/ml as described previously [37].

Isolation, passaging and plaque purification of a vaccine candidate

The CVA16 P4 strain was isolated from a clinical sample of a patient with HFMD in Hubei in 2010 and passaged serially 11 times in Vero cells at 0.1 MOI. Plaque-to-plaque purification was performed 3 times using 0.5% agarose-DMEM overlay and an identical overlay containing 0.01% of Neutral Red at 3-4 days post-infection. One of clones, referred to as CVA16-P4-L731, was passaged 3 more times in Vero cells for further characterization.

Viral RNA amplification and sequencing

Viral RNA was extracted from virus infected Vero cells using Quantum Viral RNA Mini Kit (Qiagen) following the manufacturer’s instructions. Reverse transcription-polymerase chain reactions (RT-PCR) were performed using random and CVA16-specific primers, respectively, following the manufacturers’ protocols (TaKaRa). Nine pairs of primers were used to generate overlapping PCR fragments covering the whole genome for sequencing. Sequencing was performed by Genescript Co.

Virus purification

The supernatant of infected cells was clarified by centrifugation at 5°C at 10,940 g in a Beckman JA10 rotor for 15 min and by microfiltration through 0.45 μm filter (Sartorius Intec). Viruses were concentrated by ultrafiltration using a 100 kDa tangential flow filter capsule (Sartorius Intec) to the 1/10 of the original volume. The concentrate was centrifuged through 5 ml of 20% (W/V) sucrose cushion in a Beckman SW28 rotor at 141,000 g for 6 hr at 5°C.

The viruses were dissolved in 0.1 mM CaCl\(_2\)-10 mM phosphate buffered saline (PBS) at 4°C overnight and centrifuged at 5°C on a 15-55% (W/W) of a sucrose gradient in a Beckman SW41 rotor at 288,000 g for 3 hr. The 80S- and 160S-particles, referred to as empty and full particles (EP and FP), respectively [33], were further purified by centrifugation of a virus mixture at a density of 1.31 g/ml of CsCl at 5°C in a Beckman Ti-90 Rotor at 214,197 g for 18 hr.

CaCl was removed from diluted virus fractions by centrifugation in a Beckman SW41 rotor at 288,000 g for 2 h at 5°C. The concentration and purity of viral proteins were determined by SDS-PAGE using a BSA standard (Thermal, Pierce) and by densitometry scanning with Densimeter and software Quantity One (Biorad). The total protein concentration was also determined by a BCA protein assay (Thermal, Pierce).

Transmission electron microscope (TEM)

The 200 mesh, carbon-coated copper grids were soaked in a drop of the purified particle suspension for 5 min, stained with a drop of 1% of uranyl acetate, pH 6.8 for 5 min and air-dried overnight. The grids were viewed under the TEM (FEI Tecnai G2 20 TWiN), and images were photographed.

SDS-PAGE and Western blotting

Proteins of purified CVA16 particles or infected-cell lysates were separated in a 12.5% SDS-PAGE gel and transferred onto NC membranes. The membranes were blocked overnight at 4°C in 1% BSA in PBS containing 0.05% Tween-20 (PBST). Each membrane strip was incubated for 2 hr with individual mouse serum.

A horseradish peroxidase-conjugated goat anti-mouse secondary antibody (Boster, China) was added at a dilution of 1:2000 in PBST-BSA and incubated for 30 min. The membranes were washed three times with PBST after each incubation step. The protein bands were visualized by adding DAB-substrate solution.

Enzyme-linked-immunosorbent-assay (ELISA)

Microplates (YunPeng, China) were coated with 200 ng/well of purified FP or EP in 100 μl of 0.1 M carbonate-bicarbonate buffer (pH 9.6) at 4°C overnight. The plates were washed 5 times with PBST after each incubation step. Each well was blocked with 150 μl PBST-BSA at 37°C for 1 hr. Then 100 μl of serial twofold dilutions of mouse serum in BSA-PBST were incubated at 37°C for 1 hr.

Each well was incubated with 100 μl of horseradish peroxidase-conjugated goat anti-mouse IgG (Boster, China) diluted with PBST-BSA at 1:10,000. The plates were incubated at 37°C for 1 hr. Then 100 μl of the substrate solution (Sigma) was added to each well and incubated at 37°C for 30 min. Subsequently, 50 μl of 1M H\(_2\)SO\(_4\) was added to each well, and absorbance at 450 nm wavelengths was measured in a microplate reader (Multiskan MK3, Thermo).

Neutralizing antibody assay

Mouse sera were inactivated at 56°C for 30 min, and serial 2-fold dilutions in DMEM were made. Viruses were diluted in DMEM so that 50 μl of the virus suspension contained 100 TCID\(_{50}\). Equal volumes (50 μl) of serum dilution and virus suspension were mixed and added to each well in 8 duplicates for each dilution.
After incubation at 37°C for 2 h, 1 x 10^4 RD or vero cells in 100 μl of DMEM per well were added. Cells were fixed and stained 7 days later. The virus back-titration was performed and titers were in a range of 32 to 320 TCID 50/50 μl. The neutralizing titer was calculated using the Reed-Muench method [38] and expressed as the reciprocal of the highest serum dilution at which CPE in 50% of the wells was completely inhibited.

Immunogenicity and efficacy studies in mice

The purified FP and EP were tested or untreated with commercially-supplied formaldehyde at dilution of 1:2,000 at 37°C for 48 hr. Formaldehyde was removed by ultracentrifugation of particles as described above and residual infectivity were tested in Vero cells. Each antigen were emulsified in 1 ml/ml of Al(OH)3 or 0.2 mg/ml of monophosphoryl lipid A (MPL-A). Groups of ten 6-8 week-old female Balb/C mice were immunized intramuscularly with 0.1 ml of the most recently circulating CV A16 isolates in mainland China and commercially-supplied formaldehyde at dilution of 1:2,000 at 37°C for 48 hr. Formaldehyde was removed by ultracentrifugation of particles as described above and residual infectivity were tested in Vero cells. Each antigen were emulsified in 1 ml/ml of Al(OH)3 or 0.2 mg/ml of monophosphoryl lipid A (MPL-A). Groups of ten 6-8 week-old female Balb/C mice were immunized intramuscularly with 0.1 ml of antigen at 2 week intervals.

Five mice in each group were sacrificed and the sera were collected on days 10, 21 and 35 after priming. Seven days after the second boosting, spleen lymphocytes were collected for analysis of T-cell-mediated immune responses. For challenge experiments, five mice in each group were mated with male mice on the day of first boosting.

Seven days post-second-boosting, groups of 7-16 one-day-old pups were challenged intraperitoneally with known LD50 doses of BIII and P5 strains of CVA16, respectively. The mice were observed daily for 14 days for clinical signs, limb paralyses, eye irritation, loss of balance and death.

IFN-γ Enzyme-linked immunospot (ELISPOT) Assay

The mouse IFN-γ ELISPOT kit (Dakewei Biotech) was used to determine the number of IFN-γ expressing cells in the single-cell suspension following the manufacturer’s instruction. Lymphocytes from spleen of three mice in each group were prepared using EZ-Sep Mouse Lymphocyte Separation Medium following the manufacturer’s recommendation.

Cells were diluted to 2 x 10^6/ml with Lympho-Spot serum-free medium for rodent containing stimulus, 2 μg/well of phytohemagglutinin or 0.5 μg/well of inactivated FPs and 2 x 10^5 cells were added per well. IFN-γ spot-forming cells (SFCs) were enumerated using an ELISPOT Reader (Biosys Bioreader 4000). The number of spots in triplicate wells with medium only was subtracted from the number of spots in test wells. The mean number of antigen-specific IFN-γ SFCs per million cells for each group in triplicate wells was calculated.

Results

Isolation, plaque-purification and characterization of CVA16-P4-L73

The virus was chosen from 20 plaque-purified clones as a vaccine candidate based on (1) its high similarity in the VP1 sequences with the most recently circulating CVA16 isolates in mainland China and Taiwan [30,39]; (2) high yields in Vero cells; (3) high ratio of FP to EP. Sequence (GenBank accession number, KF924762) comparison showed that it belongs to CVA16 cluster B2b (Supplementary Figure 1).

Proteins of the mixed FP and EP of virus preparations were separated by SDSPAGE, scanned by densitometry, and relative yields of individual proteins were calculated (Figure 1A).

Based on the molar ratio of VP0 (representing EP) and VP2 (representing FP), the EP and FP ratios were determined. The purities of EP and FP preparations (Figure 1B) were more than 99%. The EP consisted of VP0, VP1 and VP3, representing the empty procapsids.

The FP contained VP1, VP2, VP3 and VP4 as well traces of VP0. The FP contained less VP2 than calculated if the amounts of VP0 were fully cleaved, suggesting that some of the FP were not matured and infectious. Thus, the FPs are a mixture of provirions and mature, infectious particles [40,41].

When formalin-treated and untreated particles were compared by transmission EM (Figure 1C), defined shapes were observed for untreated FP and EP.

The formalin-treated FP appeared to have a less-dense outline than the untreated FP as also observed by Liu [33]. This image may have resulted from formalin cross-linking of viral proteins.

Immunogenicity of inactivated and non-inactivated FP and EP

Vaccine adjuvants are fundamental to stimulate an intense, durable and fast immune response in the presence of low doses and of inactivated antigens. Therefore, Al(OH)3 and MPL-A were co-administered with viral particles to investigate the roles played by them in enhancing humoral and T cell-mediated immune responses.

The FP and EP were formalin-treated and untreated for a fair comparison of immunogenicity of the four particle preparations.
Table 1: Neutralizing titers of antisera of immunized mice and protective efficacies of vaccine formulations in the pup mouse challenge model. Antisera were collected from groups 7 and 8 as indicated in Table 1. Formalin-treated L731/FPs was used for immunization of mice.

The total level of anti-CVA16 IgG in each group at day 7 post-second-boosting was determined by a semi-quantitative ELISA.

As shown in Table 1, titers were detected for all mouse groups immunized with antigens in a dose dependent manner. Higher titers were detected in the EP-coated assays than in the FP-coated assays.

No antibodies were detected using the sera from naive, PBS- and PBS-adjuvants-injected mice. If not treated with formalin, the EP induced higher antibody response than the FP. Interestingly, if treated the FP stimulated higher antibody response than the EP. Furthermore, if treated both, the FP and FP+EP mixtures, induced stronger antibody responses than the EP did. When the FP were formulated with MPL-A, the antibody response was weaker than those with Al(OH)₃.

The specificity of anti-sera from the immunized and mock-immunized mice by WB assays. Antiserum against CVA16 vaccine strain raised from immunized mouse only recognized the VP1 (Figure 2A, lanes 7 and 8) in the lysates of cells infected by CVA16-P2 and CVA16-P4 but not by EV71-Q1 or mock infected cells (Figure 2A, lanes 5 and 6).

Figure 2: Specificity of anti-sera from the immunized and mock-immunized mice by WB assays. Antiserum against Al(OH)₃-1.5μg-FP-injected or from Al(OH)₃-injected mice (A) were used to detect viral proteins in lysates of infected or mock-infected cells. GAPDH was used as the loading control. Equal amounts of FP (B and C) or EP (D) were used for blotting. Reference antibodies used are indicated (B).

Specificity of antibodies was determined by WB assays (Figure 2). As shown in Figure 2A, antiserum against CVA16 vaccine strain raised from immunized mouse only recognized the VP1 (Figure 2A, lanes 7 and 8) in the lysates of cells infected by CVA16-P2 and CVA16-P4 but not by EV71-Q1 or mock infected cells (Figure 2A, lanes 5 and 6).
The antisera recognized other structural proteins weekly under these conditions. The antisera from control mouse did not recognize the VP1 in these same lysates (Figure 2A, lanes 1-4).

In order to demonstrate the specificity of the VP1 detected in Figure 2A, viral proteins of purified CVA16 particles were detected with rabbit anti-virion, anti-VP1, anti-VP2 and mouse monoclonal antibody against VP1(Figure 2B).

When the FP was blotted (Figure 2C), sera from the FP-immunized mice recognized the VP1, VP2 and VP3 while those of the EP-immunized mice recognized the VP1 and VP3 but not the VP2. When the EP was blotted (Figure 2D), antisera of both FP- and EP-immunized mice recognized the VP1 and VP3 but not the VP0. The results showed that the antigenicity of the VP0 in the EP was poor.

The extra larger bands might represent the cross-linking of viral proteins as the formalin-treated FP and EF were used in these experiments. The neutralizing titers of sera were determined (Table 1).

It was observed that the higher titers (groups 3 to 8), or the roughly same titers (groups 14 and 15) were reached when low dose of 1.5 μg of particles was used. Dose dependent effects were recorded only in groups 9 to 12. The FP induced higher titers than the EP and the untreated preparations higher titers than the formalin-treated ones.

### Dynamics of humoral immune response

The titers of neutralizing antibody after priming and the first and second boosting at two dosages (1.5 and 4.5 μg) of Alum-inactivated-FP were compared (Table 2). The results showed that 1.5 μg-inactivated FP induced the same level of antibody post-first-boosting and much high level of antibody post-second-boosting than the 4.5 μg dosage.

**Table 2:** Comparison of titers of neutralizing antibodies after each of immunization in Balb/C mice; Sera were collected 10 days post-priming, 7 days post-first and second boosting at an interval of 14 days.

<table>
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<tr>
<th>Dose/Adjuvant</th>
<th>Neutralizing antibody titers</th>
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<tr>
<td></td>
<td>Priming</td>
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<tr>
<td>4.5 μg/Al(OH)₃</td>
<td>24</td>
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<tr>
<td>1.5 μg/Al(OH)₃</td>
<td>16</td>
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**Cross-neutralizing activity of mouse anti-sera against other strains**

As shown in Table 3A, antisera, pooled from the groups of immunized mice, cross-neutralized other isolates with different VP1 or genome sequences.

Furthermore, the prototype G10 and other subgenotypes were also well neutralized by the mouse antiserum pool or by rabbit hyperimmune antiserum (Table 3B).

Also, L731 was neutralized by human reference sera collected from patients infected with currently prevailing CVA16 strains (Table 3B).

The results suggested that the CVA16-P4-L731 vaccine candidate might provide broad-spectrum-protection against infection and disease caused by different genotypes and prevailing isolates of CVA16.
As shown in Figure 3 and Table 1, for the vaccine groups the percentage of survival ranged between 45.5-100% for BIII and 53.9-100% for P5. In contrast, groups of pups born of naïve, PBS-, Al(OH)₃, and MPL-A-injected mice fell ill and died within 3-4 days and 4-5 days post-challenging with BIII and P5, respectively.

![Figure 3: Determination of lethal doses of BIII (A) and P5 strains (B) and efficacy of vaccine formulations (C and D). One-day-old naïve mice were used for determination of LD₅₀. One-day-old pups born to immunized or mock-immunized female mice were challenged with BIII (C) and P5 (D) by IP route at dosages indicated. Percentage of protection of mice after challenging with 10,000 times of LD₅₀ of (C) homologous BIII (2.00 x10⁵ TCID₅₀) and (D) heterogonous P5 (1.33x10⁶ TCID₅₀) strains within 14 days.](image)

Specific proliferation of IFN-γ secreting T cell

The T cell mediated immune response to the vaccine strain was evaluated. As shown in Figure 4, the FP emulsified with Alum at all doses enhanced the proliferation of INF-γ secreting cells, specifically after in vitro stimulation of spleen cells with the purified vaccine strain.

The FP at a high dose co-administrated with MPL-A-Alum mixture significantly increased this innate immune response compared to administration of FP with alum alone.

![Figure 4: Proliferation of IFN-γ secreting spleen cells of immunized mice after in vitro stimulation with the FP. The purified and inactivated FP was used to stimulate the spleen cells in vitro. MPL-A, monophosphoryl lipid A. Alum, Al(OH)₃.](image)

Discussion

A bivalent EV71-CVA16 or polyvalent vaccines including other enteroviruses causing HFMD may be needed in a long term perspective of HFMD prevention and in case of the emergence of new virulent viruses with antigenic variations. Bivalent vaccine, or polyvalent vaccine in circumstances of outbreaks caused by CVA6 [12], CVA9 [8,10], CVA10 and CVA12 [15], would increase public confidence and acceptance of HFMD vaccines [16,32].

A major issue for multivalent vaccines is the balancing of immunogenicity for each of the individual viruses. Based on published reports [31,42,43] and our experience, the immunogenicity of CVA16 is much weaker than that of EV71. For EV71, cross-neutralizing activity of antibodies against different genotypes is generally high [30,44]. For CVA16, however, lower cross-neutralizing activity or one-sided cross-activity may exist between different genotypes or even within subgenotypes as indicated in this study although a report describes high cross-neutralizing activities [45]. Unlike L731, the LZ134 strain was not well neutralized by the human reference antisera, showing more than 10 times differences in titers of neutralizing antibodies against other strains. The genetic variations among subgenotypes and clusters might result in antigenic difference, making it difficult to select appropriate vaccine candidates. Our data suggest that the neutralizing epitopes determine the cross-neutralizing activity instead of genotypes, subgenotypes or clusters based on the VP1 nucleotide sequences. For example, the identity of L731 (B2b) is 79.0% and 91.8% in nt sequence with the G10 (A) and P6 (B2a) and there are 27 and zero amino acid changes in the VP1, respectively. However, the cross-neutralizing titers are the same or very close to those against itself (Table 3).

The experimental evidence has shown that CVA16-P4-L731 is a vaccine candidate with excellent immunogenicity against itself and...
other relatively closely-related enterovirus strains. The titers of neutralizing antibodies reached to 8,192 for the non-inactivated and 4,291 for inactivated FP, the highest reported so far, in a neutralization assay with an end point of complete inhibition of CPE in seven days rather than in 3 days [39,46]. The neutralizing titers were much higher than those previously reported [31,43,44].

The EP and FP of EV71, and maybe those of CVA16, are different in size, conformation, protein cleavage and density due to packaging of viral RNA [40,47,48]. Our results demonstrated that immunization with FP stimulated higher titers of neutralizing antibodies than with EP in agreement with the report Chou AH, et al. [31]. The reasons for this observation might be as follows: The EP degraded quicker than the FP; some of neutralization specific epitopes were not formed or exposed before the final cleavage of VP0 into the VP2 and VP4; or viral RNA might function as an adjuvant. We found that the FPs were better than the EPs at stimulating high levels of neutralizing antibodies. Western blot assays showed that the EPs did not induce much of the anti-VP2 antibody and none of the mouse sera recognized the unprocessed VP0. If the VP2 of CVA16 carries the neutralization specific epitopes, the non-cleaved VP0 in EPs may not induce antibodies against them.

It has been documented that formalin treatment could partially damage the neutralization specific epitopes of viral vaccines [31]. However, it was also reported that viral proteins purified from formalin-inactivated poliovirus induced neutralizing antibodies but not those from untreated virus, as the treatment protected the proteins from degradation [49]. The results presented here support the use of formalin-inactivated CVA16-EV71 bivalent vaccine against HFMD. Vaccines against HFMD formulated with adjuvants enhancing both B cell- and T cell-mediated immune responses might be more effective [50,51]. This might be achieved by use of a combination of alum and MPL-A adjuvants as shown in this study.

Tremendous observations indicate that interferon (IFN) plays a pivotal role in the antiviral immunity [47] and transcription factors (TFs), such as IRF3 and IRF7, along with NF-kB and AP1, have been found to be essential for the initiation of IFN genes transcription in human [52,53]. In addition to these TFs, which are able to directly bind to the promoters of IFN genes, some epigenetic modifiers also are also involved in these processes. For instance, histone H3K9 modifier G9a has been found to reversibly regulate the expression of IFN genes in humans [44]. As recent observation indicates that G9a is also required for the maintenance of imprinted DNA methylation via interaction with DNMT3s [54], the promoters of IFN genes are very likely to be protected from establishment of DNA methylation, a stable epigenetic mark that associated with gene suppression. Therefore, it will be a new direction to investigate the underlying mechanisms that associated with occupying and preventing gain of 5 mC DNA methylation at the promoters of IFN genes.

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Conflicts of Interests

The authors declare that they have no conflicts of interests. The information on this work was not presented previously in any meeting and was not submitted for publication elsewhere.

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