

Research Article

Structural and immunological characterization of *E. coli* derived recombinant CRM₁₉₇ protein used as carrier in conjugate vaccines

Ravi P.N. Mishra¹, Ravi S.P. Yadav¹, Christopher Jones², Salvatore Nocadello³, George Minasov³, Ludmilla A. Shuvalova³, Wayne F. Anderson³ and Akshay Goel¹

¹Biological E. Limited, Genome Valley, MN Park, Shameerpet, Hyderabad 500078, India; ²National Institute for Biological Standards and Control (NIBSC), Blanche Lane, South Mimms, Potters Bar, Hertfordshire EN6 3QG, U.K.; ³Center for Structural Genomics of Infectious Diseases, Northwestern University, Feinberg School of Medicine, Morton Bld. # 7-614 303 E. Chicago Ave., Chicago, IL 60611, U.S.A.

Correspondence: Akshay Goel (akshay.goel@biologicale.co.in)



It is established that the immunogenicity of polysaccharides is enhanced by coupling them to carrier proteins. Cross reacting material (CRM₁₉₇), a nontoxic variant of diphtheria toxin (DT) is widely used carrier protein for polysaccharide conjugate vaccines. Conventionally, CRM₁₉₇ is isolated by fermentation of *Corynebacterium diphtheriae* C7 (β ₁₉₇) cultures, which often suffers from low yield. Recently, several recombinant approaches have been reported with robust processes and higher yields, which will improve the affordability of CRM₁₉₇-based vaccines. Vaccine manufacturers require detailed analytical information to ensure that the CRM₁₉₇ meets quality standards and regulatory requirements. In the present manuscript we have described detailed structural characteristics of *Escherichia coli* based recombinant CRM₁₉₇ (rCRM₁₉₇) carrier protein. The crystal structure of the *E. coli* based rCRM₁₉₇ was found to be identical with the reported crystal structure of the C7 CRM₁₉₇ produced in *C. diphtheriae* C7 strain (Protein Data Bank (PDB) ID: 4EA0). The crystal structure of rCRM₁₉₇ was determined at 2.3 Å resolution and structure was submitted to the PDB with accession number ID 5182. This is the first report of a crystal structure of *E. coli* derived recombinant CRM₁₉₇ carrier protein. Furthermore, the rCRM₁₉₇ was conjugated to Vi polysaccharide to generate Typhoid conjugate vaccine (Vi-rCRM₁₉₇) and its immunogenicity was evaluated in Balb/C Mice. The Vi-rCRM₁₉₇ conjugate vaccine was found to generate strong primary α -Vi antibody response and also showed a booster response after subsequent vaccination in mice. Overall data suggest that *E. coli* based recombinant CRM₁₉₇ exhibits structural and immunological similarity with the C7 CRM₁₉₇ and can be used as a carrier protein in conjugate vaccine development.

Introduction

Over the past three decades, many routine childhood and adult vaccines have been developed using conjugation technology. Conjugate vaccines are developed by covalent attachment of an antigenic polysaccharide to a nontoxic carrier protein. Conjugation to a nontoxic carrier protein enhances the immunogenicity of polysaccharide antigens, enabling host defence against diseases caused by encapsulated pathogens. Conjugation transforms the T cell-independent response of polysaccharide vaccines to a T cell-dependent one. Conjugate vaccines have been demonstrated to be immunogenic and capable of inducing immunological memory and high avidity antibodies. Unlike unconjugated polysaccharides, conjugate vaccines also elicit protective responses in the immature immune system of young infants and the senescent immune system of the elderly [1-3].

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Cross reacting material (CRM₁₉₇) is widely used as a carrier protein in conjugate vaccines [3]. CRM₁₉₇ protein is a nontoxic variant of diphtheria toxin (DT, molecular weight: ~58 kDa) with a single mutation (Glycine to Glutamate substitution at position 52) that eliminates its toxicity [4,5]. The protein nonetheless retains the same immunostimulant properties as DT and it is used to generate safe and effective polysaccharides conjugate vaccines for all age groups [6,7].

CRM₁₉₇-based polysaccharides conjugate vaccines have already been developed to confer protection against important bacterial pathogens such as *Streptococcus pneumoniae* (Prenvar), *Haemophilus influenzae* type b (HibTITER, Vaxem-Hib) and *Neisseria meningitidis* serogroup A, C, Y and W-135 (e.g. Menveo and Menjugate). These vaccines have been successfully used to immunized hundreds of millions of people worldwide. Several conjugate polysaccharides vaccines using CRM₁₉₇ as carrier protein are under preclinical and clinical evaluation. These include vaccines against *Staphylococcus aureus*, *Streptococcus agalactiae* (Group B *Streptococcus*), *Salmonella* Typhi, *Salmonella* Paratyphi, *Mycobacterium tuberculosis* and several pneumococcal conjugate vaccines [3,8-14].

CRM₁₉₇ is synthesized as a single chain holoprotein, which comprises two domains fragment A (catalytic domain) and fragment B (transmembrane domain), bound together by a disulfide bridge. The B domain contains a subdomain for binding to the HB-EGF cell receptor and another subdomain for translocation into eukaryotic cells. *In vitro*, mild trypsin treatment and reduction divides the toxin into two functional moieties (the 'nicked form'). The receptor-binding subdomain of fragment B has an all- β -sheet structure, in contrast with the transmembrane subdomain containing nine α -helices. Two disulfide bridges are present in the intact holoprotein: one bridge joins Cys¹⁸⁶ to Cys²⁰¹, linking fragment A to fragment B while a second bridge joins Cys⁴⁶¹ to Cys⁴⁷¹ within fragment B [15].

Since CRM₁₉₇ is genetically nontoxic, chemical toxoiding is not required. This helps to preserve epitopes which would otherwise be lost. It has been suggested that preservation of T-helper epitopes explains the improved carrier effect of CRM₁₉₇ compared with diphtheria toxoid [13]. The lack of toxicity in CRM₁₉₇ has been examined *in vivo* and *in vitro* using lethality tests in guinea pigs, cytotoxic activity assays on HeLa cells and Vero cells and an ADP-ribosyltransferase enzymatic assay [16].

CRM₁₉₇ is classically purified from culture supernatant of the *Corynebacterium diphtheriae* C7 (β_{197}) tox negative (Tox-) strain [4]. The production of CRM₁₉₇ from its C7 strain often suffers from low yield and also requires sophisticated laboratory conditions to cultivate the *C. diphtheriae* C7 strain. More recently, CRM₁₉₇ has been produced in heterologous recombinant systems such as *Escherichia coli* and *Pseudomonas fluorescens* [17-19] with higher yield. The characterization of CRM₁₉₇ protein produced from *C. diphtheriae* C7 strain has been published [15,16]. However, limited data are available for CRM₁₉₇ produced from *E. coli* based recombinant source. Also, there is lack of information whether *E. coli* based recombinant CRM₁₉₇ is structurally and immunologically equivalent to its incipient counterpart (C7 CRM₁₉₇). In the present study, for the first time we have performed structural and immunological characterization of *E. coli* derived recombinant CRM₁₉₇ protein. The data are particularly important because the increasing global demand of polysaccharide conjugate vaccines against encapsulated bacterial pathogens highlight the pressing need for high yield processes delivering well-characterized carrier protein that meets regulatory and safety requirements. The objective of present study was to: (i) express the rCRM₁₉₇ protein from *E. coli* and its physicochemical characterization, (ii) determination of X-ray crystal structure of recombinant CRM₁₉₇ and demonstration of its structural equivalence with C7 CRM₁₉₇, (iii) development of polysaccharide conjugate vaccine using rCRM₁₉₇ and demonstration of immunological functionality of rCRM₁₉₇ as carrier protein.

Materials and methods

Cloning, expression and purification of CRM₁₉₇

The codon optimized sequence corresponding to CRM₁₉₇ gene was synthesized and cloned into pTWIN 1 vector (New England Biolabs, U.K.) using BamH1 and Sap1 restriction sites [18]. The pTWIN1_CRM₁₉₇ plasmid was transformed into chemically competent BL21-DE3 *E. coli* expression host. Transformants were screened on LB agar plates supplemented with ampicillin (100 μ g/ml). Five to ten randomly selected colonies were purified by repeated subculturing on LB + ampicillin (100 μ g/ml) plate. Finally, a few colonies were selected and a research cell bank was generated. For the expression analysis of rCRM₁₉₇ protein, clones were grown in 100-ml LB + Ampicillin (100 μ g/ml) till mid log phase (OD₆₀₀ ~1). At this stage, culture was induced with 1 mM IPTG and was further grown to reach late log phase (OD₆₀₀ ~4-5). One millilitre culture from each transformant was centrifuged and pellet was resuspended into 100- μ l SDS loading buffer and heated at 80°C for 10 min. Twenty microlitres of lysate was loaded on 8-12% Tri-Glycine gel and SDS/PAGE was run using 80 V constant current for ~2 h. Gel was stained with Coomassie Blue (make: Amresco, Cat No: 0472-25 G). The expression of rCRM₁₉₇ was confirmed by comparing a band appearing at

~58 kDa on the gel. Further, Western blot was performed using anti-CRM₁₉₇ antibody (Make: Santa Cruz Biotechnology, Cat No: SC2054). One of the clones expressing rCRM₁₉₇ protein has been used for fermentation and large-scale production of protein.

Fermentation inoculum was generated by inoculating one vial of research cell bank into seed flask containing 200-ml LB + ampicillin (100 µg/ml). Seed flask (containing growth media and inoculum) was grown to achieve OD₆₀₀ ~4 and inoculated into 20-l fermentation medium. Fermentation was performed in chemically defined medium. After fermentation was complete, cell mass was separated by centrifugation. The cell pellet was lysed by Micro fluidizer high pressure homogenization. Homogenized cell mass was centrifuged and pellet containing inclusion body protein was retained and used for rCRM₁₉₇ purification. The inclusion body was solubilized in buffer containing 6 M urea at 30°C and rCRM₁₉₇ was purified by refolding followed by ion exchange chromatography [18]. The solubilized inclusion body protein was refolded and purified by ion exchange chromatography method described by Stefan et al. [19].

C7 CRM₁₉₇ used as a reference in the studies was procured as a lyophilized powder from Sigma–Aldrich (Cat# D2189).

Endonuclease assay for rCRM₁₉₇

The assay was performed as described by Stefan et al. (2011) [19] with minor modifications. In brief, 2 µg rCRM₁₉₇ was mixed with 1 µg pUC57 plasmid DNA using endonuclease reaction buffer (10 mM Tris/HCl pH 7.6, 2.5 mM CaCl₂, 2.5 mM MgCl₂). Content was mixed by brief centrifugation (3000 g, 30 s) and incubated for different time points (0, 0.5, 1, 3, 5 and 18 h) at 25°C. The 0-h time point sample was used as a control. At each time point, reaction was terminated by the addition of 5 mM EDTA (final concentration) to each tube and frozen at –20°C. Thawed samples were mixed with DNA loading buffer and run on 1% agarose gel electrophoresis in 1× TAE buffer using 80 V constant voltage. DNA staining dye SYBR Safe gel stain (Invitrogen, Cat# S33102) was added to the agarose gel at the time of pouring. After the electrophoresis run was completed, the gel was analysed under UV transillumination (Bio-Rad ChemiDoc Gel imaging System).

SDS/PAGE and Western immunoblot analysis

Reducing and nonreducing SDS/PAGE and Western blot analyses were performed with purified rCRM₁₉₇ as per the standard protocol [20]. Four micrograms of purified rCRM₁₉₇ along with the reference CRM₁₉₇ was loaded on to 4–12% Tris-Glycine gels under reducing and nonreducing conditions. Gels were run at 80 V constant voltage until blue dye reached the bottom of the gel. SDS/PAGE molecular weight markers (New England Biolabs, Cat# P7712S) were used for molecular weight calibration. After Coomassie Brilliant Blue staining, the image was captured using transilluminator (Bio-Rad ChemiDoc system). For Western immunoblotting, 2 µg protein was separated by SDS/PAGE and transferred to nitrocellulose membrane (0.45 µm, Bio-Rad, Cat# 1620115). The blot was probed with α-CRM₁₉₇ rabbit polyclonal primary antibody (Abcam, Cat# ab151222) by incubating the membrane for 1 h. After the primary incubation, the membrane was further incubated in HRP-conjugated secondary antibody (rabbit) (make: Santa Cruz Biotechnology, Cat No: SC2054). Bands were visualized using 3,3'-diaminobenzidine (DAB) substrate and the image captured using Bio-Rad ChemiDoc gel imaging system.

Intact MS

The intact mass of rCRM₁₉₇ was determined by LC-MS (ABSCIEX Triple TOF 5600+) using the published protocol [21,22]. Protein samples were buffer exchanged by centrifugal ultrafiltration (Amicon Ultra-0.5 ml, 10 K MWCO, 14000 g, 15 min, 4°C) and analysed by LC-qToF using the manufacturer's protocol.

Amino acid composition analysis

rCRM₁₉₇ was subjected to high sensitivity amino acid analysis (AAA). The analysis was performed on samples desalted by ultrafiltration (Vivaspin 6 spin columns with a 10-kDa polyethersulfone filter: Sartorius, Cat# VS0601). Samples were resuspended in 20% acetonitrile (ACN) containing 0.1% trifluoroacetic acid (TFA) and subjected to gas phase hydrolysis (6 N HCl at 110°C). α-amino butyric acid (AABA) was run as an internal standard prior to sample analysis. Cysteine analysis was performed after performic acid oxidation followed by hydrolysis as above. Hydrolysates were analysed using a Waters Acquity UPLC system with AccQTag Ultra chemistry. Samples were analysed in duplicate and the mean reported [23,24].

Western immunoblotting with structural monoclonal antibodies against CRM₁₉₇/DT

rCRM₁₉₇ samples along with reference were run in SDS/PAGE. The gel was transferred to nitrocellulose membrane and Western immunoblotting was done using five different monoclonal antibodies (mAbs) against DT/CRM₁₉₇. These mAbs were 7F2 (Abcam, Cat. No: ab8310), 8A4 (Abcam, Cat. No: ab8308), 3B6 (Abcam, Cat No: ab8307), 1H2 (Abcam, Cat. No: ab8306) and HYB 123-09 (Statens Serum Institute, Cat. No: 66045). The SDS/PAGE and Western blot methods were as described above.

Determination of crystal structure of rCRM₁₉₇ by X-ray crystallography

Center for the Structural Genomics of Infectious Diseases (CSGID) standard protocols were used to determine the structure of CRM₁₉₇ [25-27].

Sitting drop crystallization plates were set up at room temperature and crystals of rCRM₁₉₇ were obtained by mixing 22.9 mg/ml of protein in 0.15 M sodium chloride, 0.01 M Tris/HCl buffer (pH 8.3) with the crystal screen solution of 0.2 M lithium sulfate, 0.1 M Bis-Tris buffer (pH 5.5), 25% (w/v) PEG 3350 (Qiagen, Classics II suite (G2), in a 1:1 ratio. Harvested crystals were transferred to the reservoir solution before being flash-frozen in liquid nitrogen. Diffraction data were collected at 100 K at the Life Sciences Collaborative Access Team at the Advance Photon Source, Argonne, Illinois (APS BEAMLINE 21-ID-F). Data were processed using HKL-3000 for indexing, integration and scaling. The crystal structure of recombinant CRM₁₉₇ (Protein Data Bank (PDB) ID: 5I82) was determined by molecular replacement using PDB entry 4AE0 as a search model. The structure was refined with Refmac 5 [28]. Models were displayed in Coot and manually corrected based on electron density maps [29]. All structure figures were prepared using PyMOL Molecular Graphics System, version 1.3 (Schrödinger, LLC). The structure was submitted to the PDB (PDB ID: 5I82).

Generation of typhoid conjugate vaccine (Vi-rCRM₁₉₇) using rCRM₁₉₇ and immunogenicity assessment

rCRM₁₉₇ was conjugated with Vi polysaccharide antigen to generate Vi-rCRM₁₉₇ conjugate vaccine (typhoid conjugate vaccine). The Vi polysaccharide was produced by fermentation of *Citrobacter freundii* as per method described by Rondini et al. [30]. The *C. freundii* strain was provided by Dr Laura B. Martin, GSK Vaccine Institute of Global Health, Siena, Italy. The Vi-rCRM₁₉₇ conjugate was prepared according to the method reported by Kossaczka et al. [31] and as detailed by Micoli et al. [8]. Briefly, rCRM₁₉₇ was derivatized with adipic acid dihydrazine (ADH). Then, Vi (purified from *C. freundii*) was activated with N-hydroxy succinamide and 1-ethyl-3(3-dimethylaminopropyl) carbodiimide hydrochloride and linked to the derivatized rCRM₁₉₇ protein generating Vi-rCRM₁₉₇ conjugate vaccine. The ADH derivatized rCRM₁₉₇ (rCRM₁₉₇-ADH) was mixed with activated Vi polysaccharides in 1:1 ratio and incubated at room temperature for 5–6 h. The resulting conjugate was purified by HIC (hydrophobic interaction chromatography) using phenyl sepharose (GE) resin.

The immunogenicity of Vi-rCRM₁₉₇ was performed as per procedures described by Rondini et al. [30]. Four to six weeks old female Balb/C mice were used in the experiment. A group of 30 mice were subcutaneously immunized with three doses of Vi-rCRM₁₉₇, unconjugated Vi polysaccharide and PBS (placebo control). A 100 µl dose of Vi-rCRM₁₉₇ conjugate vaccine containing 2.5 µg Vi, unconjugated Vi polysaccharide (2.5 µg) and PBS (control) were administered in mice on day 1 followed by booster dose at days 14 and 21. Fourteen days after first immunization, ten randomly selected mice from each group were terminally bled to collect the post-dose 1 sera. Remaining animals (20 each) were given the second dose of vaccine. Seven days after second immunization (day 21), ten mice from each group were terminally bled and post-dose 2 sera was collected. Finally, remaining ten mice were given the third dose of vaccine. The final sera (post-dose 3) was collected at day 28 by bleeding of remaining mice from each group. Mice sera was subjected to ELISA analysis and anti-Vi IgG response was measured as per the procedure described by Micoli et al. [8].

Results

Determination of molecular weight and identity of *E. coli* derived rCRM₁₉₇

rCRM₁₉₇ appears as a ~58 kDa band in SDS/PAGE and Western blot analyses performed under both nonreducing and reducing conditions. This is consistent with the reported molecular weight of CRM₁₉₇ of 58.4 kDa [16]. Gels run under nonreducing conditions help to evaluate possible covalent and noncovalent aggregation in the samples and to detect impurities. rCRM₁₉₇ contains two disulfide bonds, one linking chains A and B and a second within the

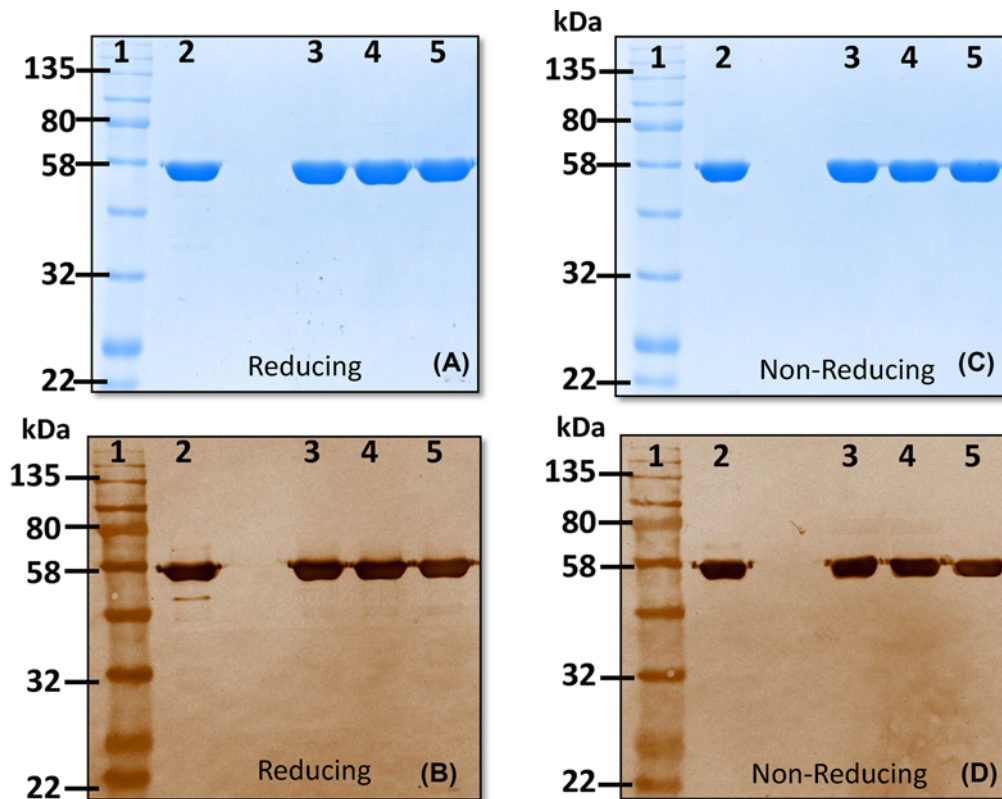


Figure 1. Determination of molecular weight, identity and purity of CRM₁₉₇ by SDS/PAGE and Western immunoblot respectively

(A,B) Reducing SDS/PAGE and Western blot analysis of three different lots of rCRM₁₉₇, and reference CRM₁₉₇. (C,D) Nonreducing SDS/PAGE and Western blot analysis of three different lots of rCRM₁₉₇ and reference CRM₁₉₇ (Lane 1: protein molecular weight marker, Lane 2: reference CRM₁₉₇, Lane 3–5: rCRM₁₉₇ Lot 1–3). Purified recombinant CRM₁₉₇ protein appeared as ~58 kDa molecular weight protein on SDS/PAGE with ≥95% purity. This molecular weight matches the expected size of the protein and with reference standard used. Anti-CRM₁₉₇ antibodies recognized rCRM₁₉₇ in Western blot, thereby confirming the protein identity as CRM₁₉₇.

B chain. Gels run in reducing conditions allow detection of the proteolytically nicked form (if any) of the protein. β-mercaptoethanol (2-βME) was used to reduce disulfide bridges during sample preparation. Under these reducing conditions, rCRM₁₉₇ (test sample) showed no visible evidence of nicked and/or degraded forms in SDS/PAGE (Figure 1A,B). Figure 1C,D summarizes the Western blot analysis of rCRM₁₉₇ and reference CRM₁₉₇ run in reducing and nonreducing conditions. The Western blot was probed with α-CRM₁₉₇ rabbit polyclonal antibody (make: Abcam; Cat No: ab151222).

Three independent manufacturing lots of rCRM₁₉₇ were evaluated in the present study and exhibited similar profiles under both nonreducing and reducing conditions, also supporting consistency of manufacturing for rCRM₁₉₇.

Intact mass determination by MS

MS is a powerful analytical tool as it measures an intrinsic property of a molecule, its mass, with very high sensitivity and therefore it is used in a wide range of applications [22]. The intact mass of rCRM₁₉₇ was obtained by deconvolution of mass/charge (m/z) ratio. Data obtained by TOF-MS showed most abundant mass of 58413 Da compared with a theoretical calculated mass of 58421 Da based on amino acid sequence (Figure 2). This result confirms the molecular weight of the purified rCRM₁₉₇ protein i.e. ~58.4 kDa. Similar data have also been reported both for a recombinant CRM₁₉₇ produced in *Pseudomonas fluorescens* and C7 CRM₁₉₇ [21,32].

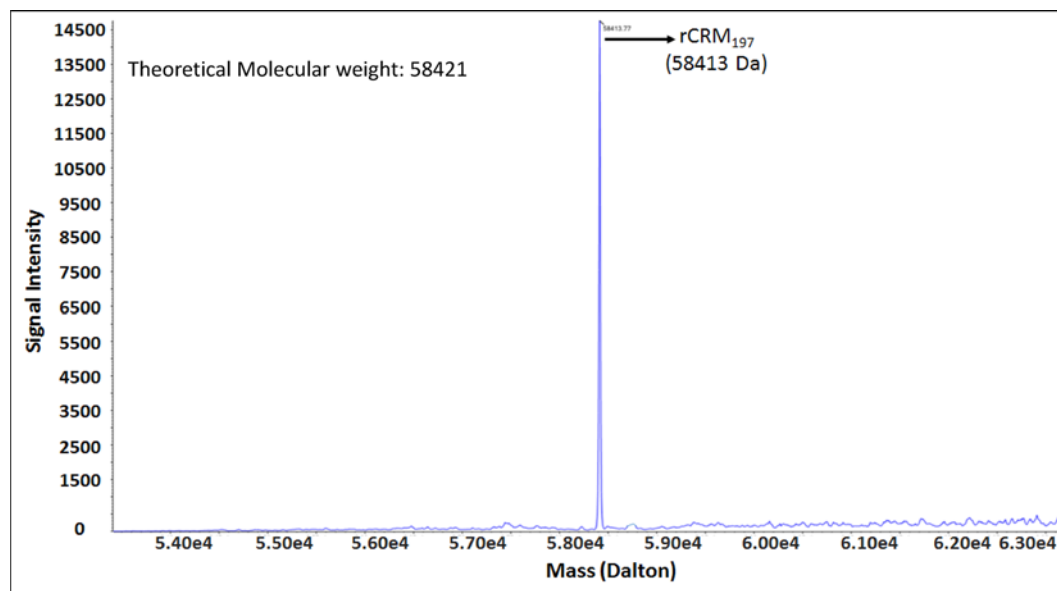


Figure 2. Determination of intact mass of rCRM₁₉₇ using MS (ESI-qTof)

The major abundant peak of 58.4 kDa has been observed which is similar to the expected/theoretical molecular weight of CRM₁₉₇ protein.

Endonuclease activity

Mild endonuclease activity is an inherent property of CRM₁₉₇ and its parent molecule DT [4] and is considered as an indicator of the correctly folded bioactive form of the protein. Incubation of CRM₁₉₇ with supercoiled plasmid DNA in the presence of divalent cations (e.g. Ca⁺⁺ and Mg⁺⁺) results in DNA cleavage [19,33]. Incubation of pUC57 plasmid DNA with rCRM₁₉₇ in a 2:1 ratio for different times and analysis of the DNA by agarose gel electrophoresis confirmed the positive nuclease activity in the protein. During incubation, rCRM₁₉₇ was able to linearize the supercoiled DNA. No endonuclease activity was seen in the 0-h time point sample but it increased with the incubation time (Figure 3). This endonuclease assay can be used as an in-process control assay for CRM₁₉₇ protein produced from insoluble inclusion bodies. Similarly, earlier studies in which endonuclease activity was used to establish the purity and correctness of refolding of rCRM₁₉₇ produced in *E. coli* is also reported [19]. Previous reports also showed that both DT and CRM₁₉₇ should have fragment A-associated nuclease activity [33].

Determination of amino acid composition

Acid hydrolysis followed by high-sensitivity AAA has been performed with rCRM₁₉₇ and compared with reference CRM₁₉₇. After acid hydrolysis, amino acid mixture was analysed by ultra performance liquid chromatography (UPLC) using the Waters AccQTag Ultra chemistry. All samples were analysed in duplicate and results expressed as an average. The amino acid composition of rCRM₁₉₇ closely matches (within 1–2% variability range) with reference and with the theoretical values of CRM₁₉₇ protein (Figure 4). AAA can be used to confirm the primary amino acid composition of protein. The overall data suggest that rCRM₁₉₇ produced in *E. coli* has similar amino acid composition as CRM₁₉₇ produced by *C. diphtheriae* C7 strain (reference CRM₁₉₇) [15].

Immunoblotting with α -CRM₁₉₇ mAbs

Three independent lots of rCRM₁₉₇ were run in SDS/PAGE and were probed with five different mAbs with specificity to different epitopes on the protein in Western immunoblotting experiments. All these mAbs recognized the protein at similar level and showed a major immune-reactive band at ~58 kDa, the expected size of the protein (Figure 5). The reference CRM₁₉₇ was recognized at similar level as rCRM₁₉₇. However, a slight difference in the recognition pattern was observed with respect to different mAbs, which suggests differential affinity of different mAbs for recognition of CRM₁₉₇. mAb mapping of CRM₁₉₇ has been found useful to assess potential nicking of A and B chains that could be seen in Western blot. A similar study was performed with different mAbs raised against CRM₁₉₇ for their ability to bind DT and six functional mutants CRM₁₉₇, CRM₁₇₆, CRM₂₂₈, CRM₁₀₀₁, CRM₄₅ and CRM₃₀ by immunoblotting

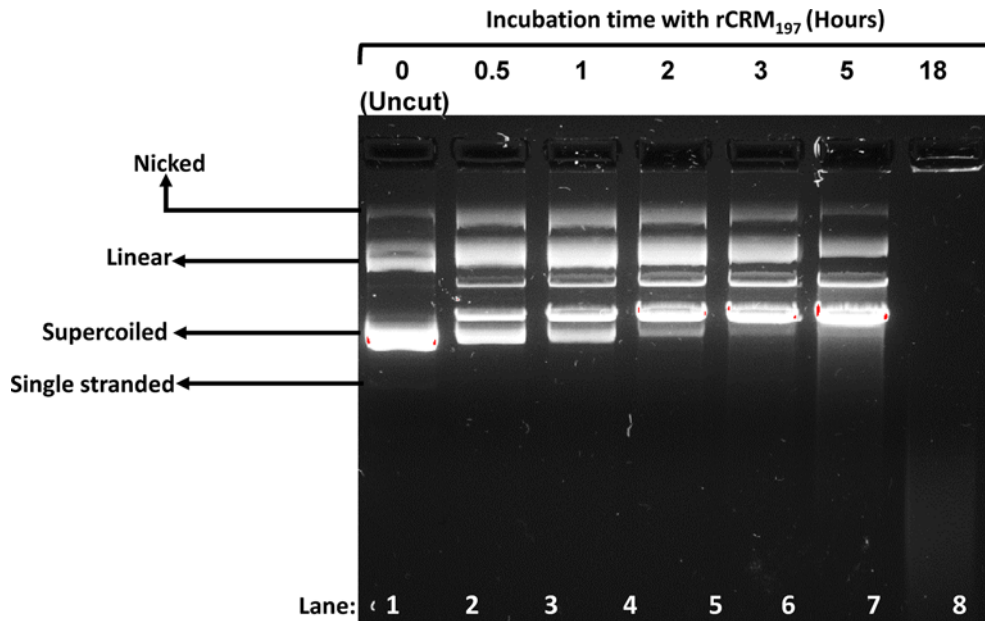


Figure 3. Demonstration of functional activity in rCRM₁₉₇ samples purified from inclusion body

pUC57 plasmid was incubated with rCRM₁₉₇ for different time points and the mix was run on 1% agarose gel. Native CRM₁₉₇ possesses a mild endonuclease activity and this assay is used to prove the restoration of biological activity and conformation integrity of the protein purified from insoluble inclusion body. The purified rCRM₁₉₇ exhibits the endonuclease activity as evident by nicking of supercoiled plasmid DNA incubated with the protein (Lanes 2–7).

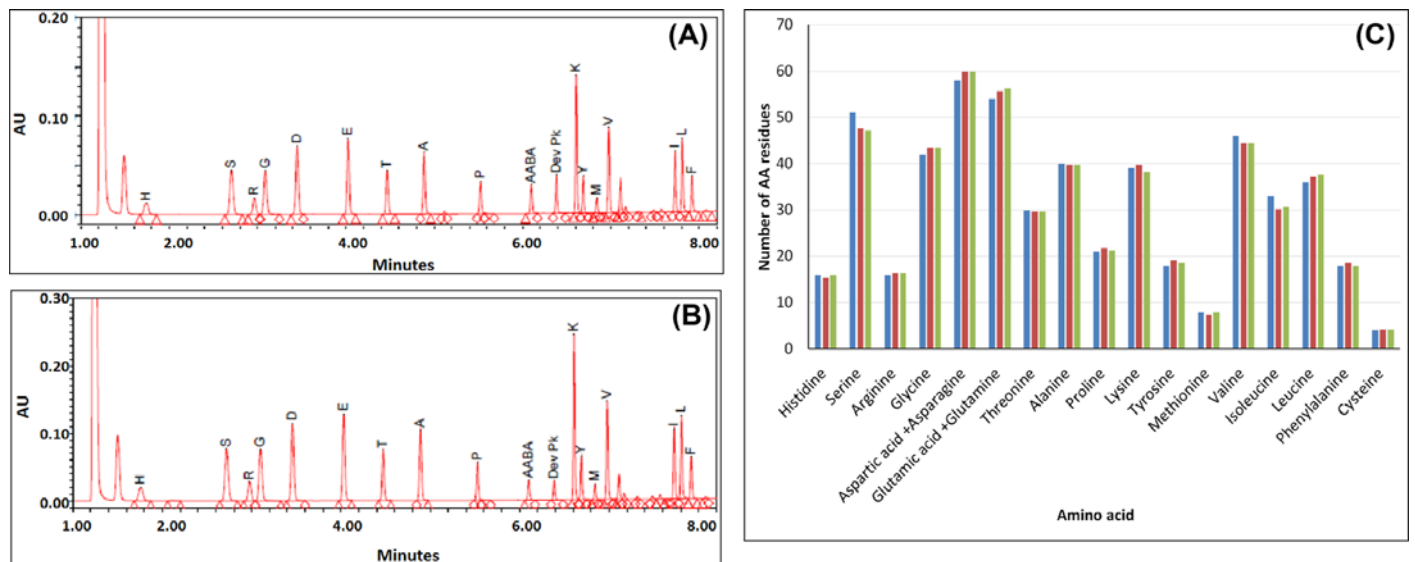


Figure 4. High-sensitivity AAA profile of rCRM₁₉₇ and CRM₁₉₇

Both the proteins showed similar elution profile during AAA chromatogram of high-sensitivity AAA of CRM₁₉₇. (A) AAA chromatogram of BioE rCRM₁₉₇ and (B) AAA chromatogram of reference CRM₁₉₇. Both the proteins showed similar elution profile. The peak data were extrapolated and plotted in bar diagram (C). Blue bar: theoretical number of amino acid residues, Red bar: experimental amino acid residues of rCRM₁₉₇ and Green bar: experimental amino acid residues of reference CRM₁₉₇.

[35]. The results showed a difference in the recognition pattern of mAbs specific to different part of the protein. Our data are of particular importance because protein is purified from insoluble inclusion body after refolding. There is potential chance of misfolding and or improper folding that might impact the conformation of the protein. Hence, the

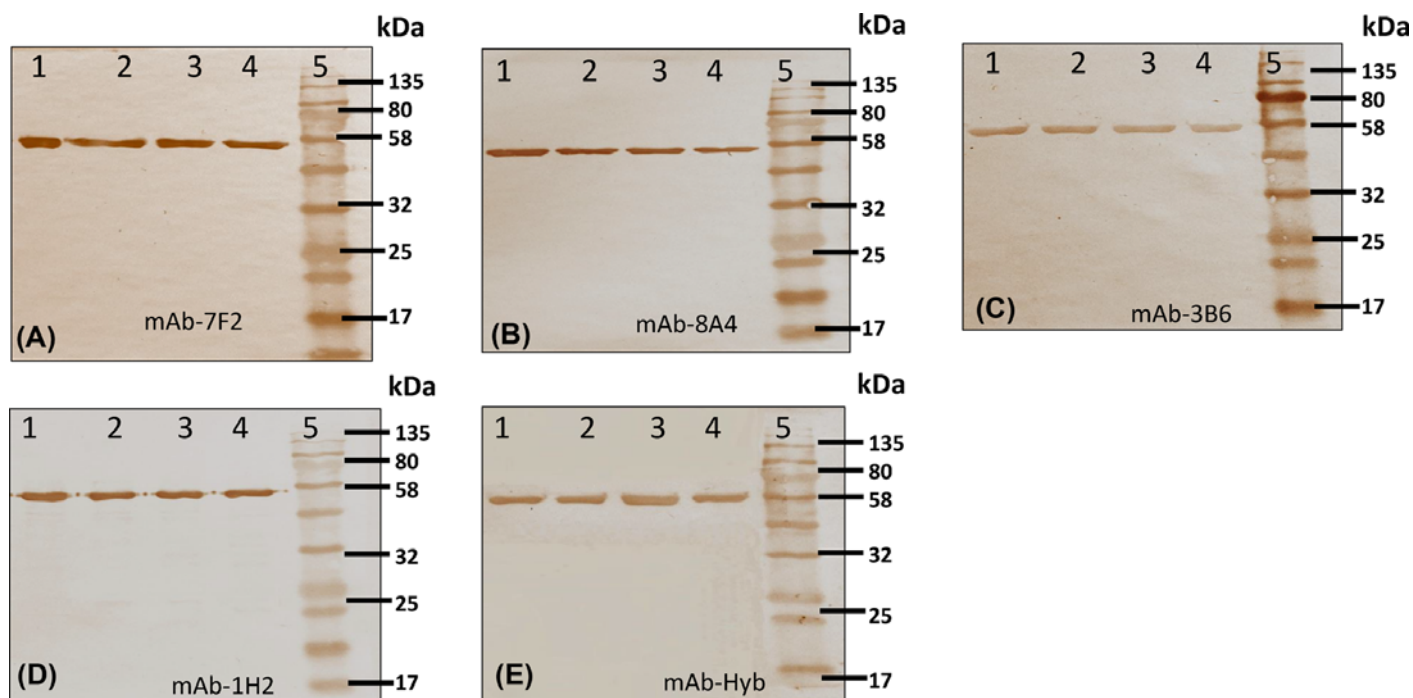


Figure 5. Mapping of rCRM₁₉₇ protein by structural mAbs against CRM₁₉₇ by Western immunoblot

All mAbs recognized rCRM₁₉₇ as single major band of ~58 kDa. This confirms the structural integrity and identity of rCRM₁₉₇ and its equivalence with reference. (A,B) Western blot using A chain specific α -CRM₁₉₇ mAb 7F2 and 8A4; (C) Western blot using B chain specific α -CRM₁₉₇ mAb 73B6; (D,E) Western blot using whole CRM₁₉₇ specific α -CRM₁₉₇ mAb 1H2 and Hyb. Lane 1: Reference CRM₁₉₇, 2-4: rCRM₁₉₇ Lot 1-3 and Lane 5: protein molecular weight marker.

data also confirm that rCRM₁₉₇ has attained the correct conformational structure after purification from insoluble inclusion bodies [35].

Determination of crystal structure of *E. coli* rCRM₁₉₇ by X-ray crystallography

The crystal structure of *E. coli* derived recombinant CRM₁₉₇ was determined at 2.3 Å resolution and the structure was submitted to the PDB with the PDB ID: 5I82. Previous studies have elucidated the structural features of DT in different conditions, such as in the *apo* conformation, at acidic pH, in complex with NAD, with dinucleotide inhibitor adenylyl 3'-5' uridine 3'-monophosphate and with an extracellular fragment of its receptor [36-39]. More recently, the structures of C7 CRM₁₉₇ in the *apo* form and in complex with the NAD hydrolysis product nicotinamide have been determined [16]. This showed that the single amino acid substitution of G52E that is the difference between CRM₁₉₇ and DT, results in an intrinsic flexibility of the active-site loop [16]. All these studies were performed on DT and CRM₁₉₇ produced by *C. diphtheriae*. However, there is very limited information available on CRM₁₉₇ protein produced from the recombinant *E. coli*. To verify if the rCRM₁₉₇ produced from insoluble inclusion body protein have the overall similar 3D structure of the protein, we determined the X-ray crystal structure of rCRM₁₉₇. Furthermore, the crystal structure of rCRM₁₉₇ confirms the presence of structural determinants that ablated the toxicity of DT and resulted in CRM₁₉₇ (G52E mutation).

The structure was determined at 2.3 Å resolution by molecular replacement using a search model, the coordinates of the *apo* form of C7 CRM₁₉₇ (PDB entry 4EA0). Data collection, refinement and the final model validation statistics for the structure of rCRM₁₉₇ (PDB ID: 5I82) have been given in Table 1. The asymmetric unit contains four molecules of rCRM₁₉₇ (chains A, B, C and D). The four chains interact as dimers (chains A-B and chains C-D) by domain swapping, with each monomer in 'open' conformation, a typical packing that has already been described for both DT and CRM₁₉₇ (Figure 6) [16]. The overall fold of each chain is similar to C7 CRM₁₉₇ (PDB: 4AE0). In fact, by superimposing the single chains or the two homodimers (chains A-B and C-D), the RMSD results were less than 0.17 Å (A and B, 428 out of 507 C α , RMSD of 0.17 Å; A and C, 424 out of 508 C α , RMSD of 0.17 Å; A and D, 426

Table 1 Refinement statistics of CRM₁₉₇ (PDB ID: 5182)

Data collection statistics	
Space group	P1
Unit cell: a, b, c (Å) α, β, γ (°)	69.21, 69.17, 127.88, 90.09, 90.01, 82.02
Wavelength (Å)	0.97872
Resolution (Å)	30.00–2.35 (2.39–2.35)*
Number of observed reflections	98137 (4888)
R _{merge} (%)	5.1 (42.2)
Completeness (%)	98.4 (97.7)
//σ(I)	18.2 (2.3)
Redundancy	2.6 (2.6)
Refinement and validation	
Resolution (Å)	29.08–2.35 (2.41–2.35)
R _{work} /R _{free} (%)	20.7 (27.2)/25.0 (32.9)
Number of atoms	
Protein	15730
Ligand/ion	21
Water	351
B-factors	
Protein/ligand/water	66.6/94.2/54.3
RMSD bond lengths (Å)	0.007
RMSD bond angles (°)	1.174
PDB ID	5182

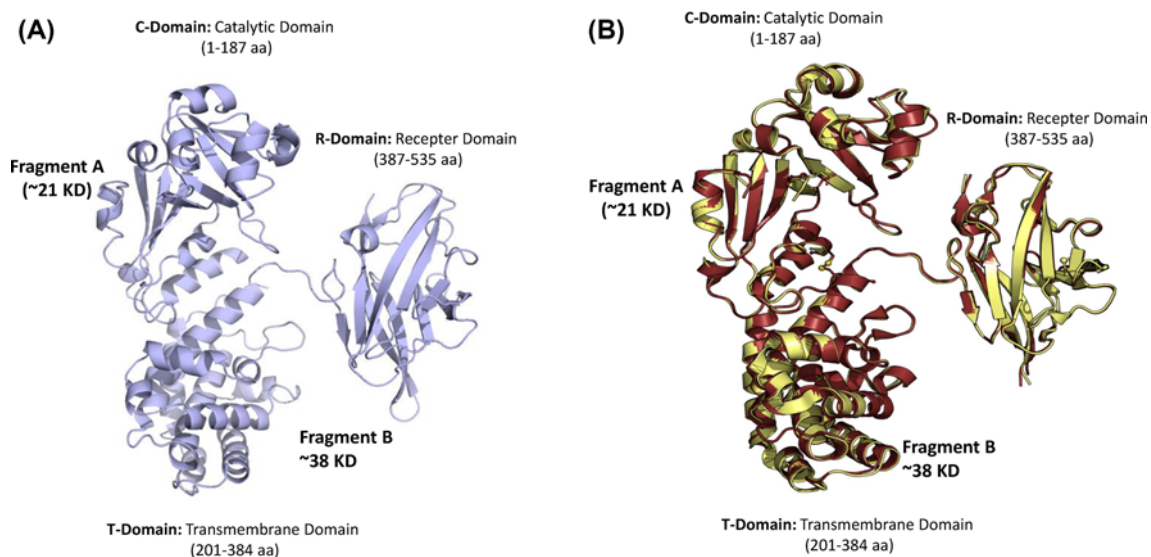


Figure 6. Crystal structure of monomeric form of *E. coli* derived rCRM197 protein (PDB ID: 5182)

(A) Superimposition of the monomer of rCRM₁₉₇ and C7 CRM₁₉₇. (B) Chain A (cartoon colored as ruby) and chain B (transparent, dark grey ribbon) of rCRM₁₉₇ are superimposed on the monomer of wild-type CRM₁₉₇ (PDB code: 4AE0, cartoon colored as yellow) and its symmetry-related molecule (transparent, yellow ribbon). (C) Catalytic domains consisting of residues 1–187, (T) Transmembrane domain consisting of residues 201–384, (R) Receptor domain consisting of residues 387–535. The dimers show the swapping of the T domains in the open conformation.

out of 504 C α , RMSD of 0.10 Å; AB and CD, 839 out of 1011 C α , RMSD of 0.17 Å) [40], demonstrating that these are very similar.

The dimers in the crystal structure of rCRM₁₉₇ and C7-CRM₁₉₇, are equivalent. By superimposing the A and B chains of rCRM₁₉₇ on to C7-CRM₁₉₇ and its symmetry-related molecule that forms the dimer, 924 out of 998 C α superimposed with RMSD of 0.38 Å (Figure 6). The overall fold of the monomer rCRM₁₉₇ is equivalent to C7-CRM₁₉₇, with an overall RMSD of 0.34 Å for 451 out of 498 superimposed C α atoms (Figure 6). Since overall structure is

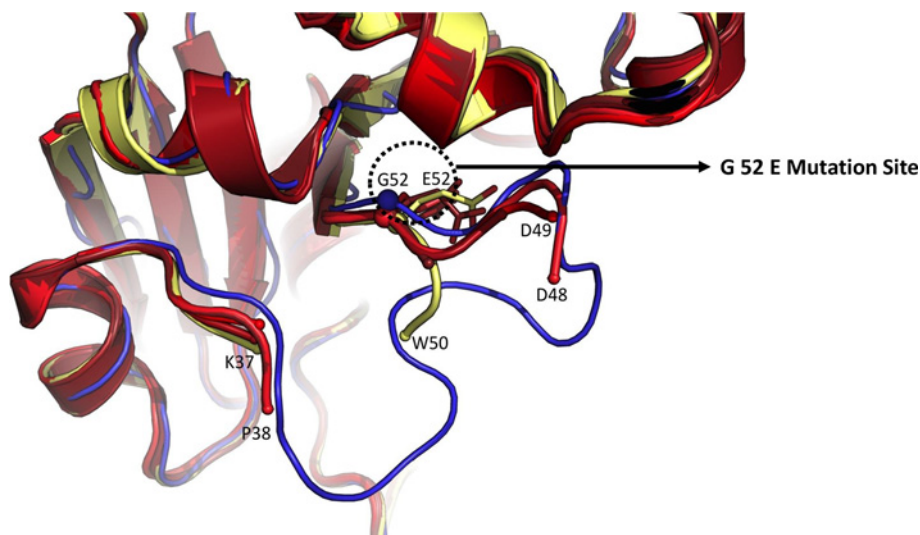


Figure 7. The glycine to glutamate mutation at amino acid position 52 (G52E) in rCRM₁₉₇.

Zoom in the active site loop CL2 where G52E mutation has taken place. The mutation creates a structural distortion in active site loop of DT thereby eliminating its toxicity (consequently CRM₁₉₇ is formed). The image depicts the structural superimposition of the DT (PDB code: 1SGK; thin blue ribbon), C7-CRM₁₉₇ (PDB code: 4AE0, yellow carton) and rCRM₁₉₇ (PDB code: 5I82: chain A as ruby cartoon, chain B as red cartoon, chain C as light red cartoon and chain D as firebrick color cartoon). The data demonstrate that rCRM₁₉₇ possesses G52E point mutation that converted DT into CRM₁₉₇. The same point mutation was also evident in C7-CRM₁₉₇.

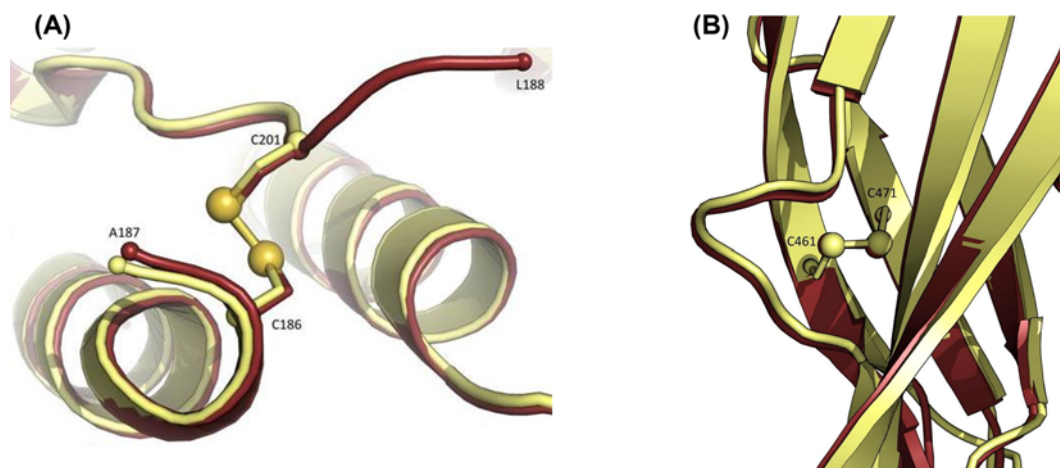


Figure 8. The presence of correct disulphide bonds in *E.coli* derived rCRM₁₉₇.

Zoom in the disulfide bond C186-C201 (A) and C461-C471 (B) in rCRM₁₉₇ (PDB ID: 5I82). Two disulfide bonds has been identified in at the expected position of the resolved structure of rCRM₁₉₇.

preserved between C7 CRM₁₉₇ and rCRM₁₉₇, it is unlikely that there will be any significant differences between these two molecules in terms of T epitopes that can potentially impact the efficacy of rCRM₁₉₇.

Furthermore, previous studies showed that the substitution G52E induces the dislocation of the active-site loop CL2 in C7-CRM₁₉₇ with respect to DT [16]. The intrinsic flexibility of CL2 results in a localized low quality electron density map in the C7-CRM₁₉₇ structure [38,39]. In our crystal structure of rCRM₁₉₇, we have also observed a low electron density in the active loop CL2, consistent with the previous results (Figure 7). This confirms the structural basis of lack of toxicity in rCRM₁₉₇.

Another important biochemical and structural features of DT, as well as C7 CRM₁₉₇, is the presence of the disulfide bond C186-C201 that maintains the A and B fragments bound together in the nicked form after the proteolytic cleavage of the chain. The structure revealed that the disulfide bond is conserved in rCRM₁₉₇ (Figure 8).

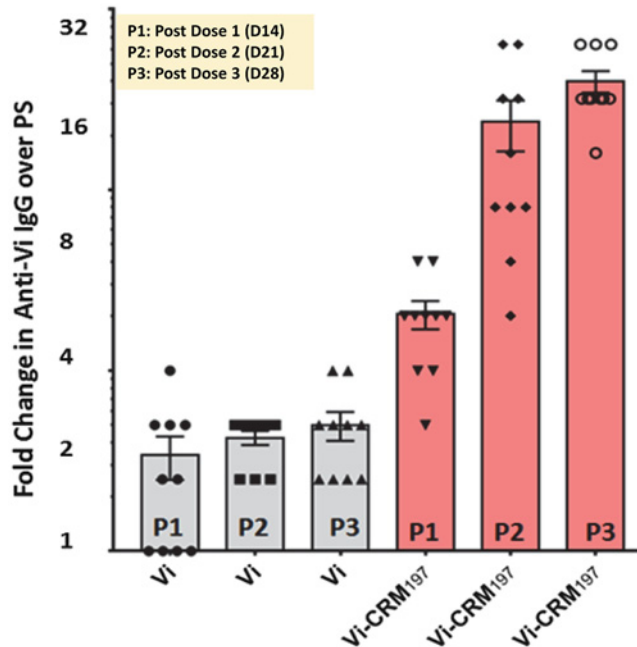


Figure 9. Immunogenicity of typhoid conjugate vaccine (Vi-rCRM₁₉₇) generated by conjugating Vi polysaccharide with rCRM₁₉₇

Balb/C mice immunized with three dose of Vi-rCRM₁₉₇ conjugate vaccine and Anti Vi-IgG was measured by ELISA. The titre of Vi specific antibody was compared with Vi PS (unconjugated Vi polysaccharide) group. There was significant increase in the Vi antibody was observed in Vi-rCRM₁₉₇ immunized mice and a booster response was also observed after second injection.

To our knowledge, this is the first reported crystal structure of *E. coli* derived rCRM₁₉₇ and first described comparison with C7-CRM₁₉₇. In conclusion, rCRM₁₉₇ conserves the overall folding, making it structurally equivalent to C7-CRM₁₉₇, which is used in many licenced conjugate vaccine such as meningitis, pneumococcal conjugate vaccine etc. [10-14].

Immunogenicity of Vi-rCRM₁₉₇ conjugate vaccine in mice

Immunogenicity of typhoid conjugate vaccine (Vi-rCRM₁₉₇) generated using *E. coli* derived rCRM₁₉₇ as carrier protein was evaluated in Balb/c mice. Mice were immunized with three doses of conjugate vaccine Vi-rCRM₁₉₇ and anti-Vi IgG immune response was measured by ELISA. Sera from mice immunized with PBS was used to generate the baseline. Vi-rCRM₁₉₇ immunized mice were able to elicit strong Anti-Vi IgG antibody response. Mice immunized with unconjugated Vi polysaccharide (Control) showed only basal level of Anti-Vi IgG. The Anti-Vi antibodies response also found to be bootable as seen in a steep increase in IgG titre after second dose of vaccine (Booster) (Figure 9). The Anti-Vi IgG response in mice immunized with Vi-rCRM₁₉₇ conjugate vaccine was found to be ~16-fold higher than the baseline response (PBS) after booster dose whereas un-conjugated Vi showed only 2 fold increase in the anti-Vi IgG over baseline. Noteworthy, the response of anti-Vi IgG in mice immunized with conjugate vaccine was 10-12 fold higher than the un-conjugated Vi.

As per the recommendation of World Health Organization-Technical Report Series (WHO-TRS) on typhoid conjugate vaccine, the conjugate vaccine should induce an immune response that is at least four-fold higher than the response induced by un-conjugated Vi, and a booster response should occur after the second dose in animals (WHO-TRS, 2013) [41]. The conjugate vaccine generated by using rCRM₁₉₇ as carrier protein comfortably meets the WHO immunogenicity criteria for typhoid conjugate vaccine, as it induces more than four-fold Anti-Vi IgG and also exhibits a booster response in mice. The data confirm the immunological functionality of rCRM₁₉₇ as carrier protein and suggests that it is able to convert T-independent response of polysaccharides in mice into T-dependent one and therefore booster antibody response is resulted. These data corroborate that rCRM₁₉₇ conjugation with Vi polysaccharide in mice can be resulted in immunogenic conjugates and therefore strengthening the suitability of rCRM₁₉₇ as a carrier protein for the development of conjugate vaccines.

Conclusion

The present study provides the detailed structural and immunological characteristics of CRM₁₉₇ carrier protein produced by heterologous expression in *E. coli*. In addition to reveal the scientific insights of rCRM₁₉₇, the data presented in this work are also important in the context that regulatory guidelines on conjugate vaccines that emphasizes the need for in-depth characterization of carrier protein used in the conjugate vaccines development. Successful and broadly marketed carbohydrate conjugate vaccines are based on just a few FDA-approved carrier proteins. First-generation of carrier proteins such as DT and tetanus toxin require detoxification with formaldehyde, potentially eliminating part of the lysine residues needed for glycan attachment, thereby potentially compromising the conjugation efficacy. CRM₁₉₇ is one of the most widely used and highly effective carrier protein for conjugate vaccine. Licensed conjugate vaccines such as HibTITER (*Haemophilus influenzae* type b associated diseases), Prevnar (pneumococcal diseases), and Menveo (meningococcal diseases) are containing CRM₁₉₇ as a carrier protein. Majority of currently marketed glycoconjugate vaccines contain CRM₁₉₇ as a carrier protein and access to high quality material will support the development and production of new conjugate vaccines. The data presented herein demonstrate for the first time structural and immunological characterization of rCRM₁₉₇ produced in *E. coli*.

In a recent study, the recombinant CRM₁₉₇ has also been expressed in *E. coli* using pET28a expression vector [42]. Unlike our study, CRM₁₉₇ was expressed in Histidine-tagged form (*His*-CRM₁₉₇) which requires a histidine removal step by enterokinase treatment. This process often poses the limitation during the scale-up of production process and inconsistencies with respect to batch to batch purity/impurity profiles of protein. The protein was purified at small scale by immobilized metal affinity chromatography (IMAC) with reasonable purity [42]. However, the in-depth analytical characterization of protein was also lacking in this study. In the present study, rCRM₁₉₇ expression was done in pTWIN expression vector, which releases the expressed protein from fusion tag by self-splicing with high fidelity. CRM₁₉₇ protein was purified at high scale (~20 l Fermentation Scale) using ion exchange chromatography and highly pure form of protein (>95%) was recovered as the final product.

Recently, we have studied the analytical comparability of five CRM₁₉₇ proteins produced in three different expression systems (*E. coli*, *P. fluorescens* and *C. diphtheriae* C7). A comprehensive physicochemical analysis of the CRM₁₉₇ molecules demonstrate that recombinant CRM₁₉₇s expressed in *E. coli* are overall highly similar to those expressed in the traditional system (*C. diphtheriae* C7) in terms of primary sequence/post-translational modifications, higher order structural integrity, apparent solubility, physical stability profile and *in vitro* antigenicity [43]. However, the detailed structural analysis and its *in vivo* antigenicity/immunological characterization was lacking. As a part of further in-depth characterization, we have determined the 3D crystal structure of *E. coli* derived rCRM₁₉₇. The rCRM₁₉₇ has shown the equivalence with C7 CRM₁₉₇ in terms of overall folding pattern, domain architecture and the presence of correct disulfide bonds. The absence of toxicity in CRM₁₉₇ relies on a point mutation at amino acid position 52 that replaces Glycine with Glutamate (G52E) and renders protein nontoxicity. In the present study, the genetic basis of nontoxicity of *E. coli* based rCRM₁₉₇ was also confirmed by X-ray crystallography. These data are important for scientific communities as well as industries working on conjugate vaccine development using CRM₁₉₇ as carrier protein. The *in vivo* functionality of *E. coli* expressed recombinant CRM₁₉₇ carrier protein has been demonstrated by conjugating it with Vi polysaccharide antigen of *Salmonella* Typhi thereby generating Vi-rCRM₁₉₇ conjugate vaccine. The data confirm that the use of rCRM₁₉₇ as a carrier protein in conjugate vaccine will be able to elicit the primary and a booster antibody response in animals. The Anti-Vi IgG elicited by Vi-rCRM₁₉₇ conjugate vaccine was more than four-fold higher than the response generated by unconjugated Vi which is one of the benchmark for a protective typhoid conjugate vaccine [41]. The comparative immunogenicity of Vi polysaccharide conjugated to other carrier protein like tetanus toxoid (TT), diphtheria toxoid (DT) and C7 CRM₁₉₇ is already published [44]. The immunogenicity of Vi-rCRM₁₉₇ showed the comparable result in terms of overall fold change in Anti-Vi IgG over Vi and booster impact. The overall data presented in our study will potentially catalyse the efforts for the production of CRM₁₉₇ protein in the recombinant system and its use as carrier protein in the conjugate vaccine development.

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Competing interests

A.G., R.P.N.M. and to R.S.P.Y. are full-time employees of Biological E. Ltd., Hyderabad, India. The other authors declare that they have no competing interests associated with the manuscript.

Author contribution

R.P.N.M., A.G., W.F.A. and C.J. conceived and designed the research. R.P.N.M., R.P.Y., G.M., S.N., and L.A.S. were responsible for execution, analysis, and interpretation of data. R.P.N.M., A.G., C.J., S.N., and G.M. wrote and reviewed the manuscript.

Abbreviations

AAA, amino acid analysis; ADH, adipic acid dihydrazine; ADP, adenosine diphosphate; CRM₁₉₇, cross reacting material 197; CSGID, Center for the Structural Genomics of Infectious Diseases; DT, diphtheria toxin; FDA, Food and Drug Administration; HB-EGF, heparin binding epidermal growth factor; HRP, horseradish peroxidase; LC-qTOF, liquid chromatography-quadrupole time-of-flight; mAb, monoclonal antibody; PDB, Protein Data Bank; TAE, Tris-acetate-EDTA; WHO-TRS, World Health Organization-Technical Report Series.

References

- 1 Goldblatt, D. (2000) Conjugate vaccines. *Clin. Exp. Immunol.* **119**, 1–3, <https://doi.org/10.1046/j.1365-2249.2000.01109.x>
- 2 Avci, F.Y., Li, X., Tsuji, M. and Kasper, D.L. (2011) A mechanism for glycoconjugate vaccine activation of the adaptive immune system and its implications for vaccine design. *Nat. Med.* **17**, 1602–1609, <https://doi.org/10.1038/nm.2535>
- 3 Pichichero, M.E. (2013) Protein carriers of conjugate vaccines: characteristics, development, and clinical trials. *Hum. Vacc. Immun.* **9**, 2505–2523, <https://doi.org/10.4161/hv.26109>
- 4 Uchida, T., Pappenheimer, A.M. and Greany, R. (1973) Diphtheria toxin and related proteins I. Isolation and properties of mutant proteins serologically related to diphtheria toxin. *J. Biol. Chem.* **248**, 3838–3844
- 5 Giannini, G., Rappuoli, R. and Ratti, G. (1984) The amino-acid sequence of two non-toxic mutants of diphtheria toxin: CRM₄₅ and CRM₁₉₇. *Nucleic Acid Res.* **12**, 4063–4069, <https://doi.org/10.1093/nar/12.10.4063>
- 6 Gupta, R.K. and Siber, G.R. (1995) Adjuvants for human vaccines—current status, problems and future prospects. *Vaccine* **13**, 1263–1276, [https://doi.org/10.1016/0264-410X\(95\)00011-0](https://doi.org/10.1016/0264-410X(95)00011-0)
- 7 Shinefield, H.R. (2010) Overview of the development and current use of CRM₁₉₇ conjugate vaccines for pediatric use. *Vaccine* **28**, 4335–4339, <https://doi.org/10.1016/j.vaccine.2010.04.072>
- 8 Micoli, F., Rondini, S., Pisoni, I., Proietti, D., Berti, F., Costantino, P. et al. (2011) Vi-CRM₁₉₇ as a new conjugate vaccine against *Salmonella* Typhi. *Vaccine* **29**, 712–720, <https://doi.org/10.1016/j.vaccine.2010.11.022>
- 9 Micoli, F., Rondini, S., Gavini, M., Lanzilao, L., Medagliani, D., Saul, A. et al. (2012) O: 2-CRM₁₉₇ conjugates against *Salmonella* paratyphi A. *PLoS ONE* **7**, e47039, <https://doi.org/10.1371/journal.pone.0047039>
- 10 Fiorino, G., Peyrin-Biroulet, L., Naccarato, P., Szabó, H., Sociale, O.R., Vetrano, S. et al. (2012) Effects of immunosuppression on immune response to pneumococcal vaccine in inflammatory bowel disease: a prospective study. *Inflamm. Bowel Dis.* **18**, 1042–1047, <https://doi.org/10.1002/ibd.21800>
- 11 Golden, A.R., Adam, H.J. and Zhan, G.G. (2016) Canadian Antimicrobial Resistance Alliance (CARA). Invasive *Streptococcus pneumoniae* in Canada, 2011–2014: characterization of new candidate 15-valent pneumococcal conjugate vaccine serotypes 22F and 33F. *Vaccine* **34**, 2527–2530, <https://doi.org/10.1016/j.vaccine.2016.03.058>
- 12 Cheng, B.L., Nielsen, T.B., Pantapalangkoor, P., Zhao, F., Lee, J.C., Montgomery, C.P. et al. (2017) Evaluation of serotypes 5 and 8 capsular polysaccharides in protection against *Staphylococcus aureus* in murine models of infection. *Hum. Vaccine Immunother.* **20**, 1–6
- 13 Carboni, F., Adamo, R., Fabbrini, M., De Ricco, R., Cattaneo, V., Brogioni, B. et al. (2017) Structure of a protective epitope of group B *Streptococcus* type III capsular polysaccharide. *Proc. Natl. Acad. Sci. U.S.A.* **114**, 5017–5022, <https://doi.org/10.1073/pnas.1701885114>
- 14 Meng, X., Ji, C., Su, C., Shen, D., Li, Y., Dong, P. et al. (2017) Synthesis and immunogenicity of PG-tb1 monovalent glycoconjugate. *Eur. J. Med. Chem.* **34**, 140–146, <https://doi.org/10.1016/j.ejmech.2017.03.058>
- 15 Bröker, M., Costantino, P., DeTora, L., McIntosh, E.D. and Rappuoli, R. (2011) Biochemical and biological characteristics of cross-reacting material 197 (CRM₁₉₇), a non-toxic mutant of diphtheria toxin: use as a conjugation protein in vaccines and other potential clinical applications. *Biologicals* **39**, 195–204, <https://doi.org/10.1016/j.biologicals.2011.05.004>
- 16 Malito, E., Bursulaya, B., Chen, C., Surdo, P.L., Picchianti, M., Balducci, E. et al. (2012) Structural basis for lack of toxicity of the diphtheria toxin mutant CRM₁₉₇. *Proc. Natl. Acad. Sci. U.S.A.* **109**, 5229–5234, <https://doi.org/10.1073/pnas.1201964109>
- 17 Patkar, A. (2012) Recombinant production of carrier proteins. *Genet. Eng. Biotech. News* **32**, <https://doi.org/10.1089/gen.32.21.23>
- 18 Goel, A., Mishra, R.P.N., Mantena, N. and Datla, M. (2017) Codon optimized polynucleotide for high level expression of CRM₁₉₇. WO 2,016,079,755 A1 (patent pending)

- 19 Stefan, A., Conti, M., Rubboli, D., Ravagli, L., Presta, E. and Hochkoepler, A. (2011) Overexpression and purification of the recombinant diphtheria toxin variant CRM₁₉₇ in *Escherichia coli*. *J. Biotechnol.* **156**, 245–252, <https://doi.org/10.1016/j.jbiotec.2011.08.024>
- 20 Sambrook, J., Fritsch, E.F. and Maniatis, T. (1989) *Molecular Cloning*, Cold Spring Harbor Laboratory Press, New York
- 21 Brady, C., Killeen, K., Taylor, W., Patkar, A. and Lees, A. (2012) Carrier protein outsourcing. *BioProcess Int.* **10**, 50–55
- 22 Mann, M., Hendrickson, R.C. and Pandey, A. (2001) Analysis of proteins and proteomes by mass spectrometry. *Annu. Rev. Biochem.* **70**, 437–473, <https://doi.org/10.1146/annurev.biochem.70.1.437>
- 23 Edman, P. (1950) Method for determination of the amino acid sequence in peptides. *Acta Chem. Scand.* **4**, 283–293, <https://doi.org/10.3891/acta.chem.scand.04-0283>
- 24 Metz, B., Jiskoot, W., Hennink, W.E., Crommelin, D.J.A. and Kerstena, G.F.A. (2003) Physicochemical and immunochemical techniques predict the quality of diphtheria toxoid vaccines. *Vaccine* **22**, 156–167, <https://doi.org/10.1016/j.vaccine.2003.08.003>
- 25 Anderson, W.F. (2009) Structural genomics and drug discovery for infectious diseases. *Infect. Disorder Drug Targets* **9**, 507, <https://doi.org/10.2174/187152609789105713>
- 26 Kim, Y., Bigelow, L., Borovilos, M., Dementieva, I., Duggan, E., Hatzos, C. et al. (2008) High-throughput protein purification for X-ray crystallography and NMR. *Adv. Protein Chem. Struct. Biol.* **75**, 85–105
- 27 Otwinowski, Z. and Minor, W. (1997) Processing of X-ray diffraction data collected in oscillation mode. *Methods Enzymol.* **276**, 307–326, [https://doi.org/10.1016/S0076-6879\(97\)76066-X](https://doi.org/10.1016/S0076-6879(97)76066-X)
- 28 Murshudov, G.N., Skubák, P., Lebedev, A.A., Pannu, N.S., Steiner, R.A., Nicholls, R.A. et al. (2011) REFMAC5 for the refinement of macromolecular crystal structures. *Acta Crystallogr. D Biol. Crystallogr.* **67**, 355–367, <https://doi.org/10.1107/S0907444911001314>
- 29 Emsley, P. and Cowtan, K. (2004) Coot: model-building tools for molecular graphics. *Acta Crystallogr. D Biol. Crystallogr.* **60**, 2126–2132, <https://doi.org/10.1107/S0907444904019158>
- 30 Rondini, S., Micoli, F., Lanzilao, L., Hale, C., Saul, A.J. and Martin, L.B. (2011) Evaluation of the immunogenicity and biological activity of the *Citrobacter freundii* Vi-CRM₁₉₇ conjugate as a vaccine for *Salmonella enterica* serovar Typhi. *Clin. Vaccine Immunol.* **18**, 460–468, <https://doi.org/10.1128/CVI.00387-10>
- 31 Kossaczka, Z., Bystrycki, S., Bryla, D.A., Shiloach, J., Robbins, J.B. and Szu, S.C. (1997) Synthesis and immunological properties of Vi and di-O-acetyl pectin protein conjugates with adipic acid dihydrazide as the linker. *Infect. Immun.* **65**, 2088–2093
- 32 Crane, D.T., Bolgiano, B. and Jones, C. (1997) Comparison of the diphtheria mutant toxin, CRM₁₉₇, with a *Haemophilus influenzae* type-b polysaccharide-CRM₁₉₇ conjugate by optical spectroscopy. *Eur. J. Biochem.* **246**, 320–327, <https://doi.org/10.1111/j.1432-1033.1997.00320.x>
- 33 Bruce, C., Baldwin, R.L., Lessnick, S.L. and Wisniewski, B.J. (1990) Diphtheria toxin and its ADP-ribosyltransferase-defective homologue CRM₁₉₇ possess deoxyribonuclease activity. *Proc. Natl. Acad. Sci. U.S.A.* **87**, 2995–2998, <https://doi.org/10.1073/pnas.87.8.2995>
- 34 Mahamad, P., Boonchird, C. and Panbangred, W. (2016) High level accumulation of soluble diphtheria toxin mutant (CRM₁₉₇) with co-expression of chaperones in recombinant *Escherichia coli*. *Appl. Microbiol. Biotechnol.* **28**, 1–2
- 35 Bigio, M., Rossi, R., Nucci, D., Antoni, G., Rappouli, R. and Ratti, G. (1987) Conformational changes in diphtheria toxoids: analysis with monoclonal antibodies. *FEBS Lett.* **218**, 271–276, [https://doi.org/10.1016/0014-5793\(87\)81060-8](https://doi.org/10.1016/0014-5793(87)81060-8)
- 36 Nocadello, S., Minasov, G., Shuvalova, L., Dubrovskaya, I., Sabini, E., Bagnoli, F. et al. (2016) Crystal structures of the components of the *Staphylococcus aureus* leukotoxin ED. *Acta Crystallogr. D Struct. Biol.* **72**, 113–120, <https://doi.org/10.1107/S2059798315023207>
- 37 Louie, G.V., Yang, W., Bowman, M.E. and Choe, S. (1997) Crystal structure of the complex of diphtheria toxin with an extracellular fragment of its receptor. *Mol. Cell* **1**, 67–78, [https://doi.org/10.1016/S1097-2765\(00\)80008-8](https://doi.org/10.1016/S1097-2765(00)80008-8)
- 38 Bell, C.E. and Eisenberg, D. (1997) Crystal structure of nucleotide-free diphtheria toxin. *Biochemistry* **36**, 481–488, <https://doi.org/10.1021/bi962214s>
- 39 Bennett, M.J., Choe, S. and Eisenberg, D. (1994) Refined structure of dimeric diphtheria toxin at 2.0 Å resolution. *Protein Sci.* **3**, 1444–1463, <https://doi.org/10.1002/pro.5560030911>
- 40 Leka, O., Vallese, F., Pirazzini, M., Berto, P., Montecucco, C. and Zanotti, G. (2014) Diphtheria toxin conformational switching at acidic pH. *FEBS J.* **281**, 2115–2122, <https://doi.org/10.1111/febs.12783>
- 41 World Health Organization (2013) Guidelines on the quality, safety and efficacy of typhoid conjugate vaccines.
- 42 Lu, X.U., Jing, Z., Rong, Y.U. and Zhiguo, S.U. (2017) Expression of CRM₁₉₇ in *E. coli* system and its application in universal influenza vaccine. *Chin. J. Process Eng.* **17**, 1054–1058, <https://doi.org/10.12034/j.issn.1009.606X.217132>
- 43 Hickey, J.M., Toprani, V.M., Kaur, K., Mishra, R.P., Goel, A., Oganessian, N. et al. (2018) Analytical comparability assessments of 5 recombinant CRM₁₉₇ proteins from different manufacturers and expression systems. *J. Pharm. Sci.* **107**, 1806–1819, <https://doi.org/10.1016/j.xphs.2018.03.002>
- 44 Arcuri, M., Di Benedetto, R., Cunningham, A.F., Saul, A., MacLennan, C.A. and Micoli, F. (2017) The influence of conjugation variables on the design and immunogenicity of a glycoconjugate vaccine against *Salmonella* Typhi. *PLoS ONE* **12**, e0189100, <https://doi.org/10.1371/journal.pone.0189100>