Effects of N-Linked Glycan on Lassa Virus Envelope Glycoprotein Cleavage, Infectivity, and Immune Response

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Abstract
Lassa virus (LASV) belongs to the Mammarenavirus genus (family Arenaviridae) and causes severe hemorrhagic fever in humans. The glycoprotein complex (GPC) contains eleven N-linked glycans that play essential roles in GPC functionalities such as cleavage, transport, receptor recognition, epitope shielding, and immune response. We used three mutagenesis strategies (asparagine to glutamine, asparagine to alanine, and serine/tyrosine to alanine mutants) to abolish individual glycan chain on GPC and found that all the three strategies led to cleavage inefficiency on the 2nd (N89), 5th (N119), or 8th (N365) glycosylation motif. To evaluate N to Q mutagenesis for further research, it was found that deletion of the 2nd (N89Q) or 8th (N365Q) glycan completely inhibited the transduction efficiency of pseudotyped particles. We further investigated the role of individual glycan on GPC-mediated immune response by DNA immunization of mice. Deletion of the individual 1st (N79Q), 3rd (N99Q), 5th (N119Q), or 6th (N167Q) glycan significantly enhanced the proportion of effector CD4+ cells, whereas deletion of the 1st (N79Q), 2nd (N89Q), 3rd (N99Q), 4th (N109Q), 5th (N119Q), 6th (N167Q), or 9th (N373Q) glycan enhanced the proportion of CD8+ effector T cells. Deletion of specific glycan improves the Th1-type immune response, and abolishment of glycan on GPC generally increases the antibody titer to the glycan-deficient GPC. However, the antibodies from either the mutant or WT GPC-immunized mice show little neutralization effect on wild-type LASV. The glycan residues on GPC provide an immune shield for the virus, and thus represent a target for the design and development of a vaccine.

Keywords Lassa virus (LASV) · Glycoprotein complex (GPC) · N-linked glycan · Immune response

Introduction
Lassa virus (LASV) belongs to genus Mammarenavirus, family Arenaviridae. The natural reservoir of LASV is Mastomys natalensis in Africa, and humans are infected through direct contact with their excreta or exposure to the aerosol. Between 300,000 and 500,000 people are infected with LASV annually, and the mortality of hospitalized patients ranges from 20% to 70% (Houlihan and Behrens 2017). The United States Centers for Disease Control and Prevention classifies the virus as a Category A bioterrorism agent, and there is currently no drug or vaccine approved by the Food and Drug Administration capable of treating or preventing Lassa fever.

The envelope glycoprotein complex (GPC) of LASV is sequentially cleaved by signal peptidases and subtilisin kexin isozyme-1 (SKI-1)/site-1 protease (S1P) enzyme during the maturation process to obtain the stable signal peptide (SSP), receptor-binding GP1, and envelope fusion protein GP2. Mammarenavirus GPC is a heavily glycosylated protein. It has been estimated that N-linked glycosylation accounts for nearly 30% of the total mass of LASV GPC (Eichler et al. 2006; Hastie et al. 2017; Watanabe et al. 2018). The LASV lineage IV Josiah strain is the most commonly used strain in the development of LASV vaccines. It has 11 N-glycosylation motifs (Asn-X-Thr/Ser, where X is any amino acid except proline) within the GPC
These 11 N-linked glycans are distributed relatively evenly on the surface of the GPC in terms of spatial conformation, with seven glycans on GP1 and four on GP2. Glycans play critical roles in many biological functions associated with the GPC, such as cleavage, folding, receptor recognition, epitope shielding, and immune response (Sommerstein et al. 2015). Investigation on hospitalized patients infected with LASV found that a small amount of neutralizing antibodies could be detected in only a few convalescents at a relatively late stage of disease course, and that the neutralizing power of these antibodies was relatively low (McCormick et al. 1986; Johnson et al. 1987; Gunther et al. 2000; Sommerstein et al. 2015; Robinson et al. 2016). On the other hand, a previous study found that the cellular immune response plays a major role in immune defense against the LASV (ter Meulen et al. 2000, 2004). We hypothesize that the large number of N-linked glycans on the surface of LASV virions shield important epitopes within GPC, which adversely thereby making recovery from Lassa fever difficult. Therefore, studying the role of the 11 N-sugar chains on the GPC of LASV with regards to the host immune response will help to identify the mechanism of immune escape employed by the virus and facilitate subsequent development of effective vaccines or therapeutic antibodies.

**Materials and Methods**

### Cells and Plasmids

HEK 293 T, HeLa, and Vero cells were cultured in Dulbecco’s modified Eagle’s medium (HyClone, Logan, UT, USA) supplemented with 10% fetal bovine serum (FBS, Gibco, Grand Island, NY, USA). The pseudotype VSV bearing the GPC of LASV (strain Josiah, GenBank accession number HQ688673.1) as well as containing the Renilla luciferase (Rluc) reporter gene were generated as previously reported with the titer of $3 \times 10^7$/mL (Wang et al. 2018). The pseudotyped viruses carrying the WT and mutant GPC were also generated and treated with DNase I (Thermo Scientific, Waltham, MA, USA) and RNase A (ST 576, Beyotime, Shanghai, China) to remove the unpackaged DNA and RNA. The genome copies of the pseudotyped viruses were quantified using qPCR, and Vero

![Fig. 1A](image1.png)

**Fig. 1** Proteolytic cleavage of the individual N-linked glycosylation motif mutants. A Schematic diagram and cartoon representation (PDB: 5VK2 (1)) of the glycosylation of LASV GPC. The precursor glycoprotein GPC was cleaved by SPase and SKI-1/S1P. The mature GPC contains SSP (1–58), GP1 (59–259), and GP2 (260–491) domains and is modified by 11 N-linked glycan chains (green). Transmembrane domains are indicated in gray. B–D HEK 293 T cells expressing the wild-type GPC and individual N-glycosylation mutants of N → Q (B), N → A (C), and S/T → A (D), respectively. The expressed proteins were separated via SDS-PAGE, and western blotting was carried out using anti-GP2 antisera. Images chosen are representative of at least three independent assays.
cells were infected with the WT and mutant pseudotyped viruses with the same genome copies.

To generate individual glycan deletion mutants, we introduced three amino acid substitutions (N → A, N → Q, and T/S → A) into the 11 N-glycosylation motifs using 33 pairs of primers (Supplementary Table S1) (Synthesized by Sangon, Shanghai, China). The recombinant plasmids underwent PCR amplification to perform site-directed mutagenesis, after which the template was removed by DpnI restriction digestion, and products were obtained via gel extraction, transformation, and monoclonal identification.

Mice

Specific pathogen-free (SPF) 6-week-old female BALB/c mice were maintained at the Laboratory Animal Center of Wuhan Institute of Virology, Chinese Academy of Sciences (CAS). All mice were fed in independent ventilated cages (IVCs), and the IVCs were kept within an SPF barrier environment for experimental animals. The feed was sterilized via Co 60 irradiation, and water was sterilized using an autoclave.

Immunization Strategy

Immunization was performed via intramuscular injection using 40 μg of each plasmid in the medial thigh of each mouse while avoiding blood vessels. To improve immunogenicity, mice were shocked with an Electro Square Porator (BTX, ECM830) using the cross method with the injection hole as the center. Six mice in each group were immunized three times over a 2-week interval, and at 10 days after the final immunization, mice were euthanized via decapitation; eyeball enucleation was conducted for blood collection.

Separation of Mouse Spleen Lymphocytes

Lymphocyte separation was performed using an EZ-Sep kit (Dakewe Biotech Co., Ltd., Beijing, China) according to the manufacturer’s instructions. Mouse spleen was soaked in Roswell Park Memorial Institute (RPMI) 1640 medium. EZ-Sep separation solution (3–4 mL) was then added to a sterile 3 cm culture dish, over which a nylon mesh was fixed with hemostatic forceps. The spleen was placed onto the mesh for grinding and the grinding solution was rapidly transferred to a 15 mL centrifuge tube along with ~500 μL of serum-free RPMI 1640 along the tube wall; the solution was centrifuged at 800 × g for 30 min. The lymphocyte layer was transferred to a new 15 mL centrifuge tube, followed by the addition of 10 mL serum-free RPMI 1640 medium and centrifugation at 250 × g for 10 min. The supernatant was then carefully removed; the cells were resuspended with 500 μL RPMI 1640 medium containing 10% FBS, and 10 μL was used for tenfold serial dilutions for cell counting. The solution was diluted to a cell density of 2 × 10^7 cells/mL, transferred to a 96-well U-shaped-bottom plate at 100 μL/well, and cultured in a cell incubator with 5% CO 2 at 37 °C.

Detection of CD4^+ and CD8^+ T Cells and Cytokines

The culture system used to stimulate spleen lymphocytes involved the application of a stimulator and co-stimulator [(anti-CD28) + Golgi blocker (BFA) + spleen lymphocytes (2 × 10^6 cells)]. Negative control, positive control, and experimental groups underwent stimulation with phosphate-buffered saline (PBS), phorbol myristate acetate/ionomycin, and polypeptide (Supplementary Table S2) (Synthesized by Bankpeptide, Heifei, China) (Vita et al. 2019), respectively. After 4.5 h in culture, the solution was centrifuged at 800 × g for 3 min; 50 μL TrueStain FeX™ (mouse anti-CD16/32; BioLegend, San Diego, CA, USA) was added to reduce non-specific fluorescent staining, incubated at 4 °C for 10 min, and centrifuged at 800 × g for 3 min. PBS solution was then used to dilute fluorescently labeled antibodies (PE-conjugated rat anti-mouse CD8α, FITC-conjugated rat anti-mouse CD4, PE-Cy™7-conjugated hamster anti-mouse CD3ε), and 7-AAD viability staining solution (BioLegend) by 1:200. They were then added (100 μL/well) for staining at 4 °C for 20 min in the dark. After washing, the cells were fixed, permeabilized, and subjected to intracellular staining. An allophycocyanin-conjugated rat anti-mouse IFN-γ antibody (1:200) was then added and incubated at 4 °C for 30 min in the dark. The cells were filtered through a 200 μm nylon mesh before being loaded onto the flow cytometer (BD FACSArıa III). Cytokines were detected using a cytometric bead array kit (BD Biosciences, Franklin Lakes, NJ, USA) according to the manufacturer’s instructions.

Antibody Titration

The sera were collected from each immunized mouse 10 day after the last immunization to determine specific IgG using cell-based ELISA. The serum was diluted 50-fold, followed by separation into eight gradient dilutions at 1:4 ratios. HeLa cells transfected with the individual glycan deletion GPC or WT GPC were served as antigens, which were blocked, washed, and incubated with sera, followed by detection with HRP-conjugated Affi-niPure Goat Anti-Mouse IgG (Proteintech, Wuhan, China).
Antibody Neutralization

Serum was diluted tenfold with FBS-free medium and then separated into six gradients at 1:2 ratios. Forty microliters of the diluted serum were mixed with 10 μL LASVpv at 37 °C for 1 h. The mixture was added to Vero cells for 1 h incubation. Neutralization activities were determined 24 h later using the Rluc assay system (Promega, Madison, WI, USA).

Statistical Analysis

All results are expressed as the mean ± standard error. A two-tailed Student’s t-test was used to determine statistical significance between groups. All statistical analyses were performed using GraphPad Prism software (ver. 6.0; San Diego, CA, USA). *P < 0.05 was considered statistically significant.

Results

Effects of N-Glycosylation Modification on GPC Cleavage

To investigate the influences of different mutations of the 11 glycosylation motifs upon GPC cleavage, the corresponding asparagine residues were individually mutated to glutamine or alanine, or the serine/tyrosine residues in the motif were mutated to alanine. Thus, 33 mutants were constructed, and western blotting was performed using antiserum against GP2. As shown in Fig. 1, all mutants of the N-glycan motifs within GP2 resulted in slight decreases (~2 kDa) in the molecular weight of the respective GP2 bands relative to the wild-type (WT) GP2 band.

We determined the effect of the N-linked glycans on protease cleavage by testing for the presence of cleaved GP2. Disruption of the N-linked glycosylation motif by substitution of asparagine with the structurally similar glutamine on the 2nd, 4th, 5th, or 8th motif (corresponding to N89, N109, N119, N365) inhibited proteolytical processing, while substituting glutamine for asparagine at the remaining seven glycosylation motifs (corresponding to N79, N99, N167, N224, N373, N390, N395) did not alter GP1–GP2 cleavage (Fig. 1B). Similarly, disruption of the glycosylation motif by substitution of asparagine with alanine on the 2nd, 4th, or 8th motif (corresponding to N89A, N109A, N365A) abolished proteolytical processing, and the mutant in the 5th or 9th (corresponding to N119A, N373A) exerted mild inhibition, whereas the other N to A mutants had no influence (Fig. 1C). Intriguingly, substitution of serine at the 8th glycosylation site with alanine (corresponding to S367A) abolished the GP1–GP2 cleavage, and the mutant in the 1st, 2nd, 5th, or 9th (corresponding to T81A, S91A, S121A, T375A) exerted mild inhibition on the cleavage efficiency (Fig. 1D). It was shown that disrupting the glycosylation motifs by introducing different mutants led to differing results, which might be due to the changes of the residue per se, rather than the loss of the specific glycan. However, all three mutants led to a decrease in cleavage efficiency on the 2nd (N89), 5th (N119), or 8th (N365) glycosylation motif, suggesting that the 2nd, 5th, or 8th N-linked glycan was indispensable for GP1–GP2 cleavage. Given the consistency in the results for the N to A and N to Q substitutions, and that the structures of N and Q are the most similar, we used the N to Q mutant for further research, which caused a minimal change to the spatial structure of LASV GPC while ensuring removal of the glycosylation-site motif.

Effects of N-Glycosylation Modification on the Infectivity of LASV Pseudotype Virus

To evaluate the influence of the individual glycan on the pseudotype virus infectivity, we produced VSV particles bearing the altered LASV GP proteins. The transduction activities were evaluated in Vero cells (Fig. 2). Disruption of the N-linked glycan by the N to Q substitution on the 2nd, 4th, 5th, or 8th glycosylation motif (corresponding to N89Q, N109Q, N119Q, N365Q) led to a significant loss of LASVpv (LASV pseudovirus) entry when compared with LASVpv packaged with WT GPC. Deletion of the 2nd or 8th N-linked glycan (corresponding to N89Q, N365Q) completely inhibited the infectivity, and deletion of the 4th

![Fig. 2 The transduction activity of pseudotyped viruses bearing mutant GPC.](image-url)
or 5th N-linked glycan (corresponding to N109Q, N119Q) led to partial inhibition. These results were in line with the protease cleavage results depicted above, indicating that efficient cleavage of premature GPC was a prerequisite for downstream function.

**Effects of N-Glycosylation Modification on Effector CD4+ T Cells and CD8+ T Cells among Spleen Lymphocytes**

The 11 N-linked glycans on LASV GP play an important role in GPC cleavage and maturation, as well as in pseudovirus infection. Additionally, glycan chain is involved in various aspects of the immune response (Bowden et al. 2009; Abraham et al. 2010; Cohen-Dvashi et al. 2015; Sommerstein et al. 2015; Watanabe et al. 2018). Therefore, we focused on how modification of the 11 N-linked glycans on LASV GP influences the immune response to LASV GPC. We employed recombinant GPC plasmids with N to Q substitution as the DNA vaccine used to immunize BALB/c mice. BALB/c mice were immunized with 13 groups (6 mice per group) of DNA vaccines, including 11 groups of mutated recombinant plasmids, pCAGGS-GPCN→Q. WT GPC, and control pCAGGS. Immunofluorescence analyses of CD3, CD4, CD8, and interferon (IFN)-γ were performed (Fig. 3). The proportions of CD3+ T cells in spleen lymphocytes in 78 BALB/c mice across the 13 vaccinated groups were within the range (~40%) of those observed during normal immune response, whereas the deletion of the 1st, 2nd, 4th, or 8th (N79Q, N89Q, N109Q, N365Q) glycan caused the proportion of CD3+ T cells to decrease slightly when compared to the WT (Fig. 3A). Moreover, abolishment of the 1st, 2nd, or 6th (N79Q, N89Q, N167Q) glycan caused the proportion of CD4+ T cells to decrease, whereas abolishment of the 5th, 9th, or 10th (N119Q, N373Q, N390Q) glycan led to the proportion of CD8+ T cells to increase (Fig. 3B, 3C).

Lymphocytes that secrete IFN-γ represent an effector subset of these cells. First, we found that immunization with the WT resulted in a significant increase in the proportion of both the effector CD4+ and CD8+ cells relative to the control plasmid vaccine group. Moreover, deletion of the 1st, 3rd, 5th or 6th (N79Q, N99Q, N119Q, N167Q) N-linked glycan on GPC significantly enhanced the proportion of effector CD4+ cells, and deletion of the remaining seven N-sugar chains had no effect on their proportion. For CD8+ effector T cells, deletion of the 1st, 2nd, 3rd, 4th 5th, 6th, 9th (N79Q, N89Q, N99Q, N109Q, N119Q, N167Q, N373Q) N-linked glycan enhanced their proportion relative to IFN-γ, and deletion of the 7th, 8th, 10th, or 11th (N224Q, N365Q, N390Q, N395Q) N-linked glycan had no effect (Fig. 3D, 3E).

![Fig. 3](image-url)  
**Fig. 3** Effects of N-glycan deletion on T cells in spleen lymphocytes. The mouse spleen lymphocytes were separated, cultured, and stained with phycoerythrin (PE)-conjugated rat anti-mouse CD8a, fluorescein isothiocyanate-conjugated rat anti-mouse CD4, PE-CyTM7-conjugated hamster anti-mouse CD3ε, and 7-AAD viability staining solution (BioLegend) at 4 °C for 20 min in the dark. A–C Effects of individual N-glycosylation deletions on CD3+ (A), CD4+ (B), and CD8+ (C) T cells in spleen lymphocytes. Effector T cells CD4+ (D) and CD8+ (E) T cells were stained with allophycocyanin-conjugated rat anti-mouse IFN-γ antibody (1:200). Each group contained five to six mice. *P < 0.05, **P < 0.01, ***P < 0.001, ****P < 0.0001, compared to WT.
Effects of N-Linked Glycans on Cytokines Secreted by Spleen Lymphocytes

Among the seven cytokines tested, interleukin (IL)-2, IFN-γ, and tumor necrosis factor (TNF)-α are associated with the Th1-type cellular immune response, IL-4, IL-6, and IL-10 with the Th2-type response, and IL-17A to the Th17-type response. Figure 4 shows that all seven cytokines, except for IL-17A, increased in WT group relative to control pCAGGS group. Specifically, deletion of the individual 3rd (N99Q) to 10th (N390Q) N-linked glycan on GPC significantly increased IL-2 levels in spleen lymphocytes relative to those in mice receiving WT GPC (Fig. 4A), whereas deletion of the individual 3rd, 5th, 6th, 8th, 9th, or 10th (N99Q, N119Q, N167Q, N365Q, N373Q, N390Q) glycan increased IFN-γ levels (Fig. 4B); TNF-α was only slightly elevated when either the 3rd (N99Q) or 9th (N373Q) glycan was abolished (Fig. 4C).

Furthermore, we found that deletion of the 1st (N79Q) or 5th (N119Q) N-linked glycan on GPC significantly increased IL-4 secretion, deletion of the 2nd, 3rd, 4th, or 8th (N89Q, N99Q, N109Q, N365Q) slightly increased IL-4 secretion, and deletion of the remaining five glycans had no effect (Fig. 4D). However, IL-6 levels in either glycan mutant motif showed no significant difference relative to levels in the WT group (Fig. 4E). Furthermore, deletion of the 3rd, 5th, 7th, 8th, 9th, or 10th (N99Q, N119Q, N224Q, N365Q, N373Q, N390Q) N-linked glycan slightly increased IL-10 levels, whereas deletion of the remaining five N-linked glycans had no effect (Fig. 4F).

When studying induction of the Th17-type immune response (Fig. 4G), we observed elevated levels of IL-17A in spleen lymphocytes from mice in the 7th, 8th, or 10th (N224Q, N365Q, N390Q) deletion mutant group relative to those in mice receiving the WT vaccine.

These results indicated that deletion of N-linked glycan on GPC, especially those at the 3rd, 5th, 8th, or 9th site.

Fig. 4 Effects of N-glycan deletions on cytokines secreted by spleen lymphocytes were labeled by using a cytometric bead array kit (BD Biosciences), and IL-2 (A), IFN-γ (B), TNF-α (C), IL-4 (D), IL-6 (E), IL-10 (F), and IL-17A (G) were determined using flow cytometry. Each group contained five to six mice. *P < 0.05, **P < 0.01, ***P < 0.001, compared to WT.
immunization, sera were collected, and the titers were determined

Figure 5 shows that antibody response to N-glycan deletion GPC showed higher titer to mutant GPC than to WT GPC, suggesting that the glycans on LASV GPC might play a role in immune escape by shielding the epitope, thus making it inaccessible to antibodies. Evaluation of which N-glycosylation site affected antibody titers revealed that deletion of N-linked glycan at the 3rd, 5th, 6th, 8th, or 9th site (N99Q, N119Q, N167Q, N365Q, N373Q) significantly increased the antibody titer. The titers obtained from these five groups could reach to approximately $17(-\log 2)$ whereas the titers from the remaining seven groups were approximately $15(-\log 2)$ (Fig. 5A–5L). Similarly, to compare the antibody titers when titrated with WT GPC, we found that deletion of the 3rd, 5th, 6th, 8th, or 9th

(N99Q, N119Q, N365Q, N373Q), primarily enhanced the induction of both Th1 and Th2 immune response. Intriguingly, deletion of the 11th (N395Q) glycan on GPC had no effect on Th1, Th2, and Th17A immune response. These findings suggest these specific glycan chains assist LASV in escaping immune response by reducing host recognition of GPC and precluding induction of the immune response.

**Effects of N-Glycosylation Modification on Antibody Titers**

We then detected antibody titers obtained from the individual glycan deletion mutant immunized serum against each mutated GPC variant using a cell-based ELISA.

**Fig. 5** Antibody response to N-glycan deletion GPC showed higher titer to mutant GPC than to WT GPC. After three rounds of immunization, sera were collected, and the titers were determined using cell-based ELISA. A–L. The titer of the indicated serum to the mutated GPC. M The titer of the indicated serum to WT GPC. *P < 0.05, **P < 0.01, ***P < 0.001, compared to WT.
(N99Q, N119Q, N167Q, N365Q, N373Q) N-linked glycan generated significantly higher antibody titers than other deletion mutants as well as WT (Fig. 5M), thereby indicating a role for the 3rd (N99), 5th (N119), 6th (N167), 8th (N365), and 9th (N373) N-glycosylation sites in the GPC in immune escape and reduced humoral immune response by shielding key GPC epitopes.

**Effects of N-Glycosylation Modification on Antibody Neutralization**

We then evaluated the neutralization ability of the serum against LASVpv infection (Fig. 6). Unfortunately, we found that the serum generated by both the WT GPC immunized group and each of the N-linked glycan deletion mutated variants showed minimal to no neutralizing effect to the LASVpv infection.

**Discussion**

In a variety of virions such as influenza A virus, human immunodeficiency virus, Ebola virus, etc., a glycan shield on the glycoprotein plays critical roles in the host immune response. The LASV GPC harbors 11 N-linked glycans that almost completely encapsulate the GPC. Therefore, modification of these chains represents a possible strategy for identifying the LASV immune-escape mechanism, given that these chains might shield key epitopes on the protein surface from immune recognition. We hypothesized that removal of these chains could expose the epitopes, thereby promoting host recognition and triggering of the immune response.

First, to investigate the role of individual N-linked glycan in GPC expression and function, we found that using three different strategies to abolish the glycosylation motif, the deletion of the 2nd (N89) or the 8th (N365) glycan led to decreased GPC cleavage and pseudotype virus infectivity. These results were in line with previous reports that commented on the importance of these two glycans in LCMV GPC function (Bonhomme et al. 2011, 2013). As the 2nd (N89) and 8th (N365) glycans are conserved in *Mammarenavirus* GPC, it was supposed that both of the glycan chains play essential roles in GPC expression and function (Abraham et al. 2010; Hastie et al. 2017). The 2nd (N89) glycan was reported to interact with H92 in the prefusion conformation, and during endocytosis, the glycan chain rotated and H92 was released, which could bind to the second receptor LAMP1 (Cohen-Dvashi et al. 2015; Hastie et al. 2017). Similarly, the 2nd (N89) glycan of the New World *Mammarenavirus* Machupo virus was reported to form a stacking interaction with F98, which was essential for receptor binding (Abraham et al. 2010; Bonhomme et al. 2013). The 8th (N365) glycan of LASV GPC was the first glycan in GP2. This glycan was reported to interact with Q232 and R235, shielding the fusion peptide at the tip of GP2, and thus contributing to the stability of the prefusion GPC (Hastie et al. 2017).

It was recently reported that the 3rd (N99) and 5th (N119) glycans on LASV GPC shield the neutralizing epitopes of the virus (Sommerstein et al. 2015; Robinson et al. 2016), and the 5th (N119), 8th (N365) and 9th (N373) glycosylation motifs were reported to be located in the epitopes of GPC (Robinson et al. 2016). Most recently, the 10th (N390) and 11th (N395) glycans were reported to occlude the conformational GPC-B epitope located at the stalk of GPC (Hastie et al. 2019). Notably, evidence from human survivor and vaccine development studies have shown that adaptive immune protection in LASV infection is probably conferred mainly by a cell-mediated immune response, especially for Type I IFN (Fisher-Hoch et al. 2000; ter Meulen et al. 2004; Geisbert et al. 2005; Botten et al. 2006; Baize et al. 2009; Mire et al. 2017; Warner et al. 2018; Ibukun 2020). It was supposed that deletion of the glycan chain would expose the epitope and thus increase the immunogenicity of GPC. We found that deletion of specific N-linked glycan residues on the GPC had no effect on the proportions of lymphocytes (CD3+/CD4++CD8−, and CD8+/CD3−) but significantly increased proportions of effector lymphocytes (IFN-γ+/CD4+ and IFN-γ+/CD8+), especially following deletion of glycan chain at the 1st, 3rd, 5th, or 6th glycosylation site (N79Q, N99Q, N119Q, N167Q). Additionally, we observed significant increases in the secretion of molecules involved in Th1 immune response (IL-2 and IFN-γ) following deletion.

**Fig. 6** N-glycan deletions showed little effect in improving the neutralization ability of the antibody from sera against the seven individual glycan deletions in GP1 (A) and four in GP2 (B).

| Serum dilution | Inhibition (%) |
|---------------|---------------|
| 1:10          | 0             |
| 1:20          | -20           |
| 1:40          | -40           |
| 1:80          | -60           |
| 1:160         | -80           |

Legend:
- **N79Q**: N-linked glycan in GP1
- **N99Q**: N-linked glycan in GP1
- **N119Q**: N-linked glycan in GP1
- **N167Q**: N-linked glycan in GP1
- **N89Q**: N-linked glycan in GP1
- **N395Q**: N-linked glycan in GP2
- **N365Q**: N-linked glycan in GP2
- **N390Q**: N-linked glycan in GP2
- **N373Q**: N-linked glycan in GP2
- **N224Q**: N-linked glycan in GP2
- **WT**: Wild-type GPC

![Fig. 6](image-url)
of chain at the 3rd, 5th, 8th, 9th, or 10th glycosylation site (N99Q, N119Q, N365Q, N373Q, N390Q), although this did not affect the secretion of Th2- and Th17-related cytokines. These results support the role of N-linked glycans in inhibiting host recognition and Th1 immune response.

Additionally, analysis of changes in antibody titers from the sera of mice immunized with GPC variants revealed general increases in titer following glycan removal, suggesting that the presence of these glycan chains promotes immune escape by shielding the antigenic epitope. Moreover, we verified that the antibodies generated by all GPC variants showed no neutralizing effect on the WT LASVpv. This indicated that the epitopes exposed by the deletion of N-linked glycans on the GPC did not generate neutralizing antibodies, or the glycan chains shield the neutralizing epitopes of the WT virus.

In summary, we found that the 3rd, 5th, 6th, 8th, and 9th N-linked glycans (N99, N119, N167, N365, N373) likely shield epitopes on the LASV GPC that reduce host cellular and humoral immune responses. These sites can potentially be used as breakthrough points to develop effective therapeutic or prophylactic antibodies against Lassa fever.

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Author Contributions XZ, GX, and WW conceived and designed the study. Experiments were performed by XZ, YL, JC, ZW, and WW. Data were analyzed by XZ, YL, and WW. XZ and WW wrote the manuscript. All authors read and approved the manuscript.

Compliance with Ethical Standards

Conflict of interest The authors declare that there are no conflicts of interest.

Animal and Human Rights Statement All mouse studies were performed according to the Regulations of the Administration of Affairs Concerning Experimental Animals in China (WIVA25201801), and the protocols were reviewed and approved by the Laboratory Animal Care and Use Committee at the Wuhan Institute of Virology, CAS.

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