Quantitation of IgG and IgM Response to Recombinant Zika Virus Proteins

Imraan Ali, James P. Chambers, Masarrat Ali and Richard C. Tamfu*

Alpha Diagnostics International, Inc., San Antonio Texas, USA

Corresponding author: Richard C Tamfu, Alpha Diagnostics International, Inc., San Antonio Texas, USA, Tel: 210-561-9515; Fax: 210-561-9544; E-mail: rtamfu@adi.com

Received date: August 11, 2017; Accepted date: August 30, 2017; Published date: September 05, 2017

Copyright: ©2017 Ali I, et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Abstract

Background: Currently, neither vaccine nor drugs are available for treatment of Zika infection. Vaccines for Flaviviruses have been developed using whole inactivated Yellow Fever virus (YFV) as well as YFV-vectored constructs expressing Dengue Envelop-PrM protein. Thus, Flavivirus proteins show promise as vaccine candidates. Using a rabbit model, the aim of this work was to evaluate the response elicited to recombinant Zika Envelop and NS1 proteins.

Methods: His tagged (carboxy terminus) recombinant proteins were expressed in E. coli, purified using Ni-NTA Agarose and imidazole elution, and used to immunize adult New Zealand white male rabbits. Envelop and NS1 antibody ELISAs were developed, and levels of IgG and IgM assessed during the course of vaccination.

Results: Zika recombinant Envelop protein elicited a robust IgG response (4500 µg/ml serum) in contrast to the NS1 protein (~100 µg/ml serum). Both proteins elicited an IgM response; although, significantly lower, i.e., ~25-50 µg/ml serum. E. coli expressed Envelop and NS1 elicited IgG reacted robustly compared with sf9 cell expressed Envelop and NS1 glycosylated homologs. Zika Anti-Envelop IgG cross-reacted with E. coli expressed YFV and Dengue Type I Envelop homologs (approximately 20% control); whereas, Zika anti-NS1 IgG cross-reacted (approximately 20% control) with E. coli expressed WNV NS1 homolog.

Discussion and Conclusion: Zika Envelop protein is unique among Flaviviruses; although, parts of it resemble West Nile, Japanese Encephalitis, and Dengue virus homologs. Although some cross-reactivity, i.e., ~25% A450 values was observed, and as with all animal model derived data there are limitations to human extrapolation, data presented here support the potential usefulness of 1) Zika recombinant Envelop protein as a vaccine candidate, and 2) Zika Envelop specific ELISA reagents.

Keywords: Zika; Flavivirus; Recombinant Env and NS1 proteins; Immune response; Protein vaccines

Introduction

Zika virus is a Flavivirus first discovered in the serum of a rhesus macaque monkey in the Ugandan Zika Forest in 1947, and subsequently in humans [1]. It is transmitted by the Aedes aegypti mosquito [2] which is endemic to the Southern, Eastern, and Mid-Eastern parts of the United States. Due to the large area inhabited by the mosquito [3], and the proximity to South America and the Caribbean where the most recent outbreak originated, it has become a major health concern in the United States [4]. Currently, there has been 5,381 confirmed cases of Zika in the US [5]. The majority of cases observed have been travel-related with the first 14 cases arising by localized transmission in Miami, Florida in July 2016 [5]. Due to the localized nature of infection, the CDC for the first time in its history issued a warning not to travel to an American neighborhood due to an outbreak of an infectious disease [6].

Zika viral infection does not initially give rise to severe symptoms with 80% of all cases being non-symptomatic, and the remaining 20% giving rise to minor flu-like symptoms [7]. Currently, no vaccine or anti-viral therapeutic agents exist for treatment of Zika infection [7].

The major concern for Zika viral infection is the increase in microcephaly observed in pregnant women in mosquito infected areas prompting the CDC to conclude Zika virus as being the causative agent [8]. Microcephaly is a lifelong condition associated with much smaller head size-giving rise to symptoms ranging from seizures, developmental delay, cognitive deficit with movement/balance, feeding, hearing, and vision problems, and in severe cases death [9]. Additionally, Guillain-Barré Syndrome (GBS) is a peripheral nervous system disorder increasingly observed in individuals in areas where Zika virus is prevalent [9]. Although the greater majority of GBS individuals recover from the most severe cases, some individuals experience long-term symptoms [8].

Materials and Methods

Recombinant protein expression and purification

cDNA for Zika virus envelop protein corresponding to amino acids 1-251 (Accession# ABL43461.1), and cDNA for Zika virus NS1 protein corresponding to amino acids 795-1148 (Accession# BAP7441.1) were optimized and synthesized for expression in E.coli. Each respective cDNA was sub-cloned into PET28A vector using Ncol and Xhol restriction sites. Following transformation of BL21 (DE3)
cells with each construct, respective colonies were screened for recombinant protein expression. Briefly, cells were grown in LB-Kanamycin at 37°C in a shaker (150 rpm) to OD_{600} of 0.5-0.6 followed by induction with 0.5 mM IPTG for recombinant protein expression for an additional 3 hours. Cells were lysed in 50 mM Tris-HCl, pH 8.0 buffer containing 0.2 mg/ml lysozyme, 0.5 mM EDTA, 1 mM PMSF, 5 mM β-mercaptoethanol and 250 mM NaCl on ice. Lysates were sonicated 3 times (30 seconds) on ice (60% full power setting). Whole cell lysate material was centrifuged at 10,000 rpm for 1 hour using a GS3 rotor to pellet inclusion bodies. Inclusion bodies were washed twice in the above buffer, and inclusion body proteins were extracted in buffer containing 8M urea. His-tagged extracted proteins were applied to a Ni-NTA Agarose column, and eluted with buffer containing 0.25M imidazole. Protein was quantitated using the BioRad dye-binding assay [10], and BSA as standard. Protein purity was assessed by SDS-PAGE, and proteomic analysis (see below).

SDS-PAGE and proteomic analysis of recombinant proteins

Electrophoresis was carried out under denaturing conditions according to the method of Laemmli [11]. Recombinant Envelop and NS1 protein (1.6 and 1.4 µg, respectively) were heated for 5 minutes at 95°C in the presence of SDS and β-mercaptoethanol, and applied to 10% Bis-Tris gels (Life Sciences Technology, Inc., Hewlett, NY). Electrophoresis was carried out at constant voltage (80V) for 1.5-2.0 hours at room temperature. Gels were stained with Coomassie Blue. The gel bands of interest were reduced in situ with TCEP [tris(2-carboxyethyl)phosphine] and alkylated in the dark with iodoacetamide prior to treatment with trypsin (Promega). The digests were analyzed by capillary HPLC-electrospray ionization tandem mass spectrometry (HPLC-ESI-MS/MS) on a Thermo Fisher Orbitrap Fusion Lumos mass spectrometer fitted with a New Objective PicoView 550 nanospray interface. On-line HPLC separation of the digests was accomplished with an RSLC NANO HPLC system (Thermo Fisher/Dionex). Precursor ions were acquired in the orbitrap from m/z 300-m/z 1500 at 120,000 mass resolutions (m/z 200); data-dependent higher-energy collision dissociation (HCD) spectra were acquired at the same time in the linear trap using the “top speed” option. Mascot (Matrix Science) was used to search the spectra against the virus subset of the UniProt database (downloaded on 4/11/2017) concatenated to a database of common protein contaminants. Cysteine carbamidomethylation was set as a fixed modification and methionine oxidation, deamidation of glutamine and asparagine, and acetylation of the protein N-terminus were considered as variable modifications; trypsin was specified as the proteolytic enzyme, with one missed cleavage allowed. Subset searching of the identified proteins was achieved by X! Tandem. cross-correlation with Mascot results and determination of protein and peptide identity probabilities was accomplished by Scaffold (Proteome Software). The thresholds for acceptance of peptide and protein assignments in Scaffold were 95% and 99%, respectively, two peptides minimum per protein (peptide and protein FDR<0.1%). Mass spectrometry analyses were conducted in the Mass Spectrometry Core Laboratory of The Department of Biochemistry, The University of Texas Health Science Center at San Antonio.

Animals

Adult, New Zealand White male (5-6 lbs.) rabbits were used for antibody production in a USDA sanctioned facility with approved IACUC protocols.

Immunization protocol

Rabbits (one for each antigen) were immunized by subcutaneous injection (3-4 sites) of recombinant Zika protein antigens: Envelop protein 50 µl (135 µg protein/ml) and NS1 protein, 50 µl (130 µg protein/ml). Due to very low antibody titers following 5 injections, NS1 dosage was doubled, i.e., 260 µg/50 µl for injections 6 and 7. A pre-immune bleed was collected prior to immunization (Day 0). Immunization was initiated on Day 1, and subsequently performed at 2, 4, 6, 8, 10, and 12 weeks. Bleeds were collected at 3, 7, 9, 14, 16, and 17 weeks, respectively. Serum was separated at room temperature by centrifugation (15 minutes at 3000 rpm), and stored at -20°C until used.

ELISA plate coating

ELISA plates were prepared by coating overnight at 4°C. For Envelop protein IgG and IgM detection, antigen was coated at a concentration of 0.5 µg/ml in PBS. For NS1 IgG and IgM detection, NS1 antigen was coated at concentrations of 0.1 and 0.5 µg/ml, respectively, in PBS. For cross-reactivity assessment (IgG only), Flavivirus Envelop and NS1 related recombinant proteins were coated at 0.2 µg/ml. Plates were washed with 1X wash buffer 3 times. Stablecoat reagent (100 µl) was added to each well, incubated at room temperature for 1 hour after which time wells were aspirated using an ELISA wash apparatus, and subsequently dried at 30°C for 3 hours.

ELISA

Zika Envelop and NS1 ELISAs were carried out as follows: serum (100 µl) was added to each well and incubated for 1 hour. Wells were washed (3 times) with wash buffer followed by addition of 100 µl horse radish peroxidase (HRP) conjugated immunoglobulins, i.e., Goat anti-Rabbit IgG or Goat ant-Rabbit IgM (Alpha Diagnostic International, San Antonio, TX), and incubated for 30 minutes at room temperature. Following washing (5 times) with wash buffer, 100 µl 3,3’,5,5’-Tetramethylbenzidine (TMB) substrate was added to each well, incubated for 15 minutes at room temperature, and reactions were terminated by addition of 100 µl Stop' solution. Absorbance at 450 nm was determined using an ELISA Wash plate reader (MPR-2100, Awareness Technologies, Westport, CT).

Generation of standard curve and quantitation of ELISA IgG/IgM

Titer strips were generated as follows: Rabbit IgG and IgM (Equitech Bio, Kerrville, TX) was coated at specific concentrations, i.e., 3, 10, 30, and 100 ng, respectively. HRP-conjugate (100 µl Goat-anti Rabbit IgG or Goat-anti Rabbit IgM) was added to strips at appropriate dilutions (IgG, 1:200; IgM, 1:100) for 30 minutes. Following washing (5 times), 100 µl TMB substrate was added to each well, incubated for 15 minutes at room temperature, and absorbance at 450 nm determined following addition of Stop' solution. Absorbance for standards was linear from 3 to 100 ng. ELISA absorbance values were all quantitated within this linear range by appropriate dilution. Concentration was determined by multiplication of respective extrapolated A_{450} values by the dilution necessary to achieve A_{450} values within the standard linear range.

Results

SDS-PAGE analysis of expressed Envelop and NS1 recombinant proteins used in this study is shown in Figure 1.
Figure 1: SDS-PAGE and sequencing analysis was carried out as previously described under ‘Materials and Methods’. **Frame A:** Zika Envelope. **Frame B:** Zika NS1.

Figure 2: NS1 and Envelop recombinant protein elicited IgG for different bleeds was determined as previously described under ‘Materials and Methods’. **Frame A:** Anti-serum raw absorbance (mean value of three determinations) values (closed squares=Envelop protein, closed triangles = NS1 protein). A Dilution factor of 1:60,000 and 1:1,500 for Envelop and NS1 recombinant protein was used, respectively. **Frame B:** Envelop (closed squares) and NS1 (closed triangles) IgG titers (mean value of 3 determinations) as a function of post-vaccination time. The titer of elicited IgG was achieved by extrapolation to a standard curve. **Inset:** Rabbit IgG Standard Curve. A rabbit IgG titer strip was prepared as described under ‘Materials and Methods’, and standard curve derived (mean value of 3 determinations, $y = 33.02x-5.82$).

Zika Envelop recombinant protein (expressed in E. coli) antiserum, i.e., IgG was evaluated for cross-reactivity with recombinant Envelop and NS1 homologs from related Flaviviruses. While we observed no cross-reactivity with West Nile virus (WNV), Chikungunya virus (CV), and Japanese Encephalitis virus (JEV) homologs, we did observe approximately 20% control $A_{450}$ values (E. coli expressed Zika Envelope elicited response) cross-reactivity with Yellow Fever (YFV) and Dengue Type 1 virus homologs. Antiserum obtained from rabbits vaccinated with E. coli Zika NS1 recombinant protein exhibited approximately 20% control $A_{450}$ values (E. coli expressed Zika NS1 elicited response) cross-reactivity with WNV recombinant NS1 homolog (Figure 4).
Assessment of Zika recombinant Envelop and NS1 IgM response

Figure 3: NS1 and Envelop recombinant protein elicited IgM for different bleeds was determined as previously described under 'Materials and Methods'. Frame A: Anti-serum raw absorbance (mean value of three determinations) values (closed squares=Envelop protein; closed triangles=NS1 protein). A Dilution factor of 1:600 and 1:300 for Envelop and NS1 recombinant protein was used, respectively. Frame B: Envelop (closed squares) and NS1 (closed triangles) IgM titers (mean value of 3 determinations) as a function of post-vaccination time. The titer of elicited IgM was achieved by extrapolation to a standard curve essentially identical to that shown above. A rabbit IgM titer strip was prepared as described under 'Materials and Methods' and standard curve generated (mean value of 3 determinations, y=33.03x-1.35, data not shown).

Cross-reactivity of related Flavivirus recombinant proteins with rabbit anti-Zika Envelop and NS1 IgG

Figure 4: ELISA plate preparation for cross-reactivity assessment was carried out as described under 'Materials and Methods'. Error bars represent 4 replicate determinations at a dilution of 1:5000 serum with all proteins coated at 0.2 μg/ml. Frame A: Flavivirus recombinant Envelop proteins. Frame B: Flavivirus recombinant NS1 proteins. The mean raw absorbance value is shown above each respective bar.

Discussion

Zika recombinant Envelop protein elicited a robust IgG response starting approximately 3 weeks post-immunization 'up' to 'out' continuing up to 110 days (~4500 μg/ml) consistent with an expected 'adaptive' albeit continuous response following 5 subsequent injections, i.e., roughly 4 months after the initial injection. In contrast, IgG elicited in response to Zika recombinant NS1 protein albeit lower was significant (~100 μg/ml at day 90 post immunization). Both Envelop and NS1 recombinant protein elicited IgM levels were observed to be lower in comparison (2 maxima of approximately 50 μg/ml at 20 and 110 days, and 10 and 25 μg/ml at 50 and 110 days post-immunization, respectively) than that observed for IgG. One possible explanation for the much lower IgM titers is inhibition or cross-reactivity, i.e., competition of immune reagents elicited to similar/identical epitopes. Future experiments involving selective precipitation of serum IgG and subsequent IgM quantitation could shed further light on this issue. If IgG competitively blocks IgM binding to the target antigen, detectable
IgM levels should increase in its absence. It is also noteworthy that a portion of the NS1 recombinant protein was truncated as determined by mass spectroscopy and could contribute to the lower NS1 elicited IgG and IgM responses. NS1 elicited antibodies could be used to discriminate animals previously vaccinated with subunit or killed vaccine which does not elicit antibodies toward NS1 protein from those with active Zika viral infection.

Flaviviruses (Zika, Dengue, West Nile, Japanese Encephalitis, and Yellow Fever) have similar genetic makeup, and the primary antigenic proteins (Envelope, prM, and NS1) share 50-70% protein sequence homology across the group with large regions in Envelope and NS1 proteins more conserved within the Flavivirus family than all others. However, only approximately 20% cross-reactivity of rabbit anti-Zika immunizations were given every 2 weeks in contrast to that used for those with active Zika viral infection. Currently a number of vaccines are in development using recombinant proteins. Although recombinant protein vaccines potentially can be less effective due to being less immunogenic as natural pathogen proteins, i.e., eliciting a less robust Tc response, such an approach offers some advantage over live, attenuated or inactivated vaccines, e.g., stability, no need to use whole organisms, genetic manipulation to increase immunogenicity, and no chance of mutation thus reversing attenuation as in live vaccines. Attenuated recombinant Yellow Fever virus containing the Zika Envelope gene is currently being produced by Sanofi Pasteur (Lyon France). Additionally, Bharat Biotech (Hyderabad, India) and Protein Sciences Corporation (Meriden, CT) are using viral surface antigen expression to induce immune responses [12].

As with all animal model derived data, there are serious limitations to human extrapolation, e.g., vaccination of rabbits was carried out using a Mycobacterium adjuvant that is highly antigenic persisting much longer than other adjuvants [13]. However, Mycobacterium adjuvants are banned for human use by the FDA due to toxicity, and can only be injected subcutaneously or intraperitoneally in animals due to complications that can arise via other routes. Alternatively, aluminum salt adjuvants are available for human vaccine development but could lead to lower antibody titers, as they are not as persistent or antigenic. Additionally, in the rabbit model described here, booster immunizations were given every 2 weeks in contrast to that used for human vaccination in which booster immunizations are typically given several months to a year apart.

Conclusion

The Envelope protein is a major viral surface protein involved in various steps of the Zika virus life cycle. IgG titers produced in response to both Zika Envelope and NS1 recombinant proteins increased with each subsequent injection, i.e., 3 through 7; the peak NS1 IgG titer being roughly 45-fold less than that observed for Envelope recombinant protein. IgM levels elicited by both Envelope and NS1 recombinant proteins were lower, i.e., approximately 82- and 180-fold less, respectively, than the IgG response to the Envelope recombinant protein. Early sequence and structural comparisons of the Zika Envelope protein with that of other Flaviviruses suggest that overall the Zika Envelope protein is unique among Flaviviruses; although, parts of it resemble its homologs in West Nile, Japanese Encephalitis, and Dengue viruses [14,15]. This is consistent with the small amount of cross-reactivity of rabbit anti-Zika Envelope elicited antibodies with related Flavivirus recombinant protein homologs. It remains to be determined if the antibodies produced are effective at limiting Zika virus transmission. Data reported here support the potential of 1) the Zika Envelop protein as a vaccine candidate [16,17], 2) usefulness of the NS1 response in discriminating prior vaccine exposure from active viral infection, and 3) usefulness of Zika specific ELISA reagents.

Acknowledgements

We dedicate this work to the memory of our dear colleague Dr. Masarrat Ali who passed away unexpectedly, April 21, 2017. We gratefully acknowledge Alpha Diagnostics International, Inc. for financial support of this work. The expert technical assistance of Sammy Pardo and Dana Molleur of the Mass Spectrometry Core Laboratory at the University of Texas Health Science Center at San Antonio is gratefully acknowledged.

References


6. Centers for Disease Control and Prevention (2016) Additional area of active Zika transmission identified in Miami Beach.


