



Published in final edited form as:

ACS Infect Dis. 2022 March 11; 8(3): 574–583. doi:10.1021/acsinfecdis.1c00585.

Virus-like Particle Display of *Vibrio cholerae* O-Specific Polysaccharide as a Potential Vaccine against Cholera

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Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acsinfecdis.1c00585>.

Scheme S1: $Q\beta$ conjugation to lactose squarate **3**; Figure S1: SELDI-TOF MS result of the conjugation of $Q\beta$ and lactose; Figure S2: The SDS-PAGE of different samples; Figure S3: SELDI-TOF MS result of conjugation of $Q\beta$ to OSP, Figure S4: The MP result of $Q\beta$ triple mutant A38K/A40C/D102C; Figure S5: The MP result of $Q\beta$ WT mutant without RNA; Figure S6: The MP result of $Q\beta$ WT mutant without RNA (a) before and (b) after $Q\beta$ conjugation to lactoside **3**; Figure S7: Mass spectrum of wild-type $Q\beta$ without RNA; Figure S8: Mass spectrum of wild-type $Q\beta$ without RNA after conjugation to lactoside; Figure S9: SELDI-TOF MS result of wild type $Q\beta$ without RNA before and after conjugation to lactoside; Figure S10: Sequence coverage of wild type $Q\beta$ conjugated with lactoside **3**; Table S1: TIC from sequencing of wild type $Q\beta$ conjugated with lactoside **3** (PDF)

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The authors declare the following competing financial interest(s): Xuefei Huang is the founder of Iaso Therapeutics Inc., which is dedicated to the development of next generation of vaccines using the bacteriophage Qbeta platform. All authors declare no other competing interests.

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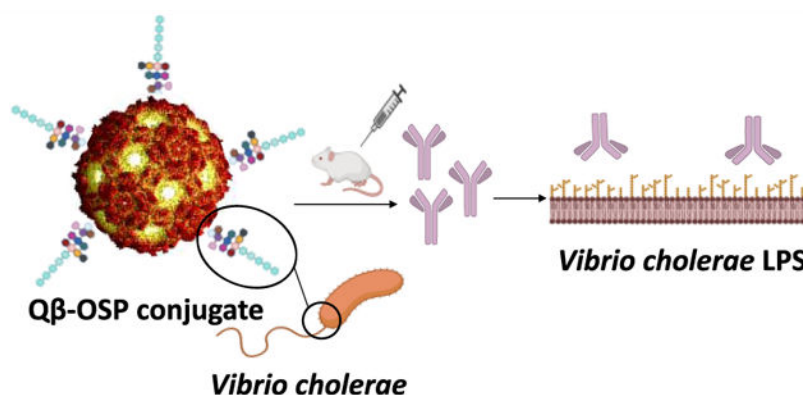
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Abstract

Vibrio cholerae, a noninvasive mucosal pathogen, is endemic in more than 50 countries. Oral cholera vaccines, based on killed whole-cell strains of *Vibrio cholerae*, can provide significant protection in adults and children for 2–5 years. However, they have relatively limited direct protection in young children. To overcome current challenges, in this study, a potential conjugate vaccine was developed by linking O-specific polysaccharide (OSP) antigen purified from *V. cholerae* O1 El Tor Inaba strain PIC018 with Q β virus-like particles efficiently via squarate chemistry. The Q β -OSP conjugate was characterized with mass photometry (MP) on the whole particle level. Pertinent immunologic display of OSP was confirmed by immunoreactivity of the conjugate with convalescent phase samples from humans with cholera. Mouse immunization with the Q β -OSP conjugate showed that the construct generated prominent and long-lasting IgG antibody responses against OSP, and the resulting antibodies could recognize the native lipopolysaccharide from *Vibrio cholerae* O1 Inaba. This was the first time that Q β was conjugated with a bacterial polysaccharide for vaccine development, broadening the scope of this powerful carrier.

Graphical Abstract



Keywords

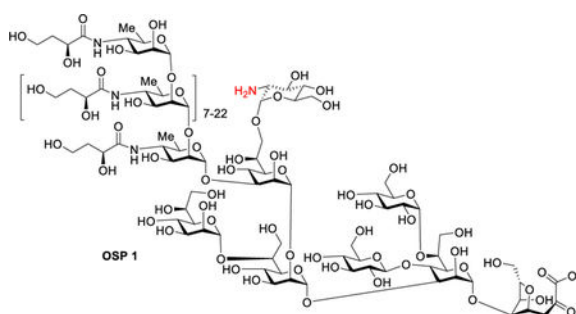
bacteriophage Q β ; mass photometry; O-specific polysaccharide; vaccine; *Vibrio cholerae*

Cholera is an acute, secretory diarrheal disease caused by the highly transmissible bacterium *Vibrio cholerae* (*V. cholerae*). *V. cholerae* are Gram-negative and highly motile bacteria with a single polar flagellum. There are more than 200 serogroups of *V. cholerae* based on the O-antigen of surface lipopolysaccharide (LPS) structures, but only serogroups O1 and O139 are capable of causing epidemic cholera. *V. cholerae* O1 has two serotypes, i.e., Ogawa and Inaba, based on the presence or absence of a methyl group on the nonreducing terminal perosamine moiety of the surface O-specific polysaccharide (OSP, O-antigen).¹ There are 2–3 million cases of cholera each year, resulting in tens of thousands of deaths annually.² Current cholera vaccines include oral killed whole cell vaccine with or without cholera toxin B subunit (CtxB), and attenuated oral cholera vaccine.³ Inclusion of the cholera vaccine into global cholera control strategies has been transformative, but current oral vaccines have the lowest level and duration of protection in young children,^{4–8} who bear a large share of global cholera burden, especially in cholera-endemic countries.^{2,3,9–11} As such, there is a need to develop new cholera vaccines that can provide high-level and long-term immunity.

Immunity protection against cholera infection targets the OSP of *V. cholerae*.^{12,13} However, as O-antigens are T cell independent B cells antigens, direct administration of the O-antigens often only leads to low titers of low-affinity IgM antibodies with limited duration of antibody responses and a lack of induction of immunological memory, rendering O-antigen-based vaccination suboptimal.^{14–16} Covalent linkage of carbohydrate to a carrier protein provides a T cell-dependent immune response by activating CD4⁺ T cells and enables memory B cell proliferation for long lasting antibody protection. Recently, we have demonstrated that self-assembled virus-like particles (VLPs) such as bacteriophage Q β could be used to conjugate with carbohydrate antigens such as the Thomsen–Nouveau (Tn) antigen, ganglioside GM2 and GD2 as potential vaccines.^{17–19} The resulting glycoconjugates were able to induce strong glycan specific IgG antibody responses. However, to date, only low molecular weight (MW generally below 2000 Da) glycans have been investigated for Q β based anticarbohydrate vaccine studies. It is not

known whether bacterial polysaccharide antigens could be conjugated with $Q\beta$ and whether such conjugates could induce strong IgG antibody responses to polysaccharides.

Herein, we report that the native *O*-specific polysaccharide (OSP) 1 of *Vibrio cholerae* O1, Inaba serotype was successfully conjugated with $Q\beta$ through squarate chemistry, which is the first time that a bacterial polysaccharide antigen is covalently linked with $Q\beta$ as a potential vaccine. This approach provides direct conjugation without prior introduction of a linker to the protein carrier. High levels of antipolysaccharide IgG antibodies were induced by the conjugate in mice, and the antibodies were effective in killing the bacteria.

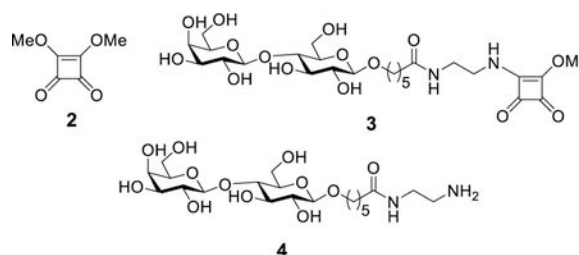


RESULTS

Conjugation of the OSP Core Antigen to $Q\beta$.

To efficiently link the OSP antigen to $Q\beta$, we built upon previous work using squarate based conjugation of polysaccharide.^{20–22} 3,4-Dimethoxy-3-cyclobutene-1,2-dione (dimethyl squarate) **2** is a unique bifunctional linker, both of whose methoxy groups are reactive with amines but under different pH conditions. The first methoxy group can be substituted by a primary amine in neutral pH and the second one is active toward amines in a basic solution. The squaramide moiety itself is stable to hydrolysis under the aqueous reaction conditions.²³ To optimize the conjugation reaction with the squarate ester chemistry, $Q\beta$ was conjugated with a squarate functionalized lactose **3**²⁴ as a model reaction. Lactoside **3** was prepared by derivatizing lactose **4**²⁴ with dimethyl squarate **2** in 76% yield (Scheme S1). In parallel, the coat protein of $Q\beta$ triple mutant A38K/A40C/D102C was expressed in *E. coli*,²⁵ which self-assembled into nanoparticles with average diameters of 28 nm consisting of 180 copies of the monomer. The $Q\beta$ triple mutant A38K/A40C/D102C was selected as the carrier as it has been shown to induce lower levels of antibodies against the carrier itself as compared to the wild type $Q\beta$, thus leading to stronger IgG antibody responses against the target antigen.²⁵ The lactoside **3** (14 equiv per monomer) was then incubated with $Q\beta$ at 22 °C for 20 h leading to the $Q\beta$ -lactose conjugate **5** (Scheme S1). In order to quantify the degree of modification, surface-enhanced laser desorption/ionization time-of-flight (SELDI-TOF) mass spectrometry analysis was performed (Figure S1). On the basis of the intensities of the mass spectrometry (MS) peaks for $Q\beta$ coat protein monomer conjugated with lactosides, it was estimated there was an average loading of 4.5 haptens per $Q\beta$ monomer, corresponding to 810 copies per $Q\beta$ capsid. Increasing the amount of **3** to 28 equiv led to an average of 8 lactosides conjugated per $Q\beta$ monomer unit (1440 copies per $Q\beta$ capsid). As each monomer of the $Q\beta$ triple mutant A38K/A40C/D102C has 9 total

free amines (8 lysines plus the free *N*-terminus), the ability to nearly fully functionalize $Q\beta$ suggests the squarate chemistry is highly efficient in promoting glycan conjugation with $Q\beta$. The 1440 copies per capsid are one of the highest ligand loading levels on $Q\beta$ reported to date.^{18,19,26,27} MS sequencing of $Q\beta$ conjugated with lactoside **3** indicated high functionalization efficiency (80–100%) of lysines (K14, K17, K61, K64, and K68) on the capsid surface and moderate efficiency (46%) of the lysine K3 that lays between monomers in the capsid compared to almost no functionalization (~1%) of K47, the least accessible lysine residue based on the crystal structure of $Q\beta$ (Table S1).



To further characterize the conjugate, gel electrophoresis analysis was carried out (Figure S2). In the presence of reducing agents, the $Q\beta$ capsid disassembled to its subunits showing bands at 14 and 28 kDa corresponding to the monomer and dimer of the coat protein respectively (Figure S2, lane 11). After conjugation, the monomer band of the $Q\beta$ coat protein shifted to about 19 kDa, correlating well with the addition of ~8 lactose units per $Q\beta$ monomer on average (lane 10).

We next explored the conjugation of *Vibrio cholerae* O1 Inaba OSP from PIC018²² with $Q\beta$ using the squarate chemistry. *V. cholerae* O1 Inaba OSP **1** was treated with dimethyl squarate **2** first, which was then incubated with $Q\beta$. To preserve the valuable material, the amount of OSP-squarate was reduced to 8 equiv per $Q\beta$ monomer (Scheme 1). After 120 h, the conjugation reaction was analyzed by SELDI-TOF, which only showed the peak for $Q\beta$ monomer at 14.1 kDa with very weak signals from the potential OSP adduct (Figure S3). The SDS-PAGE of the $Q\beta$ -OSP conjugate (Figure S2, lane 9) showed a very faint band close to the $Q\beta$ monomer MW under the reducing condition with the majority of the protein sample appearing smeared at the high MW region of the gel. The incomplete disassembly of $Q\beta$ -OSP under the reducing condition as compared to $Q\beta$ -lactose may be due to the relatively large size of the OSP (~6 kDa). The loading of multiple OSP molecules on $Q\beta$ surface may sterically impede the access of reducing agents to the capsid, hindering reduction of the disulfides resulting in multiple monomers remaining bound together. As there was little $Q\beta$ monomer observed on the gel, the low signal from the OSP conjugate observed in SELDI-TOF (Figure S3) was most likely due to the difficulty in ionizing the $Q\beta$ -OSP conjugate by MS. Thus, we needed to employ additional techniques beyond SELDI-TOF and SDS-PAGE to provide more quantitative information on the degree of OSP functionalization on $Q\beta$.

We next tested Mass Photometry (MP)²⁸ to quantify OSP functionalization of $Q\beta$. While the aforementioned SELDI-TOF and SDS-PAGE methods for VLP analysis rely on disassembly of the particles and assessing the conjugation at the individual monomeric coat protein

levels, MP measures light scattering from the intact particle, which is proportional to the mass of the scattering particles. By analyzing hundreds to thousands of particles, the mass distribution of the sample is generated. To the best of our knowledge, MP has only been applied twice to study VLPs to date.^{29,30} When we measured the Q β sample, there were two populations and the MW was reducing over time (Figure S4). This could be resulting from the instability and degradation of RNA encapsulated inside the Q β . To test this possibility, we prepared a Q β sample without RNA by cleaving the RNA chemically with lead acetate³¹ and measured the empty particles with MP. Although we still observed two populations in the sample, the MW remained stable in two different measurements of the same sample over a 3-month interval (Figure S5). This supported that the result is reproducible and that MW reduction over time observed with full Q β was likely due to RNA degradation. The population of Q β particles with smaller molecular weight observed in the MP spectrum may be the result of partially disassembled VLPs.

In order to calibrate the mass shifts in MP, we conjugated the Q β capsid without RNA with lactoside **3** as a control sample. On the basis of the MW of this conjugate obtained from MP, the lactose loading was about 7 per monomer (Figure S6), which was close to the average loading of 6 obtained from the same sample on QTOF-ESI (Figure S8) and SELDI-TOF (Figure S9). With this result, we confirmed that the mass shift in Q β sample after conjugation is due to the loading of the carbohydrate and hence the difference in mass can be used to calculate the loading.

We analyzed the Q β -OSP conjugate with MP next. Upon OSP conjugation, the mean mass shifted about 400 kDa based on the MW of intact particles (Figure 1). With the average MW of OSP at ~6000 Da, it was calculated that on average there were 68 OSP molecules per Q β capsid. The OSP conjugation and MP protocols were reproducible, giving 66–68 OSP units per Q β on two independent batches of Q β -OSP conjugates.

Immunogenicity of the Q β -OSP Conjugate.

With the Q β -OSP conjugate in hand, to analyze whether the Q β -OSP was displaying OSP in an immunologically relevant manner, we assessed the ability of convalescent plasma from humans recovering from cholera to recognize Q β -OSP. Plasma was collected and analyzed as previously described.²² As shown in Figure 2, Q β -OSP was recognized by convalescent phase plasma of humans recovering from cholera (day 7 sample), but not by acute phase plasma (day 2 sample). In comparison to Q β -OSP, there was little binding of convalescent sera to Q β by itself. Furthermore, there was no increased immune-recognition of Q β -OSP by plasma from *Salmonella* Typhi infected patients (typhoid fever) suggesting the binding of Q β -OSP was a result from cholera infection.

Next, we evaluated the ability of Q β -OSP to generate anti-OSP antibodies in animals. A group of five female Swiss-Webster (3–5 week old) mice was injected intramuscularly on days 0, 21, 42, and 265 with the Q β -OSP construct (10 μ g OSP per mouse) in the absence of any exogenous adjuvants (Figure 3a). Blood was collected from the immunized mice on days 0, 7, 28, 49, 56 and during the study as shown in Figure 3a. A control group of Swiss-Webster mice received Q β only following the same protocol. To analyze the levels of anti-OSP antibodies in the sera by enzyme-linked immunosorbent assay (ELISA), a bovine

serum albumin (BSA) conjugate of OSP was prepared to avoid the interference of anti-Q β antibodies. The anti-OSP IgG titer, the highest dilution above background that gives optical density (OD) = 0.1, was determined in pooled sera at different time points by ELISA. While there were weak IgG responses 1 week after the first immunization, after the second immunization, significantly higher levels of IgG were observed on days 28, 49, and 56 (Figures 3b and 3c). The average IgG titers reached the maximum value of 226 504 on day 56 (Figure 3d). The IgG titer from the Q β -OSP group remained at high levels over time with IgG titers still detectable at day 265. In contrast, there was no detectable anti-OSP IgG responses in the control group at any time point suggesting Q β -OSP potentially induced antibody responses against OSP. No anti-OSP IgM antibodies were detected.

In order to assess whether memory responses were generated, on day 265 post initial immunization, mice received an additional vaccination. One week (day 272) after the booster, the average anti-OSP IgG antibody levels of the mice increased over 35 times compared to those on day 265 and reached the similar level of IgG titer as day 56. These results suggest that Q β -OSP vaccination induced memory B cell responses and the anti-OSP humoral immunity could be boosted (Figure 3b).

In order to determine whether immunization with Q β -OSP would elicit antibodies recognizing the native LPS containing OSP from *V. cholerae*, ELISA analysis was also performed using *V. cholerae* O1 Inaba LPS PIC018 as the coating antigen. Sera from mice immunized with Q β -OSP had significantly higher levels of anti-LPS IgG antibodies as compared to those from mice receiving Q β alone (Figure 4). Furthermore, serum binding to Inaba LPS was significantly higher compared to binding to LPS from *E. coli*, suggesting the antibodies induced by Q β -OSP were selective toward Inaba (Figure 4).

With the ability to selectively bind native *V. cholerae* O1 Inaba LPS and OSP by the Q β -OSP induced antibodies established, we next measured the vibriocidal activities of the postimmune sera.²² In the presence of an exogenous source of complement, *V. cholerae* cells are incubated with serial serum dilutions. Anti-*V. cholerae* antibodies present in the serum sample(s) in combination with complement can lyse the live bacteria. While none of the mice from the Q β immunized group showed any vibriocidal activities, sera from 2 of the 5 Q β -OSP immunized mice were able to kill the bacteria at dilutions higher than those in mice immunized with Q β alone (Figure 5).

DISCUSSION

OSP of *V. cholerae* has been used as an antigen in conjugation with BSA and recombinant heavy chain of tetanus toxin, and synthetic hexasaccharide and synthetic hexasaccharide cluster conjugates have also been evaluated as vaccine antigens against *V. cholerae*.^{22,32-34} A virus-like particle (VLP) based approach has a number of attractive features for vaccine applications since the highly ordered organization of the protein(s) in the VLPs is well recognized via pathogen-associated molecular patterns (PAMPs).³⁵ VLPs can present antigens in an organized and polyvalent manner to cross-link B-cell receptors to induce intense cellular signaling for strong immune activation. Bacteriophage Q β VLP is a

promising platform for organized display to induce antibody responses against a target antigen.

For polysaccharide based conjugate vaccines, there are several coupling methods such as periodate activation for reductive amination, cyanation, and carbodiimide-mediated coupling.³⁶ These approaches can suffer from incompatibility with some proteins and substrates and low selectivity. Squaric acid esters are favorable linker molecules in glyco-conjugate formation between amino-saccharides and proteins due to the amine-selectivity, high reactivity at room temperature, possibility of stoichiometric modification, and recovery of high-value unreacted (oligo)saccharide.³⁷ In the current study, we have prepared the squaric acid monoester derivative of *V. cholerae* O1 OSP core antigen utilizing the core amine group of the OSP and conjugated the monoester with VLP (carrier protein). While bacteriophage Q β has been conjugated with glycan antigens,^{18,19,26,27} this is the first example of using a squarate linker for glycan conjugation with Q β VLP. The squarate chemistry was highly efficient, leading to close to full derivatization of all free amines of Q β coat proteins with a small glycan such as lactose 4.

With squarate chemistry, the polysaccharide of *V. cholerae* O1 OSP was conjugated to the Q β carrier protein via single point attachment due to the presence of single amino group per OSP molecule. The final construct may mimic native bacteria by presenting multiple OSP polysaccharide on the surface. OSP display on Q β was in an immunologically relevant manner, which was recognized by convalescent sera of cholera infected humans but not of typhoid fever patients. The Q β -OSP vaccine was immunogenic in mice, inducing persistent IgG responses against OSP. Such long-term IgG production may assist with the long-term protective goal of anticholera vaccination.³⁸

The vaccine was administrated in the absence of an exogenous adjuvant. This in part may be due to inherent adjuvant properties of Q β VLP, which can encapsulate *E. coli* RNA molecules, thus stimulating internal cellular signals via TLR7/8 in the antigen presenting cells.³⁹ The induced IgG antibodies recognized native *V. cholerae* LPS and had vibriocidal activities. Induction of vibriocidal antibodies correlate with protection against cholera,⁴⁰ and these responses largely target *V. cholerae* OSP.^{40,41} The mechanism of protection against *V. cholerae* in the intestinal lumen is currently unclear, but may involve inhibition of *V. cholerae* motility through the binding of OSP-specific antibodies.^{42,43} The induction of low levels *E. coli* LPS-specific IgG antibody responses (Figure 4) may be due to the presence of trace amounts of residual *E. coli* LPS in Q β VLP preps. Studies are ongoing to express Q β VLPs in LPS deficient *E. coli* strains.⁴⁴

Our study has several limitations. We focused on developing suitable coupling chemistry and analytical tools to synthesize and characterize the Q β -OSP conjugate. We evaluated the immunogenicity of the conjugate, and future assessments could evaluate protective efficacy in a wild type *V. cholerae* challenge assay. We did not investigate the effect of adjuvant administration on anti-OSP IgG production, or the mechanism by which induced antibodies can provide protection.

CONCLUSIONS

In this work, we have conjugated the *V. cholerae* O1 OSP polysaccharide to Q β carrier for the first time. The MP technique yielded critical information on the level of OSP loading on Q β . The Q β -OSP conjugate was recognized by sera from humans with cholera and was able to induce long lasting antibody production in the absence of adjuvant in a mice immunization study. The resulting antibodies exhibited vibriocidal activities. VLP-based display of bacterial OSP may warrant additional evaluation as a next generation anticholera vaccine.

METHODS

All chemicals were reagent grade and were used as received from the manufacturer, unless otherwise noted. Protein concentration was measured using the Coomassie Plus Protein Reagent (Bradford Assay, Pierce) with BSA as the standard. OSP and BSA-OSP was produced as previously described.²² Q β capsid was expressed recombinantly following the reported procedure.^{25,45}

Q β Conjugation to Lactoside 3 and Purification.

A stock solution of 4.8 mg/mL Q β in 0.1 M (pH 7.0) KPB buffer (208 μ L) was placed in a Millipore Amicon Ultra-0.5 (10k Da cutoff) ultrafiltration device and the content was ultrafiltered against 0.5 M pH 9.0 borate buffer for three times (at 10 °C, 7500 rcf, 10 min/run) to exchange the buffer to 0.5 M pH 9.0 borate buffer. The filtrates were discarded and the final retentate was transferred into a 0.5 mL V-shaped reaction vessel and the same buffer was added to adjust the overall volume to 200 μ . Lactose squarate **3** (0.6 mg, 0.986 mmol) was carefully added into the reaction mixture and the content of the vessel was stirred at r.t. for 20 h. SELDI-TOF-MS analysis showed that an average loading (lactose/Q β monomer ratio) of ~4.5 was achieved (Figure S1b). Another 0.6 mg (0.986 mmol) of compound **3** was added and the reaction mixture was further stirred for 72 h. SELDI-TOF-MS showed that the average loading reached ~8.0 per monomer. The reaction was worked up by ultrafiltering the reaction mixture in a Millipore Amicon Ultra-0.5 (30k Da cutoff) tube against pH 7.2 PBS (1 \times) buffer for 6 times to remove the unconjugated lactose derivatives. The final retentate was transferred into a conical tube for storage.

SELDI-TOF-MS Analysis of Samples.

The above Q β -lactose reaction mixture (1 mL) was diluted with 0.5 M pH 7.0 phosphate buffer (10 mL) and then mixed with dithiothreitol (DTT) solution (0.1 M in water, 11 mL). The mixture was incubated at 37 °C for 30 min. The above solution (1 mL) was withdrawn from the mixture for SELDI analysis²⁰ using sinapinic acid (SPA) as the matrix.

Preparation of SPA Matrix.

To SPA (5 mg) in a 1 mL Eppendorf tube was added acetonitrile (100 μ L) followed by 1% TFA aqueous solution (100 μ L). The mixture was vigorously vortexed for 20 s and then centrifuged for 10 min at 1000 rcf and the supernatant was used as the matrix solution.

MS Sequencing of Q β Lactoside 3 Conjugates.

Q β conjugated with lactoside **3** (10 μ g) was run on SDS-PAGE gel under reducing conditions. The gel was fixed and stained with Coomassie and the band corresponding to the Q β monomers was excised. In gel trypsin digestion was performed and the resulting peptides were analyzed by mass spectrometry. The resulting data was analyzed using Scaffold v5.1.0.

Q β Conjugation to OSP and Purification.

V. cholerae O1 Inaba OSP **1** (5.4 mg, 0.0009 mmol) was converted to its squarate derivative by reacting with 3,4-dimethoxy-3-cyclobutene-1,2-dione (**2**, 2.56 mg, 0.018 mmol) in 0.5 M pH 7.0 phosphate buffer as described previously.²¹ A white fluffy solid (5.4 mg, 0.00088 mmol, 98%) was obtained after workup. A stock solution of 6.7 mg/mL Q β in 0.1 M (pH 7.0) KPB buffer (240 μ L) was placed in a Millipore Amicon Ultra-0.5 (10 kDa cutoff) ultrafiltration device and the content was ultrafiltered against 0.5 M pH 9.0 borate buffer for three times (at 10 °C, for speed/rpm, time and volume of the concentrate, manufacturer's suggestions were followed) to exchange the buffer to 0.5 M pH 9.0 borate buffer. The filtrates were discarded and the final retentate was transferred into a 1 mL V-shaped reaction vessel and the same buffer was added to adjust the overall volume to 225 μ L. The squarate derivative obtained above (5.4 mg, 0.00088 mmol) was added into the vessel and the clear solution formed was stirred at r.t. After 120 h, the reaction was worked up by ultrafiltering the reaction mixture in a Millipore Amicon Ultra-4 (30 kDa cutoff) tube against pH 7.2 PBS (1 \times) buffer for 6 times to remove the unconjugated antigen. The final retentate was transferred into a conical tube for storage.

MP Procedure.

The 24 \times 50 mm microscope coverslips (Fisher Scientific, Waltham, MA) and precut 2 \times 2 silicon gasket wells (GBL103250, Sigma, MO) were cleaned and assembled as described in the literature.⁴⁶ Measurements were performed on OneMP instrument (Refeyn, Oxford, UK) at room temperature. PBS (1 \times , pH 7.2) buffer was filtered through 0.22 μ M filters before use. Ten microliters of the buffer were loaded in gasket well to focus the objective on the coverslip surface. Q β conjugates stock were diluted 100 times in PBS (1 \times , pH 7.2) buffer and added to buffer in the well. The MP video was immediately recorded after the sample loading using the AcquireMP software (Refeyn, Oxford, UK). A 1 min video was recorded for each sample, and each sample was repeated twice. Data were processed using the DiscoverMP software (Refeyn, Oxford, UK) with the threshold filter values of 5. The mass distribution was plotted as histograms with bin width of 50 kDa and fit with Gaussian peaks to obtain the average mass of different species. The contrast-to-mass calibration was performed using an unstained protein ladder (LC0725, Thermo Fisher, Waltham, MA) and empty AAV5 sample (Virovek, Hayward, CA).

Immunization.

All animal experiments were approved by and performed in accordance with the guidelines of the Institutional Animal Care and Use Committee (IACUC) of Michigan State University. The animal usage protocol number is PROTO201900423. Female Swiss-Webster (3–5 week

old) mice were used for studies ($n = 5$ for each group). Mice were injected intramuscularly on days 0, 21, and 42 with 0.06 mL Q β -OSP construct (10 μg OSP, 65 μg Q β per mouse) without an exogenous adjuvant. The control group was injected with the same amount of Q β carrier (65 μg Q β per mouse) as the experimental group. The final boost was given at day 265. Serum samples were collected on days 0 (before immunization), 7, 28, 49, 56, 86, 118, 170, 265, and 272.

Evaluation of Antibody Titers by ELISA.

The Nunc MaxiSorp flat-bottom 96-well microtiter plates were coated with 1 $\mu\text{g mL}^{-1}$ of the BSA-OSP conjugate (100 $\mu\text{L/well}$)¹ in NaHCO₃/Na₂CO₃ buffer (0.05 M, pH 9.6) containing 0.02% NaN₃ by incubation at 4 °C overnight. For ELISA study against LPS, plates were coated with 10 $\mu\text{g mL}^{-1}$ of LPS, Inaba or *E. coli*, in PBS buffer overnight at room temperature. The coated plates were washed with PBS/0.5% Tween-20 (PBST) (4 \times 200 μL) and blocked with 1% BSA in PBS (200 $\mu\text{L/well}$) at rt for 1 h. The plates were washed again with PBST (4 \times 200 μL) and incubated with serial dilutions of mouse sera in 0.1% BSA/PBS (100 $\mu\text{L/well}$, 2 wells for each dilution). The plates were incubated for 2 h at 37 °C and then washed with PBST (4 \times 200 μL). A 1:2000 dilution of HRP-conjugated goat antimouse IgG or IgM (Jackson ImmunoResearch Laboratory, 115-035-003) in 0.1% BSA/PBS (100 μL) was added to the wells respectively to determine the titers of antibodies generated. The plates were incubated for 1 h at 37 °C and then washed with PBST (4 \times 200 μL). A solution of the enzymatic substrate 3,3',5,5'-tetramethylbenzidine (TMB, 200 μL) was added to the plates (for one plate: 5 mg of TMB was dissolved in 2 mL of DMSO plus 18 mL of citric acid buffer containing 20 μL of H₂O₂). Color was allowed to develop for 15 min and then quenched by adding 50 μL of 0.5 M H₂SO₄. The readout was measured at 450 nm using a microplate reader. The titer was determined by regression analysis with log₁₀ dilution plotted with optical density and reported as the highest fold of dilution giving the optical absorbance value of 0.1 over those of the preimmune control sera. 1:5000 serum dilution was used for LPS binding study as shown in Figure 4.

Evaluation of Q β -OSP Conjugates Using Human Serum.

To assess immunoreactivity of the OSP display on the Q β conjugates, antigen-specific ELISA using sera collected from humans with cholera in Bangladesh was performed. The responses to human sera with cholera were compared to the response detected in humans with typhoid fever in Bangladesh. All samples were collected following informed consent, and human subjects work was approved by the Institutional Review Boards of the Massachusetts General Hospital and the International Centre for Diarrheal Disease Research, Bangladesh (icddr,b). Plates coated with 100 ng of Q β or Q β -OSP (based on protein mass) per well. After blocking and washing of plates, acute and convalescent phase sera from humans with cholera or typhoid (diluted 1:250 in 0.1% BSA in phosphate buffered saline-Tween) were added and incubated for 90 min at 37 °C. HRP-conjugate antihuman IgG antibody at 1:5000 dilution in 0.1% BSA in phosphate buffered saline-Tween was used to detect antigen-specific antibodies. After 90 min incubation at 37 °C, 0.55 mg/mL solution of 2,2'-azinobis (3-ethylbenzothiazoline-6-sulfonic acid) (ABTS; Sigma) with 0.03% H₂O₂ (Sigma) were added to the plates and optical density was read at 405 nm for 5 min at 30

s intervals. The maximum slope for an optical density change of 0.2 U was reported as milli-optical density units per minute (mOD/min).

Serum Vibriocidal Responses.

The vibriocidal antibody titers against *V. cholerae* O1 El Tor Inaba strain PIC018 were assessed in a microassay as previously reported.⁴⁷ The endogenous complement activity of mouse serum was inactivated by heating it for 30 min at 56 °C. 50 μ L aliquots of serial dilution of heat-inactivated sera in 0.15 M saline were added to wells of sterile 96-well tissue culture plates containing 25 μ L/well of *V. cholerae* O1 El Tor Inaba strain PIC018 (OD 0.3) in 0.15 M saline and 22% guinea pig complement. After 1 h incubation at 37 °C, 150 μ L of brain heart infusion broth was added to each well, and plates were incubated for an additional 2 h at 37 °C. The optical density of plates was then measured at 595 nm. A responder was defined as having a 4-fold increase of vibriocidal titer at day 56 compared with baseline day 0 titer.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

ACKNOWLEDGMENTS

This research was supported in part by grants from the National Institutes of Health: (R01AI146210 (XH), R01AI106878 (ETR)), the Fogarty International Center, Training Grant in Vaccine Development and Public Health (TW005572 (MK, TRB)), and Emerging Global Fellowship Award TW010362 (TRB), the Intramural Research Program of the NIH and NIDDK (PX), and Michigan State University. We would like to thank the Michigan State University RTSF Proteomics Core for their help with the sequencing of Q β -lactose conjugates.

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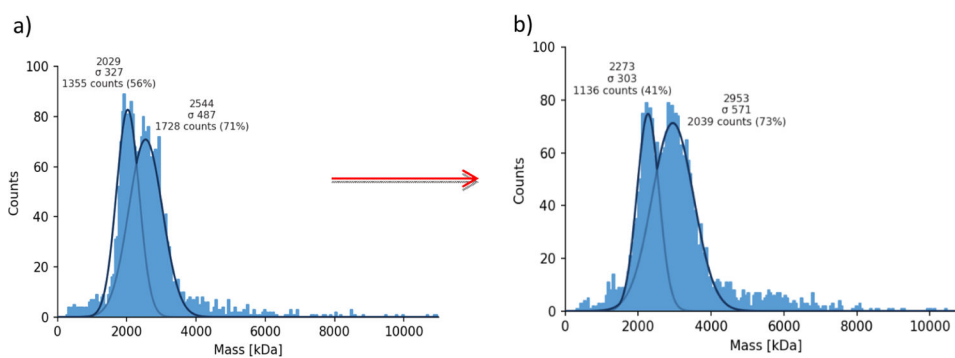


Figure 1. MP results of (a) $Q\beta$ and (b) $Q\beta$ -OSP conjugate. The right peak shifted from 2544 kDa to 2953 kDa, which suggests the conjugation of an average of 68 OSP per full $Q\beta$ capsid calculated based on the mass of the intact particle.

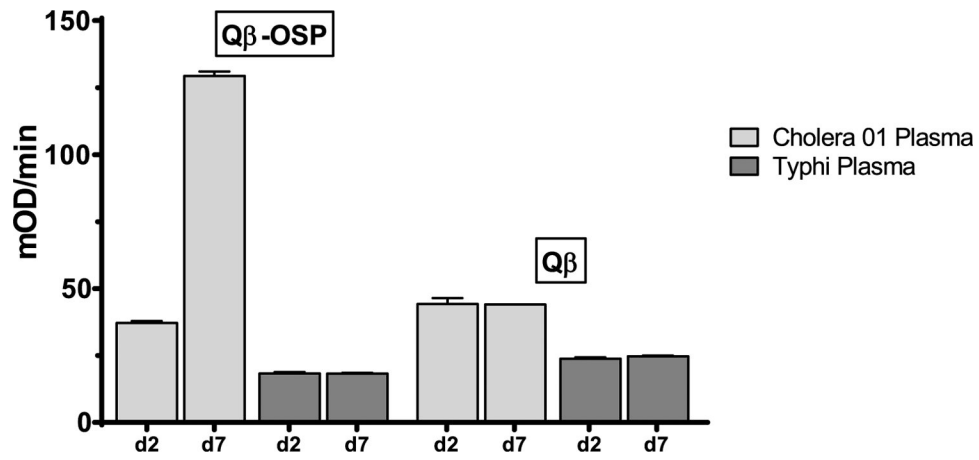


Figure 2. Immunoreactivities of human plasma toward $Q\beta$ and $Q\beta$ -OSP were measured by acute phase plasma (day 2 sample) versus convalescent phase plasma (day 7 sample) of patients with cholera versus typhoid fever in Dhaka, Bangladesh.

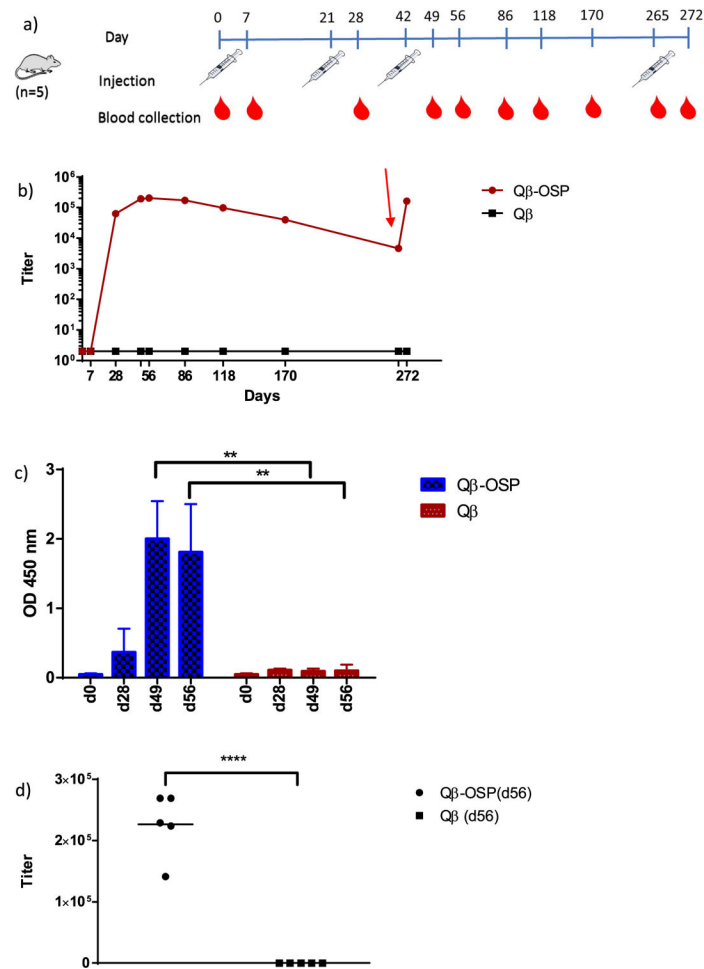


Figure 3. Evaluation of Q β -OSP immunogenicity. (a) Immunization and blood collection schedule. Each group received 3 immunizations 3 weeks apart with blood collected at day 0 and on days 7, 28, 49, 56, 86, 118, 170, 265, and 272 respectively. (b) OSP-specific IgG titer of pooled sera from Q β and Q β -OSP groups up to day 272 postimmunization. The red arrow indicates a booster injection at day 265. (c) ELISA analysis showed significant IgG binding to BSA-OSP by postimmune sera at d49 and d56 ($p = 0.0014$ and 0.0065 , respectively), compared to the control sera from mice immunized with Q β only. Each bar represents data for 5 mice at 20 000 fold of serum dilution. (d) Individual mouse serum OSP-specific IgG titer of Q β and Q β -OSP groups at day 56. The statistical significance was determined through a two-tailed t test using GraphPad Prism. ** $p < 0.0001$.

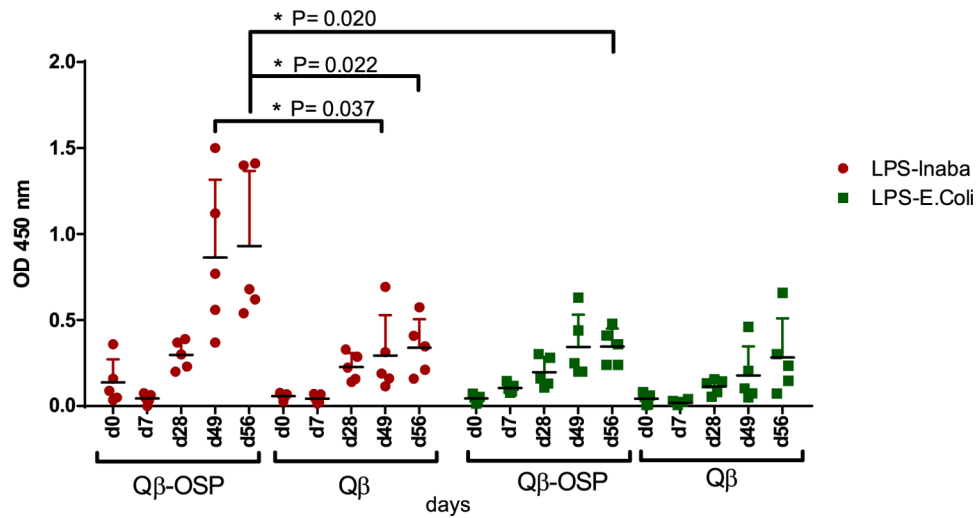


Figure 4. Binding of mouse serum immunized with Q β -OSP and Q β to LPS from Inaba vs *E. coli*. Serum binding against Inaba LPS was observed in the Q β -OSP immunized group while sera from the Q β group had lower binding. The binding to *E. coli* LPS was lower by sera from both Q β -OSP and Q β immunized mice. The statistical significance was determined through an unpaired two-tailed *t* test using GraphPad Prism. **p* < 0.05.

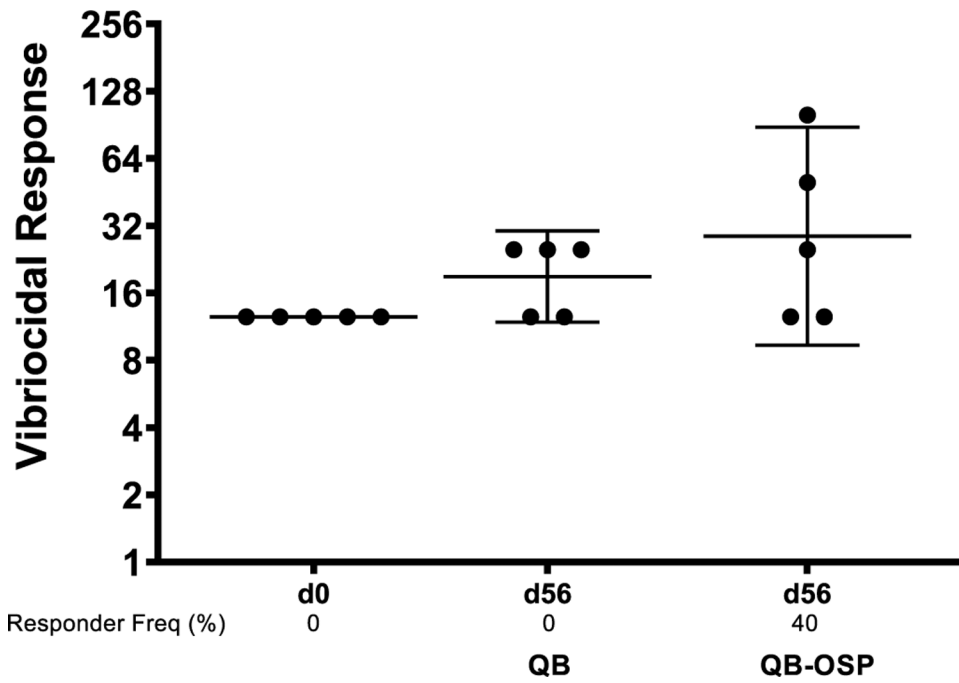
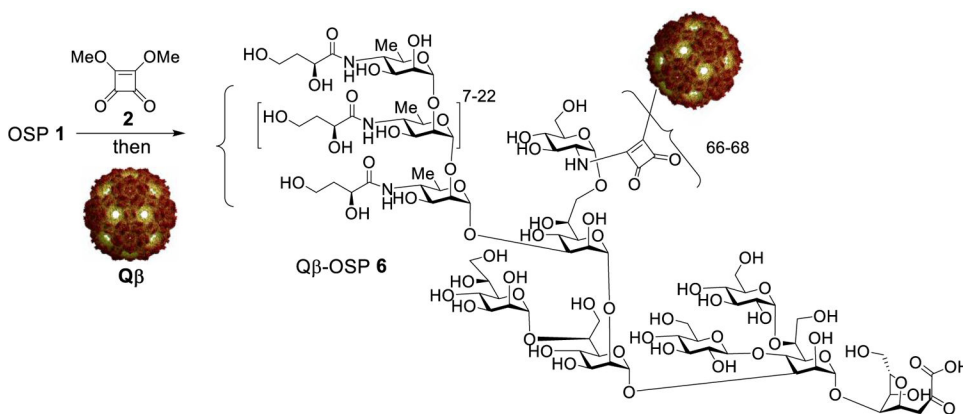


Figure 5. Vibriocidal responses in vaccine cohorts. We defined responders as having an increase in vibriocidal titer by 4-fold at day 56 than day 0.

**Scheme 1.**Conjugation of OSP 1 to Qβ VLP^a

^aOSP 1 was activated with dimethyl squarate 2, and subsequently added to a Qβ solution in 0.5 M borate buffer, pH = 9.0. After 120 h, the reaction was worked up by ultrafiltration against pH 7.2 (1×) PBS buffer.