



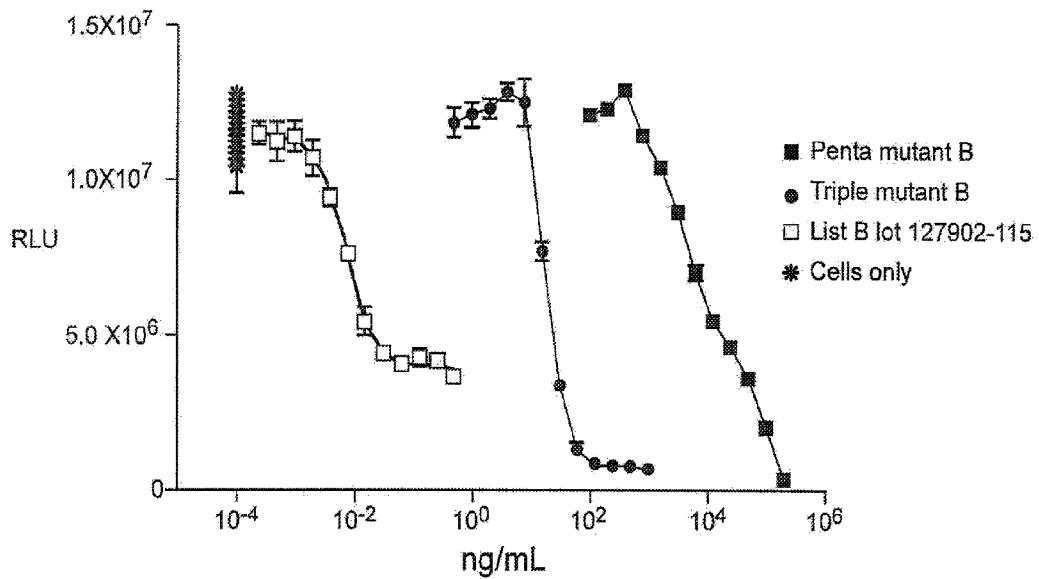
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(54) Titre : COMPOSITIONS ET METHODES RELATIVES A UNE TOXINE MUTANTE DE CLOSTRIDIUM DIFFICILE
(54) Title: COMPOSITIONS AND METHODS RELATING TO A MUTANT CLOSTRIDIUM DIFFICILE TOXIN



(57) Abrégé/Abstract:

In one aspect, the invention relates to an immunogenic composition that includes a mutant Clostridium difficile toxin A and/or a mutant Clostridium difficile toxin B. The mutant toxin may include a glucosyltransferase domain having at least one mutation and a cysteine protease domain having at least one mutation, relative to the corresponding wild-type C. difficile toxin. The mutant toxins may include at least one amino acid that is chemically crosslinked. In another aspect, the invention relates to methods and compositions for use in culturing Clostridium difficile and in producing C. difficile toxins.

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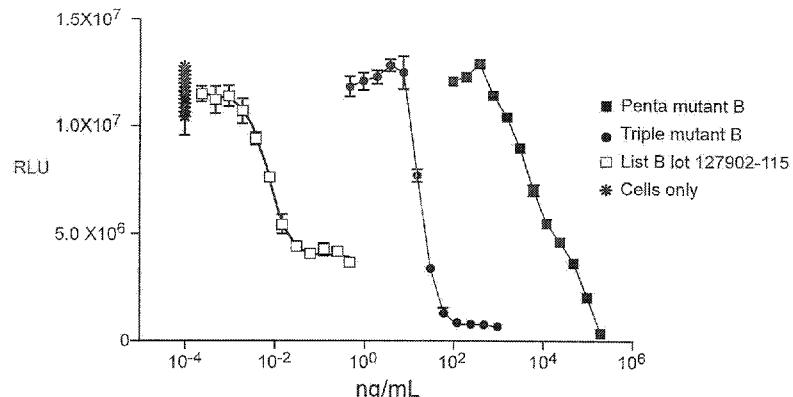
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[Continued on next page]

(54) Title: COMPOSITIONS AND METHODS RELATING TO A MUTANT CLOSTRIDIUM DIFFICILE TOXIN

FIG. 26



(57) Abstract: In one aspect, the invention relates to an immunogenic composition that includes a mutant *Clostridium difficile* toxin A and/or a mutant *Clostridium difficile* toxin B. The mutant toxin may include a glucosyltransferase domain having at least one mutation and a cysteine protease domain having at least one mutation, relative to the corresponding wild-type *C. difficile* toxin. The mutant toxins may include at least one amino acid that is chemically crosslinked. In another aspect, the invention relates to methods and compositions for use in culturing *Clostridium difficile* and in producing *C. difficile* toxins.

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COMPOSITIONS AND METHODS RELATING TO A MUTANT *CLOSTRIDIUM**DIFFICILE* TOXIN

CROSS REFERENCE TO RELATED APPLICATIONS

The present application claims the benefit of U.S. Provisional Patent Application 5 61/716,605, filed on October 21, 2012,

FIELD

The present invention is directed to compositions and methods relating to mutant *Clostridium difficile* toxins.

BACKGROUND

10 *Clostridium difficile* (*C. difficile*) is a Gram-positive anaerobic bacterium that is associated with gastrointestinal disease in humans. Colonization of *C. difficile* usually occurs in the colon if the natural gut flora is diminished by treatment with antibiotics. An infection can lead to antibiotic-associated diarrhea and sometimes pseudomembranous colitis through the secretion of the glucosylating toxins, toxin A and toxin B (308 and 270 15 kDa, respectively), which are the primary virulence factors of *C. difficile*.

Toxin A and toxin B are encoded within the 19 kb pathogenicity locus (PaLoc) by the genes *tcdA* and *tcdB*, respectively. Nonpathogenic strains of *C. difficile* have this locus replaced by an alternative 115 base pair sequence.

Both toxin A and toxin B are potent cytotoxins. These proteins are homologous 20 glucosyltransferases that inactivate small GTPases of the Rho/Rac/Ras family. The resulting disruption in signaling causes a loss of cell-cell junctions, dysregulation of the actin cytoskeleton, and/or apoptosis, resulting in the profound secretory diarrhea that is associated with *Clostridium difficile* infections (CDI).

In the last decade, the numbers and severity of *C. difficile* outbreaks in hospitals, 25 nursing homes, and other long-term care facilities increased dramatically. Key factors in this escalation include emergence of hypervirulent pathogenic strains, increased use of antibiotics, improved detection methods, and increased exposure to airborne spores in health care facilities.

Metronidazole and vancomycin represent the currently accepted standard of care 30 for the antibiotic treatment of *C. difficile* associated disease (CDAD). However, about 20% of patients receiving such treatment experience a recurrence of infection after a first episode of CDI, and up to about 50% of those patients suffer from additional

recurrences. Treatment of recurrences represents a very significant challenge, and the majority of recurrences usually occur within one month of the preceding episode.

Accordingly, there is a need for immunogenic and/or therapeutic compositions and methods thereof directed to *C. difficile*.

5 SUMMARY OF THE INVENTION

These and other objectives are provided by the invention herein.

In one aspect, the invention relates to a mutant *C. difficile* toxin A. The mutant toxin A includes a mutation at residues positions 285, 287, 700, 972, and 978 as compared to a wild-type toxin A. In one embodiment, the mutant toxin A includes SEQ 10 ID NO: 183. In one embodiment, the mutant toxin A is less cytotoxic than a corresponding wild-type toxin A. In one embodiment, the mutant toxin A includes at least one amino acid residue that is chemically modified. In one aspect, the invention relates to an isolated polypeptide that includes SEQ ID NO: 183.

In another aspect, the invention relates to a mutant *C. difficile* toxin B. The 15 mutant toxin B includes a mutation at residues 286, 288, 698, 970, and 976 as compared to a wild-type toxin B. In one embodiment, the mutant toxin B includes SEQ ID NO: 184. In one embodiment, the mutant toxin B is less cytotoxic than a corresponding wild-type toxin B. In one embodiment, the mutant toxin A includes at least one amino acid residue that is chemically modified. In one aspect, the invention 20 relates to an isolated polypeptide that includes SEQ ID NO: 184.

The invention further relates to compositions and methods for use in culturing *C. difficile* and producing *C. difficile* toxins. In one aspect, the invention relates to a culture medium including a vegetable hydrolysate and a *C. difficile* cell. In a preferred embodiment, the hydrolysate is soy hydrolysate. More preferably, the soy hydrolysate 25 is soy hydrolysate SE50MK.

In another aspect, the invention relates to a culture medium including a nitrogen source and a *C. difficile* cell. In one embodiment, the nitrogen source is a yeast extract. Preferably, the yeast extract is HY YEST 412 (Kerry Biosciences).

In a further aspect, the invention relates to a culture medium including a 30 vegetable hydrolysate, yeast extract, and a *C. difficile* cell. In one embodiment, the medium does not contain a carbon source.

In a preferred embodiment, the medium further includes a carbon source. The inventors discovered that fermentation of *C. difficile* in a culture medium including at

least one carbon source provided high OD₆₀₀ values and high toxin production yields, as compared to fermentation without a carbon source. In one embodiment, the carbon source is glucose, mannitol, fructose, and/or mannose.

5 In one embodiment, the *C. difficile* cell is not genetically modified. In another embodiment, the *C. difficile* cell is a recombinant *C. difficile* cell. In one embodiment, the *C. difficile* cell lacks an endogenous polynucleotide encoding a toxin. In another embodiment, the cell includes a constitutive promoter. In a preferred embodiment, the promoter is a *Clostridium sporogenes* ferredoxin(*fdx*) promoter. In a further embodiment, the cell does not include a native, regulated chromosomal promoter.

10 In another aspect, the invention relates to a method of culturing *C. difficile*. The method includes culturing a *C. difficile* cell in a medium. In one embodiment, the medium includes soy hydrolysate and/or yeast extract. In a preferred embodiment, the medium further includes a carbon source. Preferably, the carbon source is glucose.

15 In one embodiment, the culturing step is carried out under anaerobic conditions. In one embodiment, the *C. difficile* is grown as a seed culture. In one embodiment, the seed culture is started by inoculation from a stock culture that was grown in the medium.

20 In one embodiment, the *C. difficile* is grown as a fermentation culture. In one embodiment, the fermentation culture was inoculated from a seed culture that was grown in the medium. In an alternative aspect, the invention relates to a method of culturing *C. difficile*. The method includes culturing a *C. difficile* cell in a monoclonal antibody medium.

25 In one aspect, the invention relates to a method of producing a *C. difficile* toxin. The method includes culturing a *C. difficile* cell in a medium. The method further includes isolating a *C. difficile* toxin from said medium.

BRIEF DESCRIPTION OF DRAWINGS

Figure 1 A-H: Sequence alignment of wild-type *C. difficile* toxin A from strains 630, VPI10463, R20291, CD196, and mutant toxin A having SEQ ID NO: 4, using CLUSTALW alignment, default parameters.

5 Figure 2 A-F: Sequence alignment of wild-type *C. difficile* toxin B from strains 630, VPI10463, R20291, CD196, and mutant toxin B having SEQ ID NO: 6, using CLUSTALW alignment, default parameters.

10 Figure 3: Graph showing identification of wild-type toxin-negative *C. difficile* strains. Culture media of 13 *C. difficile* strains were tested by ELISA for toxin A. As illustrated, seven strains expressed toxin A and 6 strains did not (strains 1351, 3232, 7322, 5036, 4811 and VPI 11186).

15 Figure 4 A and B: SDS-PAGE results illustrating that triple mutant A (SEQ ID NO: 4), double mutant B (SEQ ID NO: 5), and triple mutant B (SEQ ID NO: 6) do not glucosylate Rac1 or RhoA GTPases in an *in vitro* glucosylation assays with UDP-¹⁴C-glucose; whereas 10 µg to 1 ng of wild type toxin B does glucosylate Rac1.

Figure 5: Western blot indicating abrogation of cysteine protease activity in mutant toxins A and B (SEQ ID NOs: 4 and 6, respectively), as compared to observation of cleaved fragments of wild-type toxins A and B (SEQ ID NOs: 1 and 2, respectively). See Example 13.

20 Figure 6: Graphs showing that triple mutant toxins A and B (SEQ ID NOs: 4 and 6, respectively) exhibit residual cytotoxicity when tested at high concentrations (e.g., about 100 µg/ml) by *in vitro* cytotoxicity assay in IMR-90 cells.

Figure 7: Graph showing that EC₅₀ values are similar for the triple mutant toxin B (SEQ ID NO: 6) and hepta mutant toxin B (SEQ ID NO: 8).

25 Figure 8: Graph representing results from *in vitro* cytotoxicity tests in which the ATP levels (RLUs) are plotted against increasing concentrations of the triple mutant TcdA (SEQ ID NO: 4)(top panel) and triple mutant TcdB (SEQ ID NO: 6)(bottom panel). Residual cytotoxicity of mutant toxin A and B can be completely abrogated with neutralizing antibodies specific for mutant toxin A (top panel-pAb A and mAbs A3-25 + A60-22) and mutant toxin B (bottom panel-pAb B).

30 Figure 9: Images of IMR-90 cell morphology at 72 hours post treatment. Panel A shows mock treated control cells. Panel B shows cell morphology following treatment with formalin inactivated mutant TcdB (SEQ ID NO: 6). Panel C shows cell morphology

following treatment with EDC inactivated mutant TcdB (SEQ ID NO: 6). Panel D shows cell morphology following treatment with wild-type toxin B (SEQ ID NO: 2). Panel E shows cell morphology following treatment with triple mutant TcdB (SEQ ID NO: 6). Similar results were observed for TcdA treatments.

5 Figure 10: Graph showing neutralizing antibody titers as described in Example 25 (study muCdiff2010-06).

Figure 11A-B: Graph showing neutralizing antibody titers as described in Example 26 (study muCdiff2010-07).

10 Figure 12: Graph showing neutralizing antibody responses against toxins A and B in hamsters after four immunizations as described in Example 27 (study hamC. difficile2010-02)

Figure 13A-B: Graph showing neutralizing antibody responses in hamsters after vaccination with chemically inactivated genetic mutant toxins and List Biological toxoids, as described in Example 27 (study hamC. difficile2010-02).

15 Figure 14: Survival curves for three immunized groups of hamsters as compared to the non-immunized controls, described in Example 28 (study hamC. difficile2010-02, continued).

Figure 15: Graph showing relative neutralizing antibody response against different formulations of *C. difficile* mutant toxins in hamsters (study hamC. difficile2010-03), as described in Example 29.

Figure 16A-B: Graphs showing strong relative neutralizing antibody response against chemically inactivated genetic mutant toxins A and B (SEQ ID NOs: 4 and 6, respectively) in cynomolgus macaques, as described in Example 30.

25 Figure 17: Amino acid sequences of variable regions of light (VL) and heavy (HL) chains of A3-25 mAb IgE. Signal peptide – highlighted; CDRs – italicized and underlined; Constant region – bolded and underlined (complete sequence not shown).

Figure 18: Graph showing titration of individual toxin A monoclonal antibodies in the toxin neutralization assay using ATP levels (quantified by relative light units- RLU) as an indicator of cell viability. In comparison to the toxin (4xEC₅₀) control, mAbs A80-30, A29, A65-33, A60-22 and A3-25 had increasing neutralizing effects on toxin A with concentration but not to the level of the positive rabbit anti-toxin A control. mAbs A50-10, A56-33, and A58-46 did not neutralize toxin A. The cell only control was 1 -1.5x10⁶ RLU.

Figure 19: Mapping of 8 epitope groups of toxin B mAbs by BiaCore

Figure 20A-C: Synergistic neutralizing activities of combinations of toxin A mAbs:
Adding different dilutions of neutralizing antibodies A60-22, A65-33, and A80-29 to
increasing concentrations of A3-25 mAb synergistically increased the neutralization of
5 toxin A regardless of the dilution. The RLUs of the toxin A only (4x EC₅₀) control is
illustrated (<0.3x10⁶) and cell only controls were 2-2.5 x10⁶ RLUs as depicted in graphs
shown in Figure 20B and Figure 20C.

10 Figure 21: Synergistic neutralizing activities of toxin B mAbs: Neutralization of
toxin B by mAbs 8-26, B60-2 and B59-3 is illustrated in Figure 21A. Neutralization of
15 toxin B is synergistically increased after combining B8-26 with dilutions of B59-3 (Figure
21B)

Figure 22: Western blot showing that Rac1 GTPase expression is reduced in
genetic mutant toxin B (SEQ ID NO: 6) extracts from 24 to 96 hours, but not in wild-type
toxin B (SEQ ID NO: 2) treated extracts. The blot also shows that Rac1 is glucosylated
15 in toxin B-treated extracts, but not in genetic mutant toxin B treated extracts.

Figure 23A-K: Graph representing results from *in vitro* cytotoxicity tests in which
the ATP levels (RLUs) are plotted against increasing concentrations of *C. difficile*
culture media and the hamster serum pool (■); crude toxin (culture harvest) from the
respective strain and the hamster serum pool (●); purified toxin (commercial toxin
20 obtained from List Biologicals) and the hamster serum pool (▲); crude toxin (▼),
control; and purified toxin (◆), control. The toxins from the respective strains were
added to the cells at 4xEC₅₀ values. Figure 23 shows that an immunogenic composition
including mutant TcdA (SEQ ID NO: 4) and mutant TcdB (SEQ ID NO: 6), wherein the
mutant toxins were inactivated with EDC, according to, for example, Example 29, Table
25 15, described herein, induced neutralizing antibodies that exhibited neutralizing activity
against toxins from at least the following 16 different CDC strains of *C. difficile*, in
comparison to the respective toxin only control: 2007886 (Figure 23A); 2006017 (Figure
23B); 2007070 (Figure 23C); 2007302 (Figure 23D); 2007838 (Figure 23E); 2007886
30 (Figure 23F); 2009292 (Figure 23G); 2004013 (Figure 23H); 2009141 (Figure 23I);
2005022 (Figure 23J); 2006376 (Figure 23K).

Figure 24: Illustration of an exemplary EDC/NHS inactivation of mutant *C. difficile*
toxins, resulting in at least three possible types of modifications: crosslinks, glycine
adducts, and beta-alanine adducts. Panel A illustrates crosslinking. Carboxylic

residues of triple mutant toxins are activated by the addition of EDC and NHS. The activated esters react with primary amines to form stable amide bonds, resulting in intra- and intermolecular crosslinks. Panel B illustrates formation of glycine adducts. After inactivation, residual activated esters are quenched by the addition of glycine to form 5 stable amide bonds. Panel C illustrates formation of beta-alanine adducts. Three moles of NHS can react with one mole of EDC to form activated beta-alanine. This then reacts with primary amines to form stable amide bonds.

Figure 25: Illustration of an exemplary EDC/NHS inactivation of mutant *C. difficile* toxins, resulting in at least one of the following types of modifications: (A) crosslinks, (B) 10 glycine adducts, and (C) beta-alanine adducts.

Figure 26: Graph representing results from an *in vitro* cytotoxicity assay in which the ATP levels (RLUs) (72 hr ATP) are plotted against increasing concentrations of the wild-type TcdB, commercially obtained from List Biologicals (□), triple mutant TcdB (SEQ ID NO: 86) (●), and penta mutant TcdB (SEQ ID NO: 184) (■). IMR-90 cells 15 (*) were used as control.

Figure 27: Graph showing competitive inhibition of triple mutant toxin B (SEQ ID NO: 86)-mediated cytotoxicity by penta mutant toxin B (SEQ ID NO: 184) on IMR -90 cells, (72 hr ATP assay). —●— represents penta mutant toxin B (SEQ ID NO: 184) ("PM-B"); —▲— represents triple mutant (TM) at 200 ng/mL.

Figure 28: Graph showing final OD and triple mutant toxin B titer (mg/l) following a perfusion fermentation (CDF-5126). —●— represents OD_{600 nm}; —▲— represents 20 perfusion flow rate (Fermentor volumes/2.0h); —■— represents glucose (g/L); —●— represents toxoid B (triple mutant, SEQ ID NO: 86)

Figure 29: Graph showing final OD and triple mutant toxin B titer (mg/l) results 25 from another perfusion culture (CDF-5127). —●— represents OD_{600 nm}; —▲— represents perfusion flow rate (Fermentor volumes/2.0h); —■— represents glucose (g/L); —●— represents toxoid B (triple mutant, SEQ ID NO: 86).

BRIEF DESCRIPTION OF SEQUENCES

SEQ ID NO: 1 sets forth the amino acid sequence for wild-type *C. difficile* 630 toxin A (TcdA).

SEQ ID NO: 2 sets forth the amino acid sequence for wild-type *C. difficile* 630 toxin B (TcdB).

SEQ ID NO: 3 sets forth the amino acid sequence for a mutant TcdA having a mutation at positions 285 and 287, as compared to SEQ ID NO: 1.

SEQ ID NO: 4 sets forth the amino acid sequence for a mutant TcdA having a mutation at positions 285, 287, and 700, as compared to SEQ ID NO: 1.

10 SEQ ID NO: 5 sets forth the amino acid sequence for a mutant TcdB having a mutation at positions 286 and 288, as compared to SEQ ID NO: 2.

SEQ ID NO: 6 sets forth the amino acid sequence for a mutant TcdB having a mutation at positions 286, 288, and 698, as compared to SEQ ID NO: 2.

15 SEQ ID NO: 7 sets forth the amino acid sequence for a mutant TcdA having a mutation at positions 269, 272, 285, 287, 460, 462, and 700, as compared to SEQ ID NO: 1

SEQ ID NO: 8 sets forth the amino acid sequence for a mutant TcdB having a mutation at positions 270, 273, 286, 288, 461, 463, and 698, as compared to SEQ ID NO: 2

SEQ ID NO: 9 sets forth a DNA sequence encoding a wild-type *C. difficile* 630 toxin A (TcdA).

20 SEQ ID NO: 10 sets forth a DNA sequence encoding a wild-type *C. difficile* 630 toxin B (TcdB).

SEQ ID NO: 11 sets forth a DNA sequence encoding SEQ ID NO: 3

SEQ ID NO: 12 sets forth a DNA sequence encoding SEQ ID NO: 4

SEQ ID NO: 13 sets forth a DNA sequence encoding SEQ ID NO: 5

25 SEQ ID NO: 14 sets forth a DNA sequence encoding SEQ ID NO: 6

SEQ ID NO: 15 sets forth the amino acid sequence for wild-type *C. difficile* R20291 TcdA.

SEQ ID NO: 16 sets forth a DNA sequence encoding SEQ ID NO: 15.

SEQ ID NO: 17 sets forth the amino acid sequence for wild-type *C. difficile* CD196

30 TcdA.

SEQ ID NO: 18 sets forth a DNA sequence encoding SEQ ID NO: 17.

SEQ ID NO: 19 sets forth the amino acid sequence for wild-type *C. difficile* VPI10463 TcdA.

SEQ ID NO: 20 sets forth a DNA sequence encoding SEQ ID NO: 19.

SEQ ID NO: 21 sets forth the amino acid sequence for wild-type *C. difficile* R20291 TcdB.

SEQ ID NO: 22 sets forth a DNA sequence encoding SEQ ID NO: 21.

5 SEQ ID NO: 23 sets forth the amino acid sequence for wild-type *C. difficile* CD196 TcdB.

SEQ ID NO: 24 sets forth a DNA sequence encoding SEQ ID NO: 23.

SEQ ID NO: 25 sets forth the amino acid sequence for wild-type *C. difficile* VPI10463 TcdB.

10 SEQ ID NO: 26 sets forth a DNA sequence encoding SEQ ID NO: 25.

SEQ ID NO: 27 sets forth a DNA sequence of a pathogenicity locus of wild-type *C. difficile* VPI10463.

SEQ ID NO: 28 sets forth the amino acid sequence for residues 101 to 293 of SEQ ID NO: 1.

15 SEQ ID NO: 29 sets forth the amino acid sequence for residues 1 to 542 of SEQ ID NO: 1.

SEQ ID NO: 30 sets forth the amino acid sequence for residues 101 to 293 of SEQ ID NO: 2.

20 SEQ ID NO: 31 sets forth the amino acid sequence for residues 1 to 543 of SEQ ID NO: 2.

SEQ ID NO: 32 sets forth the amino acid sequence for residues 543 to 809 of SEQ ID NO: 1.

SEQ ID NO: 33 sets forth the amino acid sequence for residues 544 to 767 of SEQ ID NO: 2.

25 SEQ ID NO: 34 sets forth the amino acid sequence for a mutant TcdA, wherein residues 101, 269, 272, 285, 287, 460, 462, 541, 542, 543, 589, 655, and 700 may be any amino acid.

SEQ ID NO: 35 sets forth the amino acid sequence for a mutant TcdB, wherein 102, 270, 273, 286, 288, 384, 461, 463, 520, 543, 544, 587, 600, 653, 698, and 751 may be 30 any amino acid.

SEQ ID NO: 36 sets forth the amino acid sequence for the variable light chain of a neutralizing antibody of *C. difficile* TcdA (A3-25 mAb).

SEQ ID NO: 37 sets forth the amino acid sequence for the variable heavy chain of a neutralizing antibody of *C. difficile* TcdA (A3-25 mAb).

SEQ ID NO: 38 sets forth the amino acid sequence for CDR1 of the variable light chain of neutralizing antibody of *C. difficile* TcdA (A3-25 mAb).

5 SEQ ID NO: 39 sets forth the amino acid sequence for CDR2 of the variable light chain of neutralizing antibody of *C. difficile* TcdA (A3-25 mAb).

SEQ ID NO: 40 sets forth the amino acid sequence for CDR3 of the variable light chain of neutralizing antibody of *C. difficile* TcdA (A3-25 mAb).

10 SEQ ID NO: 41 sets forth the amino acid sequence for CDR1 of the variable heavy chain of neutralizing antibody of *C. difficile* TcdA (A3-25 mAb).

SEQ ID NO: 42 sets forth the amino acid sequence for CDR2 of the variable heavy chain of neutralizing antibody of *C. difficile* TcdA (A3-25 mAb).

SEQ ID NO: 43 sets forth the amino acid sequence for CDR3 of the variable heavy chain of neutralizing antibody of *C. difficile* TcdA (A3-25 mAb).

15 SEQ ID NO: 44 sets forth a DNA sequence encoding SEQ ID NO: 3.

SEQ ID NO: 45 sets forth a DNA sequence encoding SEQ ID NO: 4.

SEQ ID NO: 46 sets forth a DNA sequence encoding SEQ ID NO: 5.

SEQ ID NO: 47 sets forth a DNA sequence encoding SEQ ID NO: 6.

20 SEQ ID NO: 48 sets forth the nucleotide sequence of immunostimulatory oligonucleotide ODN CpG 24555.

SEQ ID NO: 49 sets forth the amino acid sequence for the variable heavy chain of a *C. difficile* TcdB neutralizing antibody (B8-26 mAb).

SEQ ID NO: 50 sets forth the amino acid sequence for the signal peptide of the variable heavy chain of a *C. difficile* TcdB neutralizing antibody (B8-26 mAb).

25 SEQ ID NO: 51 sets forth the amino acid sequence for CDR1 of the variable heavy chain of a *C. difficile* TcdB neutralizing antibody (B8-26 mAb).

SEQ ID NO: 52 sets forth the amino acid sequence for CDR2 of the variable heavy chain of a *C. difficile* TcdB neutralizing antibody (B8-26 mAb).

30 SEQ ID NO: 53 sets forth the amino acid sequence for CDR3 of the variable heavy chain of a *C. difficile* TcdB neutralizing antibody (B8-26 mAb).

SEQ ID NO: 54 sets forth the amino acid sequence for the constant region of the variable heavy chain of a *C. difficile* TcdB neutralizing antibody (B8-26 mAb).

SEQ ID NO: 55 sets forth the amino acid sequence for the variable light chain of a *C. difficile* TcdB neutralizing antibody (B8-26 mAb).

SEQ ID NO: 56 sets forth the amino acid sequence for the signal peptide of the variable light chain of a *C. difficile* TcdB neutralizing antibody (B8-26 mAb).

5 SEQ ID NO: 57 sets forth the amino acid sequence for CDR1 of the variable light chain of a *C. difficile* TcdB neutralizing antibody (B8-26 mAb).

SEQ ID NO: 58 sets forth the amino acid sequence for CDR2 of the variable light chain of a *C. difficile* TcdB neutralizing antibody (B8-26 mAb).

10 SEQ ID NO: 59 sets forth the amino acid sequence for CDR3 of the variable light chain of a *C. difficile* TcdB neutralizing antibody (B8-26 mAb).

SEQ ID NO: 60 sets forth the amino acid sequence for the variable heavy chain of a *C. difficile* TcdB neutralizing antibody (B59-3 mAb).

SEQ ID NO: 61 sets forth the amino acid sequence for the signal peptide of the variable heavy chain of a *C. difficile* TcdB neutralizing antibody (B59-3 mAb).

15 SEQ ID NO: 62 sets forth the amino acid sequence for CDR1 of the variable heavy chain of a *C. difficile* TcdB neutralizing antibody (B59-3 mAb).

SEQ ID NO: 63 sets forth the amino acid sequence for CDR2 of the variable heavy chain of a *C. difficile* TcdB neutralizing antibody (B59-3 mAb).

20 SEQ ID NO: 64 sets forth the amino acid sequence for CDR3 of the variable heavy chain of a *C. difficile* TcdB neutralizing antibody (B59-3 mAb).

SEQ ID NO: 65 sets forth the amino acid sequence for the constant region of the variable heavy chain of a *C. difficile* TcdB neutralizing antibody (B59-3 mAb).

SEQ ID NO: 66 sets forth the amino acid sequence for the variable light chain of a *C. difficile* TcdB neutralizing antibody (B59-3 mAb).

25 SEQ ID NO: 67 sets forth the amino acid sequence for the signal peptide of the variable light chain of a *C. difficile* TcdB neutralizing antibody (B59-3 mAb).

SEQ ID NO: 68 sets forth the amino acid sequence for CDR1 of the variable light chain of a *C. difficile* TcdB neutralizing antibody (B59-3 mAb).

30 SEQ ID NO: 69 sets forth the amino acid sequence for CDR2 of the variable light chain of a *C. difficile* TcdB neutralizing antibody (B59-3 mAb).

SEQ ID NO: 70 sets forth the amino acid sequence for CDR3 of the variable light chain of a *C. difficile* TcdB neutralizing antibody (B59-3 mAb).

SEQ ID NO: 71 sets forth the amino acid sequence for the variable heavy chain of a *C. difficile* TcdB neutralizing antibody (B9-30 mAb).

SEQ ID NO: 72 sets forth the amino acid sequence for the signal peptide of the variable heavy chain of a *C. difficile* TcdB neutralizing antibody (B9-30 mAb).

5 SEQ ID NO: 73 sets forth the amino acid sequence for CDR1 of the variable heavy chain of a *C. difficile* TcdB neutralizing antibody (B9-30 mAb).

SEQ ID NO: 74 sets forth the amino acid sequence for CDR2 of the variable heavy chain of a *C. difficile* TcdB neutralizing antibody (B9-30 mAb).

10 SEQ ID NO: 75 sets forth the amino acid sequence for CDR3 of the variable heavy chain of a *C. difficile* TcdB neutralizing antibody (B9-30 mAb).

SEQ ID NO: 76 sets forth the amino acid sequence for the constant region of the variable heavy chain of a *C. difficile* TcdB neutralizing antibody (B9-30 mAb).

SEQ ID NO: 77 sets forth the amino acid sequence for the variable light chain of a *C. difficile* TcdB neutralizing antibody (B9-30 mAb).

15 SEQ ID NO: 78 sets forth the amino acid sequence for the signal peptide of the variable light chain of a *C. difficile* TcdB neutralizing antibody (B9-30 mAb).

SEQ ID NO: 79 sets forth the amino acid sequence for CDR1 of the variable light chain of a *C. difficile* TcdB neutralizing antibody (B9-30 mAb).

20 SEQ ID NO: 80 sets forth the amino acid sequence for CDR2 of the variable light chain of a *C. difficile* TcdB neutralizing antibody (B9-30 mAb).

SEQ ID NO: 81 sets forth the amino acid sequence for CDR3 of the variable light chain of a *C. difficile* TcdB neutralizing antibody (B9-30 mAb).

25 SEQ ID NO: 82 sets forth the amino acid sequence for a mutant TcdB, wherein a residue at positions 102, 270, 273, 286, 288, 384, 461, 463, 520, 543, 544, 587, 600, 653, 698, and 751 may be any amino acid.

SEQ ID NO: 83 sets forth the amino acid sequence for a mutant TcdA having a mutation at positions 269, 272, 285, 287, 460, 462, and 700, as compared to SEQ ID NO: 1, wherein the methionine at position 1 is absent.

30 SEQ ID NO: 84 sets forth the amino acid sequence for a mutant *C. difficile* toxin A having a mutation at positions 285, 287, and 700, as compared to SEQ ID NO: 1, wherein the methionine at position 1 is absent.

SEQ ID NO: 85 sets forth the amino acid sequence for a mutant *C. difficile* toxin B having a mutation at positions 270, 273, 286, 288, 461, 463, and 698, as compared to SEQ ID NO: 2, wherein the methionine at position 1 is absent.

SEQ ID NO: 86 sets forth the amino acid sequence for a mutant *C. difficile* toxin B

5 having a mutation at positions 286, 288, and 698, as compared to SEQ ID NO: 2, wherein the methionine at position 1 is absent.

SEQ ID NO: 87 sets forth the amino acid sequence for wild-type *C. difficile* 2004013 TcdA.

SEQ ID NO: 88 sets forth the amino acid sequence for wild-type *C. difficile* 2004111

10 TcdA.

SEQ ID NO: 89 sets forth the amino acid sequence for wild-type *C. difficile* 2004118 TcdA.

SEQ ID NO: 90 sets forth the amino acid sequence for wild-type *C. difficile* 2004205 TcdA.

15 SEQ ID NO: 91 sets forth the amino acid sequence for wild-type *C. difficile* 2004206 TcdA.

SEQ ID NO: 92 sets forth the amino acid sequence for wild-type *C. difficile* 2005022 TcdA.

20 SEQ ID NO: 93 sets forth the amino acid sequence for wild-type *C. difficile* 2005088 TcdA.

SEQ ID NO: 94 sets forth the amino acid sequence for wild-type *C. difficile* 2005283 TcdA.

SEQ ID NO: 95 sets forth the amino acid sequence for wild-type *C. difficile* 2005325 TcdA.

25 SEQ ID NO: 96 sets forth the amino acid sequence for wild-type *C. difficile* 2005359 TcdA.

SEQ ID NO: 97 sets forth the amino acid sequence for wild-type *C. difficile* 2006017 TcdA.

30 SEQ ID NO: 98 sets forth the amino acid sequence for wild-type *C. difficile* 2007070 TcdA.

SEQ ID NO: 99 sets forth the amino acid sequence for wild-type *C. difficile* 2007217 TcdA.

SEQ ID NO: 100 sets forth the amino acid sequence for wild-type *C. difficile* 2007302 TcdA.

SEQ ID NO: 101 sets forth the amino acid sequence for wild-type *C. difficile* 2007816 TcdA.

5 SEQ ID NO: 102 sets forth the amino acid sequence for wild-type *C. difficile* 2007838 TcdA.

SEQ ID NO: 103 sets forth the amino acid sequence for wild-type *C. difficile* 2007858 TcdA.

10 SEQ ID NO: 104 sets forth the amino acid sequence for wild-type *C. difficile* 2007886 TcdA.

SEQ ID NO: 105 sets forth the amino acid sequence for wild-type *C. difficile* 2008222 TcdA.

SEQ ID NO: 106 sets forth the amino acid sequence for wild-type *C. difficile* 2009078 TcdA.

15 SEQ ID NO: 107 sets forth the amino acid sequence for wild-type *C. difficile* 2009087 TcdA.

SEQ ID NO: 108 sets forth the amino acid sequence for wild-type *C. difficile* 2009141 TcdA.

20 SEQ ID NO: 109 sets forth the amino acid sequence for wild-type *C. difficile* 2009292 TcdA.

SEQ ID NO: 110 sets forth the amino acid sequence for wild-type *C. difficile* 2004013 TcdB.

SEQ ID NO: 111 sets forth the amino acid sequence for wild-type *C. difficile* 2004111 TcdB.

25 SEQ ID NO: 112 sets forth the amino acid sequence for wild-type *C. difficile* 2004118 TcdB.

SEQ ID NO: 113 sets forth the amino acid sequence for wild-type *C. difficile* 2004205 TcdB.

30 SEQ ID NO: 114 sets forth the amino acid sequence for wild-type *C. difficile* 2004206 TcdB.

SEQ ID NO: 115 sets forth the amino acid sequence for wild-type *C. difficile* 2005022 TcdB.

SEQ ID NO: 116 sets forth the amino acid sequence for wild-type *C. difficile* 2005088 TcdB.

SEQ ID NO: 117 sets forth the amino acid sequence for wild-type *C. difficile* 2005283 TcdB.

5 SEQ ID NO: 118 sets forth the amino acid sequence for wild-type *C. difficile* 2005325 TcdB.

SEQ ID NO: 119 sets forth the amino acid sequence for wild-type *C. difficile* 2005359 TcdB.

10 SEQ ID NO: 120 sets forth the amino acid sequence for wild-type *C. difficile* 2006017 TcdB.

SEQ ID NO: 121 sets forth the amino acid sequence for wild-type *C. difficile* 2006376 TcdB.

SEQ ID NO: 122 sets forth the amino acid sequence for wild-type *C. difficile* 2007070 TcdB.

15 SEQ ID NO: 123 sets forth the amino acid sequence for wild-type *C. difficile* 2007217 TcdB.

SEQ ID NO: 124 sets forth the amino acid sequence for wild-type *C. difficile* 2007302 TcdB.

20 SEQ ID NO: 125 sets forth the amino acid sequence for wild-type *C. difficile* 2007816 TcdB.

SEQ ID NO: 126 sets forth the amino acid sequence for wild-type *C. difficile* 2007838 TcdB.

SEQ ID NO: 127 sets forth the amino acid sequence for wild-type *C. difficile* 2007858 TcdB.

25 SEQ ID NO: 128 sets forth the amino acid sequence for wild-type *C. difficile* 2007886 TcdB.

SEQ ID NO: 129 sets forth the amino acid sequence for wild-type *C. difficile* 2008222 TcdB.

30 SEQ ID NO: 130 sets forth the amino acid sequence for wild-type *C. difficile* 2009078 TcdB.

SEQ ID NO: 131 sets forth the amino acid sequence for wild-type *C. difficile* 2009087 TcdB.

SEQ ID NO: 132 sets forth the amino acid sequence for wild-type *C. difficile* 2009141 TcdB.

SEQ ID NO: 133 sets forth the amino acid sequence for wild-type *C. difficile* 2009292 TcdB.

- 5 SEQ ID NO: 134 sets forth the amino acid sequence for wild-type *C. difficile* 014 TcdA.
- SEQ ID NO: 135 sets forth the amino acid sequence for wild-type *C. difficile* 015 TcdA.
- SEQ ID NO: 136 sets forth the amino acid sequence for wild-type *C. difficile* 020 TcdA.
- SEQ ID NO: 137 sets forth the amino acid sequence for wild-type *C. difficile* 023 TcdA.
- SEQ ID NO: 138 sets forth the amino acid sequence for wild-type *C. difficile* 027 TcdA.
- 10 SEQ ID NO: 139 sets forth the amino acid sequence for wild-type *C. difficile* 029 TcdA.
- SEQ ID NO: 140 sets forth the amino acid sequence for wild-type *C. difficile* 046 TcdA.
- SEQ ID NO: 141 sets forth the amino acid sequence for wild-type *C. difficile* 014 TcdB.
- SEQ ID NO: 142 sets forth the amino acid sequence for wild-type *C. difficile* 015 TcdB.
- SEQ ID NO: 143 sets forth the amino acid sequence for wild-type *C. difficile* 020 TcdB.
- 15 SEQ ID NO: 144 sets forth the amino acid sequence for wild-type *C. difficile* 023 TcdB.
- SEQ ID NO: 145 sets forth the amino acid sequence for wild-type *C. difficile* 027 TcdB.
- SEQ ID NO: 146 sets forth the amino acid sequence for wild-type *C. difficile* 029 TcdB.
- SEQ ID NO: 147 sets forth the amino acid sequence for wild-type *C. difficile* 046 TcdB.
- SEQ ID NO: 148 sets forth the amino acid sequence for wild-type *C. difficile* 001 TcdA.
- 20 SEQ ID NO: 149 sets forth the amino acid sequence for wild-type *C. difficile* 002 TcdA.
- SEQ ID NO: 150 sets forth the amino acid sequence for wild-type *C. difficile* 003 TcdA.
- SEQ ID NO: 151 sets forth the amino acid sequence for wild-type *C. difficile* 004 TcdA.
- SEQ ID NO: 152 sets forth the amino acid sequence for wild-type *C. difficile* 070 TcdA.
- SEQ ID NO: 153 sets forth the amino acid sequence for wild-type *C. difficile* 075 TcdA.
- 25 SEQ ID NO: 154 sets forth the amino acid sequence for wild-type *C. difficile* 077 TcdA.
- SEQ ID NO: 155 sets forth the amino acid sequence for wild-type *C. difficile* 081 TcdA.
- SEQ ID NO: 156 sets forth the amino acid sequence for wild-type *C. difficile* 117 TcdA.
- SEQ ID NO: 157 sets forth the amino acid sequence for wild-type *C. difficile* 131 TcdA.
- SEQ ID NO: 158 sets forth the amino acid sequence for wild-type *C. difficile* 001 TcdB.
- 30 SEQ ID NO: 159 sets forth the amino acid sequence for wild-type *C. difficile* 002 TcdB.
- SEQ ID NO: 160 sets forth the amino acid sequence for wild-type *C. difficile* 003 TcdB.
- SEQ ID NO: 161 sets forth the amino acid sequence for wild-type *C. difficile* 004 TcdB.
- SEQ ID NO: 162 sets forth the amino acid sequence for wild-type *C. difficile* 070 TcdB.

SEQ ID NO: 163 sets forth the amino acid sequence for wild-type *C. difficile* 075 TcdB.
SEQ ID NO: 164 sets forth the amino acid sequence for wild-type *C. difficile* 077 TcdB.
SEQ ID NO: 165 sets forth the amino acid sequence for wild-type *C. difficile* 081 TcdB.
SEQ ID NO: 166 sets forth the amino acid sequence for wild-type *C. difficile* 117 TcdB.
5 SEQ ID NO: 167 sets forth the amino acid sequence for wild-type *C. difficile* 131 TcdB.
SEQ ID NO: 168 sets forth the amino acid sequence for wild-type *C. difficile* 053 TcdA.
SEQ ID NO: 169 sets forth the amino acid sequence for wild-type *C. difficile* 078 TcdA.
SEQ ID NO: 170 sets forth the amino acid sequence for wild-type *C. difficile* 087 TcdA.
SEQ ID NO: 171 sets forth the amino acid sequence for wild-type *C. difficile* 095 TcdA.
10 SEQ ID NO: 172 sets forth the amino acid sequence for wild-type *C. difficile* 126 TcdA.
SEQ ID NO: 173 sets forth the amino acid sequence for wild-type *C. difficile* 053 TcdB.
SEQ ID NO: 174 sets forth the amino acid sequence for wild-type *C. difficile* 078 TcdB.
SEQ ID NO: 175 sets forth the amino acid sequence for wild-type *C. difficile* 087 TcdB.
SEQ ID NO: 176 sets forth the amino acid sequence for wild-type *C. difficile* 095 TcdB.
15 SEQ ID NO: 177 sets forth the amino acid sequence for wild-type *C. difficile* 126 TcdB.
SEQ ID NO: 178 sets forth the amino acid sequence for wild-type *C. difficile* 059 TcdA.
SEQ ID NO: 179 sets forth the amino acid sequence for wild-type *C. difficile* 059 TcdB.
SEQ ID NO: 180 sets forth the amino acid sequence for wild-type *C. difficile* 106 TcdA.
SEQ ID NO: 181 sets forth the amino acid sequence for wild-type *C. difficile* 106 TcdB.
20 SEQ ID NO: 182 sets forth the amino acid sequence for wild-type *C. difficile* 017 TcdB.
SEQ ID NO: 183 sets forth the amino acid sequence for a mutant TcdA having a
mutation at positions 285, 287, 700, 972, and 978 as compared to SEQ ID NO: 1.
SEQ ID NO: 184 sets forth the amino acid sequence for a mutant TcdB having a
mutation at positions 286, 288, 698, 970, and 976 as compared to SEQ ID NO: 2.
25 SEQ ID NO: 185 through SEQ ID NO: 195 each set forth the amino acid sequence for
an exemplary mutant toxin.
SEQ ID NO: 196 through SEQ ID NO: 212 each set forth the amino acid sequence for
an exemplary mutant toxin A.
SEQ ID NO: 213 through SEQ ID NO: 222 each set forth the amino acid sequence for
30 an exemplary mutant toxin B.
SEQ ID NO: 223 through SEQ ID NO: 236 each set forth the amino acid sequence for
an exemplary mutant toxin A.

SEQ ID NO: 237 through SEQ ID NO: 243 each set forth the amino acid sequence for an exemplary mutant toxin B.

SEQ ID NO: 244 through SEQ ID NO: 245 each set forth the amino acid sequence for an exemplary mutant toxin A.

5 SEQ ID NO: 246 through SEQ ID NO: 249 each set forth the amino acid sequence for an exemplary mutant toxin B.

SEQ ID NO: 250 through SEQ ID NO: 253 each set forth the amino acid sequence for an exemplary mutant toxin A.

SEQ ID NO: 254 sets forth the amino acid sequence for an exemplary mutant toxin.

10 SEQ ID NO: 255 through SEQ ID NO: 263 each set forth the amino acid sequence for an exemplary mutant toxin A.

SEQ ID NO: 264 through SEQ ID NO: 269 each set forth the amino acid sequence for an exemplary mutant toxin B.

15 SEQ ID NO: 270 through SEQ ID NO: 275 each set forth the amino acid sequence for an exemplary mutant toxin.

SEQ ID NO: 276 through SEQ ID NO: 323 each set forth the amino acid sequence for an exemplary mutant toxin A.

SEQ ID NO: 324 through SEQ ID NO: 373 each set forth the amino acid sequence for an exemplary mutant toxin B.

20 SEQ ID NO: 374 through SEQ ID NO: 421 each set forth the amino acid sequence for an exemplary mutant toxin A.

SEQ ID NO: 422 through SEQ ID NO: 471 each set forth the amino acid sequence for an exemplary mutant toxin B.

25 SEQ ID NO: 472 through SEQ ID NO: 519 each set forth the amino acid sequence for an exemplary mutant toxin A.

SEQ ID NO: 568 through SEQ ID NO: 615 each set forth the amino acid sequence for an exemplary mutant toxin B.

SEQ ID NO: 520 through SEQ ID NO: 567 each set forth the amino acid sequence for an exemplary mutant toxin A.

30 SEQ ID NO: 616 through SEQ ID NO: 663 each set forth the amino acid sequence for an exemplary mutant toxin B.

SEQ ID NO: 664 through SEQ ID NO: 711 each set forth the amino acid sequence for an exemplary mutant toxin A.

SEQ ID NO: 712 through SEQ ID NO: 761 each set forth the amino acid sequence for an exemplary mutant toxin B.

DETAILED DESCRIPTION

The inventors surprisingly discovered, among other things, a mutant *C. difficile* toxin A and toxin B, and methods thereof. The mutants are characterized, in part, by being immunogenic and exhibiting reduced cytotoxicity compared to a wild-type form of the respective toxin. The present invention also relates to immunogenic portions thereof, biological equivalents thereof, and isolated polynucleotides that include nucleic acid sequences encoding any of the foregoing.

The immunogenic compositions described herein unexpectedly demonstrated the ability to elicit novel neutralizing antibodies against *C. difficile* toxins and they may have the ability to confer active and/or passive protection against a *C. difficile* challenge. The novel antibodies are directed against various epitopes of toxin A and toxin B. The inventors further discovered that a combination of at least two of the neutralizing monoclonal antibodies can exhibit an unexpectedly synergistic effect in respective *in vitro* neutralization of toxin A and toxin B.

The inventive compositions described herein may be used to treat, prevent, decrease the risk of, decrease occurrences of, decrease severity of, and/or delay the onset of a *C. difficile* infection, *C. difficile* associated disease (CDAD), syndrome, condition, symptom, and/or complication thereof in a mammal, as compared to a mammal to which the composition was not administered.

Moreover, the inventors discovered a recombinant asporogenic *C. difficile* cell that can stably express the mutant *C. difficile* toxin A and toxin B, and novel methods for producing the same.

25 Immunogenic Compositions

In one aspect, the invention relates to an immunogenic composition that includes a mutant *C. difficile* toxin. The mutant *C. difficile* toxin includes an amino acid sequence having at least one mutation in a glucosyltransferase domain and at least one mutation in a cysteine protease domain, relative to the corresponding wild-type *C. difficile* toxin.

The term "wild-type," as used herein, refers to the form found in nature. For example, a wild-type polypeptide or polynucleotide sequence is a sequence present in an organism that can be isolated from a source in nature and which has not been intentionally modified by human manipulation. The present invention also relates to

isolated polynucleotides that include nucleic acid sequences encoding any of the foregoing. In addition, the present invention relates to use of any of the foregoing compositions to treat, prevent, decrease the risk of, decrease severity of, decrease occurrences of, and/or delay the onset of a *C. difficile* infection, *C. difficile* associated 5 disease, syndrome, condition, symptom, and/or complication thereof in a mammal, as compared to a mammal to which the composition is not administered, as well as methods for preparing said compositions.

As used herein, an "immunogenic composition" or "immunogen" refers to a composition that elicits an immune response in a mammal to which the composition is 10 administered.

An "immune response" refers to the development of a beneficial humoral (antibody mediated) and/or a cellular (mediated by antigen-specific T cells or their secretion products) response directed against a *C. difficile* toxin in a recipient patient. The immune response may be humoral, cellular, or both.

15 The immune response can be an active response induced by administration of an immunogenic composition, an immunogen. Alternatively, the immune response can be a passive response induced by administration of antibody or primed T-cells.

The presence of a humoral (antibody-mediated) immune response can be determined, for example, by cell-based assays known in the art, such as a neutralizing 20 antibody assay, ELISA, etc.

A cellular immune response is typically elicited by the presentation of polypeptide epitopes in association with Class I or Class II MHC molecules to activate antigen-specific CD4 + T helper cells and/or CD8 + cytotoxic T cells. The response may also involve activation of monocytes, macrophages, NK cells, basophils, dendritic cells, 25 astrocytes, microglia cells, eosinophils or other components of innate immunity. The presence of a cell-mediated immunological response can be determined by proliferation assays (CD4 + T cells) or CTL (cytotoxic T lymphocyte) assays known in the art.

In one embodiment, an immunogenic composition is a vaccine composition. As used herein, a "vaccine composition" is a composition that elicits an immune response 30 in a mammal to which the composition is administered. The vaccine composition may protect the immunized mammal against subsequent challenge by an immunizing agent or an immunologically cross-reactive agent. Protection can be complete or partial with

regard to reduction in symptoms or infection as compared to a non-vaccinated mammal under the same conditions.

The immunogenic compositions described herein are cross-reactive, which refers to having a characteristic of being able to elicit an effective immune response (e.g., 5 humoral immune response) against a toxin produced by another *C. difficile* strain that is different from the strain from which the composition is derived. For example, the immunogenic compositions (e.g., derived from *C. difficile* 630) described herein may elicit cross-reactive antibodies that can bind to toxins produced by multiple strains of *C. difficile* (e.g., toxins produced by *C. difficile* R20291 and VPI10463). See, for example, 10 Example 37. Cross-reactivity is indicative of the cross-protection potential of the bacterial immunogen, and vice versa.

The term "cross-protective" as used herein refers to the ability of the immune response induced by an immunogenic composition to prevent or attenuate infection by a different bacterial strain or species of the same genus. For example, an immunogenic 15 composition (e.g., derived from *C. difficile* 630) described herein may induce an effective immune response in a mammal to attenuate a *C. difficile* infection and/or to attenuate a *C. difficile* disease caused by a strain other than 630 (e.g., *C. difficile* R20291) in the mammal.

Exemplary mammals in which the immunogenic composition or immunogen 20 elicits an immune response include any mammals, such as, for example, mice, hamsters, primates, and humans. In a preferred embodiment, the immunogenic composition or immunogen elicits an immune response in a human to which the composition is administered.

As described above, toxin A (TcdA) and toxin B (TcdB) are homologous 25 glucosyltransferases that inactivate small GTPases of the Rho/Rac/Ras family. The action of TcdA and TcdB on mammalian target cells depends on a multistep mechanism of receptor-mediated endocytosis, membrane translocation, autoproteolytic processing, and monoglucosylation of GTPases. Many of these functional activities have been ascribed to discrete regions within the primary sequence of the toxins, and the toxins 30 have been imaged to show that these molecules are similar in structure.

The wild-type gene for TcdA has about 8130 nucleotides that encode a protein having a deduced molecular weight of about 308-kDa, having about 2710 amino acids. As used herein, a wild-type *C. difficile* TcdA includes a *C. difficile* TcdA from any wild-

type *C. difficile* strain. A wild-type *C. difficile* TcdA may include a wild-type *C. difficile* TcdA amino acid sequence having at least about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, preferably about 98%, more preferably about 99% or most preferably about 100% identity to SEQ ID NO: 1 (full length) when optimally aligned, such as by the 5 programs GAP or BESTFIT using default gap weights.

In a preferred embodiment, the wild-type *C. difficile* TcdA includes an amino acid sequence set forth in SEQ ID NO: 1, which describes the wild-type amino acid sequence for TcdA from *C. difficile* strain 630 (also disclosed in GenBank accession number YP_001087137.1 and/or CAJ67494.1). *C. difficile* strain 630 is known in the 10 art as being a PCR-ribotype 012 strain. SEQ ID NO: 9 describes the wild-type gene for TcdA from *C. difficile* strain 630, which is also disclosed in GenBank accession number NC_009089.1.

Another example of a wild-type *C. difficile* TcdA includes an amino acid sequence set forth in SEQ ID NO: 15, which describes the wild-type amino acid 15 sequence for TcdA from *C. difficile* strain R20291 (also disclosed in GenBank accession number YP_003217088.1). *C. difficile* strain R20291 is known in the art as being a hypervirulent strain and a PCR-ribotype 027 strain. The amino acid sequence for TcdA from *C. difficile* strain R20291 has about 98% identity to SEQ ID NO:1. SEQ ID NO: 16 describes the wild-type gene for TcdA from *C. difficile* strain R20291, which is also 20 disclosed in GenBank accession number NC_013316.1.

An additional example of a wild-type *C. difficile* TcdA includes an amino acid sequence set forth in SEQ ID NO: 17, which describes the wild-type amino acid sequence for TcdA from *C. difficile* strain CD196 (also disclosed in GenBank accession number CBA61156.1). CD196 is a strain from a recent Canadian outbreak, and it is 25 known in the art as a PCR-ribotype 027 strain. The amino acid sequence for TcdA from *C. difficile* strain CD196 has about 98% identity to SEQ ID NO: 1, and has about 100% identity to TcdA from *C. difficile* strain R20291. SEQ ID NO: 18 describes the wild-type gene for TcdA from *C. difficile* strain CD196, which is also disclosed in GenBank accession number FN538970.1.

30 Further examples of an amino acid sequence for a wild-type *C. difficile* TcdA include SEQ ID NO: 19, which describes the wild-type amino acid sequence for TcdA from *C. difficile* strain VPI10463 (also disclosed in GenBank accession number CAA63564.1). The amino acid sequence for TcdA from *C. difficile* strain VPI10463 has

about 100% (99.8%) identity to SEQ ID NO: 1. SEQ ID NO: 20 describes the wild-type gene for TcdA from *C. difficile* strain VPI10463, which is also disclosed in GenBank accession number X92982.1.

Additional examples of a wild-type *C. difficile* TcdA include TcdA from wild-type 5 *C. difficile* strains obtainable from the Centers for Disease Control and Prevention (CDC, Atlanta, GA). The inventors discovered that the amino acid sequence of TcdA from wild-type *C. difficile* strains obtainable from the CDC include at least about 99.3% to 100% identity, when optimally aligned, to amino acid residues 1 to 821 of SEQ ID NO: 1 (TcdA from *C. difficile* 630). See Table 1.

10 The inventors also discovered that the amino acid sequence of TcdA from wild-type *C. difficile* strains may include at least about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, to about 100% identity, when optimally aligned (e.g., when full, length sequences are optimally aligned) to SEQ ID NO: 1.

15 Table 1: wild-type *C. difficile* strains obtained from CDC and the percent identity of amino acid residues 1-821 of TcdA from the respective wild-type *C. difficile* strain to amino acid residues 1-821 of SEQ ID NO: 1, when optimally aligned.

Table 1: Wild-type *C. difficile* Strains from CDC

<i>C. difficile</i> Strain ID	Approximate % Amino Acid Identity to Residues 1-821 of SEQ ID NO: 1
2004111	100
2004118	99.6
2004205	100
2004206	100
2005325	99.3
2005359	99.6
2006017	100
2007070	100
2007302	100
2007816	99.3
2007838	99.6
2007886	99.6
2008222	100
2009078	100
2009087	100
2009141	100
2009292	99.6

Accordingly, in one embodiment, the wild-type *C. difficile* TcdA amino acid sequence includes a sequence of at least about 500, 600, 700, or 800 contiguous residues, which has at least about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 5 preferably about 98%, more preferably about 99%, or most preferably about 100% identity to a sequence of equal length between residues 1 to 900 of SEQ ID NO: 1 when optimally aligned, such as by the programs GAP or BESTFIT using default gap weights. Examples include strains described above (e.g., R20291, CD196, etc) and those listed in Table 1.

10 In another embodiment, the wild-type *C. difficile* TcdA amino acid sequence includes a sequence having at least about 90%, 91%, 92%, 93%, 94%, 95%, 96%, preferably about 97%, preferably about 98%, more preferably about 99% or most

preferably about 100% identity to any sequence selected from SEQ ID NOs: 87-109 when optimally aligned. See Table 1-a.

Table 1-a: Wild-type <i>C. difficile</i> Strains	
<i>C. difficile</i> Strain ID	Toxin A, SEQ ID NO:
2004013	SEQ ID NO: 87
2004111	SEQ ID NO: 88
2004118	SEQ ID NO: 89
2004205	SEQ ID NO: 90
2004206	SEQ ID NO: 91
2005022	SEQ ID NO: 92
2005088	SEQ ID NO: 93
2005283	SEQ ID NO: 94
2005325	SEQ ID NO: 95
2005359	SEQ ID NO: 96
2006017	SEQ ID NO: 97
2006376	N/A
2007070	SEQ ID NO: 98
2007217	SEQ ID NO: 99
2007302	SEQ ID NO: 100
2007816	SEQ ID NO: 101
2007838	SEQ ID NO: 102
2007858	SEQ ID NO: 103
2007886	SEQ ID NO: 104
2008222	SEQ ID NO: 105
2009078	SEQ ID NO: 106
2009087	SEQ ID NO: 107
2009141	SEQ ID NO: 108
2009292	SEQ ID NO: 109
001	SEQ ID NO: 148
002	SEQ ID NO: 149
003	SEQ ID NO: 150

012 (004)	SEQ ID NO: 151
014	SEQ ID NO: 134
015	SEQ ID NO: 135
017	
020	SEQ ID NO: 136
023	SEQ ID NO: 137
027	SEQ ID NO: 138
029	SEQ ID NO: 139
046	SEQ ID NO: 140
053	SEQ ID NO: 168
059	SEQ ID NO: 178
070	SEQ ID NO: 152
075	SEQ ID NO: 153
077	SEQ ID NO: 154
078	SEQ ID NO: 169
081	SEQ ID NO: 155
087	SEQ ID NO: 170
095	SEQ ID NO: 171
106	SEQ ID NO: 180
117	SEQ ID NO: 156
126	SEQ ID NO: 172
131	SEQ ID NO: 157
SE844	SEQ ID NO: 196
12087	SEQ ID NO: 197
K14	SEQ ID NO: 198
BI6	SEQ ID NO: 199
BI17	SEQ ID NO: 200
CH6230	SEQ ID NO: 201
SE881	SEQ ID NO: 202

The wild-type gene for TcdB has about 7098 nucleotides that encode a protein with a deduced molecular weight of about 270 kDa, having about 2366 amino acids. As used herein, a wild-type *C. difficile* TcdB includes a *C. difficile* TcdB from any wild-type *C. difficile* strain. A wild-type *C. difficile* TcdB may include a wild-type amino acid sequence having at least about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, preferably about 98%, more preferably about 99% or most preferably about 100% identity to SEQ ID NO: 2 when optimally aligned, such as by the programs GAP or BESTFIT using default gap weights. In a preferred embodiment, the wild-type *C. difficile* TcdB includes an amino acid sequence set forth in SEQ ID NO: 2, which 5 describes the wild-type amino acid sequence for TcdB from *C. difficile* strain 630 (also disclosed in GenBank accession number YP_001087135.1 and/or CAJ67492). SEQ ID NO: 10 describes the wild-type gene for TcdB from *C. difficile* strain 630, which is also disclosed in GenBank accession number NC_009089.1.

Another example of a wild-type *C. difficile* TcdB includes an amino acid 15 sequence set forth in SEQ ID NO: 21, which describes the wild-type amino acid sequence for TcdB from *C. difficile* strain R20291 (also disclosed in GenBank accession number YP_003217086.1 and/or CBE02479.1). The amino acid sequence for TcdB from *C. difficile* strain R20291 has about 92% identity to SEQ ID NO: 2. SEQ ID NO: 22 describes the wild-type gene for TcdB from *C. difficile* strain R20291, which is also 20 disclosed in GenBank accession number NC_013316.1.

An additional example of a wild-type *C. difficile* TcdB includes an amino acid sequence set forth in SEQ ID NO: 23, which describes the wild-type amino acid sequence for TcdB from *C. difficile* strain CD196 (also disclosed in GenBank accession number YP_003213639.1 and/or CBA61153.1). SEQ ID NO: 24 describes the wild- 25 type gene for TcdB from *C. difficile* strain CD196, which is also disclosed in GenBank accession number NC_013315.1. The amino acid sequence for TcdB from *C. difficile* strain CD196 has about 92% identity to SEQ ID NO: 2.

Further examples of an amino acid sequence for a wild-type *C. difficile* TcdB include SEQ ID NO: 25, which describes the wild-type amino acid sequence for TcdB from *C. difficile* strain VPI10463 (also disclosed in GenBank accession number P18177 and/or CAA37298). The amino acid sequence for TcdB from *C. difficile* strain VPI10463 has 100% identity to SEQ ID NO: 2. SEQ ID NO: 26 describes the wild-type gene for

TcdB from *C. difficile* strain VPI10463, which is also disclosed in GenBank accession number X53138.1.

Additional examples of a wild-type *C. difficile* TcdB include TcdB from wild-type *C. difficile* strains obtainable from the Centers for Disease Control and Prevention (CDC, Atlanta, GA). The inventors discovered that the amino acid sequence of TcdB from wild-type *C. difficile* strains obtainable from the CDC include at least about 96% to 100% identity, when optimally aligned, to amino acid residue 1 to 821 of SEQ ID NO: 2 (TcdB from *C. difficile* 630). See Table 2.

Table 2: wild-type *C. difficile* strains obtained from CDC and the % identity of amino acid residues 1-821 of TcdB from the respective wild-type *C. difficile* strain to amino acid residues 1-821 of SEQ ID NO: 2, when optimally aligned.

Table 2: Wild-type <i>C. difficile</i> Strains from CDC	
<i>C. difficile</i> Strain ID	Approximate % Amino Acid Identity to Residues 1-821 of SEQ ID NO: 2
2004013	96.0
2004111	100
2004118	96.0
2004206	100
2005022	100
2005325	96.7
2007302	100
2007816	96.7
2008222	100
2009078	100
2009087	100
2009141	100

Accordingly, in one embodiment, a wild-type *C. difficile* TcdB amino acid sequence includes a sequence of at least about 500, 600, 700, or 800 contiguous residues, which has at least about 90%, 91%, 92%, 93%, 94%, 95%, 96%, preferably about 97%, preferably about 98%, more preferably about 99% or most preferably about 100% identity to a sequence of equal length between residues 1 to 900 of SEQ ID NO: 2 when optimally aligned, such as by the programs GAP or BESTFIT using default gap

weights. Examples include strains described above (e.g., R20291, CD196, etc) and those listed in Table 2.

In another embodiment, the wild-type *C. difficile* TcdB amino acid sequence includes a sequence having at least about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 5 preferably about 97%, preferably about 98%, more preferably about 99% or most preferably about 100% identity to any sequence selected from SEQ ID NOs: 110-133 when optimally aligned. See Table 2-a.

Table 2-a Wild-type *C. difficile* Strains

<i>C. difficile</i> Strain ID	Toxin B, SEQ ID NO:
2004013	SEQ ID NO: 110
2004111	SEQ ID NO: 111
2004118	SEQ ID NO: 112
2004205	SEQ ID NO: 113
2004206	SEQ ID NO: 114
2005022	SEQ ID NO: 115
2005088	SEQ ID NO: 116
2005283	SEQ ID NO: 117
2005325	SEQ ID NO: 118
2005359	SEQ ID NO: 119
2006017	SEQ ID NO: 120
2006376	SEQ ID NO: 121
2007070	SEQ ID NO: 122
2007217	SEQ ID NO: 123
2007302	SEQ ID NO: 124
2007816	SEQ ID NO: 125
2007838	SEQ ID NO: 126
2007858	SEQ ID NO: 127
2007886	SEQ ID NO: 128
2008222	SEQ ID NO: 129
2009078	SEQ ID NO: 130
2009087	SEQ ID NO: 131
2009141	SEQ ID NO: 132
2009292	SEQ ID NO: 133
001	SEQ ID NO: 158
002	SEQ ID NO: 159
003	SEQ ID NO: 160
012 (004)	SEQ ID NO: 161
014	SEQ ID NO: 141

015	SEQ ID NO: 142
017	SEQ ID NO: 182
020	SEQ ID NO: 143
023	SEQ ID NO: 144
027	SEQ ID NO: 145
029	SEQ ID NO: 146
046	SEQ ID NO: 147
053	SEQ ID NO: 173
059	SEQ ID NO: 179
070	SEQ ID NO: 162
075	SEQ ID NO: 163
077	SEQ ID NO: 164
078	SEQ ID NO: 174
081	SEQ ID NO: 165
087	SEQ ID NO: 175
095	SEQ ID NO: 176
106	SEQ ID NO: 181
117	SEQ ID NO: 166
126	SEQ ID NO: 177
131	SEQ ID NO: 167

The genes for toxins A and B (*tcdA* and *tcdB*) are part of a 19.6-kb genetic locus (the pathogenicity locus, PaLoc) that includes 3 additional small open-reading frames (ORFs), *tcdD*, *tcdE*, and *tcdC*, and may be considered useful for virulence. The PaLoc is known to be stable and conserved in toxigenic strains. It is present at the same 5 chromosomal integration site in all toxigenic strains that have been analyzed to date. In nontoxigenic strains, the pathogenicity locus (PaLoc) is not present. Accordingly, a characteristic of the wild-type *C. difficile* strains described herein is the presence of a pathogenicity locus. Another preferred characteristic of the wild-type *C. difficile* strains described herein is the production of both TcdA and TcdB.

10 In one embodiment, the wild-type *C. difficile* strain is a strain having a pathogenicity locus that is at least about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, preferably about 98%, more preferably about 99% or most preferably about 100% identical to that of *C. difficile* 630 or VPI10463. The total pathogenicity locus sequence of *C. difficile* VPI10463, is registered at the EMBL database with the sequence 15 accession number X92982, also shown in SEQ ID NO: 26. Strains in which the PaLoc is identical to that of the reference strain VPI10463 are referred to as toxinotype 0. Strains of toxinotypes I-VII, IX, XII-XV, and XVIII-XXIV produce both TcdA and TcdB despite variations in their toxin genes.

At the N-terminus of the toxins, the glucosyltransferase domain is located. The 20 glucosyltransferase activity of the toxins is associated with the cytotoxic function of the toxins. Without being bound by mechanism or theory, the glucosyltransferase activity in both toxins is believed to catalyze the monoglucosylation of small GTP-binding proteins in the Rho/Rac/Ras superfamily. After glucosylation of these GTP binding proteins, cellular physiology is modified dramatically, resulting in a loss of structural integrity and 25 disruption of essential signaling pathways of the host cells infected by the toxins. The Asp-Xaa-Asp (DXD) motif, which is involved with manganese, uridine diphosphate (UDP), and glucose binding, is a typical characteristic for the glucosyltransferase domain. Without being bound by mechanism or theory, it is believed that residues critical for catalytic activity, such as the DXD motif, do not vary between a TcdB from a 30 known "historical" strain, such as 630, and a TcdB from a hypervirulent strain, such as R20291. The DXD motif is located at residues 285 to 287 of a wild-type *C. difficile* TcdA, according to the numbering of SEQ ID NO: 1, and at residues 286 to 288 of a wild-type *C. difficile* TcdB, according to the numbering of SEQ ID NO: 2.

Global alignment algorithms (e.g., sequence analysis programs) are known in the art and may be used to optimally align two or more amino acid toxin sequences to determine if the toxin includes a particular signature motif (e.g., DXD in the glucosyltransferase domain, DHC in the cysteine protease domain described below, etc.). The optimally aligned sequence(s) are compared to a respective reference sequence (e.g., SEQ ID NO:1 for TcdA or SEQ ID NO: 2 for TcdB) to determine the existence of the signature motif. “Optimal alignment” refers to an alignment giving the highest percent identity score. Such alignment can be performed using known sequence analysis programs. In one embodiment, a CLUSTAL alignment (such as CLUSTALW) under default parameters is used to identify suitable wild-type toxins by comparing the query sequence against the reference sequence. The relative numbering of the conserved amino acid residues is based on the residue numbering of the reference amino acid sequence to account for small insertions or deletions (for example, five amino acids of less) within the aligned sequence.

As used herein, the term “according to the numbering of” refers to the numbering of the residues of a reference sequence when the given amino acid or polynucleotide sequence is compared to the reference sequence. In other words, the number or residue position of a given polymer is designated with respect to the reference sequence rather than by the actual numerical position of the residue within the given amino acid or polynucleotide sequence.

For example, a given amino acid sequence, such as that of a hypervirulent wild-type *C. difficile* strain, can be aligned to a reference sequence (e.g., such as that of a historical wild-type *C. difficile* strain, e.g., 630) by introducing gaps, if necessary, to optimize residue matches between the two sequences. In these cases, although the gaps are present, the numbering of the residue in the given amino acid or polynucleotide sequence is made with respect to the reference sequence to which it has been aligned. As used herein, a “reference sequence” refers to a defined sequence used as a basis for a sequence comparison.

Unless stated otherwise, all references herein to amino acid positions of a TcdA refer to the numbering of SEQ ID NO: 1. Unless stated otherwise, all references herein to amino acid positions of a TcdB refer to the numbering of SEQ ID NO: 2.

The glucosyltransferase domain of TcdA, as used herein, may begin at exemplary residue 1, 101, or 102, and may end at exemplary residue 542, 516, or 293

of a wild-type *C. difficile* TcdA, e.g., SEQ ID NO: 1. Any minimum residue position may be combined with a maximum residue position between residues 1 and 542 of TcdA to define a sequence for the glucosyltransferase domain as long as the DXD motif region is included. For example, in one embodiment, the glucosyltransferase domain of TcdA 5 includes SEQ ID NO: 27, which is identical to residues 101-293 of SEQ ID NO: 1, and it includes the DXD motif region. In another embodiment, the glucosyltransferase domain of TcdA includes SEQ ID NO: 28, which is identical to residues 1-542 of SEQ ID NO: 1.

The glucosyltransferase domain of TcdB, as used herein, may begin at exemplary residue 1, 101, or 102, and may end at exemplary residue 543, 516, or 293 10 of a wild-type *C. difficile* TcdB, e.g., SEQ ID NO: 2. Any minimum residue position may be combined with a maximum residue position between residues 1 and 543 of TcdB to define a sequence for the glucosyltransferase domain as long as the DXD motif region is included. For example, in one embodiment, the glucosyltransferase domain of TcdB includes SEQ ID NO: 29, which is identical to residues 101-293 of SEQ ID NO: 2, and it 15 includes the DXD motif region. In another embodiment, the glucosyltransferase domain of TcdB includes SEQ ID NO: 30, which is identical to residues 1-543 of SEQ ID NO: 2.

Without being bound to theory or mechanism, it is believed that the N-terminus of TcdA and/or TcdB is cleaved by an autoproteolytic process for the glucosyltransferase domain to be translocated and released into the host cell cytosol, where it can interact 20 with Rac/Ras/Rho GTPases. Wild-type *C. difficile* TcdA has been shown to be cleaved between L542 and S543. Wild-type *C. difficile* TcdB has been shown to be cleaved between L543 and G544.

The cysteine protease domain is associated with the autocatalytic proteolytic activity of the toxin. The cysteine protease domain is located downstream of the 25 glucosyltransferase domain and may be characterized by the catalytic triad aspartate, histidine, and cysteine (DHC), e.g., D589, H655, and C700 of a wild-type TcdA, and D587, H653, and C698 of a wild-type TcdB. Without being bound by mechanism or theory, it is believed that the catalytic triad is conserved between a toxin from a “historical” strain, such as 630, and a TcdB from a hypervirulent strain, such as R20291.

The cysteine protease domain of TcdA, as used herein, may begin at exemplary residue 543, and may end at exemplary residue 809 769, 768, or 767 of a wild-type TcdA, e.g., SEQ ID NO: 1. Any minimum residue position may be combined with a maximum residue position between 543 and 809 of a wild-type TcdA to define a

sequence for the cysteine protease domain as long as the catalytic triad DHC motif region is included. For example, in one embodiment, the cysteine protease domain of TcdA includes SEQ ID NO: 32, which has the DHC motif region located at residues 47, 113, and 158 of SEQ ID NO: 32, which respectively correspond to D589, H655, and

5 C700 of a wild-type TcdA according to the numbering of SEQ ID NO: 1. SEQ ID NO: 32 is identical to residues 543 to 809 of SEQ ID NO: 1, TcdA.

The cysteine protease domain of TcdB, as used herein, may begin at exemplary residue 544, and may end at exemplary residue 801, 767, 755, or 700 of a wild-type TcdB, e.g., SEQ ID NO: 2. Any minimum residue position may be combined with a

10 maximum residue position between 544 and 801 of a wild-type TcdB to define a sequence for the cysteine protease domain as long as the catalytic triad DHC motif region is included. For example, in one embodiment, the cysteine protease domain of TcdB includes SEQ ID NO: 33, which includes the DHC motif region located at residues 44, 110, and 115 of SEQ ID NO: 33, which respectively correspond to D587, H653, and

15 C698 of a wild-type TcdB according to the numbering of SEQ ID NO: 2. SEQ ID NO: 33 is identical to residues 544 to 767 of SEQ ID NO: 2, TcdB. In another embodiment, the cysteine protease domain of TcdB includes residues 544-801 of SEQ ID NO: 2, TcdB.

Mutant toxin

In the present invention, the immunogenic composition includes a mutant *C. difficile* toxin. The term "mutant," as used herein, refers to a molecule that exhibits a structure or sequence that differs from the corresponding wild-type structure or sequence, e.g., by having crosslinks as compared to the corresponding wild-type structure and/or by having at least one mutation, as compared to the corresponding wild-type sequence when optimally aligned, such as by the programs GAP or BESTFIT

20 using default gap weights. The term "mutant" as used herein further includes a molecule that exhibits a functional property (e.g., abrogated glucosyltransferase and/or abrogated cysteine protease activity) that differs from the corresponding wild-type molecule.

25

A *C. difficile* toxin from any of the wild-type strains described above may be used as a source from which a mutant *C. difficile* toxin is produced. Preferably, *C. difficile* 630 is the source from which a mutant *C. difficile* toxin is produced.

The mutation may involve a substitution, deletion, truncation or modification of the wild type amino acid residue normally located at that position. Preferably, the

mutation is a non-conservative amino acid substitution. The present invention also contemplates isolated polynucleotides that include nucleic acid sequences encoding any of the mutant toxins described herein.

A “non-conservative” amino acid substitution, as used herein, refers to an 5 exchange of an amino acid from one class for an amino acid from another class, according to the following Table 3:

Table 3: Amino Acid Classes	
Class	Amino acid
Nonpolar:	Ala (A), Val (V), Leu (L), Ile (I), Pro (P), Met (M), Phe (F), Trp (W)
Uncharged polar:	Gly (G), Ser (S), Thr (T), Cys (C), Tyr (Y), Asn (N), Gln (Q)
Acidic:	Asp (D), Glu (E)
Basic:	Lys (K), Arg (R), His (H)

Examples of a non-conservative amino acid substitution include a substitution wherein an aspartic acid residue (Asp, D) is replaced by an alanine residue (Ala, A). 10 Other examples include replacing an aspartic acid residue (Asp, D) with an asparagine residue (Asn, N); replacing an arginine (Arg, R), glutamic acid (Glu, E), lysine (Lys, K), and/or histidine (His, H) residue with an alanine residue (Ala, A).

A conservative substitution refers to an exchange between amino acids from the same class, for example, according to Table 3.

15 The mutant toxins of the invention may be prepared by techniques known in the art for preparing mutations, such as, for example, site-directed mutagenesis, mutagenesis using a mutagen (e.g., UV light), etc. Preferably, site-directed mutagenesis is used. Alternatively, a nucleic acid molecule having an objective sequence may be directly synthesized. Such chemical synthesis methods are known in 20 the art.

In the present invention, the mutant *C. difficile* toxin includes at least one mutation in a glucosyltransferase domain, relative to the corresponding wild-type *C. difficile* toxin. In one embodiment, the glucosyltransferase domain includes at least two mutations. Preferably, the mutation decreases or abrogates glucosyltransferase 25 enzyme activity of the toxin, as compared to the glucosyltransferase enzyme activity of the corresponding wild-type *C. difficile* toxin.

Exemplary amino acid residues in a glucosyltransferase domain of TcdA that may undergo a mutation include at least one of the following, or any combination thereof: W101, D269, R272, D285, D287, E460, R462, S541, and L542, as compared to a wild-type *C. difficile* TcdA, according to the numbering of SEQ ID NO: 1. Further 5 exemplary amino acid residues that may undergo a mutation include E514, S517, and W519, as compared to a wild-type *C. difficile* TcdA, according to the numbering of SEQ ID NO: 1.

Exemplary mutations in a glucosyltransferase domain of TcdA include at least one of the following, or any combination thereof: W101A, D269A, R272A, D285A, 10 D287A, E460A, R462A, S541A, and L542G, as compared to a wild-type *C. difficile* TcdA. In a preferred embodiment, the glucosyltransferase domain of TcdA includes a L542G mutation, as compared to a wild-type *C. difficile* TcdA. In another preferred embodiment, the glucosyltransferase domain of TcdA includes a D285A and a D287A mutation, as compared to a wild-type *C. difficile* TcdA.

15 Exemplary amino acid residues in a glucosyltransferase domain of TcdB that may undergo a mutation include at least one of the following, or any combination thereof: W102, D270, R273, D286, D288, N384, D461, K463, W520, and L543, as compared to a wild-type *C. difficile* toxin B, according to the numbering of SEQ ID NO: 2. Further exemplary amino acid residues that may undergo a mutation include E515, 20 S518, and W520, as compared to a wild-type *C. difficile* toxin B, according to the numbering of SEQ ID NO: 2.

Exemplary mutations in a glucosyltransferase domain of TcdB include at least one of the following, or any combination thereof: W102A, D270A, D270N, R273A, D286A, D288A, N384A, D461A, D461R, K463A, K463E, W520A, and L543A, as 25 compared to a wild-type *C. difficile* TcdB. In a preferred embodiment, the glucosyltransferase domain of TcdB includes a L543A, as compared to a wild-type *C. difficile* TcdB. In another preferred embodiment, the glucosyltransferase domain of TcdB includes a D286A and a D288A mutation, as compared to a wild-type *C. difficile* TcdB.

30 Any of the mutations described herein above may be combined with a mutation in a cysteine protease domain. In the present invention, the mutant *C. difficile* toxin includes at least one mutation in a cysteine protease domain, relative to the corresponding wild-type *C. difficile* toxin. Preferably, the mutation decreases or

abrogates cysteine protease activity of the toxin, as compared to the cysteine protease activity of the corresponding wild-type *C. difficile* toxin.

Exemplary amino acid residues in a cysteine protease domain of TcdA that may undergo a mutation include at least one of the following, or any combination thereof:

- 5 S543, D589, H655, and C700, as compared to a wild-type *C. difficile* TcdA, according to the numbering of SEQ ID NO: 1. Exemplary mutations in a glucosyltransferase domain of TcdA include at least one of the following, or any combination thereof: S543A, D589A, D589N, H655A, C700A, as compared to a wild-type *C. difficile* TcdA. In a preferred embodiment, the cysteine protease domain of TcdA includes a C700A mutation, as compared to a wild-type *C. difficile* TcdA.
- 10

Exemplary amino acid residues in a cysteine protease domain of TcdB that may undergo a mutation include at least one of the following, or any combination thereof: G544, D587, H653, and C698, as compared to a wild-type *C. difficile* TcdB, according to the numbering of SEQ ID NO: 2. Exemplary mutations in a glucosyltransferase domain of TcdB include at least one of the following, or any combination thereof: G544A, D587A, D587N, H653A, C698A, as compared to a wild-type *C. difficile* TcdB. In a preferred embodiment, the cysteine protease domain of TcdB includes a C698A mutation, as compared to a wild-type *C. difficile* TcdB. Additional amino acid residues in a cysteine protease domain of TcdB that may undergo a mutation include: K600 and/or R751, as compared to a wild-type TcdB. Exemplary mutations include K600E and/or R751E.

Accordingly, the inventive mutant *C. difficile* toxin includes a glucosyltransferase domain having a mutation and a cysteine protease domain having a mutation, relative to the corresponding wild-type *C. difficile* toxin. In one embodiment, the mutant toxin

- 25 includes at least one mutation in the glucosyltransferase domain and at least one mutation in the cysteine protease domain. In a preferred embodiment, a mutant toxin A includes at least a D285, D287, and C700 mutation. In a preferred embodiment, a mutant toxin B includes at least a D286, D288, and C698 mutation. The mutant toxins may include any further mutation, individually or in combination, described herein.

- 30 An exemplary mutant *C. difficile* TcdA includes a glucosyltransferase domain including SEQ ID NO: 29 having an amino acid substitution at positions 285 and 287, and a cysteine protease domain comprising SEQ ID NO: 32 having an amino acid substitution at position 158, relative to the corresponding wild-type *C. difficile* toxin A.

For example, such a mutant *C. difficile* TcdA includes the amino acid sequence set forth in SEQ ID NO: 4, wherein the initial methionine is optionally not present. In another embodiment, the mutant mutant *C. difficile* toxin A includes the amino acid sequence set forth in SEQ ID NO: 84.

5 Further examples of a mutant *C. difficile* toxin A include the amino acid sequence set forth in SEQ ID NO: 7, which has a D269A, R272A, D285A, D287A, E460A, R462A, and C700A mutation, as compared to SEQ ID NO: 1, wherein the initial methionine is optionally not present. In another embodiment, the mutant mutant *C. difficile* toxin A includes the amino acid sequence set forth in SEQ ID NO: 83.

10 Another exemplary mutant TcdA includes SEQ ID NO: 34, wherein the residue at positions 101, 269, 272, 285, 287, 460, 462, 541, 542, 543, 589, 655, and 700 may be any amino acid.

15 In some embodiments, the mutant *C. difficile* toxin exhibits decreased or abrogated autoproteolytic processing as compared to the corresponding wild-type *C. difficile* toxin. For example, a mutant *C. difficile* TcdA may include a mutation at one of the following residues, or any combination thereof: S541, L542 and/or S543, as compared to the corresponding wild-type *C. difficile* TcdA. Preferably, the mutant *C. difficile* TcdA includes at least one of the following mutations, or any combination thereof: S541A, L542G, and S543A, as compared to the corresponding wild-type *C. difficile* TcdA.

20 Another exemplary mutant *C. difficile* TcdA includes a S541A, L542, S543 and C700 mutation, as compared to the corresponding wild-type *C. difficile* TcdA.

25 An exemplary mutant *C. difficile* toxin B includes a glucosyltransferase domain comprising SEQ ID NO: 31 having an amino acid substitution at positions 286 and 288, and a cysteine protease domain comprising SEQ ID NO: 33 having an amino acid substitution at position 155, relative to the corresponding wild-type *C. difficile* toxin B. For example, such a mutant *C. difficile* TcdB includes the amino acid sequence set forth in SEQ ID NO: 6, wherein the initial methionine is optionally not present. In another embodiment, the mutant mutant *C. difficile* toxin A includes the amino acid sequence set forth in SEQ ID NO: 86.

30 Further examples of a mutant *C. difficile* TcdB include the amino acid sequence set forth in SEQ ID NO: 8, which has a D270A, R273A, D286A, D288A, D461A, K463A, and C698A mutation, as compared to SEQ ID NO: 2. SEQ ID NO: 8 wherein the initial

methionine is optionally not present. In another embodiment, the mutant *C. difficile* toxin A includes the amino acid sequence set forth in SEQ ID NO: 85.

Another exemplary mutant TcdB includes SEQ ID NO: 35, wherein the residue at positions 101, 269, 272, 285, 287, 460, 462, 541, 542, 543, 589, 655, and 700 may be 5 any amino acid.

As another example, a mutant *C. difficile* TcdB may include a mutation at positions 543 and/or 544, as compared to the corresponding wild-type *C. difficile* TcdB. Preferably, the mutant *C. difficile* TcdB includes a L543 and/or G544 mutation, as compared to the corresponding wild-type *C. difficile* TcdB. More preferably, the mutant 10 *C. difficile* TcdB includes a L543G and/or G544A mutation, as compared to the corresponding wild-type *C. difficile* TcdB.

Another exemplary mutant *C. difficile* TcdB includes a L543G, G544A and C698 mutation, as compared to the corresponding wild-type *C. difficile* TcdB.

In one aspect, the invention relates to an isolated polypeptide having a mutation 15 at any position from amino acid residue 1 to 1500 according to the numbering of SEQ ID NO: 2, to define an exemplary mutant *C. difficile* toxin B. For example, in one embodiment, the isolated polypeptide includes a mutation between amino acids residues 830 and 990 of SEQ ID NO: 2. Exemplary positions for mutations include positions 970 and 976 according to the numbering of SEQ ID NO: 2. Preferably, the 20 mutation between residues 830 and 990 is a substitution. In one embodiment, the mutation is a non-conservative substitution wherein an Asp (D) and/or a Glu (E) amino acid residue is replaced by an amino acid residue that is not neutralized upon acidification, such as, for example, lysine (K), arginine (R), and histidine (H). Exemplary mutations include: E970K, E970R, E970H, E976K, E976R, E976H of SEQ ID NO: 2, to 25 define a mutant *C. difficile* toxin B.

In one embodiment, the isolated polypeptide includes the following substitutions D286A/D288A/C698A/E970K/E976K (SEQ ID NO: 184). E970 and E976 are conserved in toxin B from all *C. difficile* strains observed (see Table 2-a, SEQ ID NOs: 110-30 133), except in ribotype 078 and ribotype 126 strains (see Table 35 and Table 37). In toxin B from ribotype 078 and ribotype 126 strains, there is a glycine-970 (G970) instead of glutamate-970. Accordingly, in one embodiment, the isolated polypeptide includes a mutation at G970 and E976, such as, for example, G970K and E976K. The

mutant toxins described above and herein may exhibit reduced cytotoxicity as compared to the corresponding wild-type toxin. See Examples 8 and 15).

In another aspect, the invention relates to an isolated polypeptide having a mutation at any position from amino acid residue 1 to 1500 according to the numbering

5 of SEQ ID NO: 1, to define an exemplary mutant *C. difficile* toxin A. For example, in one embodiment, the isolated polypeptide includes a mutation between amino acids residues 832 and 992 of SEQ ID NO: 1. Exemplary positions for mutations include positions 972 and 978 according to the numbering of SEQ ID NO: 1. Preferably, the mutation between residues 832 and 992 is a substitution. In one embodiment, the 10 mutation is a non-conservative substitution wherein an Asp (D) and/or a Glu (E) amino acid residue is replaced by an amino acid residue that is not neutralized upon acidification, such as, for example, lysine (K), arginine (R), and histidine (H). Exemplary mutations include: D972K, D972R, D972H, D978K, D978R, D978H of SEQ ID NO: 1, to define a mutant *C. difficile* toxin A.

15 In one embodiment, the isolated polypeptide includes the following substitutions D285A/D287A/C700A/D972K/D978K (SEQ ID NO: 183). D972 and D978 residues are conserved in toxin A from all *C. difficile* strains assessed (see Table 1-a, SEQ ID NOs: 87-109). The mutant toxins described above and herein may exhibit reduced cytotoxicity as compared to the corresponding wild-type toxin.

20 The following describes additional exemplary mutant toxins. In one embodiment, the mutant toxin TcdA includes (i) SEQ ID NO: 185 ; (ii) a polypeptide of SEQ ID NO:185 having at least 90%, 92%, 93%, 95%, 98%, 99% or 100% identity to SEQ ID NO:185 ;or (iii) a fragment of at least 250, 280 or 300 amino acids of SEQ ID NO:185. In another embodiment, the mutant toxin TcdB includes (iv) SEQ ID NO:186;(v) a 25 polypeptide of SEQ ID NO:186 having at least 80%, 85%, 90%, 92%, 93%, 95%, 98%, 99% or 100% identity to SEQ ID NO:186; or (vi) a fragment of at least 250, 280 or 300 amino acids of SEQ ID NO:186.

In one embodiment, the mutant toxin TcdA consists of less than 600, 675, 650, 625, 600, 575, 550, 525, 500, 475, 450, 425, 400, 375, 350, 325, 300, 275, 250, or 225 30 amino acids. In one embodiment, the mutant toxin consists of less than 800, 775, 750, 725, 700, 675, 650, 625, 600, 575, 550, or 525 amino acids. In one embodiment, the mutant toxin includes at least 200, 225, 250, 270, 280, 300 or 310 amino acids of SEQ ID NO:185 or at least 200, 225, 250, 270, 280, 300 or 310 amino acids of a polypeptide

having at least 80%, 85%, 90%, 92%, 95%, 98%, 99%, or 100% identity to SEQ ID NO:185. In one embodiment the mutant toxin includes at least 400, 425, 450, 475, 500, 525, 550, 575, 600 or 610 amino acids of SEQ ID NO: 186 or at least 400, 425, 450, 475, 500, 525, 550, 575, 600 or 610 amino acids of a polypeptide having at least 80%, 5 85%, 90%, 92%, 95%, 98%, 99%, or 100% identity to SEQ ID NO:186.

In one embodiment, the mutant toxin includes a fusion protein that includes A) (i) SEQ ID NO:185 ; (ii) a polypeptide of SEQ ID NO:185 having at least 90%, 92%, 93%, 95%, 98%, 99% or 100% identity to SEQ ID NO:185 ;or (iii) a fragment of at least 250, 10 280 or 300 amino acids of SEQ ID NO:185 fused to B) (iv) SEQ ID NO:186;(v) a polypeptide of SEQ ID NO:186 having at least 80%, 85%, 90%, 92%, 93%, 95%, 98%, 99% or 100% identity to SEQ ID NO:186; or (vi) a fragment of at least 250, 280 or 300 15 amino acids of SEQ ID NO:186. In a further embodiment the N-terminus of SEQ ID NO: 185 is adjacent to the C-terminus of SEQ ID NO: 186. In a further embodiment the N-terminus of SEQ ID NO: 185 is adjacent to the N-terminus of SEQ ID NO: 186. In a further embodiment the C-terminus of SEQ ID NO:185 is adjacent to the C-terminus of SEQ ID NO:186. Further examples of a mutant toxin include a polypeptide having any one of the following amino acid sequences SEQ ID NO: 187, SEQ ID NO: 188, SEQ ID NO: 189, SEQ ID NO: 190, SEQ ID NO: 191, SEQ ID NO: 192, SEQ ID NO: 193, SEQ ID NO: 194, and SEQ ID NO: 195.

20 In another embodiment, the mutant toxin includes a fusion and/or hybrid polypeptide that includes any combination of amino acid sequences selected from any of the following: SEQ ID NO: 223, SEQ ID NO: 224, SEQ ID NO: 225, SEQ ID NO: 226, SEQ ID NO: 227, SEQ ID NO: 228, SEQ ID NO: 229, SEQ ID NO: 230, SEQ ID NO: 231, SEQ ID NO: 232, SEQ ID NO: 233, SEQ ID NO: 234, SEQ ID NO: 235, SEQ ID NO: 236, SEQ ID NO: 237, SEQ ID NO: 238, SEQ ID NO: 239, SEQ ID NO: 240, SEQ ID NO: 241, SEQ ID NO: 242, and SEQ ID NO: 243. For example, in one embodiment, the mutant toxin includes an amino acid sequence as set forth in any one of the following: SEQ ID NO: 254, SEQ ID NO: 270, SEQ ID NO: 271, SEQ ID NO: 272, SEQ ID NO: 273, SEQ ID NO: 274, or SEQ ID NO: 275.

30 Another example of a mutant toxin includes a fragment of a wild-type toxin. A “fragment” mutant toxin TcdA as used herein refers to a peptide sequence that has less consecutive amino acids total than the corresponding wild-type *C. difficile* toxin TcdA sequence, e.g., a sequence that includes less than 2710 consecutive amino acids total.

The fragment mutant toxin TcdA may further include a mutation of an amino acid residue as described herein. A “fragment” mutant toxin TcdB as used herein refers to a peptide sequence that has less consecutive amino acids total than the corresponding wild-type *C. difficile* toxin TcdB sequence, e.g., a sequence that includes less than 2366 consecutive amino acids total. The fragment mutant toxin TcdB may further include a mutation of an amino acid residue as described herein. Such exemplary mutant toxin sequences include SEQ ID NO: 28, SEQ ID NO: 29, SEQ ID NO: 30, SEQ ID NO: 31, SEQ ID NO: 32, SEQ ID NO: 33. In one embodiment, the mutant toxin TcdA includes at least 200, 300, 400, 500, 600, 700, 800, 900, 1000, 1100, 1200, 1300, 1400, 1500, 1600, 1700, 1800, 1900, or 2000 consecutive amino acids of a wild-type mutant toxin A, as described herein, e.g., SEQ ID NO: 1. In another embodiment, a mutant toxin includes a fragment of a genetically mutated toxin A described herein. In one embodiment, the mutant toxin TcdB includes at least 200, 300, 400, 500, 600, 700, 800, 900, 1000, 1100, 1200, 1300, 1400, 1500, 1600, 1700, 1800, 1900, or 2000 consecutive amino acids of a wild-type mutant toxin B, as described herein, e.g., SEQ ID NO: 2. In another embodiment, a mutant toxin includes a fragment of a genetically mutated toxin B described herein. In one embodiment, the mutant toxin includes at most 3000, 2710, 2500, 2400, 2366, 2000, 1900, 1800, 1700, 1600, 1500, 1400, 1300, 1200, 1100, 1000, 900, 800, 700, 600, 500, 400, or 300 consecutive amino acids. Any minimum value may be combined with any maximum value to define a suitable range. In another embodiment, the mutant toxin is fused to at least one other peptide or at least one other mutant toxin in a manner that results in the production of a hybrid molecule.

Additional exemplary fragment mutant toxin TcdA includes a polypeptide having any one of the amino acid sequences of SEQ ID NO: 374 through SEQ ID NO: 421. Further exemplary fragment mutant toxin TcdA includes a polypeptide having any one of the amino acid sequences of SEQ ID NO: 472 through SEQ ID NO: 519. Additional exemplary mutant toxin TcdB includes a polypeptide having any one of the amino acid sequences of SEQ ID NO: 422 through SEQ ID NO: 471. Further exemplary fragment mutant toxin TcdB includes a polypeptide having any one of the amino acid sequences of SEQ ID NO: 568 through SEQ ID NO: 615.

The following describes further exemplary mutant toxins. In one embodiment, the mutant toxin includes a TcdA which includes or consists of at least 3, at least 4, or at least 5 mutations at amino acid residues selected from the group consisting of: W101,

D287, E514, D285, S517, W519, and C700, e.g., according to the numbering of SEQ ID NO: 1. In additional embodiments; the TcdA mutants include or consist of at least 3, at least 4, or at least 5 mutations selected from the group consisting of: W101A, D287A, E514Q, D285A, S517A, W519A, and C700A substitutions and a W101 deletion.

5 Another exemplary mutant toxin TcdA includes the amino acid substitutions W101, D287A, and E514Q, e.g., according to the numbering of SEQ ID NO: 1. A further embodiment provides a TcdA protein including or consisting of the amino acid substitutions W101A, D287A, E514Q, and W519A. Another specific embodiment of the invention is a TcdA protein including or consisting of the amino acid substitutions

10 W101A, D287A, E514Q, W519A, and C700A.

In another embodiment, the mutant toxin TcdA includes the mutations W101A, D287A, E514Q and D285A, e.g., according to the numbering of SEQ ID NO: 1. In another embodiment, the mutant toxin TcdA includes the mutations W101A, D287A, E514Q and S517A.

15 In another embodiment, a further mutation may be added to the mutant TcdA, e.g. a C700A mutation. In additional embodiments, up to 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20 further mutations may be added to any of the mutant toxin TcdA embodiments described herein.

Further examples of a mutant toxin TcdA includes the amino acid sequence as 20 set forth in SEQ ID NO: 203, SEQ ID NO: 204, SEQ ID NO: 205, SEQ ID NO: 206, SEQ ID NO: 207, SEQ ID NO: 208, SEQ ID NO: 209, or SEQ ID NO: 211. In another embodiment, the mutant toxin TcdA includes the amino acid sequence SEQ ID NO: 210. In yet another embodiment, the mutant toxin TcdA includes a mutation at positions 25 W101, D287, E514, W519 and C700; wherein W101 is replaced with any amino acid except tryptophan, D287 is replaced with any amino acid but aspartic acid, E514 is replaced with any amino acid but glutamic acid, W519 is replaced with any amino acid but tryptophan and C700 is replaced with any amino acid but cysteine, as set forth in SEQ ID NO: 212.

Additional examples of a mutant toxin TcdA include a polypeptide having an 30 amino acid sequence that is at least 85%, at least 90%, at least 95%, at least 97%, at least 98%, or at least 99% identical to the original reference sequence (e.g. SEQ ID NO: 203, SEQ ID NO: 204, SEQ ID NO: 205, SEQ ID NO: 206, SEQ ID NO: 207, SEQ ID NO: 208, SEQ ID NO: 209, SEQ ID NO: 210, SEQ ID NO: 211 or SEQ ID NO:

212 or SEQ ID NO: 82, SEQ ID NO: 83, SEQ ID NO: 84, SEQ ID NO: 85 or SEQ ID NO: 86).

In a further embodiment, mutant toxin TcdA includes mutations at at most than 12 amino acid residues, at most than 11 amino acid residues, at most than 10 amino acid residues, at most than 9 amino acid residues, at most than 8 amino acid residues, at most than 7 amino acid residues, at most than 6 amino acid residues, at most than 5 amino acid residues, at most than 4 amino acid residues, at most than 3 amino acid residues, at most than 2 amino acid residues, or 1 amino acid residue, for example, relative to SEQ ID NO: 203, SEQ ID NO: 204, SEQ ID NO: 205, SEQ ID NO: 206, SEQ ID NO: 207, SEQ ID NO: 208, SEQ ID NO: 209, SEQ ID NO: 210, SEQ ID NO: 211 and SEQ ID NO: 212 or SEQ ID NO: 82, SEQ ID NO: 83, SEQ ID NO: 84, SEQ ID NO: 85 or SEQ ID NO: 86.

The following describes further exemplary mutant toxins. In one embodiment, the mutant toxin is a mutant TcdB that that includes at least 3, at least 4, or at least 5 mutations at amino acid residues selected from the group consisting of: W102, D288, E515, D286, S518, W520, and C698, according to the numbering of SEQ ID NO: 2. In another embodiment, the mutant toxin TcdB includes at least 3, at least 4, or at least 5 mutations selected from the group consisting of: W102A, D288A, E515Q, D286A, S518A, W520A, and C698A substitutions and a W102 deletion. In one embodiment, the mutant toxin TcdB includes the amino acid sequence as set forth in SEQ ID NO: 213, SEQ ID NO: 214, SEQ ID NO: 215, SEQ ID NO: 216, SEQ ID NO: 217, SEQ ID NO: 218, SEQ ID NO: 219, SEQ ID NO: 220, or SEQ ID NO: 221. In yet another embodiment, the mutant toxin TcdB includes a mutation at positions W102, D288, E515, W520 and C698; wherein W102 is replaced with any amino acid except tryptophan, D288 is replaced with any amino acid but aspartic acid, E515 is replaced with any amino acid but glutamic acid, W520 is replaced with any amino acid but tryptophan and C698 is replaced with any amino acid but cysteine, as set forth in SEQ ID NO: 222.

Additional examples of a mutant toxin TcdB include a polypeptide having an amino acid sequence that is at least 85%, at least 90%, at least 95%, at least 97%, at least 98%, or at least 99% identical to the original reference sequence SEQ ID NO: 213, SEQ ID NO: 214, SEQ ID NO: 215, SEQ ID NO: 216, SEQ ID NO: 217, SEQ ID NO: 218, SEQ ID NO: 219, SEQ ID NO: 220, SEQ ID NO: 221, or SEQ ID NO: 222.

In a further embodiment, mutant toxin TcdB includes mutations at at most than 12 amino acid residues, at most than 11 amino acid residues, at most than 10 amino acid residues, at most than 9 amino acid residues, at most than 8 amino acid residues, at most than 7 amino acid residues, at most than 6 amino acid residues, at most than 5 amino acid residues, at most than 4 amino acid residues, at most than 3 amino acid residues, at most than 2 amino acid residues, or 1 amino acid residue, for example, relative to SEQ ID NO: 213, SEQ ID NO: 214, SEQ ID NO: 215, SEQ ID NO: 216, SEQ ID NO: 217, SEQ ID NO: 218, SEQ ID NO: 219, SEQ ID NO: 220, SEQ ID NO: 221, and/or SEQ ID NO: 222.

10 Additional exemplary mutant toxin TcdA includes a polypeptide having any one of the amino acid sequences of SEQ ID NO: 276 through SEQ ID NO: 323. Additional exemplary mutant toxin TcdB includes a polypeptide having any one of the amino acid sequences of SEQ ID NO: 324 through SEQ ID NO: 373.

15 The following describes further exemplary mutant toxins. In one embodiment, the mutant toxin TcdA includes an amino acid sequence: (a) having 50% or more identity (e.g. 60%, 65%, 70%, 75%, 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 98.5%, 99%, 99.5%, 99.8%, 99.9%, or 100%) to SEQ ID NO: 224 or SEQ ID NO: 245; and/or b) that is a fragment of at least "n" consecutive amino acids of SEQ ID NO: 224 or SEQ ID NO: 245, wherein "n" is 7 or more (e.g. 8, 10, 12, 14, 16, 18, 20, 25, 20 30, 35, 40, 45, 50, 60, 70, 80, 90, 100, 150, 250, 300, 400, 500, 540, or more).or of a polypeptide having 80% or more identity to SEQ ID NO: 224 or SEQ ID NO: 245 and that comprises an epitope of SEQ ID NO: 224 and/or SEQ ID NO: 245.

25 In one embodiment, the mutant toxin TcdA includes the amino acid sequence set forth in SEQ ID NO: 223, SEQ ID NO: 224, SEQ ID NO: 225, SEQ ID NO: 226, SEQ ID NO: 227, SEQ ID NO: 228, SEQ ID NO: 229, SEQ ID NO: 230, SEQ ID NO: 231, SEQ ID NO: 232, SEQ ID NO: 233, SEQ ID NO: 234, SEQ ID NO: 235, or SEQ ID NO: 236. Additional exemplary embodiments of a mutant toxin TcdA includes the amino acid sequence set forth in any of the following: SEQ ID NO: 244, SEQ ID NO: 245, SEQ ID NO: 250, SEQ ID NO: 251, SEQ ID NO: 252, SEQ ID NO: 253, SEQ ID NO: 254, 30 SEQ ID NO: 255, SEQ ID NO: 256, SEQ ID NO: 257, SEQ ID NO: 258, SEQ ID NO: 259, SEQ ID NO: 260, SEQ ID NO: 261, SEQ ID NO: 262, and/or SEQ ID NO: 263.

35 In one embodiment, the mutant toxin TcdB includes an amino acid sequence (a) having 80% or more identity to SEQ ID NO: 238 or SEQ ID NO: 247; and/or b) that is a

fragment of at least 7 consecutive amino acids of SEQ ID NO: 238 or SEQ ID NO: 247, or of a polypeptide having 80% or more identity to SEQ ID NO: 238 or SEQ ID NO: 247 and that comprises an epitope of SEQ ID NO: 238 or SEQ ID NO: 247.

In one embodiment, the mutant toxin TcdB includes an amino acid sequence: (a) 5 having 50% or more identity (e.g. 60%, 65%, 70%, 75%, 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 98.5%, 99%, 99.5%, 99.8%, 99.9%, or more) to SEQ ID NO: 238 or SEQ ID NO: 247; and/or (b) that is a fragment of at least "n" consecutive amino acids of SEQ ID NO: 238 or SEQ ID NO: 247, or of a polypeptide having 50% or more identity to SEQ ID NO: 238 or SEQ ID NO: 247, wherein "n" is 7 or more (e.g. 8, 10, 12, 14, 16, 18, 20, 25, 30, 35, 40, 45, 50, 60, 70, 80, 90, 100, 150, 250, 300, 400, 500, 540, or more).

In one embodiment, the mutant toxin TcdB includes the amino acid sequence set forth in SEQ ID NO: 237, SEQ ID NO: 238, SEQ ID NO: 239, SEQ ID NO: 240, SEQ ID NO: 241, SEQ ID NO: 242, or SEQ ID NO: 243. Additional exemplary embodiments of 15 a mutant toxin TcdB includes the amino acid sequence set forth in any of the following: SEQ ID NO: 246, SEQ ID NO: 247, SEQ ID NO: 248, SEQ ID NO: 249, SEQ ID NO: 264, SEQ ID NO: 265, SEQ ID NO: 266, SEQ ID NO: 267, SEQ ID NO: 268, and/or SEQ ID NO: 269.

Additional exemplary mutant toxin TcdA includes a polypeptide having any one of 20 the amino acid sequences of SEQ ID NO: 520 through SEQ ID NO: 567. Additional exemplary mutant toxin TcdB includes a polypeptide having any one of the amino acid sequences of SEQ ID NO: 616 through SEQ ID NO: 663.

The following describes yet additional exemplary mutant toxins. In one embodiment, the mutant toxin TcdA includes a polypeptide having any one of the amino 25 acid sequences of SEQ ID NO: 664 through SEQ ID NO: 711. Additional exemplary mutant toxin TcdB includes a polypeptide having any one of the amino acid sequences of SEQ ID NO: 712 through SEQ ID NO: 761.

The polypeptides of the invention may include an initial methionine residue, in some cases as a result of a host cell-mediated process. Depending on, for example, 30 the host cell used in a recombinant production procedure and/or the fermentation or growth conditions of the host cell, it is known in the art that the N-terminal methionine encoded by the translation initiation codon may be removed from a polypeptide after

translation in cells or the N-terminal methionine may remain present in the isolated polypeptide.

Accordingly, in one aspect, the invention relates to an isolated polypeptide including the amino acid sequence set forth in SEQ ID NO: 4, wherein the initial methionine (at position 1) is optionally not present. In one embodiment, the initial methionine of SEQ ID NO: 4 is absent. In one aspect, the invention relates to an isolated polypeptide including the amino acid sequence set forth in SEQ ID NO: 84, which is identical to SEQ ID NO: 4, but for an absence of the initial methionine.

In another aspect, the isolated polypeptide includes the amino acid sequence set forth in SEQ ID NO: 6, wherein the initial methionine (at position 1) is optionally not present. In one embodiment, the initial methionine of SEQ ID NO: 6 is absent. In one aspect, the invention relates to an isolated polypeptide including the amino acid sequence set forth in SEQ ID NO: 86, which is identical to SEQ ID NO: 6, but for an absence of the initial methionine.

In a further aspect, the isolated polypeptide includes the amino acid sequence set forth in SEQ ID NO: 7, wherein the initial methionine (at position 1) is optionally not present. In one embodiment, the invention relates to an isolated polypeptide including the amino acid sequence set forth in SEQ ID NO: 83, which is identical to SEQ ID NO: 7, but for an absence of the initial methionine. In yet another aspect, the isolated polypeptide includes the amino acid sequence set forth in SEQ ID NO: 8, wherein the initial methionine (at position 1) is optionally not present. In one embodiment, the isolated polypeptide includes the amino acid sequence set forth in SEQ ID NO: 85, which is identical to SEQ ID NO: 8, but for an absence of the initial methionine.

In one aspect, the invention relates to an immunogenic composition including SEQ ID NO: 4, wherein the initial methionine (at position 1) is optionally not present. In another aspect, the invention relates to an immunogenic composition including SEQ ID NO: 6, wherein the initial methionine (at position 1) is optionally not present. In a further aspect, the invention relates to an immunogenic composition including SEQ ID NO: 7, wherein the initial methionine (at position 1) is optionally not present. In yet another aspect, the invention relates to an immunogenic composition including SEQ ID NO: 8, wherein the initial methionine (at position 1) is optionally not present.

In another aspect, the invention relates to an immunogenic composition including SEQ ID NO: 83. In one aspect, the invention relates to an immunogenic composition

including SEQ ID NO: 84. In one aspect, the invention relates to an immunogenic composition including SEQ ID NO: 85. In another aspect, the invention relates to an immunogenic composition including SEQ ID NO: 86.

Cytotoxicity

In addition to generating an immune response in a mammal, the immunogenic compositions described herein also have reduced cytotoxicity compared to the corresponding wild-type *C. difficile* toxin. Preferably, the immunogenic compositions are 5 safe and have minimal (e.g., about a 6-8 \log_{10} reduction) to no cytotoxicity, relative to the cytotoxicity of a respective wild-type toxin, for administration in mammals.

As used herein, the term cytotoxicity is a term understood in the art and refers to apoptotic cell death and/or a state in which one or more usual biochemical or biological functions of a cell are aberrantly compromised, as compared to an identical cell under 10 identical conditions but in the absence of the cytotoxic agent. Toxicity can be quantitated, for example, in cells or in mammals as the amount of an agent needed to induce 50% cell death (i.e., EC₅₀ or ED₅₀, respectively) or by other methods known in the art.

Assays for indicating cytotoxicity are known in the art, such as cell rounding 15 assays (see, for example, Kuehne et al. *Nature*. 2010 Oct 7;467(7316):711-3). The action of TcdA and TcdB causes cells to round (e.g., lose morphology) and die, and such a phenomenon is visible by light microscopy. See, for example, Figure 9.

Additional exemplary cytotoxicity assays known in the art include glucosylation assays relating to phosphorimaging of Ras labeled with [¹⁴C]glucose assays (as 20 described in Busch et al., *J Biol Chem*. 1998 Jul 31;273(31):19566-72), and preferably the *in vitro* cytotoxicity assay described in the Examples below wherein EC₅₀ may refer to a concentration of an immunogenic composition that exhibits at least about 50% of 25 cytopathogenic effect (CPE) in a cell, preferably a human diploid fibroblast cell (e.g., IMR90 cell (ATCC CCL-186TM), as compared to an identical cell under identical conditions in the absence of the toxin. The *in vitro* cytotoxicity assay may also be used to assess the concentration of a composition that inhibits at least about 50% of a wild-type *C. difficile* toxin-induced cytopathogenic effect (CPE) in a cell, preferably a human 30 diploid fibroblast cell (e.g., IMR90 cell (ATCC CCL-186TM), as compared to an identical cell under identical conditions in the absence of the toxin. Additional exemplary cytotoxicity assays include those described in Doern et al., *J Clin Microbiol*. 1992 Aug;30(8):2042-6. Cytotoxicity can also be determined by measuring ATP levels in cells treated with toxin. For example, a luciferase based substrate such as CELLTITERGLO[®] (Promega) may be used, which emits luminescence measured as a

relative light unit (RLU). In such an assay, cell viability may be directly proportional to the amount of ATP in the cells or the RLU values.

In one embodiment, the cytotoxicity of the immunogenic composition is reduced by at least about 1000, 2000, 3000, 4000, 5000-, 6000-, 7000-, 8000-, 9000-, 10000-, 5 11000-, 12000-, 13000-fold, 14000-fold, 15000-fold, or more, as compared to the corresponding wild-type *C. difficile* toxin. See, for example, Table 20.

In another embodiment, the cytotoxicity of the immunogenic composition is reduced by at least about $2\log_{10}$, more preferably by about $3\log_{10}$, and most preferably by about $4\log_{10}$ or more, relative to the corresponding wild-type toxin under identical 10 conditions. For example, a mutant *C. difficile* TcdB may have an EC₅₀ value of about 10^{-9} g/ml as measured in a standard cytopathic effect assay (CPE), as compared to an exemplary wild-type *C. difficile* TcdB which may have an EC₅₀ value of at least about 10^{-12} g/ml. See, for example, Tables 7A, 7B, 8A and 8B in the Examples section below.

In yet another embodiment, the cytotoxicity of the mutant *C. difficile* toxin has an 15 EC₅₀ of at least about 50 µg/ml, 100 µg/ml, 200 µg/ml, 300 µg/ml, 400 µg/ml, 500 µg/ml, 600 µg/ml, 700 µg/ml, 800 µg/ml, 900 µg/ml, 1000 µg/ml or greater, as measured by, for example, an *in vitro* cytotoxicity assay, such as one described herein. Accordingly, in a preferred embodiment, the immunogenic compositions and mutant toxins are biologically safe for administration to mammals.

20 Without being bound by mechanism or theory, a TcdA having a D285 and D287 mutation, as compared to a wild-type TcdA, and a TcdB having a D286 and a D288 mutation, as compared to a wild-type TcdB, were expected to be defective in glycosyltransferase activity and therefore defective in inducing a cytopathic effect. In addition, a toxin having a mutation in the DHC motif was expected to be defective in 25 autocatalytic processing, and therefore be without any cytotoxic effects.

However, the inventors surprisingly discovered, among other things, that 30 exemplary mutant TcdA having SEQ ID NO: 4 and exemplary mutant TcdB having SEQ ID NO: 6 unexpectedly exhibited cytotoxicity (albeit significantly reduced from wild-type *C. difficile* 630 toxins) despite exhibiting dysfunctional glucosyltransferase activity and dysfunctional cysteine protease activity. Without being bound by mechanism or theory, the mutant toxins are believed to effect cytotoxicity through a novel mechanism. Nevertheless, the exemplary mutant TcdA having SEQ ID NO: 4 and exemplary mutant TcdB having SEQ ID NO: 6 were surprisingly immunogenic. See Examples below.

Crosslinking

Although chemical crosslinking of a wild-type toxin has a potential to fail in inactivating the toxin, the inventors further discovered that chemically crosslinking at least one amino acid of a mutant toxin further reduced cytotoxicity of the mutant toxin, 5 relative to an identical mutant toxin lacking chemical crosslinks, and relative to the corresponding wild-type toxin. Preferably, the mutant toxin is purified before contact with the chemical crosslinking agent.

Moreover, despite a potential of chemical crosslinking agents to alter useful epitopes, the inventors surprisingly discovered that a genetically modified mutant *C. difficile* toxin having at least one amino acid chemically crosslinked resulted in 10 immunogenic compositions that elicited multiple neutralizing antibodies or binding fragments thereof. Accordingly, epitopes associated with neutralizing antibody molecules were unexpectedly retained following chemical crosslinking.

Crosslinking (also referred to as "chemical inactivation" or "inactivation" herein) is 15 a process of chemically joining two or more molecules by a covalent bond. The terms "crosslinking reagents," "crosslinking agents," and "crosslinkers" refer to molecules that are capable of reacting with and/or chemically attaching to specific functional groups (primary amines, sulhydryls, carboxyls, carbonyls, etc) on peptides, polypeptides, and/or proteins. In one embodiment, the molecule may contain two or more reactive 20 ends that are capable of reacting with and/or chemically attaching to specific functional groups (primary amines, sulhydryls, carboxyls, carbonyls, etc) on peptides, polypeptides, and/or proteins. Preferably, the chemical crosslinking agent is water-soluble. In another preferred embodiment, the chemical crosslinking agent is a 25 heterobifunctional crosslinker. In another embodiment, the chemical crosslinking agent is not a bifunctional crosslinker. Chemical crosslinking agents are known in the art.

In a preferred embodiment, the crosslinking agent is a zero-length crosslinking agent. A "zero-length" crosslinker refers to a crosslinking agent that will mediate or produce a direct crosslink between functional groups of two molecules. For example, in 30 the crosslinking of two polypeptides, a zero-length crosslinker will result in the formation of a bridge, or a crosslink between a carboxyl group from an amino acid side chain of one polypeptide, and an amino group of another polypeptide, without integrating extrinsic matter. Zero-length crosslinking agents can catalyze, for example, the

formation of ester linkages between hydroxyl and carboxyl moieties, and/or the formation of amide bonds between carboxyl and primary amino moieties.

Exemplary suitable chemical crosslinking agents include formaldehyde; formalin; acetaldehyde; propionaldehyde; water-soluble carbodiimides (RN=C=NR'), which 5 include 1-Ethyl-3-(3-Dimethylaminopropyl)-Carbodiimide (EDC), 1-Ethyl-3-(3-Dimethylaminopropyl)-Carbodiimide Hydrochloride, 1-Cyclohexyl-3-(2-morpholiny-(4-ethyl)carbodiimide metho-p-toluenesulfonate (CMC), N,N'-dicyclohexylcarbodiimide (DCC), and N,N'-diisopropylcarbodiimide (DIC), and derivatives thereof; and N-hydroxysuccinimide (NHS); phenylglyoxal; and/or UDP-dialdehyde.

10 Preferably, the crosslinking agent is EDC. When a mutant *C. difficile* toxin polypeptide is chemically modified by EDC (e.g., by contacting the polypeptide with EDC), in one embodiment, the polypeptide includes (a) at least one crosslink between a side chain of an aspartic acid residue of the polypeptide and a side chain of a lysine residue of the polypeptide. In one embodiment, the polypeptide includes (b) at least 15 one crosslink between a side chain of a glutamic acid residue of the polypeptide and a side chain of a lysine residue of the polypeptide. In one embodiment, the polypeptide includes (c) at least one crosslink between the carboxyl group at the C-terminus of the polypeptide and the amino group of the N-terminus of the polypeptide. In one embodiment, the polypeptide includes (d) at least one crosslink between the carboxyl 20 group at the C-terminus of the polypeptide and a side chain of a lysine residue of the polypeptide. In one embodiment, the polypeptide includes (e) at least one crosslink between a side chain of an aspartic acid residue of the polypeptide and a side chain of a lysine residue of a second isolated polypeptide. In one embodiment, the polypeptide includes (f) at least one crosslink between a side chain of a glutamic acid residue of the 25 polypeptide and a side chain of a lysine residue of a second isolated polypeptide. In one embodiment, the polypeptide includes (g) at least one crosslink between the carboxyl group at the C-terminus of the polypeptide and the amino group of the N-terminus of a second isolated polypeptide. In one embodiment, the polypeptide includes (h) at least one crosslink between the carboxyl group at the C-terminus of the 30 polypeptide and a side chain of a lysine residue of a second isolated polypeptide. See, for example, Figure 24 and Figure 25.

The "second isolated polypeptide" refers to any isolated polypeptide that is present during the reaction with EDC. In one embodiment, the second isolated

polypeptide is a mutant *C. difficile* toxin polypeptide having an identical sequence as the first isolated polypeptide. In another embodiment, the second isolated polypeptide is a mutant *C. difficile* toxin polypeptide having a different sequence from the first isolated polypeptide.

5 In one embodiment, the polypeptide includes at least two modifications selected from the (a)-(d) modifications. In an exemplary embodiment, the polypeptide includes (a) at least one crosslink between a side chain of an aspartic acid residue of the polypeptide and a side chain of a lysine residue of the polypeptide and (b) at least one crosslink between a side chain of a glutamic acid residue of the polypeptide and a side chain of a lysine residue of the polypeptide. In a further embodiment, the polypeptide includes at least three modifications selected from the (a)-(d) modifications. In yet a further embodiment, the polypeptide includes the (a), (b), (c), and (d) modifications.

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When more than one mutant polypeptide is present during chemical modification by EDC, in one embodiment, the resulting composition includes at least one of any of 15 the (a)-(h) modifications. In one embodiment, the composition includes at least two modifications selected from the (a)-(h) modifications. In a further embodiment, the composition includes at least three modifications selected from the (a)-(h) modifications. In yet a further embodiment, the composition includes at least four modifications selected from the (a)-(h) modifications. In another embodiment, the composition 20 includes at least one of each of the (a)-(h) modifications.

In an exemplary embodiment, the resulting composition includes (a) at least one crosslink between a side chain of an aspartic acid residue of the polypeptide and a side chain of a lysine residue of the polypeptide; and (b) at least one crosslink between a side chain of a glutamic acid residue of the polypeptide and a side chain of a lysine residue of the polypeptide. In one embodiment, the composition further includes (c) at least one crosslink between the carboxyl group at the C-terminus of the polypeptide and the amino group of the N-terminus of the polypeptide; and (d) at least one crosslink between the carboxyl group at the C-terminus of the polypeptide and a side chain of a lysine residue of the polypeptide.

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30 In another exemplary embodiment, the resulting composition includes (e) at least one crosslink between a side chain of an aspartic acid residue of the polypeptide and a side chain of a lysine residue of a second isolated polypeptide; (f) at least one crosslink between a side chain of a glutamic acid residue of the polypeptide and a side chain of a

lysine residue of a second isolated polypeptide; **(g)** at least one crosslink between the carboxyl group at the C-terminus of the polypeptide and the amino group of the N-terminus of a second isolated polypeptide; and **(h)** at least one crosslink between the carboxyl group at the C-terminus of the polypeptide and a side chain of a lysine residue 5 of a second isolated polypeptide.

In a further exemplary embodiment, the resulting composition includes **(a)** at least one crosslink between a side chain of an aspartic acid residue of the polypeptide and a side chain of a lysine residue of the polypeptide; **(b)** at least one crosslink between a side chain of a glutamic acid residue of the polypeptide and a side chain of a 10 lysine residue of the polypeptide; **(e)** at least one crosslink between a side chain of an aspartic acid residue of the polypeptide and a side chain of a lysine residue of a second isolated polypeptide; and **(f)** at least one crosslink between a side chain of a glutamic acid residue of the polypeptide and a side chain of a lysine residue of a second isolated polypeptide.

15 In a preferred embodiment, the chemical crosslinking agent includes formaldehyde, more preferably, an agent including formaldehyde in the absence of lysine. Glycine or other appropriate compound with a primary amine can be used as the quencher in crosslinking reactions. Accordingly, in another preferred embodiment, the chemical agent includes formaldehyde and use of glycine.

20 In yet another preferred embodiment, the chemical crosslinking agent includes EDC and NHS. As is known in the art, NHS may be included in EDC coupling protocols. However, the inventors surprisingly discovered that NHS may facilitate in further decreasing cytotoxicity of the mutant *C. difficile* toxin, as compared to the corresponding wild-type toxin, as compared to a genetically mutated toxin, and as 25 compared to a genetically mutated toxin that has been chemically crosslinked by EDC. See, for example, Example 22. Accordingly, without being bound by mechanism or theory, a mutant toxin polypeptide having a beta-alanine moiety linked to a side chain of at least one lysine residue of the polypeptide (e.g., resulting from a reaction of the mutant toxin polypeptide, EDC, and NHS) may facilitate in further decreasing 30 cytotoxicity of the mutant toxin, as compared to, for example, a *C. difficile* toxin (wild-type or mutant) wherein a beta-alanine moiety is absent.

Use of EDC and/or NHS may also include use of glycine or other appropriate compound with a primary amine as the quencher. Any compound having a primary

amine may be used as a quencher, such as, for example glycine methyl ester and alanine. In a preferred embodiment, the quencher compound is a non-polymeric hydrophilic primary amine. Examples of a non-polymeric hydrophilic primary amine include, for example, amino sugars, amino alcohols, and amino polyols. Specific 5 examples of a non-polymeric hydrophilic primary amine include glycine, ethanolamine, glucamine, amine functionalized polyethylene glycol, and amine functionalized ethylene glycol oligomers.

In one aspect, the invention relates to a mutant *C. difficile* toxin polypeptide having at least one amino acid side chain chemically modified by EDC and a non- 10 polymeric hydrophilic primary amine, preferably glycine. The resulting glycine adducts (e.g., from a reaction of triple mutant toxins treated with EDC, NHS, and quenched with glycine) may facilitate in decreasing cytotoxicity of the mutant toxin as compared to the corresponding wild-type toxin.

In one embodiment, when a mutant *C. difficile* toxin polypeptide is chemically 15 modified by EDC and glycine, the polypeptide includes at least one modification when the polypeptide is modified by EDC (e.g., at least one of any of the (a)-(h) modifications described above), and at least one of the following exemplary modifications: (i) a glycine moiety linked to the carboxyl group at the C-terminus of the polypeptide; (j) a glycine moiety linked to a side chain of at least one aspartic acid residue of the 20 polypeptide; and (k) a glycine moiety linked to a side chain of at least one glutamic acid residue of the polypeptide. See, for example, Figure 24 and Figure 25.

In one embodiment, at least one amino acid of the mutant *C. difficile* TcdA is 25 chemically crosslinked and/or at least one amino acid of the mutant *C. difficile* TcdB is chemically crosslinked. Any of the mutant toxins described herein may be chemically crosslinked. In another embodiment, at least one amino acid of SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 7, and/or SEQ ID NO: 8 is chemically crosslinked. In one embodiment, at least one amino acid residue of a polypeptide having the amino acid sequence of any of SEQ ID NOs: 1 through SEQ ID NO: 761 is crosslinked. In another embodiment, at least one amino acid residue of a polypeptide having the amino acid 30 sequence of any of SEQ ID NOs: 183 through SEQ ID NO: 761 includes a modification as described above, e.g., any of the (a)-(k) modifications, such as (a) at least one crosslink between a side chain of an aspartic acid residue of the polypeptide and a side chain of a lysine residue of the polypeptide.

For example, the at least one amino acid may be chemically crosslinked by an agent that includes a carbodiimide, such as EDC. Carbodiimides may form a covalent bond between free carboxyl (e.g., from the side chains of aspartic acid and/or glutamic acid) and amino groups (e.g., in the side chain of lysine residues) to form stable amide bonds.

As another example, the at least one amino acid may be chemically crosslinked by an agent that includes NHS. NHS ester-activated crosslinkers may react with primary amines (e.g., at the N-terminus of each polypeptide chain and/or in the side chain of lysine residues) to yield an amide bond.

10 In another embodiment, the at least one amino acid may be chemically crosslinked by an agent that includes EDC and NHS. For example, in one embodiment, the invention relates to an isolated polypeptide having the amino acid sequence set forth in SEQ ID NO: 4, wherein the methionine residue at position 1 is optionally not present, wherein the polypeptide includes at least one amino acid side chain chemically modified by EDC and NHS. In another embodiment, the invention relates to an isolated polypeptide having the amino acid sequence set forth in SEQ ID NO: 6, wherein the methionine residue at position 1 is optionally not present, wherein the polypeptide includes at least one amino acid side chain chemically modified by EDC and NHS. In yet another embodiment, the invention relates to an isolated polypeptide having the amino acid sequence set forth in SEQ ID NO: 84, SEQ ID NO: 86, SEQ ID NO: 83, SEQ ID NO: 85, SEQ ID NO: 7, or SEQ ID NO: 8. The polypeptide is modified by contacting the polypeptide with EDC and NHS. See, for example, Figure 24 and Figure 25.

25 When a mutant *C. difficile* toxin polypeptide is chemically modified by (e.g., by contacting) EDC and NHS, in one embodiment, the polypeptide includes at least one modification when the polypeptide is modified by EDC (e.g., at least one of any of the (a)-(h) modifications described above), and (I) a beta-alanine moiety linked to a side chain of at least one lysine residue of the polypeptide.

In another aspect, the invention relates to a mutant *C. difficile* toxin polypeptide wherein the polypeptide includes at least one amino acid side chain chemically modified by EDC, NHS, and a non-polymeric hydrophilic primary amine, preferably glycine. In one embodiment, the polypeptide includes at least one modification when the polypeptide is modified by EDC (e.g., at least one of any of the (a)-(h) modifications described above), at least one modification when the polypeptide is modified by glycine

(e.g., at least one of any of the (i)-(k) modifications described above), and (l) a beta-alanine moiety linked to a side chain of at least one lysine residue of the polypeptide. See, for example, Figure 24 and Figure 25.

In one aspect, the invention relates to a mutant *C. difficile* toxin polypeptide, 5 wherein a side chain of at least one lysine residue of the polypeptide is linked to a beta-alanine moiety. In one embodiment, a side chain of a second lysine residue of the polypeptide is linked to a side chain of an aspartic acid residue and/or to a side chain of a glutamic acid residue. The “second” lysine residue of the polypeptide includes a lysine residue of the polypeptide that is not linked to a beta-alanine moiety. The side 10 chain of an aspartic acid and/or the side chain of a glutamic acid to which the second lysine residue is linked may be that of the polypeptide to form an intra-molecular crosslink, or that of a second polypeptide to form an inter-molecular crosslink. In another embodiment, a side chain of at least one aspartic acid residue and/or a side chain of at least one glutamic acid residue of the polypeptide is linked to a glycine 15 moiety. The aspartic acid residue and/or the glutamic acid residue that is linked to a glycine moiety is not also linked to a lysine residue.

In another aspect, the invention relates to a mutant *C. difficile* toxin wherein at least one amino acid side chain of a wild-type *C. difficile* toxin is chemically modified. In one embodiment, at least one amino acid side chain of a wild-type *C. difficile* toxin A 20 and/or at least one amino acid side chain of a wild-type *C. difficile* toxin B is chemically modified by EDC. For example, in one embodiment, TcdA (SEQ ID NO: 1) and/or TcdB (SEQ ID NO: 2) is chemically modified by EDC. In another embodiment, the wild-type toxin is chemically modified by EDC and NHS. In one embodiment, the mutant toxin includes a chemically modified wild-type toxin A, wherein the wild-type toxin A is any 25 one described in Table 1-a. In another embodiment, the mutant toxin includes a chemically modified wild-type toxin B, wherein the wild-type toxin B is any one described in Table 2-a.

As yet another example of a chemically crosslinked mutant *C. difficile* toxin polypeptide, the at least one amino acid may be chemically crosslinked by an agent that 30 includes formaldehyde. Formaldehyde may react with the amino group of an N-terminal amino acid residue and the side-chains of arginine, cysteine, histidine, and lysine. Formaldehyde and glycine may form a Schiff-base adduct, which may attach to primary

N-terminal amino groups, arginine, and tyrosine residues, and to a lesser degree asparagine, glutamine, histidine, and tryptophan residues.

A chemical crosslinking agent is said to reduce cytotoxicity of a toxin if the treated toxin has less toxicity (e.g., about 100%, 99%, 95%, 90%, 80%, 75%, 60%, 5 50%, 25%, or 10% less toxicity) than untreated toxin under identical conditions, as measured, for example, by an *in vitro* cytotoxicity assay, or by animal toxicity.

Preferably, the chemical crosslinking agent reduces cytotoxicity of the mutant *C. difficile* toxin by at least about a 2- \log_{10} reduction, more preferably about a 3- \log_{10} reduction, and most preferably about a 4- \log_{10} or more, relative to the mutant toxin 10 under identical conditions but in the absence of the chemical crosslinking agent. As compared to the wild-type toxin, the chemical crosslinking agent preferably reduces cytotoxicity of the mutant toxin by at least about a 5- \log_{10} reduction, about a 6- \log_{10} reduction, about a 7- \log_{10} reduction, about an 8- \log_{10} reduction, or more.

In another preferred embodiment, the chemically inactivated mutant *C. difficile* 15 toxin exhibits EC₅₀ value of greater than or at least about 50 $\mu\text{g}/\text{ml}$, 100 $\mu\text{g}/\text{ml}$, 200 $\mu\text{g}/\text{ml}$, 300 $\mu\text{g}/\text{ml}$, 400 $\mu\text{g}/\text{ml}$, 500 $\mu\text{g}/\text{ml}$, 600 $\mu\text{g}/\text{ml}$, 700 $\mu\text{g}/\text{ml}$, 800 $\mu\text{g}/\text{ml}$, 900 $\mu\text{g}/\text{ml}$, 1000 $\mu\text{g}/\text{ml}$ or greater, as measured by, for example, an *in vitro* cytotoxicity assay, such as one described herein.

Reaction conditions for contacting the mutant toxin with the chemical crosslinking 20 agent are within the scope of expertise of one skilled in the art, and the conditions may vary depending on the agent used. However, the inventors surprisingly discovered optimal reaction conditions for contacting a mutant *C. difficile* toxin polypeptide with a chemical crosslinking agent, while retaining functional epitopes and decreasing cytotoxicity of the mutant toxin, as compared to the corresponding wild-type toxin.

25 Preferably, the reaction conditions are selected for contacting a mutant toxin with the crosslinking agent, wherein the mutant toxin has a minimum concentration of about 0.5, 0.75, 1.0, 1.25, 1.5, 1.75, 2.0 mg/ml to a maximum of about 3.0, 2.5, 2.0, 1.5, or 1.25 mg/ml . Any minimum value may be combined with any maximum value to define a range of suitable concentrations of a mutant toxin for the reaction. Most preferably, the 30 mutant toxin has a concentration of about 1.0 – 1.25 mg/ml for the reaction.

In one embodiment, the agent used in the reaction has a minimum concentration of about 1 mM, 2 mM, 3 mM, 4 mM, 5 mM, 10 mM, 15 mM, 20 mM, 30 mM, 40 mM, or 50 mM, and a maximum concentration of about 100 mM, 90 mM, 80 mM, 70 mM, 60

mM, or 50 mM. Any minimum value may be combined with any maximum value to define a range of suitable concentrations of the chemical agent for the reaction.

In a preferred embodiment wherein the agent includes formaldehyde, the concentration used is preferably any concentration between about 2 mM to 80 mM, most preferably about 40 mM. In another preferred embodiment wherein the agent includes EDC, the concentration used is preferably any concentration between about 1.3 mM to about 13 mM, more preferably about 2 mM to 3 mM, most preferably about 2.6 mM. In one embodiment, the concentration of EDC is at most 5 g/L, 4 g/L, 3 g/L, 2.5 g/L, 2 g/L, 1.5 g/L, 1.0 g/L, 0.5 g/L based on the total reaction volume, preferably at most 1 g/L, more preferably at most 0.5 g/L.

Exemplary reaction times in which the mutant toxin is contacted with the chemical crosslinking agent include a minimum of about 0.5, 1, 2, 3, 4, 5, 6, 12, 24, 36, 48, or 60 hours, and a maximum of about 14 days, 12 days, 10 days, 7 days, 5 days, 3 days, 2 days, 1 day, or 12 hours, 11, 10, 9, 8, 7, 6, 5, 4, 3, 2, or 1 hour. Any minimum value may be combined with any maximum value to define a range of suitable reaction times.

In a preferred embodiment, the step of contacting the mutant toxin with the chemical crosslinking agent occurs for a period of time that is sufficient to reduce cytotoxicity of the mutant *C. difficile* toxin to an EC₅₀ value of at least about 1000 µg/ml in a suitable human cell, e.g., IMR-90 cells, in a standard *in vitro* cytotoxicity assay, as compared to an identical mutant toxin in the absence of the crosslinking agent. More preferably, the reaction step is carried out for a time that is at least twice as long, and most preferably at least three times as long or more, as the period of time sufficient to reduce the cytotoxicity of the mutant toxin to an EC₅₀ value of at least about 1000 µg/ml in a suitable human cell. In one embodiment, the reaction time does not exceed about 168 hours (or 7 days).

For example, in one embodiment wherein the agent includes formaldehyde, the mutant toxin is preferably contacted with the agent for about 12 hours, which was shown to be an exemplary period of time that was sufficient to reduce cytotoxicity of the mutant *C. difficile* toxin to an EC₅₀ value of at least about 1000 µg/ml in a suitable human cell, e.g., IMR-90 cells, in a standard *in vitro* cytotoxicity assay, as compared to an identical mutant toxin in the absence of the crosslinking agent. In a more preferred embodiment, the reaction is carried out for about 48 hours, which is at least about three

times as long as a sufficient period of time for the reaction. In such an embodiment, the reaction time is preferably not greater than about 72 hours.

In another embodiment wherein the agent includes EDC, the mutant toxin is preferably contacted with the agent for about 0.5 hours, more preferably at least about 1 hour, or most preferably about 2 hours. In one embodiment, the mutant toxin is contacted with EDC for at most about 5 hours, preferably at most about 3 hours, more preferably at most about 2 hours. In such an embodiment, the reaction time is preferably not greater than about 6 hours.

Exemplary pH at which the mutant toxin is contacted with the chemical crosslinking agent include a minimum of about pH 5.5, 6.0, 6.5, 7.0, or 7.5, and a maximum of about pH 8.5, 8.0, 7.5, 7.0, or 6.5. Any minimum value may be combined with any maximum value to define a range of suitable pH. Preferably, the reaction occurs at pH 6.5 to 7.5, preferably at pH 7.0.

Exemplary temperatures at which the mutant toxin is contacted with the chemical crosslinking agent include a minimum of about 2°C, 4°C, 10°C, 20°C, 25°C, or 37°C, and a maximum temperature of about 40°C, 37°C, 30°C, 27°C, 25°C, or 20°C. Any minimum value may be combined with any maximum value to define a range of suitable reaction temperature. Preferably, the reaction occurs at about 20°C to 30°C, most preferably at about 25°C.

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The immunogenic compositions described above may include one mutant *C. difficile* toxin (A or B). Accordingly, the immunogenic compositions can occupy separate vials (e.g., a separate vial for a composition including mutant *C. difficile* toxin A and a 5 separate vial for a composition including mutant *C. difficile* toxin B) in the preparation or kit. The immunogenic compositions may be intended for simultaneous, sequential, or separate use.

In another embodiment, the immunogenic compositions described above may include both mutant *C. difficile* toxins (A and B). Any combination of mutant *C. difficile* 10 toxin A and mutant *C. difficile* toxin B described may be combined for an immunogenic composition. Accordingly, the immunogenic compositions can be combined in a single vial (e.g., a single vial containing both a composition including mutant *C. difficile* TcdA and a composition including mutant *C. difficile* TcdB). Preferably, the immunogenic compositions include a mutant *C. difficile* TcdA and a mutant *C. difficile* TcdB.

15 For example, in one embodiment, the immunogenic composition includes SEQ ID NO: 4 and SEQ ID NO: 6, wherein at least one amino acid of each of SEQ ID NO: 4 and SEQ ID NO: 6 is chemically crosslinked. In another embodiment, the immunogenic composition includes a mutant *C. difficile* toxin A, which includes SEQ ID NO: 4 or SEQ ID NO: 7, and a mutant *C. difficile* toxin B, which comprises SEQ ID NO: 6 or SEQ ID 20 NO: 8, wherein at least one amino acid of each of the mutant *C. difficile* toxins is chemically crosslinked.

In another embodiment, the immunogenic composition includes any sequence selected from SEQ ID NO: 4, SEQ ID NO: 84, and SEQ ID NO: 83, and any sequence selected from SEQ ID NO: 6, SEQ ID NO: 86, and SEQ ID NO: 85. In another 25 embodiment, the immunogenic composition includes SEQ ID NO: 84 and an immunogenic composition including SEQ ID NO: 86. In another embodiment, the immunogenic composition includes SEQ ID NO: 83 and an immunogenic composition including SEQ ID NO: 85. In another embodiment, the immunogenic composition includes SEQ ID NO: 84, SEQ ID NO: 83, SEQ ID NO: 86, and SEQ ID NO: 85.

30 In another embodiment, the immunogenic composition includes a polypeptide having any one sequence selected from SEQ ID NO: 1 to SEQ ID NO: 761, and a second polypeptide having any one sequence selected from SEQ ID NO: 1 to SEQ ID NO: 761.

It is understood that any of the inventive compositions, for example, immunogenic compositions including a mutant toxin A and/or mutant toxin B, can be combined in different ratios or amounts for therapeutic effect. For example, the mutant *C. difficile* TcdA and mutant *C. difficile* TcdB can be present in a immunogenic

5 composition at a ratio in the range of 0.1:10 to 10:0.1, A:B. In another embodiment, for example, the mutant *C. difficile* TcdB and mutant *C. difficile* TcdA can be present in a immunogenic composition at a ratio in the range of 0.1:10 to 10:0.1, B:A. In one preferred embodiment, the ratio is such that the composition includes a greater total amount of a mutant TcdB than a total amount of mutant TcdA.

10 In one aspect, an immunogenic composition is capable of binding to a neutralizing antibody or binding fragment thereof. Preferably, the neutralizing antibody or binding fragment thereof is one described herein below. In one exemplary embodiment, an immunogenic composition is capable of binding to an anti-toxin A antibody or binding fragment thereof, wherein the anti-toxin A antibody or binding

15 fragment thereof includes a variable light chain having the amino acid sequence of SEQ ID NO: 36 and a variable heavy chain having the amino acid sequence of SEQ ID NO: 37. For example, the immunogenic composition may include a mutant *C. difficile* TcdA, SEQ ID NO: 4, or SEQ ID NO: 7. As another example, the immunogenic composition may include SEQ ID NO: 84 or SEQ ID NO: 83.

20 In another exemplary embodiment, an immunogenic composition is capable of binding to an anti-toxin B antibody or binding fragment thereof, wherein the anti-toxin B antibody or binding fragment thereof includes a variable light chain of B8-26 and a variable heavy chain of B8-26. For example, the immunogenic composition may include a mutant *C. difficile* TcdB, SEQ ID NO: 6, or SEQ ID NO: 8. As another example, the

25 immunogenic composition may include SEQ ID NO: 86 or SEQ ID NO: 85.

Recombinant cell

In another aspect, the invention relates to a recombinant cell or progeny thereof.

In one embodiment, the cell or progeny thereof includes a polynucleotide encoding a mutant *C. difficile* TcdA and/or a mutant *C. difficile* TcdB.

5 In another embodiment, the recombinant cell or progeny thereof includes a nucleic acid sequence that encodes a polypeptide having at least about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, preferably about 98%, more preferably about 99% or most preferably about 100% identity to any of SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 7, or SEQ ID NO: 8, when optimally aligned, such as by the programs GAP or

10 BESTFIT using default gap weights. In one embodiment, the recombinant cell or progeny thereof includes a nucleic acid sequence that encodes a polypeptide having at least about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, preferably about 98%, more preferably about 99% or most preferably about 100% identity to any of SEQ ID NOs: 1 through SEQ ID NO: 761.

15 In another embodiment, the recombinant cell or progeny thereof includes a nucleic acid sequence that encodes a polypeptide having at least about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, preferably about 98%, more preferably about 99% or most preferably about 100% identity to any of SEQ ID NO: 84, SEQ ID NO: 86, SEQ ID NO: 83, or SEQ ID NO: 85, when optimally aligned, such as by the programs GAP or

20 BESTFIT using default gap weights.

 In an additional embodiment, the recombinant cell or progeny thereof includes nucleic acid sequence having at least about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, preferably about 98%, more preferably about 99% or most preferably about 100% identity to any of SEQ ID NO: 11, SEQ ID NO: 12, SEQ ID NO: 13, SEQ ID NO: 14, SEQ ID NO: 44, SEQ ID NO: 45, SEQ ID NO: 46, or SEQ ID NO: 47, when optimally aligned, such as by the programs GAP or BESTFIT using default gap weights. The recombinant cell may be derived from any cell useful in the recombinant production of a polypeptide of the present invention, e.g., a prokaryote or a eukaryote. Preferably, the recombinant cell is derived from any cell that is suitable for expressing heterologous nucleic acid sequences greater than about 5000, 6000, preferably about 7000, and more preferably about 8000 nucleotides or more. The prokaryotic host cell may be any gram-negative or gram-positive bacterium. In exemplary embodiments, the prokaryotic host cell lacks an endogenous polynucleotide encoding a toxin and/or spore.

Gram-negative bacteria include, but are not limited to, *Campylobacter*, *E. coli*, *Flavobacterium*, *Fusobacterium*, *Helicobacter*, *Hlyobacter*, *Neisseria*, *Pseudomonas*, *Salmonella*, and *Ureaplasma*. For example, the recombinant cell may be derived from a *Pseudomonas fluorescens* cell, as described in US Patent application publication 5 2010013762, paragraphs [0201]-[0230].

Gram-positive bacteria include, but are not limited to, *Bacillus*, *Clostridium*, *Enterococcus*, *Geobacillus*, *Lactobacillus*, *Lactococcus*, *Oceanobacillus*, *Staphylococcus*, *Streptococcus*, and *Streptomyces*. Preferably, the cell is derived from a *C. difficile* cell.

10 The inventors identified strains of wild-type *C. difficile* that lack an endogenous polynucleotide encoding a *C. difficile* toxin. The strains lacking endogenous toxin A and B genes include the following strains, which are available through the American Type Culture Collection (ATCC) (Manassas, VA): *C. difficile* 1351 (ATCC 43593TM), *C. difficile* 3232 (ATCC BAA-1801TM), *C. difficile* 7322 (ATCC 43601TM), *C. difficile* 5036 15 (ATCC 43603TM), *C. difficile* 4811 (ATCC 43602TM), and *C. difficile* VPI 11186 (ATCC 700057TM).

Accordingly, in one embodiment, the recombinant *C. difficile* cell is derived from a strain described herein. Preferably, the recombinant *C. difficile* cell or progeny thereof is derived from the group consisting of *C. difficile* 1351, *C. difficile* 5036, and *C. difficile* 20 VPI 11186. More preferably, the recombinant *C. difficile* cell or progeny thereof is derived from a *C. difficile* VPI 11186 cell.

In a preferred embodiment, the sporulation gene of the recombinant *C. difficile* cell or progeny thereof is inactivated. Spores may be infective, highly resistant, and facilitate the persistence of *C. difficile* in aerobic environments outside of the host. 25 Spores may also contribute to survival of *C. difficile* inside the host during antimicrobial therapy. Accordingly, a *C. difficile* cell lacking a sporulation gene is useful to produce a safe immunogenic composition for administration to mammals. In addition, use of such cells facilitates safety during manufacturing, e.g., safety to protect the facility, future products, and staff.

30 Examples of sporulation genes for targeted inactivation include, *inter alia*, *spo0A*, *spoIIIE*, σ^E , σ^G , and σ^K . Preferably, the *spo0A* gene is inactivated.

Methods of inactivating a *C. difficile* sporulation gene are known in the art. For example, a sporulation gene may be inactivated by targeted insertion of a selectable

marker, such as, an antibiotic resistance marker. See, for example, Heap et al., *J Microbiol Methods*. 2010 Jan;80(1):49-55; Heap et al., *J. Microbiol. Methods*, 2007 Sept; 70(3):452-464; and Underwood et al., *J Bacteriol.* 2009 Dec;191(23):7296-305. See also, for example, Minton et al., WO2007/148091, entitled, "DNA Molecules and 5 Methods," from pages 33-66, or the corresponding US publication US 20110124109 A1, paragraphs [00137]-[0227].

Compositions relating to Culturing a *C. difficile* Cell

The invention further relates to compositions and methods for use in culturing *C. difficile* and producing *C. difficile* toxins. In one aspect, the invention relates to a culture medium comprising a nitrogen source and a *C. difficile* cell.

5 Suitable culture medium nitrogen sources include: HY-SOY (Quest), AMI-SOY (Quest), NZ-SOY (Quest), NZ-SOY BL4 (Quest), NZ-SOY BL7 (Quest), SHEFTONE D (Sheffield), SE50M (DMV), SE50 (DMV), SE % MK (DMV), SOY PEPTONE (Gibco), BACTO-SOYTON (Difco), NUTRISOY 2207 (ADM), BAKES NUTRISOY (ADM) NUTRISOY FLOUR, Soybean meal, BACTO-YEAST EXTRACT (Difco) YEAST

10 EXTRACT (Gibco), HY-VEST 412 yeast extract(Quest), HY-VEST 441 yeast extract (Quest), HY-VEST 444 yeast extract (Quest), HY-VEST 455 yeast extract (Quest) BACTO-MALT EXTRACT (Difco), Corn Steep, and PROFLO (Traders).

In one aspect, the invention relates to a culture medium including a vegetable hydrolysate and a *C. difficile* cell. Any vegetable hydrolysate may be suitable.

15 Examples of suitable vegetable hydrolysates are cottonseed hydrolysate, pea hydrolysate, and soy hydrolysate.

In a preferred embodiment, the vegetable hydrolysate is soy hydrolysate. Preferably, the soy hydrolysate is SE50MK (Friesland-Campagna). Other examples of soy products that may be used in the invention, and their sources, include:

20 Tekniscience: Soy Peptone A1, Soy Peptone A2, Soy Peptone A3, Plant Peptone E1, Plant Peptone ET1, and Wheat Peptone E1; Quest: HY-Soy, HY-Soy T, AMI-Soy, NZ-Soy, NZ-Soy BLA, and NZ-Soy BL7; DMV: SE50M, SE70M, SE50MK, SE50MK-NK (Friesland-Campagna), WGE80BT, WGE80M, CNE50M, and SE70BT; Marcor: Soy Peptone Type AB, Soy Peptone Type AC, Soy Peptone Type SL, Soy Peptone Type II, and Soy Peptone Type F; Oxoid: Vegetable Peptone and Vegetable Peptone No. 1; Gibco: Soy Peptone; and Difco: Bacsoytone.

25 Concentrations of the vegetable hydrolysate in the culture medium can range, for example, between a minimal value of 5, 10, 20, 30, 40, or 50 g/L to a maximal value of 200, 150, 100, or 75 g/L. Any minimum value can be combined with any maximum value to define a suitable range. In a preferred embodiment, the concentration of vegetable hydrolysate in the culture medium is between 10-50 g/L, most preferably about 30 g/L. The concentration of vegetable hydrolysate in the culture medium described herein may be based on the total volume of the culture medium.

In another aspect, the invention relates to a culture medium including a yeast extract (e.g., as a nitrogen source) and a *Clostridium difficile* cell. Most preferably, the yeast extract is HY YEST 412 (Kerry Biosciences).

Concentrations of the yeast extract in the culture medium can range, for 5 example, between a minimal value of 5, 10, 20, 30, 40, or 50 g/L to a maximal value of 200, 150, 100, or 75 g/L. Any minimum value can be combined with any maximum value to define a suitable range. In a preferred embodiment, the concentration of yeast extract in the culture medium is between 10-50 g/L, most preferably about 20 g/L. The concentration of yeast extract in the culture medium described herein may be based on 10 the total volume of the culture medium.

The inventors discovered that growth of *C. difficile* can be supported in a culture medium including a vegetable hydrolysate without yeast extract, and in a culture medium that includes yeast extract without a vegetable hydrolysate. The culture of cells and/or production of toxin resulting from a culture medium that includes both a 15 vegetable hydrolysate and yeast extract, however, has been observed to be higher than the yields resulting from a culture medium including a vegetable hydrolysate in the absence of yeast extract, and higher than the yields resulting from a culture medium including yeast extract in the absence of a vegetable hydrolysate.

Accordingly, the inventors discovered that a combination of a vegetable 20 hydrolysate and yeast extract helps to support maximal growth of *C. difficile* and/or production of toxin. In one aspect, the invention relates to a culture medium including a vegetable hydrolysate, yeast extract, and a *C. difficile* cell. The vegetable hydrolysate can be any suitable vegetable hydrolysate known in the art as described above. Preferably, the hydrolysate is soy hydrolysate. More preferably, the soy hydrolysate is 25 SE50MK (Friesland-Campaigna). In a preferred embodiment, the yeast extract is HY YEST 412.

In one embodiment, the medium does not include a carbon source. The inventors observed that culture medium including soy hydrolysate / yeast extract in the absence of a carbon source achieves OD₆₀₀ values of 2-3 and toxin production yields of 30 10-15 mg/L.

However, a culture medium that includes a carbon source was surprisingly observed to increase the culture of *C. difficile* cells and production of toxin, as compared to a medium in the absence of a carbon source. In addition, the inventors surprisingly

discovered that production of the mutant toxin was not markedly inhibited by culturing the *C. difficile* cell under anaerobic conditions including a carbon source in the presence of a medium including soy hydrolysate. Accordingly, in a preferred embodiment, the medium further includes a carbon source. More preferably, in one embodiment, the culture medium includes a carbon source, wherein the medium further includes a recombinant *C. difficile* cell. Even more preferably, the recombinant *C. difficile* cell includes a constitutive promoter. In a preferred embodiment, the promoter is a *Clostridium sporogenes* ferredoxin (*fdx*) promoter. In another preferred embodiment, the recombinant *C. difficile* cell does not include a regulated chromosomal promoter that may be negatively regulated by, for example, glucose repression. In a most preferred embodiment, the *C. difficile* cell is a recombinant cell or progeny thereof as described above. Any carbon source may be used in the culture medium. Suitable carbon sources include glucose, dextrose, mannitol, fructose, and/or mannose. Preferably, the carbon source in the culture medium is glucose. In a preferred embodiment, the culture medium does not include arabinose, xylose, sucrose, lactose, maltose, glycerol, rhamnose, and/or galactose.

Concentrations of a carbon source (e.g., glucose, mannitol, fructose, and/or mannose) in the culture medium can range, for example, between a minimal value of 1, 5, 10, 15, 20, 30, 40, 50, or 60 g/L to a maximal value of 150, 100, 90, 80, 75, 70, 50, 40, or 30 g/L. Any minimum value can be combined with any maximum value to define a suitable range. In a preferred embodiment, the concentration of a carbon source in the culture medium is between 5-70 g/L. In one embodiment, the concentration of a carbon source in the culture medium is about 10 g/L. In another embodiment, the concentration of a carbon source in the culture medium is greater than 10 g/L. In another embodiment, the concentration of a carbon source (e.g., glucose) in the culture medium is 45-75 g/L, preferably 55-65 g/L more preferably 55-65 g/L. In another embodiment, the concentration of a carbon source in the culture medium is about 60 g/L. The concentration of carbon source in the culture medium described herein may be based on the total volume of the culture medium.

In one embodiment, the culture medium further includes a chloramphenicol derivative. Exemplary chloramphenicol derivatives include any one of thiamphenicol, florfenicol, chloramphenicol succinate, and fluoramphenicol. Preferably, the culture medium includes thiamphenicol. Without being bound by mechanism or theory, the

chloramphenicol derivative is believed to help in preventing plasmid loss during fermentation and in increasing production of cells and/or toxin, as compared to a culture medium in the absence of a chloramphenicol derivative.

Concentrations of a chloramphenicol derivative in the culture medium can range,

5 for example, between a minimal value of 5, 10, 15, 20, or 30 mg/L to a maximal value of 100, 75, 50, or 40 mg/L. In another embodiment, the concentration of chloramphenicol derivative in the culture medium can range, for example between a minimal value of 0.5 g/L, 1 g/L, 1.5 g/L, 2 g/L, 2.5 g/L, 3 g/L, 3.5 g/L, 4 g/L, 4.5 g/L, or 5 g/L to a maximal value of 10 g/L, 9.5 g/L, 9 g/L, 8.5 g/L, 8 g/L, 7.5 g/L, 7 g/L, 6.5 g/L, 6 g/L, 5.5 g/L, 5 g/L.

10 Any minimum value can be combined with any maximum value to define a suitable range. In a preferred embodiment, the concentration of a chloramphenicol derivative in the culture medium is between 5-20 mg/L, most preferably about 15 mg/L. In another preferred embodiment, the concentration of a chloramphenicol derivative in the culture medium is between 1 g/L-10 g/L, preferably between 1g/L-5 g/L, most preferably about 15 3 g/L. The concentration of chloramphenicol derivative in the culture medium described herein may be based on the total volume of the culture medium.

In one embodiment, the culture medium further includes a cell protectant, such as polyethylene glycol, a polyvinyl alcohol or a pluronic polyol. In one embodiment, the culture medium includes polyethylene glycol, such as polyethylene glycol 2000 (PPG 2000).

Concentrations of a cell protectant, such as polyethylene glycol, in the culture medium can range, for example, between a minimal value of 0.01, 0.05, 0.10, 0.15, 0.20, 0.25, 0.30, 0.35, 0.40, 0.45, 0.50, 0.55, 0.60, 0.65, 0.70, 0.75, 0.80, 0.85, 0.90, 0.95, or 1 ml/L to a maximal value of 2, 1, 0.90, 0.80, 0.70, 0.60, 0.50, 0.40, 0.30, 0.20 25 ml/L. Any minimum value can be combined with any maximum value to define a suitable range. In a preferred embodiment, the concentration of a the cell protectant, such as polyethylene glycol in the culture medium is between 0.01 ml/L to 0.050, most preferably about 0.25 ml/L. The concentration of cell protectant in the culture medium described herein may be based on the total volume of the culture medium.

30 In a preferred embodiment, the culture medium includes a vegetable hydrolysate, yeast extract, and a carbon source, wherein the vegetable hydrolysate is soy hydrolysate, wherein the yeast extract is HY YEST 412 (Kerry Biosciences), and wherein the carbon source is glucose, mannitol, fructose, and/or mannose. In a most

preferred embodiment, the culture medium includes soy hydrolysate SE50MK, HY YEST 412 yeast extract, glucose, and thiampenicol, at pH 7.

In one preferred embodiment, the culture medium includes 30 g/L soy hydrolysate SE50MK, 20 g/L HY YEST 412 yeast extract, 10 g/L glucose, and 15 mg/L thiampenicol, at pH 7. In one preferred embodiment, the culture medium includes about 30 g/L soy hydrolysate SE50MK, about 20 g/L HY YEST 412 yeast extract, about 10 g/L glucose, and about 15 mg/L thiampenicol, at pH 7. The culture medium may further include about 0.25 ml/L PPG 2000. In another embodiment, the culture medium may further include dextrose.

10 In another preferred embodiment, the culture medium includes 30 g/L soy hydrolysate SE50MK, 20 g/L HY YEST 412 yeast extract, 60 g/L glucose, and 15 mg/L thiampenicol, at pH 7. In another preferred embodiment, the culture medium includes about 30 g/L soy hydrolysate SE50MK, about 20 g/L HY YEST 412 yeast extract, about 60 g/L glucose, and about 15 mg/L thiampenicol, at pH 7. The culture medium may

15 further include about 0.25 ml/L PPG 2000. In another embodiment, the culture medium may further include dextrose.

In one embodiment, the culture medium further includes polypropylene glycol 2000 (PPG 2000). The culture medium may include about 0.05 mL PPG 2000 to about 1 ml PPG 2000/L medium, preferably the culture medium includes about 0.25 ml/L PPG

20 2000/L medium.

The culture medium described herein may be suitable for culturing any *C. difficile* cell. In one embodiment, the cell is not genetically modified. In another embodiment, the cell is genetically modified, such as a recombinant cell or progeny thereof as described above. In one embodiment, the *C. difficile* cell lacks an endogenous

25 polynucleotide encoding a toxin. In a preferred embodiment, the *C. difficile* cell is derived from VPI 11186.

In another embodiment, the cell includes a constitutive promoter. In a preferred embodiment, the promoter is a *Clostridium sporogenes* ferredoxin (*fdx*) promoter. In another preferred embodiment, the cell does not include a regulated chromosomal

30 promoter that may be negatively regulated by, for example, glucose repression. In a most preferred embodiment, the *C. difficile* cell is a recombinant cell or progeny thereof as described above.

The inventors also discovered that a monoclonal antibody medium supports growth of *C. difficile*. Accordingly, in one aspect, the invention relates to a culture medium that includes a monoclonal antibody medium. In one embodiment, the medium is SFM4MAbTM medium (Thermo Scientific). The medium was shown to give OD₆₀₀ values of about 10 and toxin production yield was about 40 mg/L.

5 In one embodiment, the pH of the culture medium can range, for example, between a minimal value of 6.0, 6.1, 6.2, 6.3, 6.4, 6.5, 6.6, 6.7, 6.8, 6.9, or 7.0 to a maximal value of 8.0, 7.9, 7.8, 7.7, 7.6, 7.5, 7.4, 7.3, 7.2, or 7.1. Any minimum value can be combined with any maximum value to define a suitable range. In a preferred 10 embodiment, the pH of the culture medium is between 6.0 to 8.0, more preferably, between 6.5 to 7.5. Most preferably, the pH is 7.0.

Suitable pH titrants are known in the art, including, for example, NH₄OH, Na₂CO₃, and NaOH. Preferably, the pH titrant is NaOH.

15 The culture medium may include phosphate-containing ingredients such as Na₂HPO₄, KH₂PO₄. However, in a preferred embodiment, the culture medium does not include a phosphate-containing ingredient.

Methods relating to Culturing a *C. difficile* Cell

In another aspect, the invention relates to a method of culturing *C. difficile*. The method includes culturing a *C. difficile* cell in a medium as described above.

In one embodiment, growth of *C. difficile* according to the methods of the invention proceeds in at least two phases: seed growth and fermentation. A seed culture is first grown by inoculation from a stock culture, e.g., a working cell bank. The seed is used either to inoculate a second seed culture or to inoculate a relatively large fermentation culture. As is understood in the art, the number of seed cultures used may depend, for example, on the size and volume of the fermentation step.

Accordingly, in one aspect, the invention relates to a method of culturing *C. difficile*. The method includes culturing a *C. difficile* cell in a first culture medium under conditions that facilitate growth of the cell; inoculating a second culture medium with all or a portion of said first medium after said first culturing; culturing said inoculated second medium under conditions that facilitate growth of the cell. The method may further include isolating a *C. difficile* toxin from said second medium. In one embodiment, the *C. difficile* is grown in a first culture medium referred to as a seed culture. In one embodiment, the seed culture includes a culture medium as described above and an inoculation from a stock culture that was grown in the medium.

The seed growth phase (or phases) are generally carried out to scale-up the quantity of the microorganism from a stored culture, so that it can be used as an inoculant for the fermentation phase. The seed growth phase can also be carried out to allow relatively dormant microbes in stored cultures to become rejuvenated and to grow into actively growing cultures.

The volume and quantity of viable cells used to inoculate the fermentation culture can be controlled more accurately if taken from an actively growing culture (e.g., a seed culture), rather than if taken from a stored culture.

In addition, more than one (e.g., two or three) seed growth phases can be used to scale-up the quantity of *C. difficile* for inoculation of the fermentation medium. Alternatively, growth of *C. difficile* in the fermentation phase can proceed directly from the stored culture by direct inoculation, if desired.

The first culture medium includes a culture medium as described above. For example, in one embodiment, the first culture medium includes a vegetable hydrolysate and a *C. difficile* cell. Preferably, the vegetable hydrolysate is soy hydrolysate, most

preferably soy hydrolysate SE50MK. In another embodiment, the first culture medium includes a yeast extract and a *C. difficile* cell. Preferably, the yeast extract is HY YEST 412. In a further embodiment, the first culture medium includes a vegetable hydrolysate and a yeast extract. In yet a further embodiment, the first culture medium further

5 includes a carbon source. Preferably, the carbon source is glucose. In one embodiment, the culture medium includes a recombinant *C. difficile* cell when a carbon source is included, wherein the recombinant cell includes a constitutive promoter. In another embodiment, the first culture medium further includes a chloramphenicol derivative. Preferably, the culture medium includes thiamphenicol.

10 Concentrations of the vegetable hydrolysate in the first culture medium (e.g., seed culture medium) can range, for example, between a minimal value of 5, 10, 20, 30, 40, or 50 g/L to a maximal value of 200, 150, 100, or 75 g/L. Any minimum value can be combined with any maximum value to define a suitable range. In a preferred embodiment, the concentration of vegetable hydrolysate in the culture medium is 15 between 10-50 g/L, most preferably about 30 g/L. The concentration of vegetable hydrolysate in the culture medium described herein may be based on the total volume of the culture medium.

20 Concentrations of the yeast extract in the first culture medium (e.g., seed culture medium) can range, for example, between a minimal value of 5, 10, 20, 30, 40, or 50 g/L to a maximal value of 200, 150, 100, or 75 g/L. Any minimum value can be combined with any maximum value to define a suitable range. In a preferred embodiment, the concentration of yeast extract in the culture medium is between 10-50 g/L, most preferably about 20 g/L. The concentration of yeast extract in the culture medium described herein may be based on the total volume of the culture medium.

25 Concentrations of a carbon source in the first culture medium (e.g., seed culture medium) can range, for example, between a minimal value of 1, 5, 10, 15, or 20 g/L to a maximal value of 100, 75, 50, 40, or 30 g/L. Any minimum value can be combined with any maximum value to define a suitable range. In a preferred embodiment, the concentration of a carbon source in the culture medium is between 5-20 g/L, most 30 preferably about 10 g/L.

In one embodiment, the culture medium further includes a chloramphenicol derivative selected from the group consisting of thiamphenicol, florfenicol, chloramphenicol succinate, and fluoramphenicol. Preferably, the culture medium

includes thiamphenicol. Concentrations of a chloramphenicol derivative in the first culture medium (e.g., seed culture medium) can range, for example, between a minimal value of 5, 10, 15, 20, or 30 mg/L to a maximal value of 100, 75, 50, or 40 mg/L. Any minimum value can be combined with any maximum value to define a suitable range.

5 In a preferred embodiment, the concentration of a chloramphenicol derivative in the culture medium is between 5-20 mg/L, most preferably about 15 mg/L.

In one embodiment, the pH of the first culture medium can range, for example, between a minimal value of 6.0, 6.1, 6.2, 6.3, 6.4, 6.5, 6.6, 6.7, 6.8, 6.9, or 7.0 to a maximal value of 8.0, 7.9, 7.8, 7.7, 7.6, 7.5, 7.4, 7.3, 7.2, or 7.1. Any minimum value 10 can be combined with any maximum value to define a suitable range. In a preferred embodiment, the pH of the first culture medium is between 6.0 to 8.0, more preferably, between 6.5 to 7.5. Most preferably, the pH is 7.0.

In a preferred embodiment, the first culture medium includes soy hydrolysate, yeast extract HY YEST 412, glucose, thiamphenicol, at pH 7. More preferably, the 15 culture medium includes 30 g/L soy hydrolysate SE50MK, 20 g/L yeast extract HY YEST 412, 10 g/L glucose, 15 mg/L thiamphenicol, at pH 7.

To start the fermentation phase, a portion or all of a seed culture containing *C. difficile* may be used to inoculate fermentation culture medium. An appropriate concentration of seed culture to use to inoculate fermentation media can be determined 20 by those of skill in this art and can range, for example, from 0.1-10%. As specific examples, concentrations of 0.5, 1, 5, 5.5, 6, 6.25, 6.5, 7, 8, 9, 10% can be used.

Fermentation may be used to produce the maximum amount of cells in a large-scale anaerobic environment. In one embodiment, the *C. difficile* is grown as a 25 fermentation culture. In one embodiment, the fermentation culture was inoculated from a seed culture that was grown in the medium.

The second culture medium includes a culture medium as described above. For example, in one embodiment, the second culture medium includes a vegetable hydrolysate and a *C. difficile* cell. Preferably, the vegetable hydrolysate is soy hydrolysate, most preferably soy hydrolysate SE50MK. In another embodiment, the 30 second culture medium includes a yeast extract and a *C. difficile* cell. Preferably, the yeast extract is HY YEST 412. In a further embodiment, the second culture medium includes a vegetable hydrolysate and a yeast extract. In yet a further embodiment, the second culture medium further includes a carbon source. Preferably, the carbon source

is glucose. In one embodiment, the culture medium includes a recombinant *C. difficile* cell when a carbon source is included, wherein the recombinant cell includes a constitutive promoter. In another embodiment, the second culture medium further includes a chloramphenicol derivative. Preferably, the culture medium includes

5 thiamphenicol.

Concentrations of the vegetable hydrolysate in the second culture medium can range, for example, between a minimal value of 5, 10, 20, 30, 40, or 50 g/L to a maximal value of 200, 150, 100, or 75 g/L. Any minimum value can be combined with any maximum value to define a suitable range. In a preferred embodiment, the

10 concentration of vegetable hydrolysate in the culture medium is between 10-50 g/L, most preferably about 30 g/L. The concentration of vegetable hydrolysate in the culture medium described herein may be based on the total volume of the culture medium.

Concentrations of the yeast extract in the second culture medium can range, for example, between a minimal value of 5, 10, 20, 30, 40, or 50 g/L to a maximal value of

15 200, 150, 100, or 75 g/L. Any minimum value can be combined with any maximum value to define a suitable range. In a preferred embodiment, the concentration of yeast extract in the culture medium is between 10-50 g/L, most preferably about 20 g/L.

Concentrations of a carbon source in the second culture medium can range, for example, between a minimal value of 10, 20, 30, 40, 50, or 60 g/L to a maximal value of

20 150, 100, 90, 80, or 70 g/L. Any minimum value can be combined with any maximum value to define a suitable range. In a preferred embodiment, the concentration of a carbon source in the culture medium is between 50-70 g/L, most preferably about 60 g/L. In one embodiment, the concentration of a carbon source in the second culture medium is greater than the concentration of the carbon source in the first culture

25 medium. The concentration of carbon source in the culture medium described herein may be based on the total volume of the culture medium.

In one embodiment, the culture medium further includes a chloramphenicol derivative selected from the group consisting of thiamphenicol, florfenicol, chloramphenicol succinate, and fluoramphenicol. Preferably, the culture medium

30 includes thiamphenicol. Concentrations of a chloramphenicol derivative in the second culture medium can range, for example, between a minimal value of 5, 10, 15, 20, or 30 mg/L to a maximal value of 100, 75, 50, or 40 mg/L. In another embodiment, the concentration of chloramphenicol derivative in the culture medium can range, for

example between a minimal value of 0.5 g/L, 1 g/L, 1.5 g/L, 2 g/L, 2.5 g/L, 3 g/L, 3.5 g/L, 4 g/L, 4.5 g/L, or 5 g/L to a maximal value of 10 g/L, 9.5 g/L, 9 g/L, 8.5 g/L, 8 g/L, 7.5 g/L, 7 g/L, 6.5 g/L, 6 g/L, 5.5 g/L, 5 g/L. Any minimum value can be combined with any maximum value to define a suitable range. In a preferred embodiment, the

5 concentration of a chloramphenicol derivative in the culture medium is between 5-20 mg/L, most preferably about 15 mg/L. In another preferred embodiment, the concentration of a chloramphenicol derivative in the culture medium is between 1 g/L-10 g/L, preferably between 1g/L-5 g/L, most preferably about 3 g/L. The concentration of chloramphenicol derivative in the culture medium described herein may be based on the
10 total volume of the culture medium.

In one embodiment, the culture medium further includes a cell protectant, such as polyethylene glycol, a polyvinyl alcohol or a pluronic polyol. In one embodiment, the culture medium includes polyethylene glycol, such as polyethylene glycol 2000 (PPG 2000).

15 Concentrations of a cell protectant, such as polyethylene glycol, in the culture medium can range, for example, between a minimal value of 0.01, 0.05, 0.10, 0.15, 0.20, 0.25, 0.30, 0.35, 0.40, 0.45, 0.50, 0.55, 0.60, 0.65, 0.70, 0.75, 0.80, 0.85, 0.90, 0.95, or 1 ml/L to a maximal value of 2, 1, 0.90, 0.80, 0.70, 0.60, 0.50, 0.40, 0.30, 0.20 ml/L. Any minimum value can be combined with any maximum value to define a
20 suitable range. In a preferred embodiment, the concentration of a the cell protectant, such as polyethylene glycol in the culture medium is between 0.01 ml/L to 0.050, most preferably about 0.25 ml/L. The concentration of cell protectant in the culture medium described herein may be based on the total volume of the culture medium.

In one embodiment, the pH of the second culture medium can range, for
25 example, between a minimal value of 6.0, 6.1, 6.2, 6.3, 6.4, 6.5, 6.6, 6.7, 6.8, 6.9, or 7.0 to a maximal value of 8.0, 7.9, 7.8, 7.7, 7.6, 7.5, 7.4, 7.3, 7.2, or 7.1. Any minimum value can be combined with any maximum value to define a suitable range. In a preferred embodiment, the pH of the second culture medium is between 6.0 to 8.0, more preferably, between 6.5 to 7.5. Most preferably, the pH is 7.0.

30 In a preferred embodiment, the second culture medium includes soy hydrolysate, yeast extract HY YEST 412, glucose, thiamphenicol, at pH 7. More preferably, the culture medium includes 30 g/L soy hydrolysate, 20 g/L yeast extract HY YEST 412, 60 g/L glucose, 15 mg/L thiamphenicol, at pH 7.

In one embodiment, the culturing is carried out under anaerobic conditions. In one embodiment, the culturing steps of the methods of the invention (both seed and fermentation) are carried out under anaerobic conditions, although aerobic conditions for either of these phases may be used as well.

5 Approaches to anaerobic culture of bacteria, such as *C. difficile*, are known in the art and can employ, for example, nitrogen gas or a mixture of nitrogen and hydrogen gases. The gas may be bubbled through the medium (e.g., sparging) during fermentation or passed through the area above the liquid in a culture chamber (e.g., the chamber headspace).

10 Culturing of the *C. difficile* cell can be carried out in an anaerobic chamber at approximately $30\pm1^\circ\text{ C}$, $31\pm1^\circ\text{ C}$, $32\pm1^\circ\text{ C}$, $33\pm1^\circ\text{ C}$, $34\pm1^\circ\text{ C}$, $35\pm1^\circ\text{ C}$, $36\pm1^\circ\text{ C}$, $37\pm1^\circ\text{ C}$, $38\pm1^\circ\text{ C}$, or $39\pm1^\circ\text{ C}$, preferably about $37\pm1^\circ\text{ C}$. The culturing can be carried out for times ranging, for example, between a minimal value of 1, 2, 3, 4, 7, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, or 24 hours to a maximal value of 9 days, 8 days, 15 7 days, 6 days, 5 days, 4 days, 3 days, 48 hours, or 36 hours, or 24 hours. Any minimum value can be combined with any maximum value to define a suitable range. In a preferred embodiment, the culturing of the cell occurs for 11 to 48 hours, most preferably about 24 hours. In another embodiment, the culturing of the cell occurs for 5 to 25 hours, 10 to 20 hours, preferably about 15 hours. Growth can be monitored by 20 measuring the optical density (O.D.) of the medium.

In one embodiment, the method further includes isolating *C. difficile* toxins from said medium. *C. difficile* toxins can be isolated and/or purified from cultures using known protein purification methods. The purified toxins can then, for example, be inactivated by chemically inactivation treatment.

25 In an alternative aspect, the invention relates to a method of culturing *C. difficile*. The method includes culturing a *C. difficile* cell in a monoclonal antibody medium.

30 In another aspect, the invention relates to a method for production of a *C. difficile* toxin. The method includes culturing a *C. difficile* bacterium in a medium, as described above, under conditions to permit production of the toxin. The method further includes isolating the *C. difficile* toxin from the medium.

In one embodiment, the method includes culturing a *C. difficile* cell in a first medium under conditions that facilitate growth of *C. difficile*; inoculating a second medium with all or a portion of said first medium after said culturing; culturing said

inoculated second medium under conditions that facilitate growth of *C. difficile* and toxin production; and isolating a *C. difficile* toxin from said second medium. In one embodiment, culturing of the first or second media including *C. difficile* is carried out under anaerobic conditions.

5 In one embodiment, the *C. difficile* cell is cultured in a continuous culture system. In another embodiment, the invention relates to a method of culturing *C. difficile* in a perfusion culture. Surprisingly, the inventors discovered that a recombinant *C. difficile* cell expressing a mutant toxin may be successfully cultured in an anaerobic continuous culture and in an anaerobic perfusion culture. An advantage of a continuous system
10 and/or a perfusion system is that fresh media may be added continuously. In addition, toxin byproducts from production may be removed during production while maintaining cell viability in the system.

The continuous culture system may include providing fresh medium to the cells while simultaneously removing spent medium and cells from the bioreactor. The
15 continuous culture may include a perfusion culture, in which the liquid outflow contains culture medium that is substantially free of cells or includes a substantially lower cell concentration than that in the bioreactor. In a perfusion culture, cells can be retained by, for example, filtration, ultrasonic filtration, centrifugation, or sedimentation.

In one embodiment, the spent media is removed and filtered, which prevents
20 cells from being removed from the bioreactor. The filter may be a cross-flow filter and/or a tangential flow filter. In one embodiment, said filtration system comprises a hollow fiber filter. In another embodiment, the cells are prevented from being removed from the bioreactor by a centrifugation step. In another embodiment, the cells are prevented from being removed from the bioreactor by an ultrasonic filtration step. In
25 another embodiment, the cells are prevented from being removed from the bioreactor via a sedimentation system. In another embodiment, said filtration system comprises a flat-sheet cassette.

In yet another embodiment, the perfusion system comprises a hollow fiber filter that will retain cells, but not the desired product. The cells are recycled back into the
30 bioreactor and the spent media containing the desired product is passed through a desired molecular weight cut-off filter. The filter will retain the desired product. Waste products not retained by the filter can be disposed or recycled.

Method of Producing a Mutant *C. difficile* Toxin

In one aspect, the invention relates to a method of producing a mutant *C. difficile* toxin. In one embodiment, the method includes culturing any recombinant cell or progeny thereof described above, under suitable conditions to express a polypeptide.

5 The method further includes a step of isolating the toxin from the medium.

In another embodiment, the method includes culturing a recombinant cell or progeny thereof under suitable conditions to express a polynucleotide encoding a mutant *C. difficile* toxin, wherein the cell includes the polynucleotide encoding the mutant *C. difficile* toxin, and wherein the mutant includes a glucosyltransferase domain 10 having at least one mutation and a cysteine protease domain having at least one mutation, relative to the corresponding wild-type *Clostridium difficile* toxin. In one embodiment, the cell lacks an endogenous polynucleotide encoding a toxin.

15 In a further embodiment, the method includes culturing a recombinant *C. difficile* cell or progeny thereof under suitable conditions to express a polynucleotide encoding a mutant *C. difficile* toxin, wherein the cell includes the polynucleotide encoding the mutant *C. difficile* toxin and the cell lacks an endogenous polynucleotide encoding a *C. difficile* toxin.

20 In another aspect, the invention relates to a method of producing a mutant *C. difficile* toxin. The method includes the steps of: (a) contacting a *C. difficile* cell with a recombinant *Escherichia coli* cell, wherein the *C. difficile* cell lacks an endogenous 25 polynucleotide encoding a *C. difficile* toxin and the *E. coli* cell includes a polynucleotide that encodes a mutant *C. difficile* toxin; (b) culturing the *C. difficile* cell and the *E. coli* cell under suitable conditions for transfer of the polynucleotide from the *E. coli* cell to the *C. difficile* cell; (c) selecting the *C. difficile* cell comprising the polynucleotide encoding the mutant *C. difficile* toxin; (d) culturing the *C. difficile* cell of step (c) under suitable 30 conditions to express the polynucleotide; and (e) isolating the mutant *C. difficile* toxin.

35 In the inventive method, the recombinant *E. coli* cell includes a heterologous polynucleotide that encodes the mutant *C. difficile* toxin, described herein. The polynucleotide may be DNA or RNA. In one exemplary embodiment, the polynucleotide that encodes the mutant *C. difficile* toxin is codon-optimized for *E. coli* codon usage. Methods for codon-optimizing a polynucleotide are known in the art.

In one embodiment, the polynucleotide includes a nucleic acid sequence that is at least about 60%, 65%, 70%, 75%, 80%, 85%, 86%, 87%, 88%, 89%, 90%, 91%,

92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% identical to a polynucleotide encoding a mutant *C. difficile* TcdA, as described above. In one embodiment, the polynucleotide includes a nucleic acid sequence that is at least about 60%, 65%, 70%, 75%, 80%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 5 98%, 99%, or 100% identical to a polynucleotide encoding a polypeptide having any one sequence selected from SEQ ID NO: 1 to SEQ ID NO: 761. An exemplary polynucleotide encoding a mutant *C. difficile* toxin A includes SEQ ID NO: 11, SEQ ID NO: 12, SEQ ID NO: 44, and SEQ ID NO: 45.

In another embodiment, the polynucleotide includes a nucleic acid sequence that 10 is at least about 60%, 65%, 70%, 75%, 80%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% identical to a polynucleotide encoding a mutant *C. difficile* TcdB, as described above. An exemplary polynucleotide encoding a mutant *C. difficile* toxin B includes SEQ ID NO: 13, SEQ ID NO: 14, SEQ ID NO: 46, and SEQ ID NO: 47. In another embodiment, the polynucleotide encodes SEQ 15 ID NO: 83, SEQ ID NO: 84, SEQ ID NO: 85, or SEQ ID NO: 86.

In one embodiment, the *E. coli* cell that includes the heterologous polynucleotide is an *E. coli* cell that stably hosts the heterologous polynucleotide, which encodes the mutant *C. difficile* toxin. Exemplary *E. coli* cells include a cell selected from the group consisting of MAX Efficiency® Stbl2™ *E. coli* Competent Cells (Invitrogen, Carlsbad, 20 CA), One Shot® Stbl3™ Chemically Competent *E. coli* (Invitrogen, Carlsbad, CA), ElectroMAX™ Stbl4™ *E. coli* Competent Cells (Invitrogen, and *E. coli* CA434. In a preferred embodiment, the *E. coli* cloning host cell is not DH5α. More preferably, the *E. coli* cloning host cell is a MAX Efficiency® Stbl2™ *E. coli* Competent Cell.

The inventive method further includes a step of culturing the *C. difficile* cell and 25 the *E. coli* cell under suitable conditions for transfer of the polynucleotide from the *E. coli* cell to the *C. difficile* cell, resulting in a recombinant *C. difficile* cell. In a preferred embodiment, the culture conditions are suitable for transfer of the polynucleotide from the *E. coli* cell (the donor cell) into the *C. difficile* cell (the recipient cell), and resulting in a genetically stable inheritance.

30 Most preferably, the culture conditions are suitable for bacterial conjugation, which are known in the art. “Conjugation” refers to a particular process of transferring a polynucleotide in which a unidirectional transfer of a polynucleotide (e.g., from a bacterial plasmid) occurs from one bacterium cell (i.e., the “donor”) to another (i.e., the

“recipient”). The conjugation process involves donor cell-to-recipient cell contact.

Preferably, the donor *E. coli* cell is an *E. coli* CA434 cell.

Exemplary suitable (conjugation) conditions for transferring of the polynucleotide from the *E. coli* cell to the *C. difficile* cell include growing liquid cultures of *C. difficile* in

5 brain heart infusion broth (BHI; Oxoid) or Schaedlers anaerobic broth (SAB; Oxoid). In another embodiment, solid *C. difficile* cultures may be grown on fresh blood agar (FBA) or BHI agar. Preferably, the *C. difficile* is grown at 37°C in an anaerobic environment (e.g., 80% N₂, 10% CO₂, and 10% H₂ [vol/vol]). In one embodiment, the suitable condition includes growing the *E. coli* aerobically in Luria-Bertani (LB) broth or on LB 10 agar at 37°C. For conjugative transfer to *C. difficile*, an exemplary suitable condition includes growing *E. coli* anaerobically on FBA. Antibiotics may be included in the liquid and solid media as is known in the art. Examples of such antibiotics include cycloserine (250 µg/ml), cefoxitin (8 µg/ml), chloramphenicol (12.5 µg/ml), thiampenicol (15 µg/ml), and erythromycin (5 µg/ml).

15 The inventive method additionally includes a step of selecting the resulting recombinant *C. difficile* cell that includes the polynucleotide encoding the mutant *C. difficile* toxin. In an exemplary embodiment, the recombinant *C. difficile* cell is a recipient of the polynucleotide encoding the mutant *C. difficile* toxin from the recombinant *E. coli* cell via conjugation.

20 The inventive method includes a step of culturing the recombinant cell or progeny thereof under suitable conditions to express the polynucleotide encoding the mutant *C. difficile* toxin, resulting in production of a mutant *C. difficile* toxin. Suitable conditions for a recombinant cell to express the polynucleotide include culture conditions suitable for growing a *C. difficile* cell, which are known in the art. For example, suitable conditions 25 may include culturing the *C. difficile* transformants in brain heart infusion broth (BHI; Oxoid) or Schaedlers anaerobic broth (SAB; Oxoid). In another embodiment, solid *C. difficile* cultures may be grown on FBA or BHI agar. Preferably, the *C. difficile* is grown at 37°C in an anaerobic environment (e.g., 80% N₂, 10% CO₂, and 10% H₂ [vol/vol]).

In one embodiment, the inventive method includes a step of isolating the

30 resulting mutant *C. difficile* toxin. Methods of isolating a protein from *C. difficile* are known in the art.

In another embodiment, the method includes a step of purifying the resulting mutant *C. difficile* toxin. Methods of purifying a polypeptide, such as chromatography, are known in the art.

In an exemplary embodiment, the method further includes a step of contacting 5 the isolated mutant *Clostridium difficile* toxin with a chemical crosslinking agent described above. Preferably, the agent includes formaldehyde, ethyl- 3-(3-dimethylaminopropyl) carbodiimide, or a combination of EDC and NHS. Exemplary reaction conditions are described above and in the Examples section below.

In another aspect, the invention relates to an immunogenic composition including 10 a mutant *C. difficile* toxin described herein, produced by any method, preferably by any of the methods described above.

Antibodies

Surprisingly, the inventive immunogenic compositions described above elicited novel antibodies *in vivo*, suggesting that the immunogenic compositions include a preserved native structure (e.g., a preserved antigenic epitope) of the respective wild-type *C. difficile* toxin and that the immunogenic compositions include an epitope. The antibodies produced against a toxin from one strain of *C. difficile* may be capable of binding to a corresponding toxin produced by another strain of *C. difficile*. That is, the antibodies and binding fragments thereof may be “cross-reactive,” which refers to the ability to react with similar antigenic sites on toxins produced from multiple *C. difficile* strains. Cross-reactivity also includes the ability of an antibody to react with or bind an antigen that did not stimulate its production, i.e., the reaction between an antigen and an antibody that was generated against a different but similar antigen.

In one aspect, the inventors surprisingly discovered monoclonal antibodies having a neutralizing effect on *C. difficile* toxins, and methods of producing the same.

The inventive antibodies can neutralize *C. difficile* toxin cytotoxicity *in vitro*, inhibit binding of *C. difficile* toxin to mammalian cells, and/or can neutralize *C. difficile* toxin enterotoxicity *in vivo*. The present invention also relates to isolated polynucleotides that include nucleic acid sequences encoding any of the foregoing. In addition, the present invention relates to use of any of the foregoing compositions to treat, prevent, decrease the risk of, decrease severity of, decrease occurrences of, and/or delay the onset of a *C. difficile* infection, *C. difficile* associated disease, syndrome, condition, symptom, and/or complication thereof in a mammal, as compared to a mammal to which the composition is not administered, as well as methods for preparing said compositions.

The inventors further discovered that a combination of at least two of the neutralizing monoclonal antibodies can exhibit an unexpectedly synergistic effect in respective neutralization of TcdA or TcdB. Anti-toxin antibodies or binding fragments thereof can be useful in the inhibition of a *C. difficile* infection.

An “antibody” is a protein including at least one or two heavy (H) chain variable regions (abbreviated herein as VH), and at least one or two light (L) chain variable regions (abbreviated herein as VL). The VH and VL regions can be further subdivided into regions of hypervariability, termed “complementarity determining regions” (“CDR”), interspersed with regions that are more conserved, termed “framework regions” (FR). The extent of the framework region and CDRs has been precisely defined (see, Kabat,

E. A., et al. *Sequences of Proteins of Immunological Interest*, Fifth Edition, U.S.

Department of Health and Human Services, NIH Publication No. 91-3242, 1991, and

Chothia, C. et al., *J. Mol. Biol.* 196:901-917, 1987). The term "antibody" includes intact immunoglobulins of types IgA, IgG, IgE, IgD, IgM (as well as subtypes thereof), wherein the light chains of the immunoglobulin may be of types kappa or lambda.

The antibody molecules can be full-length (e.g., an IgG1 or IgG4 antibody). The antibodies can be of the various isotypes, including: IgG (e.g., IgG1, IgG2, IgG3, IgG4), IgM, IgA1, IgA2, IgD, or IgE. In one preferred embodiment, the antibody is an IgG isotype, e.g., IgG1. In another preferred embodiment, the antibody is an IgE antibody.

10 In another embodiment, the antibody molecule includes an "antigen-binding fragment" or "binding fragment," as used herein, which refers to a portion of an antibody that specifically binds to a toxin of *C. difficile* (e.g., toxin A). The binding fragment is, for example, a molecule in which one or more immunoglobulin chains is not full length, but which specifically binds to a toxin.

15 Examples of binding portions encompassed within the term "binding fragment" of an antibody include (i) a Fab fragment, a monovalent fragment consisting of the VL, VH, CL and CH1 domains; (ii) a F(ab')₂ fragment, a bivalent fragment comprising two Fab fragments linked by a disulfide bridge at the hinge region; (iii) a Fd fragment consisting of the VH and CH1 domains; (iv) a Fv fragment consisting of the VL and VH domains of a single arm of an antibody, (v) a dAb fragment (Ward et al., *Nature* 341:544-546, 20
1989), which consists of a VH domain; and (vi) an isolated complementarity determining region (CDR) having sufficient framework to specifically bind, e.g., an antigen binding portion of a variable region.

25 A binding fragment of a light chain variable region and a binding fragment of a heavy chain variable region, e.g., the two domains of the Fv fragment, VL and VH, can be joined, using recombinant methods, by a synthetic linker that enables them to be made as a single protein chain in which the VL and VH regions pair to form monovalent molecules (known as single chain Fv (scFv); see e.g., Bird et al. (1988) *Science* 242:423-426; and Huston et al. (1988) *Proc. Natl. Acad. Sci. USA* 85:5879-5883). Such 30 single chain antibodies are also encompassed within the term "binding fragment" of an antibody. These antibody portions are obtained using techniques known in the art, and the portions are screened for utility in the same manner as are intact antibodies.

As used herein, an antibody that “specifically binds” to or is “specific” for a particular polypeptide or an epitope on a particular polypeptide is an antibody that binds to that particular polypeptide or epitope on a particular polypeptide without substantially binding to any other polypeptide or polypeptide epitope. For example, when referring to 5 a biomolecule (e.g., protein, nucleic acid, antibody, etc.) that “specifically binds” to a target, the biomolecule binds to its target molecule and does not bind in a significant amount to other molecules in a heterogeneous population of molecules that include the target, as measured under designated conditions (e.g. immunoassay conditions in the case of an antibody). The binding reaction between the antibody and its target is 10 determinative of the presence of the target in the heterogeneous population of molecules. For example, “specific binding” or “specifically binds” refers to the ability of an antibody or binding fragment thereof to bind to a wild-type and/or mutant toxin of *C. difficile* with an affinity that is at least two-fold greater than its affinity for a non-specific antigen.

15 In an exemplary embodiment, the antibody is a chimeric antibody. A chimeric antibody can be produced by recombinant DNA techniques known in the art. For example, a gene encoding the Fc constant region of a murine (or other species) monoclonal antibody molecule can be digested with restriction enzymes to remove the region encoding the murine Fc, and the equivalent portion of a gene encoding a human 20 Fc constant region is substituted. A chimeric antibody can also be created by recombinant DNA techniques where DNA encoding murine variable regions can be ligated to DNA encoding the human constant regions.

25 In another exemplary embodiment, the antibody or binding fragment thereof is humanized by methods known in the art. For example, once murine antibodies are obtained, a CDR of the antibody may be replaced with at least a portion of a human CDR. Humanized antibodies can also be generated by replacing sequences of the murine Fv variable region that are not directly involved in antigen binding with equivalent sequences from human Fv variable regions. General methods for generating humanized antibodies are known in the art.

30 For example, monoclonal antibodies directed toward *C. difficile* TcdA or *C. difficile* TcdB can also be produced by standard techniques, such as a hybridoma technique (see, e.g., Kohler and Milstein, 1975, *Nature*, 256: 495-497). Briefly, an immortal cell line is fused to a lymphocyte from a mammal immunized with *C. difficile*

TcdA, *C. difficile* TcdB, or a mutant *C. difficile* toxin described herein, and the culture supernatants of the resulting hybridoma cells are screened to identify a hybridoma producing a monoclonal antibody that binds to *C. difficile* TcdA or *C. difficile* TcdB. Typically, the immortal cell line is derived from the same mammalian species as the 5 lymphocytes. Hybridoma cells producing a monoclonal antibody of the invention are detected by screening the hybridoma culture supernatants for antibodies that bind *C. difficile* TcdA or *C. difficile* TcdB using an assay, such as ELISA. Human hybridomas can be prepared in a similar way.

As an alternative to producing antibodies by immunization and selection, 10 antibodies of the invention may also be identified by screening a recombinant combinatorial immunoglobulin library with a *C. difficile* TcdA, *C. difficile* TcdB, or a mutant *C. difficile* toxin described herein. The recombinant antibody library may be an scFv library or an Fab library, for example. Moreover, the inventive antibodies described herein may be used in competitive binding studies to identify additional anti- 15 TcdA or anti-TcdB antibodies and binding fragments thereof. For example, additional anti-TcdA or anti-TcdB antibodies and binding fragments thereof may be identified by screening a human antibody library and identifying molecules within the library that competes with the inventive antibodies described herein in a competitive binding assay.

In addition, antibodies encompassed by the present invention include 20 recombinant antibodies that may be generated by using phage display methods known in the art. In phage display methods, phage can be used to display antigen binding domains expressed from a repertoire or antibody library (e.g., human or murine). Phage expressing an antigen binding domain that binds to an immunogen described herein (e.g., a mutant *C. difficile* toxin) can be selected or identified with antigen, e.g., 25 using labeled antigen.

Also within the scope of the invention are antibodies and binding fragments thereof in which specific amino acids have been substituted, deleted, or added. In particular, preferred antibodies have amino acid substitutions in the framework region, such as to improve binding to the antigen. For example, a selected, small number of 30 acceptor framework residues of the immunoglobulin chain can be replaced by the corresponding donor amino acids. Preferred locations of the substitutions include amino acid residues adjacent to the CDR, or which are capable of interacting with a CDR. Criteria for selecting amino acids from the donor are described in U.S. Pat. No.

5,585,089 (e.g., columns 12-16). The acceptor framework can be a mature human antibody framework sequence or a consensus sequence.

As used herein, a "neutralizing antibody or binding fragment thereof" refers to a respective antibody or binding fragment thereof that binds to a pathogen (e.g., a *C. difficile* TcdA or TcdB) and reduces the infectivity and/or an activity of the pathogen (e.g., reduces cytotoxicity) in a mammal and/or in cell culture, as compared to the pathogen under identical conditions in the absence of the neutralizing antibody or binding fragment thereof. In one embodiment, the neutralizing antibody or binding fragment thereof is capable of neutralizing at least about 70%, 75%, 80%, 85%, 90%, 95%, 99%, or more of a biological activity of the pathogen, as compared to the biological activity of the pathogen under identical conditions in the absence of the neutralizing antibody or binding fragment thereof.

As used herein, the term "anti-toxin antibody or binding fragment thereof" refers to an antibody or binding fragment thereof that binds to the respective *C. difficile* toxin (e.g., a *C. difficile* toxin A or toxin B). For example, an anti-toxin A antibody or binding fragment thereof refers to an antibody or binding fragment thereof that binds to TcdA.

The antibodies or binding fragments thereof described herein may be raised in any mammal, wild-type and/or transgenic, including, for example, mice, humans, rabbits, and goats.

When an immunogenic composition described above is one that has been previously administered to a population, such as for vaccination, the antibody response generated in the subjects can be used to neutralize toxins from the same strain and from a strain that did not stimulate production of the antibody. See, for example, Example 37, which shows studies relating to cross-reactivity, generated by the immunogenic composition, between the 630 strain and toxins from various wild-type *C. difficile* strains.

In one aspect, the invention relates to an antibody or binding fragment thereof specific to *C. difficile* TcdA. Monoclonal antibodies that specifically bind to TcdA include A65-33; A60-22; A80-29 and/or, preferably, A3-25.

In one aspect, the invention relates to an antibody or binding fragment thereof specific to a TcdA from any wild type *C. difficile* strain, such as those described above, e.g., to SEQ ID NO: 1. In another aspect, the invention relates to an antibody or binding fragment thereof specific to an immunogenic composition described above. For

example, in one embodiment, the antibody or binding fragment thereof is specific to an immunogenic composition that includes SEQ ID NO: 4 or SEQ ID NO: 7. In another embodiment, the antibody or binding fragment thereof is specific to an immunogenic composition that includes SEQ ID NO: 4 or SEQ ID NO: 7, wherein at least one amino acid of SEQ ID NO: 4 or SEQ ID NO: 7 is crosslinked by formaldehyde, EDC, NHS, or a combination of EDC and NHS. In another embodiment, the antibody or binding fragment thereof is specific to an immunogenic composition that includes SEQ ID NO: 84 or SEQ ID NO: 83.

Antibodies or binding fragments thereof having a variable heavy chain and 10 variable light chain regions that are at least about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, preferably about 98%, more preferably about 99% or most preferably about 100% identity to the variable heavy and light chain regions of A65-33; A60-22; A80-29 and/or, preferably, A3-25 can also bind to TcdA.

In one embodiment, the antibody or antigen binding fragment thereof includes a 15 variable heavy chain region including an amino acid sequence at least about 60%, 65%, 70%, 75%, 80%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% identical to a variable heavy chain region amino acid sequence of A3-25 as set forth in SEQ ID NO: 37.

In another embodiment, the antibody or antigen binding fragment thereof 20 includes a variable light chain region including an amino acid sequence at least about 60%, 65%, 70%, 75%, 80%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% identical to a variable light chain region amino acid sequence of A3-25 as set forth in SEQ ID NO: 36.

In yet a further aspect, the antibody or antigen binding fragment thereof includes 25 a variable heavy chain region including an amino acid sequence at least about 60%, 65%, 70%, 75%, 80%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% identical to a variable heavy chain region amino acid sequence set forth in SEQ ID NO: 37, and a variable light chain region including an amino acid sequence at least about 60%, 65%, 70%, 75%, 80%, 85%, 86%, 87%, 88%, 30 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% identical to a variable light chain region amino acid sequence set forth in SEQ ID NO: 36.

In another embodiment, antibodies or binding fragments thereof having 35 complementarity determining regions (CDRs) of variable heavy chains and/or variable

light chains of A65-33; A60-22; A80-29 and/or, preferably, A3-25 can also bind to TcdA. The CDRs of the variable heavy chain region of A3-25 are shown in Table 4, below.

Table 4: Variable Heavy Chain CDR Amino Acid Sequences

Clone	Chain	CDR	Amino Acid Sequence	SEQ ID NO:
A3-25	Heavy	CDR1	GFTFTNYWMN	41
		CDR2	EIRLKSHNYATHFAESVKG	42
		CDR3	DYYGNPAFVY	43

The CDRs of the variable light chain region of A3-25 are shown in Table 5, 5 below.

Table 5: Variable Light Chain CDR Amino Acid Sequences

Clone	Chain	CDR	Amino Acid Sequence	SEQ ID NO:
A3-25	Light	CDR1	RSSQSLIHSNGNTYLH	38
		CDR2	KVSNRFS	39
		CDR3	SQTTYFPYT	40

In one embodiment, the antibody or binding fragment thereof includes amino acid sequences of the heavy chain complementarity determining regions (CDRs) as shown in SEQ ID NOs: 41 (CDR H1), 42 (CDR H2) and 43 (CDR H3), and/or the amino acid 10 sequences of the light chain CDRs as shown in SEQ ID NOs: 38 (CDR L1), 39 (CDR L2) and 40 (CDR L3).

In one exemplary embodiment, the antibody or binding fragment thereof specific to *C. difficile* toxin A specifically binds to an epitope within the N-terminal region of TcdA e.g., an epitope between amino acids 1-1256 of a TcdA, according to the numbering of 15 SEQ ID NO: 1.

In a preferred embodiment, the antibody or binding fragment thereof specific to *C. difficile* toxin A specifically binds to an epitope within the C-terminal region of toxin A, e.g., an epitope between amino acids 1832 to 2710 of a TcdA, according to the numbering of SEQ ID NO: 1. Examples include A3-25; A65-33; A60-22; A80-29.

20 In yet another embodiment, the antibody or binding fragment thereof specific to *C. difficile* toxin A specifically binds to an epitope within the “translocation” region of *C. difficile* toxin A, e.g., an epitope that preferably includes residues 956-1128 of a TcdA,

according to the numbering of SEQ ID NO: 1, such as an epitope between amino acids 659-1832 of a TcdA, according to the numbering of SEQ ID NO: 1.

In another aspect, the invention relates to an antibody or binding fragment thereof specific to *C. difficile* TcdB. For example, the antibody or binding fragment thereof may be specific to a TcdB from any wild type *C. difficile* strain, such as those described above, e.g., to SEQ ID NO: 2. In another aspect, the invention relates to an antibody or binding fragment thereof specific to an immunogenic composition described above. For example, in one embodiment, the antibody or binding fragment thereof is specific to an immunogenic composition that includes SEQ ID NO: 6 or SEQ ID NO: 8.

10 In another embodiment, the antibody or binding fragment thereof is specific to an immunogenic composition that includes SEQ ID NO: 6 or SEQ ID NO: 8, wherein at least one amino acid of SEQ ID NO: 6 or SEQ ID NO: 8 is crosslinked by formaldehyde, EDC, NHS, or a combination of EDC and NHS. In another embodiment, the antibody or binding fragment thereof is specific to an immunogenic composition that includes SEQ 15 ID NO: 86 or SEQ ID NO: 85.

Monoclonal antibodies that specifically bind to TcdB include antibodies produced by the B2-31; B5-40, B70-2; B6-30; B9-30; B59-3; B60-2; B56-6; and/or, preferably, B8-26 clones described herein.

20 Antibodies or binding fragments thereof that can also bind to TcdB include those having a variable heavy chain and variable light chain regions that are at least about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, preferably about 98%, more preferably about 99% or most preferably about 100% identity to the variable heavy and light chain regions of B2-31; B5-40, B70-2; B6-30; B9-30; B59-3; B60-2; B56-6, preferably B8-26, B59-3, and/or B9-30.

25 In one embodiment, the antibody or antigen binding fragment thereof includes a variable heavy chain region including an amino acid sequence at least about 60%, 65%, 70%, 75%, 80%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% identical to a variable heavy chain region amino acid sequence of A3-25 as set forth in SEQ ID NO: 49.

30 In one embodiment, the antibody or antigen binding fragment thereof includes a variable heavy chain region including an amino acid sequence at least about 60%, 65%, 70%, 75%, 80%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%,

97%, 98%, 99%, or 100% identical to a variable heavy chain region amino acid sequence of A3-25 as set forth in SEQ ID NO: 60.

In one embodiment, the antibody or antigen binding fragment thereof includes a variable heavy chain region including an amino acid sequence at least about 60%, 65%, 5 70%, 75%, 80%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% identical to a variable heavy chain region amino acid sequence of A3-25 as set forth in SEQ ID NO: 71.

In another embodiment, the antibody or antigen binding fragment thereof includes a variable light chain region including an amino acid sequence at least about 10 60%, 65%, 70%, 75%, 80%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% identical to a variable light chain region amino acid sequence of A3-25 as set forth in SEQ ID NO: 55.

In another embodiment, the antibody or antigen binding fragment thereof includes a variable light chain region including an amino acid sequence at least about 15 60%, 65%, 70%, 75%, 80%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% identical to a variable light chain region amino acid sequence of A3-25 as set forth in SEQ ID NO: 66.

In another embodiment, the antibody or antigen binding fragment thereof includes a variable light chain region including an amino acid sequence at least about 20 60%, 65%, 70%, 75%, 80%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% identical to a variable light chain region amino acid sequence of A3-25 as set forth in SEQ ID NO: 77.

The amino acid sequence for the variable heavy chain of a neutralizing antibody of *C. difficile* TcdB (B8-26 mAb) is set forth in SEQ ID NO: 49. See Table 25-a.

Table 25-a: Variable Heavy Chain Amino Acid Sequences

Clone	Region	Amino Acid Sequence	SEQ ID NO:
B8-26	Signal peptide	MGWSCIILFLVATATGVHS	50
	Variable heavy chain	QVQLQQPGAEVLVKPG PVKLSCKAS GYSFTSYWMN WVKQRPGRGLEWIG RIDPSNSEIYYNQKF KDKATLTVDKSSSTAYIQLSSL TSEDSAVYYCAS GHYGSIFAY WGQGTTLVSS	49
	CDR1	GYSFTSYWMN	51
	CDR2	RIDPSNSEIYYNQKF	52
	CDR3	GHYGSIFAY	53
	Constant region (IgG1)	AKTTPPSVYPLAPGNSK	54

The amino acid sequence for the variable light chain of a neutralizing antibody of *C. difficile* TcdB (B8-26 mAb) is set forth in SEQ ID NO: 55. See Table 25-b.

Table 25-b: Variable Light (κ) Chain Amino Acid Sequences			
Clone	Region	Amino Acid Sequence	SEQ ID NO:
B8-26	Signal peptide	MRFQVQVLGLLLLWISGAQCD	56
	Variable light chain	VQITQSPSYLAASPGETITINC RASKSISKYLA WYQEKGKTNKLLLY SGSTLQS GIPS RFSGSRSRGTDFTLISSLEPEDSAMYYC QQHNEYPLT FGAGTKLELKRADAAPTVSIFPPSSEEFQ	55
	CDR1	RASKSISKYLA	57
	CDR2	SGSTLQS	58
	CDR3	QQHNEYPLT	59

In one embodiment, the antibody or binding fragment thereof includes amino acid sequences of the heavy chain CDRs as shown in SEQ ID NOs: 51 (CDR H1), 52 (CDR H2) and 53 (CDR H3), and/or the amino acid sequences of the light chain CDRs as shown in SEQ ID NOs: 57 (CDR L1), 58 (CDR L2) and 59 (CDR L3).

5

The amino acid sequence for the variable heavy chain of a neutralizing antibody of *C. difficile* TcdB (B59-3 mAb) is set forth in SEQ ID NO: 60. See Table 26-a.

Table 26-a: Variable Heavy Chain Amino Acid Sequences

Clone	Region	Amino Acid Sequence	SEQ ID NO:
B59-3	Signal peptide	MGWSYIILFLVATATDVHS	61
	Variable heavy chain	QVQLQQPGAEVLVKPGASVKLS CKAS GYTFTSYWMH WVKQRPGQGLEWIG VINPSNGRSTYSEKF KTTATVTVDKSSSTAYMQL SILTSEDSAVYYCAR AYYSTSYYAMDY WGQGTSVTVSS	60
	CDR1	GYTFTSYWMH	62
	CDR2	VINPSNGRSTYSEKF	63
	CDR3	AYYSTSYYAMDY	64
	Constant region (IgG1)	AKTTPPSVYPLAPGNSK	65

The amino acid sequence for the variable light chain of a neutralizing antibody of 5 *C. difficile* TcdB (B59-3 mAb) is set forth in SEQ ID NO: 66. See Table 26-b.

Table 26-b: Variable Light (κ) Chain Amino Acid Sequences

Clone	Region	Amino Acid Sequence	SEQ ID NO:
B59-3	Signal peptide	MKLPVRLLVLMFWIPASSSD	67
	Variable light chain	VLMTQSPLSLPVSLGDQASIS C RSSQNIVHSNGNTYLE WYLQKPGQSPKLLIY KVSNRFS GVPDRFSGSGSGTYFTLKISRVEAEDLGVYYC FQGSHFPFT FGTGTKLEIKRADAAPTVSIFPPSSEEFQ	66
	CDR1	RSSQNIVHSNGNTYLE	68
	CDR2	KVSNRFS	69
	CDR3	FQGSHFPFT	70

In one embodiment, the antibody or binding fragment thereof includes amino acid sequences of the heavy chain CDRs as shown in SEQ ID NOs: 62 (CDR H1), 63 (CDR H2) and 64 (CDR H3), and/or the amino acid sequences of the light chain CDRs as shown in SEQ ID NOs: 68 (CDR L1), 69 (CDR L2) and 70 (CDR L3).

5 The amino acid sequence for the variable heavy chain of a neutralizing antibody of *C. difficile* TcdB (B9-30 mAb) is set forth in SEQ ID NO: 71. See Table 27-a.

Table 27-a: Variable Heavy Chain Amino Acid Sequences

Clone	Region	Amino Acid Sequence	SEQ ID NO:
B9-30	Signal peptide	MGWSCIILFLVATATGVHS	72
	Variable heavy chain	QVQLQQPGAEVVKPGAPVKLS CKAS GYPFTNYWMN WVKQRPGRGLEWIG RIDPSNSEIYYNQKF KDKATLTVDKSSSTAYIQLSSLTSEDSAVYY CAS GHYGSIFAY WGQGTTLTVSS	71
	CDR1	GYPFTNYWMN	73
	CDR2	RIDPSNSEIYYNQKF	74
	CDR3	GHYGSIFAY	75
	Constant region (IgG1)	AKTTPPSVYPLAPGNSK	76

The amino acid sequence for the variable light chain of a neutralizing antibody of *C. difficile* TcdB (B9-30 mAb) is set forth in SEQ ID NO: 77. See Table 27-b.

Table 27-b: Variable Light (κ) Chain Amino Acid Sequences			
Clone	Region	Amino Acid Sequence	SEQ ID NO:
B9-30	Signal peptide	MRFQVQVLGLLLLWISGAQCD	78
	Variable light chain	VQITQSPSYLAASPGETITINC RASKSISKYLA WYQEKGPKTNKLLIY SGSTLQS GIPS RFSGSRSRGTDFTLISSLEPEDSAMYYC QQHNEYPLT FGAGTKLELKRADAAPTVSIFPPSSEEFQ	77
	CDR1	RASKSISKYLA	79
	CDR2	SGSTLQS	80
	CDR3	QQHNEYPLT	81

In one embodiment, the antibody or binding fragment thereof includes amino acid sequences of the heavy chain CDRs as shown in SEQ ID NOs: 73 (CDR H1), 74 (CDR H2) and 75 (CDR H3), and/or the amino acid sequences of the light chain CDRs as shown in SEQ ID NOs: 79 (CDR L1), 80 (CDR L2) and 81 (CDR L3).

In one aspect, the invention relates to an antibody or binding fragment thereof specific to a wild type *C. difficile* TcdB from any *C. difficile* strain, such as those described above, e.g., to SEQ ID NO: 2. In another aspect, the invention relates to an antibody or binding fragment thereof specific to an immunogenic composition described above. For example, in one embodiment, the antibody or binding fragment thereof is specific to an immunogenic composition that includes SEQ ID NO: 6 or SEQ ID NO: 8. In another embodiment, the antibody or binding fragment thereof is specific to an immunogenic composition that includes SEQ ID NO: 6 or SEQ ID NO: 8, wherein at least one amino acid of SEQ ID NO: 6 or SEQ ID NO: 8 is crosslinked by formaldehyde, EDC, NHS, or a combination of EDC and NHS.

Antibodies or binding fragments thereof having a variable heavy chain and variable light chain regions that are at least about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, preferably about 98%, more preferably about 99% or most preferably about 100% identity to the variable heavy and light chain regions of B2-31; B5-40, B70-2; B6-30; B9-30; B59-3; B60-2; B56-6; and/or, preferably, B8-26 can also bind to TcdB.

In one embodiment, the antibody or antigen binding fragment thereof includes a variable heavy chain region including an amino acid sequence at least about 60%, 65%, 70%, 75%, 80%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% identical to a variable heavy chain region amino acid sequence of B8-26 (SEQ ID NO: 49).

5 In another embodiment, the antibody or antigen binding fragment thereof includes a variable light chain region including an amino acid sequence at least about 60%, 65%, 70%, 75%, 80%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% identical to a variable light chain region amino acid sequence of B8-26 (SEQ ID NO: 55).

10 In yet a further aspect, the antibody or antigen binding fragment thereof includes a variable heavy chain region including an amino acid sequence at least about 60%, 65%, 70%, 75%, 80%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% identical to a variable heavy chain region amino acid sequence of B8-26 (SEQ ID NO: 49), and a variable light chain region including an amino acid sequence at least about 60%, 65%, 70%, 75%, 80%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% identical to a variable light chain region amino acid sequence of B8-26 (SEQ ID NO: 55).

15 In another embodiment, antibodies or binding fragments thereof having CDRs of variable heavy chains and/or variable light chains of B2-31; B5-40, B70-2; B6-30; B9-30; B59-3; B60-2; B56-6; and/or, preferably, B8-26 can also bind to TcdB.

In one embodiment, the antibody or binding fragment thereof includes amino acid sequences of the heavy chain complementarity determining regions (CDRs) of B8-26, and/or the amino acid sequences of the light chain CDRs of B8-26.

20 25 In a preferred embodiment, the antibody or binding fragment thereof specific to *C. difficile* toxin B specifically binds to an epitope within the N-terminal region of toxin B, e.g., an epitope between amino acids 1-1256 of a TcdB, according to the numbering of SEQ ID NO: 2. Examples include B2-31; B5-40; B8-26; B70-2; B6-30; and B9-30.

30 In an exemplary embodiment, the antibody or binding fragment thereof specific to *C. difficile* toxin B specifically binds to an epitope within the C-terminal region of toxin B, e.g., an epitope between amino acids 1832 to 2710 of a TcdB, according to the numbering of SEQ ID NO: 2.

In yet another embodiment, the antibody or binding fragment thereof specific to *C. difficile* toxin B specifically binds to an epitope within the “translocation” region of *C. difficile* toxin B, e.g., an epitope that preferably includes residues 956-1128 of a TcdB, according to the numbering of SEQ ID NO: 2, such as an epitope between amino acids 5 659-1832 of a TcdB. Examples include B59-3; B60-2; and B56-6.

Combinations of antibodies

The anti-toxin antibody or binding fragment thereof can be administered in combination with other anti- *C. difficile* toxin antibodies (e.g., other monoclonal antibodies, polyclonal gamma-globulin) or a binding fragment thereof. Combinations 10 that can be used include an anti-toxin A antibody or binding fragment thereof and an anti-toxin B antibody or binding fragment thereof.

In another embodiment, a combination includes an anti-toxin A antibody or binding fragment thereof and another anti-toxin A antibody or binding fragment thereof. Preferably, the combination includes a neutralizing anti-toxin A monoclonal antibody or 15 binding fragment thereof and another neutralizing anti-toxin A monoclonal antibody or binding fragment thereof. Surprisingly, the inventors discovered that such a combination resulted in a synergistic effect in neutralization of toxin A cytotoxicity. For example, the combination includes a combination of at least two of the following neutralizing anti-toxin A monoclonal antibodies: A3-25; A65-33; A60-22; and A80-29. 20 More preferably, the combination includes A3-25 antibody and at least one of the following neutralizing anti-toxin A monoclonal antibodies: A65-33; A60-22; and A80-29. Most preferably, the combination includes all four antibodies: A3-25; A65-33; A60-22; and A80-29.

In a further embodiment, a combination includes an anti-toxin B antibody or binding fragment thereof and another anti-toxin B antibody or binding fragment thereof. Preferably, the combination includes a neutralizing anti-toxin B monoclonal antibody or 25 binding fragment thereof and another neutralizing anti-toxin B monoclonal antibody or binding fragment thereof. Surprisingly, the inventors discovered that such a combination resulted in a synergistic effect in neutralization of toxin B cytotoxicity. More preferably, the combination includes a combination of at least two of the following neutralizing anti-toxin B monoclonal antibodies: B8-26; B9-30 and B59-3. Most 30 preferably, the combination includes all three antibodies: B8-26; B9-30 and B59-3.

In yet another embodiment, a combination includes an anti-toxin B antibody or binding fragment thereof and another anti-toxin B antibody or binding fragment thereof. As stated previously, the inventors discovered that a combination of at least two of the neutralizing monoclonal antibodies can exhibit an unexpectedly synergistic effect in
5 respective neutralization of toxin A and toxin B.

In another embodiment, the agents of the invention can be formulated as a mixture, or chemically or genetically linked using art recognized techniques thereby resulting in covalently linked antibodies (or covalently linked antibody fragments), having both anti-toxin A and anti-toxin B binding properties. The combined formulation
10 may be guided by a determination of one or more parameters such as the affinity, avidity, or biological efficacy of the agent alone or in combination with another agent.

Such combination therapies are preferably additive and/or synergistic in their therapeutic activity, e.g., in the inhibition, prevention (e.g., of relapse), and/or treatment of *C. difficile* - related diseases or disorders. Administering such combination therapies
15 can decrease the dosage of the therapeutic agent (e.g., antibody or antibody fragment mixture, or cross-linked or genetically fused bispecific antibody or antibody fragment) needed to achieve the desired effect.

It is understood that any of the inventive compositions, for example, an anti-toxin A and/or anti-toxin B antibody or binding fragment thereof, can be combined in different
20 ratios or amounts for therapeutic effect. For example, the anti-toxin A and anti-toxin B antibody or respective binding fragment thereof can be present in a composition at a ratio in the range of 0.1:10 to 10:0.1, A:B. In another embodiment, the anti-toxin A and anti-toxin B antibody or respective binding fragment thereof can be present in a composition at a ratio in the range of 0.1:10 to 10:0.1, B:A.

In another aspect, the invention relates to a method of producing a neutralizing antibody against a *C. difficile* TcdA. The method includes administering an immunogenic composition as described above to a mammal, and recovering the antibody from the mammal. In a preferred embodiment, the immunogenic composition includes a mutant *C. difficile* TcdA having SEQ ID NO: 4, wherein at least one amino
30 acid of the mutant *C. difficile* TcdA is chemically crosslinked, preferably by formaldehyde or 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide. Exemplary neutralizing antibodies against TcdA that may be produced include A65-33; A60-22; A80-29 and/or A3-25.

In yet another aspect, the invention relates to a method of producing a neutralizing antibody against a *C. difficile* TcdB. The method includes administering an immunogenic composition as described above to a mammal, and recovering the antibody from the mammal. In a preferred embodiment, the immunogenic composition 5 includes a mutant *C. difficile* TcdB having SEQ ID NO: 6, wherein at least one amino acid of the mutant *C. difficile* TcdB is chemically crosslinked, preferably by formaldehyde or 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide. Exemplary neutralizing antibodies against TcdB that may be produced include B2-31; B5-40, B70-2; B6-30; B9-30; B59-3; B60-2; B56-6; and/or B8-26.

10

Formulations

Compositions of the present invention (such as, e.g., compositions including a mutant *C. difficile* toxin, immunogenic compositions, antibodies and/or antibody binding fragments thereof described herein) may be in a variety of forms. These include, for 5 example, semi-solid and solid dosage forms, suppositories, liquid forms, such as liquid solutions (e.g., injectable and infusible solutions), dispersions or suspensions, liposomes, and/or dried form, such as, for example, lyophilized powder form, freeze-dried form, spray-dried form, and/or foam-dried form. For suppositories, binders and carriers include, for example, polyalkylene glycols or triglycerides; such suppositories 10 can be formed from mixtures containing the inventive compositions. In an exemplary embodiment, the composition is in a form that is suitable for solution in, or suspension in, liquid vehicles prior to injection. In another exemplary embodiment, the composition is emulsified or encapsulated in liposomes or microparticles, such as polylactide, polyglycolide, or copolymer.

15 In a preferred embodiment, the composition is lyophilized and extemporaneously reconstituted prior to use.

In one aspect, the present invention relates to pharmaceutical compositions that include any of the compositions described herein (such as, e.g., compositions including a mutant *C. difficile* toxin, immunogenic compositions, antibodies and/or antibody 20 binding fragments thereof described herein), formulated together with a pharmaceutically acceptable carrier. "Pharmaceutically acceptable carriers" include any solvents, dispersion media, stabilizers, diluents, and/or buffers that are physiologically suitable.

Exemplary stabilizers include carbohydrates, such as sorbitol, mannitol, starch, 25 dextran, sucrose, trehalose, lactose, and/or glucose; inert proteins, such as albumin and/or casein; and/or other large, slowly metabolized macromolecules, such as polysaccharides such as chitosan, polylactic acids, polyglycolic acids and copolymers (such as latex functionalized SEPHAROSE™ agarose, agarose, cellulose, etc/), amino acids, polymeric amino acids, amino acid copolymers, and lipid aggregates (such as oil 30 droplets or liposomes). Additionally, these carriers may function as immunostimulating agents (i.e., adjuvants).

Preferably, the composition includes trehalose. Preferred amounts of trehalose (% by weight) include from a minimum of about 1%, 2%, 3%, or 4% to a maximum of about 10%, 9%, 8%, 7%, 6%, or 5%. Any minimum value can be combined with any maximum value to define a suitable range. In one embodiment, the composition 5 includes about 3% - 6% trehalose, most preferably, 4.5% trehalose, for example, per 0.5 mL dose.

Examples of suitable diluents include distilled water, saline, physiological phosphate-buffered saline, glycerol, alcohol (such as ethanol), Ringer's solutions, dextrose solution, Hanks' balanced salt solutions, and/or a lyophilization excipient. 10

Exemplary buffers include phosphate (such as potassium phosphate, sodium phosphate); acetate (such as sodium acetate); succinate (such as sodium succinate); glycine; histidine; carbonate, Tris (tris(hydroxymethyl)aminomethane), and/or bicarbonate (such as ammonium bicarbonate) buffers. Preferably, the composition includes tris buffer. Preferred amounts of tris buffer include from a minimum of about 1 15 mM, 5 mM, 6 mM, 7 mM, 8 mM, 9 mM, 10 mM to a maximum of about 100 mM, 50 mM, 20 mM, 19 mM, 18 mM, 17 mM, 16 mM, 15 mM, 14 mM, 13 mM, 12 mM, or 11 mM. Any minimum value can be combined with any maximum value to define a suitable range. In one embodiment, the composition includes about 8 mM to 12 mM tris buffer, most preferably, 10 mM tris buffer, for example, per 0.5 mL dose.

20 In another preferred embodiment, the composition includes histidine buffer. Preferred amounts of histidine buffer include from a minimum of about 1 mM, 5 mM, 6 mM, 7 mM, 8 mM, 9 mM, 10 mM to a maximum of about 100 mM, 50 mM, 20 mM, 19 mM, 18 mM, 17 mM, 16 mM, 15 mM, 14 mM, 13 mM, 12 mM, or 11 mM. Any minimum value can be combined with any maximum value to define a suitable range. In one 25 embodiment, the composition includes about 8 mM to 12 mM histidine buffer, most preferably, 10 mM histidine buffer, for example, per 0.5 mL dose.

In yet another preferred embodiment, the composition includes phosphate buffer. Preferred amounts of phosphate buffer include from a minimum of about 1 mM, 5 mM, 6 mM, 7 mM, 8 mM, 9 mM, 10 mM to a maximum of about 100 mM, 50 mM, 20 mM, 19 30 mM, 18 mM, 17 mM, 16 mM, 15 mM, 14 mM, 13 mM, 12 mM, or 11 mM. Any minimum value can be combined with any maximum value to define a suitable range. In one embodiment, the composition includes about 8 mM to 12 mM phosphate buffer, most preferably, 10 mM phosphate buffer, for example, per 0.5 mL dose.

The pH of the buffer will generally be chosen to stabilize the active material of choice, and can be ascertainable by those in the art by known methods. Preferably, the pH of the buffer will be in the range of physiological pH. Thus, preferred pH ranges are from about 3 to about 8; more preferably, from about 6.0 to about 8.0; yet more preferably, from about 6.5 to about 7.5; and most preferably, at about 7.0 to about 7.2.

5 In some embodiments, the pharmaceutical compositions may include a surfactant. Any surfactant is suitable, whether it is amphoteric, non-ionic, cationic or anionic. Exemplary surfactants include the polyoxyethylene sorbitan esters surfactants (e.g., TWEEN ®), such as polysorbate 20 and/or polysorbate 80; polyoxyethylene fatty ethers derived from lauryl, cetyl, stearyl and oleyl alcohols (known as Brij surfactants), such as triethyleneglycol monolauryl ether (Brij 30); Triton X 100TM, or t-octylphenoxyethoxyethanol; and sorbitan esters (commonly known as the SPANs), such as sorbitan trioleate (SpanTM 85) and sorbitan monolaurate, and combinations thereof. Preferred surfactants include polysorbate 80 (polyoxyethylene sorbitan 10 monooleate).

15

Preferred amounts of polysorbate 80 (% by weight) include from a minimum of about 0.001%, 0.005%, or 0.01%, to a maximum of about 0.010%, 0.015%, 0.025%, or 1.0%. Any minimum value can be combined with any maximum value to define a suitable range. In one embodiment, the composition includes about 0.005% - 0.015% 20 polysorbate 80, most preferably, 0.01% polysorbate 80.

In an exemplary embodiment, the immunogenic composition includes trehalose and phosphate 80. In another exemplary embodiment, the immunogenic composition includes tris buffer and polysorbate 80. In another exemplary embodiment, the immunogenic composition includes histidine buffer and polysorbate 80. In yet another 25 exemplary embodiment, the immunogenic composition includes phosphate buffer and polysorbate 80.

In one exemplary embodiment, the immunogenic composition includes trehalose, tris buffer and polysorbate 80. In another exemplary embodiment, the immunogenic composition includes trehalose, histidine buffer and polysorbate 80. In yet another 30 exemplary embodiment, the immunogenic composition includes trehalose, phosphate buffer and polysorbate 80.

The compositions described herein may further include components of petroleum, animal, vegetable, or synthetic origin, for example, peanut oil, soybean oil, and/or mineral oil. Examples include glycols such as propylene glycol or polyethylene glycol.

5 In some embodiments, the pharmaceutical composition further includes formaldehyde. For example, in a preferred embodiment, a pharmaceutical composition that further includes formaldehyde has an immunogenic composition, wherein the mutant *C. difficile* toxin of the immunogenic composition has been contacted with a chemical crosslinking agent that includes formaldehyde. The amount of formaldehyde
10 present in the pharmaceutical composition may vary from a minimum of about 0.001%, 0.002%, 0.003%, 0.004%, 0.005%, 0.006%, 0.007%, 0.008%, 0.009%, 0.010%, 0.013%, or 0.015%, to a maximum of about 0.020%, 0.019%, 0.018%, 0.017% 0.016%, 0.015%, 0.014%, 0.013%, 0.012% 0.011% or 0.010%. Any minimum value can be combined with any maximum value to define a suitable range. In one embodiment, the
15 pharmaceutical composition includes about 0.010% formaldehyde.

In some alternative embodiments, the pharmaceutical compositions described herein do not include formaldehyde. For example, in a preferred embodiment, a pharmaceutical composition that does not include formaldehyde has an immunogenic composition, wherein at least one amino acid of the mutant *C. difficile* toxin is
20 chemically crosslinked by an agent that includes EDC. More preferably, in such an embodiment, the mutant *C. difficile* toxin has not been contacted with a chemical crosslinking agent that includes formaldehyde. As another exemplary embodiment, a pharmaceutical composition that is in a lyophilized form does not include formaldehyde.

25 In another embodiment, the compositions described herein may include an adjuvant, as described below. Preferred adjuvants augment the intrinsic immune response to an immunogen without causing conformational changes in the immunogen that may affect the qualitative form of the immune response.

Exemplary adjuvants include 3 De-O-acylated monophosphoryl lipid A (MPL™) (see GB 2220211 (GSK)); an aluminum hydroxide gel such as Alhydrogel™ (Brenntag
30 Biosector, Denmark); aluminum salts (such as aluminum hydroxide, aluminum phosphate, aluminum sulfate), which may be used with or without an immunostimulating agent such as MPL or 3-DMP, QS-21, polymeric or monomeric amino acids such as polyglutamic acid or polylysine.

Yet another exemplary adjuvant is an immunostimulatory oligonucleotide such as a CpG oligonucleotide (see, e.g., WO 1998/040100, WO2010/067262), or a saponin and an immunostimulatory oligonucleotide, such as a CpG oligonucleotide (see, e.g., WO 00/062800). In a preferred embodiment, the adjuvant is a CpG oligonucleotide,

5 most preferably a CpG oligodeoxynucleotides (CpG ODN). Preferred CpG ODN are of the B Class that preferentially activate B cells. In aspects of the invention, the CpG ODN has the nucleic acid sequence 5' T*C*G*T*C*G*T*T*T*T*C*G*G*T*G*C*T*T*T 3' (SEQ ID NO: 48) wherein * indicates a phosphorothioate linkage. The CpG ODN of this sequence is known as CpG 24555, which is described in WO2010/067262. In a

10 preferred embodiment, CpG 24555 is used together with an aluminium hydroxide salt such as AlhydrogelTM.

A further class of exemplary adjuvants include saponin adjuvants, such as StimulonTM (QS-21, which is a triterpene glycoside or saponin, Aquila, Framingham, Mass.) or particles generated therefrom such as ISCOMs (immune stimulating complexes) and ISCOMATRIX[®] adjuvant. Accordingly, the compositions of the present invention may be delivered in the form of ISCOMs, ISCOMS containing CTB, liposomes or encapsulated in compounds such as acrylates or poly(DL-lactide-co-glycoside) to form microspheres of a size suited to adsorption. Typically, the term "ISCOM" refers to immunogenic complexes formed between glycosides, such as 15 triterpenoid saponins (particularly Quil A), and antigens which contain a hydrophobic region. In a preferred embodiment, the adjuvant is an ISCOMATRIX adjuvant.

Other exemplary adjuvants include RC-529, GM-CSF and Complete Freund's Adjuvant (CFA) and Incomplete Freund's Adjuvant (IFA).

Yet another class of exemplary adjuvants is glycolipid analogues including N-25 glycosylamides, N-glycosylureas and N-glycosylcarbamates, each of which is substituted in the sugar residue by an amino acid.

Optionally, the pharmaceutical composition includes two or more different adjuvants. Preferred combinations of adjuvants include any combination of adjuvants including, for example, at least two of the following adjuvants: alum, MPL, QS-21, 30 ISCOMATRIX, CpG, and AlhydrogelTM. An exemplary combination of adjuvants includes a combination of CpG and AlhydrogelTM.

Alternatively, in one embodiment, the composition is administered to the mammal in the absence of an adjuvant.

Compositions described herein can be administered by any route of administration, such as, for example, parenteral, topical, intravenous, mucosal, oral, subcutaneous, intraarterial, intracranial, intrathecal, intraperitoneal, intranasal, intramuscular, intradermal, infusion, rectal, and/or transdermal routes for prophylactic and/or therapeutic applications. In a preferred embodiment, the route of administration of the composition is parenteral, more preferably, intramuscular administration. Typical intramuscular administration is performed in the arm or leg muscles.

Compositions described herein can be administered in combination with therapies that are at least partly effective in prevention and/or treatment of *C. difficile* infection. For example, a composition of the invention may be administered before, concurrently with, or after biotherapy; probiotic therapy; stool implants; immunotherapy (such as intravenous immunoglobulin); and/or an accepted standard of care for the antibiotic treatment of *C. difficile* associated disease (CDAD), such as metronidazole and/or vancomycin.

A composition of the present invention relating to toxin A and toxin B may be administered to the mammal in any combination. For example, an immunogenic composition including a mutant *C. difficile* TcdA may be administered to the mammal before, concurrently with, or after administration of an immunogenic composition including a mutant *C. difficile* TcdB. Conversely, an immunogenic composition including a mutant *C. difficile* TcdB may be administered to the mammal before, concurrently with, or after administration of an immunogenic composition including a mutant *C. difficile* TcdA.

In another embodiment, a composition including an anti-toxin A antibody or binding fragment thereof may be administered to the mammal before, concurrently with, or after administration of a composition including an anti-toxin B antibody or binding fragment thereof. Conversely, a composition including an anti-toxin B antibody or binding fragment thereof may be administered to the mammal before, concurrently with, or after administration of a composition including an anti-toxin A antibody or binding fragment thereof.

In a further embodiment, a composition of the present invention may be administered to the mammal before, concurrently with, or after administration of a pharmaceutically acceptable carrier. For example, an adjuvant may be administered before, concurrently with, or after administration of a composition including a mutant *C.*

difficile toxin. Accordingly, a composition of the present invention and a pharmaceutically acceptable carrier can be packaged in the same vial or they can be packaged in separate vials and mixed before use. The compositions can be formulated for single dose administration and/or multiple dose administration.

5

Methods of Protecting and/or Treating *C. difficile* infection in a Mammal

In one aspect, the invention relates to a method of inducing an immune response to a *C. difficile* toxin in a mammal. The method includes administering an effective amount of a composition described herein to the mammal. For example, the method 5 may include administering an amount effective to generate an immune response to the respective *C. difficile* toxin in the mammal.

In an exemplary embodiment, the invention relates to a method of inducing an immune response to a *C. difficile* TcdA in a mammal. The method includes administering an effective amount of an immunogenic composition that includes a 10 mutant *C. difficile* TcdA to the mammal. In another exemplary embodiment, the invention relates to a method of inducing an immune response to a *C. difficile* TcdB in a mammal. The method includes administering an effective amount of an immunogenic composition that includes a mutant *C. difficile* TcdB to the mammal.

In a further embodiment, the method includes administering an effective amount 15 of an immunogenic composition that includes a mutant *C. difficile* TcdA and an effective amount of an immunogenic composition that includes a mutant *C. difficile* TcdB to the mammal. In additional aspects, the compositions described herein may be used to treat, prevent, decrease risk of, decrease severity of, decrease occurrences of, and/or delay onset of a *C. difficile* infection, *C. difficile* associated disease, syndrome, 20 condition, symptom, and/or complication thereof in a mammal, as compared to a mammal to which the composition is not administered. The method includes administering an effective amount of the composition to the mammal.

Three clinical syndromes caused by *C. difficile* infection are recognized, based 25 on the severity of the infection. The most severe form is pseudomembranous colitis (PMC), which is characterized by profuse diarrhea, abdominal pain, systemic signs of illness, and a distinctive endoscopic appearance of the colon.

Antibiotic-associated colitis (AAC) is also characterized by profuse diarrhea, abdominal pain and tenderness, systemic signs (e.g., fever), and leukocytosis. Intestinal 30 injury in AAC is less severe than in PMC, the characteristic endoscopic appearance of the colon in PMC is absent, and mortality is low.

Finally, antibiotic-associated diarrhea (AAD, which is also known as *C. difficile* associated diarrhea (CDAD) is a relatively mild syndrome, and is characterized by mild

to moderate diarrhea, lacking both large intestinal inflammation (as characterized by, e.g., abdominal pain and tenderness) and systemic signs of infection (e.g., fever).

These three distinct syndromes typically occur in an increasing order of frequency. That is, PMC typically occurs less frequently than AAC, and AAD is typically 5 the most frequent clinical presentation of *C. difficile* disease.

A frequent complication of *C. difficile* infection is recurrent or relapsing disease, which occurs in up to 20% of all subjects who recover from *C. difficile* disease. Relapse may be characterized clinically as AAD, AAC, or PMC. Patients who relapse once are more likely to relapse again.

10 As used herein, conditions of a *C. difficile* infection include, for example, mild, mild-to-moderate, moderate, and severe *C. difficile* infection. A condition of *C. difficile* infection may vary depending on presence and/or severity of symptoms of the infection.

Symptoms of a *C. difficile* infection may include physiological, biochemical, histologic and/or behavioral symptoms such as, for example, diarrhea; colitis; colitis with 15 cramps, fever, fecal leukocytes, and inflammation on colonic biopsy; pseudomembranous colitis; hypoalbuminemia; anasarca; leukocytosis; sepsis; abdominal pain; asymptomatic carriage; and/or complications and intermediate pathological phenotypes present during development of the infection, and combinations thereof, etc. Accordingly, for example, administration of an effective amount of the 20 compositions described herein may, for example treat, prevent, decrease risk of, decrease severity of, decrease occurrences of, and/or delay onset of diarrhea; abdominal pain, cramps, fever, inflammation on colonic biopsy, hypoalbuminemia, anasarca, leukocytosis, sepsis, and/or asymptomatic carriage, etc., as compared to a mammal to which the composition was not administered.

25 Risk factors of a *C. difficile* infection may include, for example, present or immediate future use of an antimicrobial (any antimicrobial agent with an antibacterial spectrum and/or activity against anaerobic bacteria are encompassed, including, for example, antibiotics that cause disruption of normal colonic microbiota, e.g., clindamycin, cephalosporins, metronidazole, vancomycin, fluoroquinolones (including 30 levofloxacin, moxifloxacin, gatifloxacin, and ciprofloxacin), linezolid, etc.); present or immediate future withdrawal of prescribed metronidazole or vancomycin; present or immediate future admission to a healthcare facility (such as a hospital, chronic care facility, etc.) and healthcare workers; present or immediate future treatment with proton-

pump inhibitors, H2 antagonists, and/or methotrexate, or a combination thereof; present or risk of gastrointestinal diseases, such as inflammatory bowel disease; past, present, or immediate future gastrointestinal surgery or gastrointestinal procedure on the mammal; past or present recurrence of a *C. difficile* infection and/or a CDAD, e.g.,

5 patients who have had a *C. difficile* infection and/or a CDAD once or more than once; and humans aged at least about 65 and above.

In the methods described herein, the mammal may be any mammal, such as, for example, mice, hamsters, primates, and humans. In a preferred embodiment, the mammal is a human. According to the present invention, the human may include

10 individuals who have exhibited a *C. difficile* infection, *C. difficile* associated disease, syndrome, condition, symptom, and/or complication thereof; individuals who are presently exhibiting a *C. difficile* infection, *C. difficile* associated disease, syndrome, condition, symptom, and/or complication thereof; and individuals who are at risk of a *C. difficile* infection, *C. difficile* associated disease, syndrome, condition, symptom, and/or

15 complication thereof.

Examples of individuals who have shown symptoms of *C. difficile* infection include individuals who have shown or are showing symptoms described above; individuals who have had or are having a *C. difficile* infection and/or a *C. difficile* associated disease (CDAD); and individuals who have a recurrence of a *C. difficile* infection and/or CDAD.

Examples of patients who are at risk of a *C. difficile* infection include individuals at risk of or are presently undergoing planned antimicrobial use; individuals at risk of or are presently undergoing withdrawal of prescribed metronidazole or vancomycin; individuals who are at risk of or are presently undergoing a planned admission to a

25 healthcare facility (such as a hospital, chronic care facility, etc.) and healthcare workers; and/or individuals at risk of or are presently undergoing a planned treatment with proton-pump inhibitors, H2 antagonists, and/or methotrexate, or a combination thereof; individuals who have had or are undergoing gastrointestinal diseases, such as inflammatory bowel disease; individuals who have had or are undergoing

30 gastrointestinal surgery or gastrointestinal procedures; and individuals who have had or are having a recurrence of a *C. difficile* infection and/or a CDAD, e.g., patients who have had a *C. difficile* infection and/or a CDAD once or more than once; individuals who

are about 65 years old or older. Such at-risk patients may or may not be presently showing symptoms of a *C. difficile* infection.

In asymptomatic patients, prophylaxis and/or treatment can begin at any age (e.g., at about 10, 20, or 30 years old). In one embodiment, however, it is not necessary to begin treatment until a patient reaches at least about 45, 55, 65, 75, or 85 years old. For example, the compositions described herein may be administered to an asymptomatic human who is aged 50-85 years.

In one embodiment, the method of preventing, decreasing risk of, decreasing severity of, decreasing occurrences of, and/or delaying onset of a *C. difficile* infection, *C. difficile* associated disease, syndrome, condition, symptom, and/or complication thereof in a mammal includes administering an effective amount of a composition described herein to a mammal in need thereof, a mammal at risk of, and/or a mammal susceptible to a *C. difficile* infection. An effective amount includes, for example, an amount sufficient to prevent, decrease risk of, decrease severity of, decrease occurrences of, and/or delay onset of a *C. difficile* infection, *C. difficile* associated disease, syndrome, condition, symptom, and/or complication thereof in a mammal, as compared to a mammal to which the composition is not administered. Administration of an effective amount of the compositions described herein may, for example, prevent, decrease risk of, decrease severity of, decrease occurrences of, and/or delay onset of diarrhea; abdominal pain, cramps, fever, inflammation on colonic biopsy, hypoalbuminemia, anasarca, leukocytosis, sepsis, and/or asymptomatic carriage, etc., as compared to a mammal to which the composition was not administered. In a preferred embodiment, the method includes administering an effective amount of an immunogenic composition described herein to the mammal in need thereof, the mammal at risk of, and/or the mammal susceptible to a *C. difficile* infection.

In an additional embodiment, the method of treating, decreasing severity of, and/or delaying onset of a *C. difficile* infection, *C. difficile* associated disease, syndrome, condition, symptom, and/or complication thereof in a mammal includes administering an effective amount of a composition described herein to a mammal suspected of, or presently suffering from a *C. difficile* infection. An effective amount includes, for example, an amount sufficient to treat, decrease severity of, and/or delay the onset of a *C. difficile* infection, *C. difficile* associated disease, syndrome, condition,

symptom, and/or complication thereof in a mammal, as compared to a mammal to which the composition is not administered.

Administration of an effective amount of the composition may improve at least one sign or symptom of *C. difficile* infection in the subject, such as those described below. Administration of an effective amount of the compositions described herein may, for example, decrease severity of and/or decrease occurrences of diarrhea; decrease severity of and/or decrease occurrences of abdominal pain, cramps, fever, inflammation on colonic biopsy, hypoalbuminemia, anasarca, leukocytosis, sepsis, and/or asymptomatic carriage, etc., as compared to a mammal to which the composition was not administered. Optionally, the presence of symptoms, signs, and/or risk factors of an infection is determined before beginning treatment. In a preferred embodiment, the method includes administering an effective amount of an antibody and/or binding fragment thereof described herein to the mammal suspected of, or presently suffering from, a *C. difficile* infection.

Accordingly, an effective amount of the composition refers to an amount sufficient to achieve a desired effect (e.g., prophylactic and/or therapeutic effect) in the methods of the present invention. For example, the amount of an immunogen for administration may vary from a minimum of about 1 μ g, 5 μ g, 25 μ g, 50 μ g, 75 μ g, 100 μ g, 200 μ g, 500 μ g, or 1 mg to a maximum of about 2 mg, 1 mg, 500 μ g, 200 μ g per injection. Any minimum value can be combined with any maximum value to define a suitable range. Typically about 10, 20, 50 or 100 μ g per immunogen is used for each human injection.

The amount of a composition of the invention administered to the subject may depend on the type and severity of the infection and/or on the characteristics of the individual, such as general health, age, sex, body weight and tolerance to drugs. It may also depend on the degree, severity, and type of disease. An effective amount may also vary depending upon factors, such as route of administration, target site, physiological state of the patient, age of the patient, whether the patient is human or an animal, other therapies administered, and whether treatment is prophylactic or therapeutic. The skilled artisan will be able to determine appropriate amounts depending on these and other factors.

An effective amount may include one effective dose or multiple effective doses (such as, for example, 2, 3, 4 doses, or more) for use in the methods herein. Effective dosages may need to be titrated to optimize safety and efficacy.

A combination of amount and frequency of dose adequate to accomplish

5 prophylactic and/or therapeutic uses is defined as a prophylactically- or therapeutically-effective regimen. In a prophylactic and/or therapeutic regimen, the composition is typically administered in more than one dosage until a sufficient immune response has been achieved. Typically, the immune response is monitored and repeated dosages are given if the immune response starts to wane.

10 The compositions may be administered in multiple dosages over a period of time. Treatment can be monitored by assaying antibody, or activated T-cell or B-cell responses to the therapeutic agent (e.g., the immunogenic composition including a mutant *C. difficile* toxin) over time. If the response falls, a booster dosage is indicated.

EXAMPLES

Example 1: Identification of toxin-negative *C. difficile* strains

To identify *C. difficile* strains lacking toxin (A and B) genes and toxin expression, 13 *C. difficile* strains were tested. Culture media of 13 *C. difficile* strains were tested by 5 ELISA for toxin A. Seven strains expressed toxin A: *C. difficile* 14797-2, *C. difficile* 630, *C. difficile* BDMS, *C. difficile* W1194, *C. difficile* 870, *C. difficile* 1253, and *C. difficile* 2149. See Figure 3.

Six strains did not express toxin A and lacked the entire pathogenicity locus: *C. difficile* 1351 (ATCC 43593TM), *C. difficile* 3232 (ATCC BAA-1801TM), *C. difficile* 7322 10 (ATCC 43601TM), *C. difficile* 5036 (ATCC 43603TM), *C. difficile* 4811 (4 ATCC 3602TM), and *C. difficile* VPI 11186 (ATCC 700057TM). VPI 11186 was selected based upon its effectiveness to take up plasmid DNA by conjugation.

The same 13 strains were tested in a multiplex PCR assay using primers outside of the pathogenicity locus (PaLoc; Braun et al., *Gene*. 1996 Nov 28;181(1-2):29-38.). 15 The PCR results demonstrated the DNA from the 6 toxin A negative strains by ELISA did not amplify any genes from the PaLoc (tcdA-tcdE). The PaLoc flanking sequences (*cdd3* and *cdu2*) were present (data not shown).

Example 2: Inactivation of sporulation pathway in *C. difficile* VPI 11186

Knocking-out the spore-forming function of the *C. difficile* production strain 20 facilitates large scale fermentation in a safe manufacturing environment. The ClosTron system was used to create an asporogenic *C. difficile* strain. See Heap et al., *J Microbiol Methods*. 2009 Jul;78(1):79-85. The ClosTron system allows targeted gene inactivation with a group II intron for site directed insertional inactivation of a *spo0A1* clostridial gene. The toxin-minus production strain VPI11186 was subjected to 25 sporulation inactivation by the ClosTron technology. Erythromycin resistant mutants were selected and the presence of the insertional cassette was confirmed by PCR (not shown). The inability of two independent clones to form spores was confirmed.

Example 3: Genetic modification of toxin A and B genes to inactivate cytotoxicity function

Full-length mutant toxins A and B open reading frames (ORFs) based on strain 630Δ genome sequences were designed for custom synthesis at Blue Heron Biotech.

5 See, for example, SEQ ID NOs: 9-14. The active site for the glucosyltransferase activity responsible for cellular toxicity was altered by two allelic substitutions: D285A/D287A (see SEQ ID NO: 3) for toxin A, and D286A/D288A (see SEQ ID NO: 5) for toxin B. Two nucleotides were mutated in each aspartate (D) codon to create the codon for alanine (A). See, for example, SEQ ID NOs: 9-14. In addition, a pair of vectors 10 expressing mutant toxins lacking cysteine residues was constructed following custom synthesis at Blue Heron Biotech. Seven cysteine residues from mutant toxin A and 9 cysteine residues from mutant toxin B were replaced with alanine. The substitutions include catalytic cysteines of the A and B toxin autocatalytic protease. Also, silent mutations were introduced where necessary to eliminate restriction enzyme sites used 15 for vector construction.

Example 4: pMTL84121*fdx* Expression Vector

The plasmid shuttle vector used for *C. difficile* mutant toxin antigen expression was selected from the pMTL8000-series modular system developed by the Minton lab (see Heap et al., *J Microbiol Methods*. 2009 Jul;78(1):79-85). The chosen vector

20 pMTL84121*fdx* contains the *C. difficile* plasmid pCD6 Gram+ replicon, the *catP* (chloramphenicol/ thiamphenicol) selectable marker, the p15a Gram- replicon and *tra* function, and the *C. sporogenes* ferredoxin promoter (*fdx*) and distal multiple cloning site (MCS). Empirical data suggested that the low-copy number p15a replicon conferred greater stability in *E. coli* than the ColE1 alternative. The *fdx* promoter was selected as it 25 yielded higher expression than other promoters tested in experiments with CAT reporter constructs (e.g. *tcdA*, *tcdB*; or heterologous *tetR* or *xylR*) (data not shown).

Example 5: Cloning the modified toxin ORFs into pMTL84121*fdx*

Full-length mutant toxin A and B open reading frames (ORFs) based on strain 630Δ genome sequences were subcloned using pMTL84121*fdx* vector multiple cloning 30 *NdeI* and *BglII* sites using standard molecular biology techniques. To facilitate cloning, the ORFs were flanked by a proximal *NdeI* site containing the start codon and a *BglII* site just downstream of the stop codon.

Example 6: Site directed mutagenesis of TcdA to create a triple mutant

The catalytic cysteine residue of the autocatalytic protease domain was substituted (i.e., C700A for TcdA and C698A for TcdB) in SEQ ID NOs: 3 and 5, i.e., in each of the “double mutants.” For mutagenesis of mutant toxin A, a 2.48kb *Nde*I-*Hind*III fragment from the TcdA D285A/D287A expression plasmid was subcloned into pUC19 (cut with same) and site-directed mutagenesis was performed on this template. Once the new alleles were confirmed by DNA sequence analysis, the modified *Nde*I-*Hind*III fragments were reintroduced into the expression vector pMTL84121*fdx* to create the “triple mutants,” i.e., SEQ ID NO: 4 and SEQ ID NO: 6.

10 Example 7: Site directed mutagenesis of TcdB to create a triple mutant

For mutagenesis of mutant toxin B, a 3.29 kb *Nde*I-*Eco*NI fragment from the mutant toxin B plasmid was modified and reintroduced. As the *Eco*NI site is not present in available cloning vectors a slightly larger 3.5kb *Nde*I-*Eco*RV fragment was subcloned into pUC19 (prepared with *Nde*I-*Sma*I). After mutagenesis, the modified internal 3.3kb *Nde*I-*Eco*NI fragment was excised and used to replace the corresponding mutant toxin B expression vector pMTL84121*fdx* fragment. As the cloning efficiency of this directional strategy was found to be quite low, an alternative strategy for introducing the C698A allele involving replacement of a 1.5 kb *Dra*III was attempted in parallel. Both strategies independently yielded the desired recombinants.

20 Example 8: Creating additional mutant toxin variants by site-directed mutagenesis

At least twelve different *C. difficile* mutant toxin variants were constructed. Allelic substitutions were introduced into N-terminal mutant toxin gene fragments by site directed mutagenesis (Quickchange® kit). Recombinant toxins were also engineered as reference controls to evaluate the capacity of this plasmid-based system to generate protein quantitatively equivalent in biological activity to native toxins purified from wild-type *C. difficile* strains. In this case, allelic substitutions were introduced to revert the original glucosyltransferase substitutions. In addition, a pair of cysteineless mutant toxin vectors was constructed following custom synthesis at Blue Heron Biotech.

30 The twelve toxin variants include (1) a mutant *C. difficile* toxin A having a D285A/D287A mutation (SEQ ID NO: 3); (2) a mutant *C. difficile* toxin B having a D286A/D288A mutation (SEQ ID NO: 5); (3) a mutant *C. difficile* toxin A having a D285A/D287A C700A mutation (SEQ ID NO: 4); (4) a mutant *C. difficile* toxin B having

a D286A/D288A C698A mutation (SEQ ID NO: 6); (5) a recombinant toxin A having SEQ ID NO: 1; (6) a recombinant toxin B having SEQ ID NO: 2; (7) a mutant *C. difficile* toxin A having a C700A mutation; (8) a mutant *C. difficile* toxin B having a C698A mutation; (9) a mutant *C. difficile* toxin A having a C700A C597S, C1169S, C1407S, 5 C1623S, C2023S, and C2236S mutation; (10) a mutant *C. difficile* toxin B having a C698A C395S, C595S, C824S, C870S, C1167S, C1625S, C1687S, and C2232S mutation; (11) a mutant *C. difficile* toxin A having a D285A, D287A, C700A, D269A, R272A, E460A, and R462A mutation (SEQ ID NO: 7); and (12) a mutant *C. difficile* toxin B having a D270A, R273A, D286A, D288A, D461A, K463A, and C698A mutation (SEQ 10 ID NO: 8)

Penta mutant toxins were also constructed by site directed mutagenesis and by using the same materials and methods as described above, e.g., in Examples 1-7. The penta mutant for toxin B included the following substitutions D286A/D288A C698A/E970K/E976K (SEQ ID NO: 184). The penta mutant toxin B was expressed in VPI 15 11186 *spo0A*-negative cells as described above. A western blot using mAb# B8-26 (SEQ ID NO: 49), which is specific to an N-terminal epitope of toxin B, was done to confirm expression of the penta mutant toxin B. In the second, third, and fourth lanes from the left, 50 ng, 30 ng, and 10 ng of purified triple mutant B (SEQ ID NO: 86), respectively, were used as a reference protein. In the fifth and sixth lanes from the left, a 1:100 dilution and a 1:1000 dilution, 20 respectively, of the penta mutant toxin B cell lysate concentrate was assessed. The estimated amount of protein in the penta mutant toxin B concentration is about 1000 ug/mL. As shown on the blot, the penta mutant toxin B (concentrated) exhibits a protein band of expected size at about 270 kD.

Example 9: Stability of Transformants

Rearranged plasmids were obtained with the commonly-used DH5 \square *E. coli* lab 25 strain. In contrast, transformations using the Invitrogen Stbl2TM *E. coli* host yielded slow-growing full-length mutant toxin recombinants after three days of growth at 30°C on LB chloramphenicol (25 μ g/ml) plates. Lower cloning efficiencies were obtained with related Stbl3TM and Stbl4TM *E. coli* strains, although these lines were found to be stable 30 for plasmid maintenance. Transformants were subsequently propagated in agar or in liquid culture under chloramphenicol selection at 30°C. The use of LB (Miller's) media was also found to improve the recovery and growth of transformants compared with animal-free tryptone-soy based media.

Example 10: Transformation of *C. difficile* with pMTL84121^{fdx} encoding wild-type or genetic mutant toxin genes

Transformation of *C. difficile* by *E. coli* conjugal transfer was done essentially as described in Heap et al., *Journal of Microbiological Methods*, 2009. 78(1): p. 79-85. *E. coli* host CA434 was transformed with pMTL84121 ^{fdx} encoding wild type or variant mutant toxin genes. *E. coli* host CA434 is the intermediate to mobilize expression plasmids into the *C. difficile* production strain VPI 11186 spo0A1. CA434 is a derivative of *E. coli* HB101. This strain harbors the Tra+ Mob+ R702 conjugative plasmid which confers resistance to Km, Tc, Su, Sm/Spe, and Hg (due to Tn1831). Chemically competent or electrocompetent CA434 cells were prepared and expression vector transformants were selected on Miller's LB CAM plates at 30°C. Slow growing colonies appearing after 3 days were picked and amplified in 3 mL LB chloramphenicol cultures until mid-log phase (~24h, 225 rpm, orbital shaker at 30°C). *E. coli* cultures were harvested by low speed (5,000g) centrifugation to avoid breaking pili, and cell pellets were resuspended gently with a wide-bore transfer pipette in 1 mL PBS. Cells were concentrated by low speed centrifugation. Most of the PBS was removed by inversion and the drained pellets were transferred into the anaerobic chamber and resuspended with 0.2 mL of *C. difficile* culture, spotted onto BHIS agar plates and left to grow for 8h or overnight. In the case of mutant toxin A transformants, better results were achieved with overnight conjugation. Cell patches were scraped into 0.5 mL PBS and 0.1 mL was plated on BHIS selection media supplemented with 15 µg/mL thiamphenicol (more potent analog of chloramphenicol) and D-cycloserine / cefoxitin to kill *E. coli* donor cells. Transformants appearing 16-24 h later were purified by re-streaking onto a new BHIS (plus supplements) plate and subsequent cultures were tested for expression of recombinant toxins or mutant toxins. Glycerol permanents and seed stocks were prepared from clones showing good expression. Plasmid minipreps were also prepared from 2 mL cultures using a modified Qiagen kit procedure in which cells were pretreated with lysozyme (not essential). The *C. difficile* miniprep DNA was used as a template for PCR sequencing to verify clone integrity. Alternatively, plasmid maxiprep DNA was prepared from *E. coli* Stbl2™ transformants and sequenced.

Example 11: *C. difficile* expression analysis of the toxin A and B triple mutant (SEQ ID NOs: 4 and 6, respectively) and hepta B mutant (SEQ ID NO: 8)

Transformants were grown either in 2 mL cultures (for routine analysis) or in vent-capped flasks (for time course experiments). Samples (2 mL) were centrifuged 5 briefly (10,000 rpm, 30s) to concentrate the cells: supernatants were decanted and concentrated 10x (Amicon-ultraTM 30k); pellets were drained and frozen at -80°C. Cell pellets were thawed on ice, resuspended in 1mL lysis buffer (Tris-HCl pH7.5; 1mM EDTA, 15% glycerol) and sonicated (1x 20s burst with microtip). The lysate was centrifuged at 4°C and supernatant was concentrated 5-fold. Samples of supernatant 10 and lysate were combined with sample buffer and heat treated (10 min, 80°C) before loading onto duplicate 3-8% Tris-acetate SDS-PAGE gels (Invitrogen). One gel was stained with Coomassie, the second was electroblotted for western analysis. Toxin A-specific and Toxin B-specific rabbit antisera (Fitgerald; Biodesign) were used to detect mutant toxin A and B variants. Expression of the hepta mutant toxin B (SEQ ID NO: 8) 15 was also confirmed by western blot hybridization.

Example 12: Abrogation of glucosyltransferase activity of the mutant toxins

Genetic double mutant (DM) toxins A and B (SEQ ID NOs: 3 and 5, respectively) and triple mutant (TM) toxins A and B (SEQ ID NOs: 4 and 6, respectively) did not transfer ¹⁴C-glucose to 10 µg of RhoA, Rac1 and Cdc42 GTPases in *in vitro* 20 glucosylation assays in the presence of UDP-¹⁴C-glucose [30 µM], 50 mM HEPES, pH 7.2, 100 mM KCl, 4 mM MgCl₂, 2 mM MnCl₂, 1 mM DTT, and 0.1 µg/µL BSA. However, wild-type A and B toxin controls (having SEQ ID NOs: 1 and 2, respectively) transferred ¹⁴C-glucose to GTPases efficiently at a low dose of 10 and 1 ng each (and lower- data not shown) (Figures 4A and 4B), even in the presence of 100 µg of mutant toxin (Figure 25 4B) indicating at least 100,000-fold reduction compared to respective wild-type toxins. Similar results were detected for Cdc42 GTPase (data not shown).

Specifically, in Fig. 4B, wild-type toxin A and toxin B (1ng) or triple mutant toxin A and triple mutant toxin B (100 µg) were incubated with RhoA GTPase in the presence of UDP-¹⁴C-glucose for 2 hr at 30°C. As illustrated, 1 ng of wild-type TcdA and TcdB 30 transferred ¹⁴C-glucose to RhoA but 100 µg of triple mutant toxin A and triple mutant toxin B did not. When 1 ng of wild-type TcdA or TcdB was spiked into the reaction with respective 100 µg of triple mutant toxin A or triple mutant toxin B, glucosylation of RhoA was detected, indicating the lack of glucosylation inhibitors. The sensitivity of detection

for the glucosylation activity was established to be 1 ng of wild-type toxin in a background of 100 µg mutant toxin (ratio of 1:100,000). The results show that the mutations in the active site of the glucosyltransferase in the triple mutant toxin A and triple mutant toxin B reduced any measurable (less than 100,000-fold lower activity
5 compared to the activity of the respective wild-type toxins) glucosyltransferase activity. A similar assay was also developed to quantify glucosyltransferase activity by TCA precipitation of glucosylated GTPases.

Example 13: Abrogation of Auto-catalytic Cysteine Protease Activity

The function of auto-catalytic cleavage was abrogated in the triple genetic mutants A and B (TM) (SEQ ID NOs: 4 and 6, respectively) when the cysteine protease cleavage site was mutated. As illustrated in Figure 5, the wild type (*wt*) toxins A and B (SEQ ID NOs: 3 and 5, respectively) are cleaved in the presence of inositol-6-phosphate. The double mutant toxins A and B (SEQ ID NOs: 3 and 5, respectively) are also cleaved in the presence of inositol-6-phosphate (data not shown), similar to that for wild-type. Toxin A (SEQ ID NO: 3) is cleaved from 308 kDa into 2 fragments of 245 and 60 kDa. Toxin B (SEQ ID NO: 5) is cleaved from 270 kDa into two fragments of 207 and 63 kDa. The triple genetic mutants A and B (TM) (SEQ ID NOs: 4 and 6, respectively) remain unaffected at 308 and 270 kDa respectively, even in the presence of inositol-6-phosphate. See Figure 5. Therefore, the cysteine protease activity was inactivated by genetic modification.

More specifically, in figure 5, one µg of triple mutant A and triple mutant B were incubated for 90 minutes at room temperature ($21 \pm 5^\circ\text{C}$) in parallel with wild-type TcdA and TcdB from List Biologicals. The cleavage reaction was performed in 20 µL volume in Tris-HCl, pH 7.5, 2mM DTT in the presence or absence of inositol-6-phosphate (10mM for TcdA and 0.1 mM for Tcd B). The entire reaction volume was then loaded on a 3 -8% SDS/PAGE; the protein bands were visualized by silver staining. As illustrated, wt Tcd A and TcdB were cleaved into two protein bands of 245kD and 60kD and 207kD and 63 kD, respectively, in the presence of inositol-6-phosphate. The triple mutant toxin A and triple mutant toxin B were not cleaved, thus confirming the C700A mutation in triple mutant toxin A and C698A mutation in triple mutant toxin B blocked cleavage.

Example 14: Residual Cytotoxicity of Triple Mutant Toxins A and B (SEQ ID NOs: 4 and 6, respectively)

The genetic mutant toxins were evaluated for their cytotoxicity by an *in vitro* cytotoxicity assay in IMR90 cells, a human diploid lung fibroblast cell line. These cells 5 are sensitive to both toxin A and B. As an alternative preferred embodiment, Vero normal kidney cells from *Cercopithecus aethiops* may be used in the cytotoxicity assay since they were observed to have reasonable sensitivities to toxin A and B. Preferably, HT-29 human colorectal adenocarcinoma cells are not used in the cytotoxicity assay because they have shown significantly decreased sensitivities to the 10 toxins, as compared to the Vero and IMR90 cell lines. See, for example, Table 6 below.

Table 6: Cell Line Sensitivities to Toxins A and B*				
Cell line	Toxin 50 µg/ml	Cells/well	EC ₅₀ (pg/ml)	
			48 hours	72 hours
Vero (ATCC CCL-81™)	A	10000	1816	244
	B	10000	62	29
IMR90 (ATCC CCL-186TM)	A	10000	1329	1152
	B	10000	14	13
HT-29 (ATCC HTB-38™)	A	10000	>1E6	>1E6
	B	10000	11089	53313

**In vitro* cytotoxicity assay was performed by measuring cellular ATP using luciferase-based substrate, CellTiter-Glo® (Promega, Madison, WI)

Serially diluted genetic mutant toxin or *wt* toxin samples were added to the cell monolayers grown in 96-well tissue culture plates. After incubation at 37°C for 72h, the plates were evaluated for metabolically active cells by measuring cellular ATP levels by 15 addition of luciferase based CellTiterGlo® reagent (Promega, Madison, WI) generating luminescence expressed as relative luminescence units (RLUs). High RLUs show that the cells are viable, low RLUs show that the cells are not metabolically active and are dying. The level of cytotoxicity, expressed as EC₅₀, is defined as the amount of *wt* toxin or genetic mutant toxin that elicits a 50% reduction in RLUs compared to levels in cell 20 culture controls (details of this assay are provided below). As shown in Figure 6, Tables 7A, and Table 8A, the EC₅₀ values of TcdA and TcdB were about 0.92 ng/mL and 0.009 ng/mL, respectively. The EC₅₀ values of triple mutant toxin A and triple mutant toxin B

were about 8600 ng/mL and 74 ng/mL, respectively. Despite an approximate 10,000-fold reduction in cytotoxicity relative to *wt* toxins, both genetic mutant toxins still demonstrated low residual levels of cytotoxicity. This residual cytotoxicity could be blocked by neutralizing antitoxin monoclonal antibodies indicating that it was specific to 5 the triple mutant toxins but not likely related to the known enzymatic activities of the *wt* toxins (glucosylation or autoproteolysis).

Both *wt* toxins exhibit potent *in vitro* cytotoxicity, with small amounts of the toxins being sufficient to cause various effects on mammalian cells such as cell rounding (cytopathic effect or CPE) and lack of metabolic activity (as measured by ATP levels). 10 Consequently, two *in vitro* assays (a CPE or cell rounding assay and an ATP assay) have been developed to verify that no residual cytotoxicity in the mutant toxin drug substances remains. The results are expressed as EC₅₀, which is the amount of toxin or mutant toxin that causes 1) 50% of the cells to develop CPE or 2) 50% reduction in ATP levels as measured in relative light units.

15 In the CPE assay, a sample of drug substance is serially diluted and incubated with IMR90 cells, which are observed for a potential cytopathic effect. The CPE assay is scored on a scale of 0 (normal cells) to 4 (~100 % cell rounding) and a score of 2 (~50% cell rounding) is defined as EC₅₀ value of the test sample. This method is used for testing of mutant toxin drug substance at the concentration of 1000 µg /mL, which is the 20 maximal tolerable concentration that can be tested in this assay without matrix interference. Consequently, no detectable cytotoxicity is reported as EC₅₀ >1000 µg/ml.

The ATP assay is based on measurement of the amount of luminescence signal generated from ATP, which is proportional to the number of metabolically active cells. The maximal tolerable concentration that can be tested in this assay without assay 25 interference is about 200 µg/mL. Therefore no detectable cytotoxicity in this assay is reported as EC₅₀ >200 µg/mL.

Different concentrations of mutant toxin A and B were added to cells in parallel with toxin controls. The endpoints of the assay were cell viability determined by cellular 30 ATP levels using the CellTiter-Glo® (Promega). The degree of luminescence is proportional to ATP levels or viable cell number.

The *in vitro* cytotoxicity (EC₅₀) of wild type (wt) toxin A was 920 pg/mL and 9 pg/mL for toxin B. The *in vitro* cytotoxicity (EC₅₀) of mutant toxin A (SEQ ID NO: 4) was 8600 ng/mL and 74 ng/mL for mutant toxin B (SEQ ID NO: 6). Although these values represent reductions of 9348 and 8222-fold, respectively, residual cytotoxicity was 5 detected in both mutant toxins.

In other words, the cytotoxicity of triple mutant toxins A and B (SEQ ID NOs: 4 and 6, respectively) was significantly reduced in the *in vitro* cytotoxicity assay in IMR-90 cells relative to the cytotoxicity of wt toxins A and B (SEQ ID NOs: 1 and 2, respectively). As illustrated in Figure 6, although both triple mutant toxins exhibited 10 significant reduction in cytotoxicity (10⁴ fold) relative to the wt toxin, residual cytotoxicity was observed at higher concentrations of both triple mutant toxins.

Furthermore, the residual cytotoxicity of each triple mutant toxin could be completely neutralized (e.g., at least a 6-8 log₁₀ reduction in toxicity, relative to the wild-type toxin toxicity) by the toxin specific antibodies. See Example 16, below.

15 Cell culture assays are more sensitive for detection of cytotoxicity than *in vivo* animal models. When delivered by either i.p. or i.v routes in the mouse lethal challenge model, the *wt* TcdA has an LD₅₀ of ~50 ng per mouse while the *wt* TcdB is more potent with an LD₅₀ of ~5 ng per mouse. In contrast, the cell culture based *in vitro* assays described above have EC₅₀ values of 100 pg per well for *wt* TcdA and 2 pg per well for 20 *wt* TcdB.

Example 15: Residual Cytotoxicity of the Genetic Hepta Mutant Toxin B (SEQ ID NO: 8), and Cytotoxicity of Penta Mutant Toxin B (SEQ ID NO: 184)

As illustrated in Figure 7, the EC₅₀ values are similar for the triple mutant toxin B (SEQ ID NO: 6) (20.78 ng/mL) and hepta mutant toxin B (SEQ ID NO: 8) (35.9 ng/mL)

5 mutants indicating that the four additional mutations to further modify the glucosyltransferase active site and GTPase substrate binding site did not further reduce the cytotoxicity of the genetic mutant toxins. The EC₅₀ values were also similar for the double mutant toxin B (SEQ ID NO: 5) as they are for the triple and hepta mutant toxins (data not shown). This observation suggests the mechanism for cytotoxicity of the 10 mutant toxins is surprisingly independent of the glucosyltransferase and substrate recognition mechanism.

The penta mutant toxin B (SEQ ID NO: 184) was evaluated for its cytotoxicity by an *in vitro* ATP cytotoxicity assay in IMR90 cells, as described above. See, e.g., Example 14. As shown in Figure 26, the cytotoxicity of the penta mutant toxin B was 15 greatly reduced, as compared to the wild-type toxin B (e.g., SEQ ID NO: 2), which was obtained commercially from List Biologicals), and as compared to the triple mutant toxin B (SEQ ID NO: 86). The “cells only” control relates to IMR-90 cells. See Table 62 below, showing the respective EC₅₀ values.

Table 62

Sample	EC ₅₀	Fold-reduction in EC ₅₀
Penta Mutant B (lysate concentrate)	11.5 µg/mL	1,642,857
Triple Mutant B (Purified)	18.9 ng/mL	2700
List Toxin B lot127902-115	7.0 pg/mL	1

20

Moreover, the penta mutant toxin B (“PM-B”)(SEQ ID NO: 184) was assessed for competitive inhibition of cytotoxicity that is mediated by the triple mutant toxin B (SEQ ID NO: 86) on IMR-90 cells. See Figure 27. To prepare the samples, the triple mutant toxin B (TM B)(SEQ ID NO: 86) was kept at a constant concentration of 200 ng/mL (10 25 x EC₅₀) in all wells. A 2-fold serially diluted penta mutant toxin B starting at 5 µg/mL was added to the wells containing the triple mutant toxin B. The samples were then transferred to a 96-well plate containing IMR-90 cells, then incubated for 72 hours. An ATP assay was completed, as described above, e.g., in Example 14. As can be shown

in Figure 27, the penta mutant toxin B competitively inhibited cytotoxicity of the triple mutant toxin B.

Example 16: Residual Cytotoxicity of Triple Mutant Toxins A and B (SEQ ID NOs: 4 and 6, respectively)

To further evaluate the nature of the residual cytotoxicity, the mutant toxins (SEQ ID NOs: 4 and 6) were mixed and incubated with their respective neutralizing antibodies 5 before and the mixture was added to IMR90 cell monolayer.

The results (Figure 8) showed that the residual cytotoxicity of mutant toxin A and B (SEQ ID NOs: 4 and 6, respectively) can be completely abrogated with neutralizing antibodies specific for mutant toxin A (top panel, Figure 8) and mutant toxin B (bottom panel, Figure 8). Increasing concentrations of mutant toxin A (top panel) and B (bottom 10 panel) were incubated with rabbit anti-toxin polyclonal (pAb, 1:10 dilution) or murine monoclonal antibodies (1:50 dilution from a stock containing 3.0 mg IgG/mL) before adding to IMR90 cells. After 72-hr treatment incubation with IMR90 cells at 37°C, CellTiter-Glo® substrate was added and the relative light units (RLU) were measured in a spectrophotometer with the luminescence program to measure ATP levels. The lower 15 the ATP level, the higher the toxicity. Controls included TcdA and TcdB added at 4 times their corresponding EC₅₀ values.

Published reports suggest that mutations in the glucosyltransferase or autocatalytic protease domain of the toxins result in complete inactivation of the toxicity. However, our data do not agree with these published reports and this could be 20 attributed to increased concentrations of the highly purified mutant toxins tested in our studies as opposed to crude culture lysates in published reports; increased time points at which cell rounding of mutant toxin-treated cells was observed (e.g., 24 hours, 48 hours, 72 hours, or 96 hours) as opposed to observations made in less than 12 hours; use of cell lines that exhibit significantly higher sensitivities to toxins in present 25 cytotoxicity assays in contrast to HT-29 human colorectal adenocarcinoma cells in cytotoxicity assays disclosed in published reports; and/or to an unknown activity or process, other than glycosylation, that could be driving the residual toxicity of the mutant toxins.

Example 17: Novel Mechanism of Cytotoxicity of Genetic Mutant Toxins

To investigate the mechanism of residual cytotoxicity of the genetic mutant toxins, IMR-90 cells were treated with wt toxin B (SEQ ID NO: 2) or genetic mutant toxin B (SEQ ID NO: 6), and glucosylation of Rac1 GTPase was studied with time of treatment. Samples were collected from 24 to 96 hours and cell extracts were prepared. Glucosylated Rac1 is distinguished from non-glucosylated Rac1 by western blots with two antibodies to Rac1. One antibody recognizes both forms of Rac1 (23A8) and the other (102) only recognizes non-glucosylated Rac1. As illustrated in Figure 22, for toxin B, the total Rac1 levels stayed unchanged over time with majority of the Rac1 being glucosylated. Treatment with the genetic mutant toxin B (SEQ ID NO: 6), on the other hand, resulted in significant reduction of total Rac1, however, the Rac1 was non-glucosylated at all time points. This shows that Rac1 level was negatively affected by the treatment with the genetic mutant toxin, but not by wt toxin. As illustrated in Figure 22, the level of actin was similar in toxin and genetic mutant toxin B treated cells and similar to mock treated cells at indicated time points. This showed that the genetic mutant toxins exerted cytotoxicity by a mechanism which is different than the wild-type toxin-driven glucosylation pathway.

Example 18: Chemical treatment of genetic mutant toxins

Although the genetically modified mutant toxins showed a 4-log reduction in cytotoxic activity is preferred, further reduction (2 to 4 logs) in cytotoxic activity was considered. Two chemical inactivation strategies have been evaluated.

5 The first method uses formaldehyde and glycine to inactivate the mutant toxins. Formaldehyde inactivation occurs by forming a Schiff base (imine) between formaldehyde and primary amines on the protein. The Schiff bases can then react with a number of amino acid residues (Arg, His, Trp, Tyr, Gln, Asn) to form either intra- or intermolecular crosslinks. This crosslinking fixates the structure of the protein rendering 10 it inactive. In addition, formaldehyde can react with glycine to form a Schiff base. The glycyl Schiff base can then react with the amino acid residues to form intermolecular protein-glycine crosslinks. Formaldehyde reduced the cytotoxic activity of the genetic mutant toxins to below detectable limits (reduction in cytotoxicity $>8 \log_{10}$ for triple mutant B (SEQ ID NO: 6) and $>6 \log_{10}$ for triple mutant A (SEQ ID NO: 4). However, 15 reversion was observed over time when the formaldehyde-inactivated (FI) triple mutant toxins were incubated at 25°C. The cytotoxic reversion can be prevented by addition of a low amount of formaldehyde (0.01-0.02%) into the FI-triple mutant toxins storage solution. See Example 23.

Another method uses 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide (EDC)/N-20 hydroxysuccinimide (NHS) treatment to generate inactivated mutant toxins. In this method, EDC/NHS reacts with carboxylic groups on the protein to form activated esters. The activated esters can then react with primary amines on the protein to form stable amide bonds. As with the formaldehyde reaction, this reaction results in intra- and intermolecular crosslinks. The amide bond formed by treatment with EDC/NHS is more 25 stable and non-reversible than the labile imine bond formed by formalin inactivation. In addition to crosslinks formed by the reaction of activated esters with primary amines on the polypeptide, both glycine and beta-alanine adducts can be formed. Without being bound by mechanism or theory, glycine adducts are produced when glycine is added to quench unreacted activated esters. The amine of glycine reacts with the activated ester 30 on the polypeptide to form stable amide bonds. Without being bound by mechanism or theory, beta-alanine adducts are formed by the reaction of activated beta-alanine with primary amines on the polypeptide. This reaction results in stable amide bonds.

Activated beta-alanine is produced by the reaction of three moles of NHS with one mole of EDC.

To achieve the 2-4 logs reduction of cytotoxic activity relative to the genetically modified mutant toxins (6-8 logs, relative to native toxins), the chemically inactivated 5 mutant toxins should have EC₅₀ values of $\geq 1000 \mu\text{g/mL}$. In addition to reduction in cytotoxic activity, it would be advantageous to retain key epitopes as determined by dot-blot analysis. To date, a number of reaction conditions have been identified that meet both the reduction cytotoxicity and epitope recognition criteria. Several batches of inactivated mutant toxins have been prepared for animal studies and analytical data 10 from a few representative batches is shown in Tables 7A and 7B, Table 8A and 8B.

Table 7A: Chemically Inactivated Mutant Toxin A is Safe and Antigenic					
Sample #	Toxin Sample ID	Treatment	CPE EC ₅₀	Reduction in toxicity	Reactivities to mAbs
			$\mu\text{g/mL}$	Log Scale	
1	Mutant TcdA (SEQ ID NO: 4) L44905-160A	Formalin	>1000	6.4	Medium/high
2	Mutant TcdA (SEQ ID NO: 4) L44166-166	EDC	>1000	6.4	High
3	Mutant TcdA (SEQ ID NO: 4) L44905-170A	Formalin	>1000	6.4	Low
CONTROLS					
4	TcdA wt (from List Bio)	none	390 pg/mL	1	High
5	TcdB wt (from List Bio)	none	3.90 pg/mL	Not applicable	None
6	rMutant TcdA TM Genetic L36901-79 (SEQ ID NO: 4)	none	12.5 $\mu\text{g/mL}$	4.5	High
7	Toxoid A List Bio	Formalin	Not Done	---	Low

Table 7B: Chemically inactivated Mutant Toxin A is Safe and Antigenic

Sample #	Toxin Sample ID	Treatment	Reactivity with mAb (dot blot, nondenaturing conditions)					
			N-terminal mAb#6	Mid- Domain mAb # 102	C-terminal (neut)			
					A80- 29	A3- 25	A60- 22	A65- 33
1	Mutant TcdA (SEQ ID NO: 4) L44905-160A	Formalin	++	++	++++	++	++++	++++
2	Mutant TcdA (SEQ ID NO: 4) L44166-166	EDC	++++	++++	++++	++++	++++	++++
3	Mutant TcdA (SEQ ID NO: 4) L44905-170A	Formalin	+	+	++	++	++	+
CONTROLS								
4	TcdA wt (from List Bio)	none	++++	+++	++++	++++	++++	++++
5	TcdB wt (from List Bio)	none	—	—	—	—	—	—
6	rMutant TcdA TM Genetic L36901-79 (SEQ ID NO: 4)	none	++++	++++	++++	++++	++++	++++
7	Toxoid A List Bio	Formalin	—	—	+	—	++	+

List =List Biologicals; CPE= cytopathic effect assay; EC₅₀ = the lowest concentration

where 50% of the cells show cytotoxicity; mAbs= monoclonal antibodies; neut=

neutralizing; ND= not done; TM= active site and cleavage mutant ("triple mutant")

Table 8A: Chemically Inactivated Mutant Toxin B is Safe and Antigenic

Sample #	Toxin Sample ID	Treatment	CPE EC ₅₀	Reduction in toxicity	Reactivities to mAbs
			µg/mL	Log Scale	
1	Mutant TcdB L44905-182 (SEQ ID NO: 6)	Formalin	>1000	8.4	Medium/high
2	Mutant TcdB L34346-38A (SEQ ID NO: 6)	EDC	>1000	8.4	High
3	Mutant TcdB L44905-170B (SEQ ID NO: 6)	Formalin	>1000	8.4	Low
CONTROLS					
4	TcdA wt (from List Bio)	none	390 pg/mL	Not applicable	None
5	TcdB wt (from List Bio)	none	3.90 pg/mL	1	High
6	rMutant toxin B TM Genetic (SEQ ID NO: 6) L34346-022	none	69 ng/mL	4.2	High
7	Toxoid A List	Formalin	Not done	---	Medium

Table 8B: Chemically Inactivated Mutant Toxin B is Safe and Antigenic

Sample #	Toxin Sample ID	Treatment	Reactivity with mAb (dot blot, nondenaturing conditions)			
			N-terminal aa 1-543		Mid-/C-terminal aa 544-2366	
			B8-26	B9-30	B56-6	B59-3
1	Mutant TcdB (SEQ ID NO: 6) L44905-160A	Formalin	+++	+++	++	++
2	Mutant TcdB (SEQ ID NO: 6) L44166-166	EDC	++++	++++	++++	++++
3	Mutant TcdB (SEQ ID NO: 6) L44905-170A	Formalin	++	+	+/-	-
CONTROLS						
4	TcdA wt (from List Bio)	none	—	—	—	—
5	TcdB wt (from List Bio)	none	++++	+++	++++	++++
6	rMutant TcdB TM Genetic L34346-022 (SEQ ID NO: 6)	none	++++	++++	++++	++++
7	Toxoid B List	Formalin	+++	+++	+++	+++

List =List Biologicals; CPE= cytopathic effect assay; EC₅₀ = the concentration where 50% of the cells show cytotoxicity; mAbs= monoclonal antibodies; neut= neutralizing; ND= not

5 done; TM= active site and cleavage mutant ("triple mutant")

Example 19: Fermentation and Purification

Fermentations were initiated from a frozen source of a recombinant *Clostridium difficile* including an *fdx* promoter described above. The frozen stocks were a cell suspension made to an OD₆₀₀=2.0 and 20% glycerol. A starter culture was inoculated 5 with 0.2 mL of the culture stock into 500mL SHYG10 medium (30 g/L soy hydrolysate SE50MK, 20 g/L yeast extract HY YEST 412, 10 g/L glucose, 15 mg/L thiampenicol , pH7). The medium was contained in a 500mL vented bottle. The inoculation was performed in a conventional biosafety cabinet, the bottle was flushed with nitrogen, and the bottle was then incubated static (vents closed) for ~ 16-18 hours at 37°C in a 10 conventional incubator.

A ten liter bioreactor containing 8L of SHYG60 medium (30 g/L soy hydrolysate SE50MK, 20 g/L yeast extract HY YEST 412, 60 g/L glucose, 15 mg/L thiampenicol, pH7) was used for the fermentation phase. The 500mL contents of the inoculum bottle were inoculated to the fermentor which was operated at 37°C, 400 rpm (1.47 m/s) with 15 0.1 vvm nitrogen sparge. The pH was controlled at 7.0 by auto-addition of 5N NaOH. Fermentation was typically run for ~ 24hours to reach peak potency. Growth during the course of the fermentation was monitored by OD₆₀₀ readings. Samples taken for mutant toxin quantitation were spun at ~5000xg and the resulting pellets were frozen at -70°C. The cell pellet was then defrosted and resuspended in a buffer consisting of 20mM Tris, 20 3mM NaCl, 0.5 mM EDTA, pH 6.5 and sonicated at an amplitude of 40 for 20 seconds. The resulting cell lysate was spun at 5,000xg for 10 min. The supernatant was combined with loading buffer and reducing agent and run on a 3.8% Tris-acetate PAGE gel at 150 volts for 50-55 min versus authentic toxin standards. The gel was stained overnight, then destained and quantitated on a scanning densitometer. Typically, OD₆₀₀ 25 values of ~ 10-12 were observed with toxin values of 80-120 mg/L.

The following table is an example of fermentation data for mutant toxin A.

Table 63		
Elapsed fermentation time (hrs)	OD ₆₀₀	Toxin yield (mg/L)
1	0.32	
3	1.16	
5	3.55	33
7	5.56	53
9	7.11	63
11	8.64	81
24	11.14	123

An example of fermentation data for mutant toxin B is presented in the table below.

Table 64		
Elapsed fermentation time (hrs)	OD ₆₀₀	Toxin yield (mg/L)
1	0.73	
3	1.18	
5	4.54	30
7	5.93	40
9	7.05	48
11	8.44	62
24	10.03	94

5 Modifications of the composition and methods for culturing the recombinant *C. difficile* cell and/or production of the mutant toxins have been tested and are within the scope of the invention. For example, although SE50MK (and variously SE50MK-NK, both sourced from Friesland-Campaigna) was the preferred choice for nitrogen source, a variety of other soy hydrolysates from other manufacturers were identified that worked
10 well in the process. A culture medium including soy hydrolysate in the absence of yeast extract provided 30-40% of the expected yield.

Yeast extracts from alternative manufacturers were shown to support equivalent yields. A culture medium including yeast extract in the absence of soy hydrolysate provided 60-70% of the expected yield.

Use of a culture medium based on soy hydrolysate/yeast extract, in the absence of a carbon source, yielded about 2-3 OD₆₀₀ and about 10-15 mg/L toxin. Although glucose was a preferred carbon source for culturing, equivalent results were obtained with mannitol. A screen of carbon sources in bottles, indicated that *C. difficile* may also utilize fructose and mannose which would also be expected to support high OD/yield, as compared to a culture medium in the absence of a carbon source. The following carbon sources did not appear to support optimal growth: arabinose, xylose, sucrose, lactose, maltose, glycerol, rhamnose and galactose.

In addition, extending the fermentation time to 48 hours (requiring addition of more glucose) did not appear to substantially improve yield.

Further, fermentation at pH 6.5 and 7.5 gave yields in the expected range of 80-120 mg/L. Fermentation at more extreme pH (pH 6.0 or 8.0) still gave the expected OD₆₀₀ values, but reduced yields (40-60 mg/L) of toxin.

Fermentation using a culture medium in the absence of thiamphenicol resulted in a loss of plasmid, e.g., about 10-20% for a plasmid encoding mutant toxin B, and about 30-40% for a plasmid encoding mutant toxin A. Accordingly, fermentation using a culture medium in the absence of a chloramphenicol derivative was feasible.

Alternative modes of operating fermentation were also tested and are within the scope of the invention. For example, fermentation may be run at 400 rpm or less and nitrogen overlay may be used, both techniques of which were tried and used successfully.

Moreover, a monoclonal antibody medium SFM4MAb was tested and was shown to give about 10 OD₆₀₀ of cells and about 40 mg/L of mutant toxin.

Lastly, addition of a phosphate-containing ingredient to the fermentation appeared to reduce the production of toxin, as compared to culture medium in the absence of the phosphate-containing ingredient.

At the end of fermentation, the fermenter is cooled. The cell slurry is recovered by continuous centrifugation and re-suspended in the appropriate buffer. Lysis of the cell suspension is achieved by high-pressure homogenization. For mutant toxin A, the homogenate is flocculated and the flocculated solution undergoes continuous

centrifugation. This solution is filtered and then transferred for downstream processing. For mutant toxin B, the homogenate is clarified by continuous centrifugation, and then transferred for downstream processing.

Mutant toxin A (SEQ ID NO: 4) is purified using two chromatographic steps 5 followed by a final buffer exchange. The clarified lysate is loaded onto a hydrophobic interaction chromatography (HIC) column and the bound mutant toxin is eluted using a sodium citrate gradient. The product pool from the HIC column is then loaded on a cation exchange (CEX) column and the bound mutant toxin A is eluted using a sodium chloride gradient. The CEX pool containing purified mutant toxin A is exchanged into 10 the final buffer by diafiltration. The purified mutant toxin A is exchanged into the final drug substance intermediate buffer by diafiltration. After diafiltration, the retentate is filtered through a 0.2 micron filter prior to chemically inactivation to a final drug substance. The protein concentration is targeted to 1-3 mg/mL.

Mutant toxin B (SEQ ID NO: 6) is purified using two chromatographic steps 15 followed by a final buffer exchange. The clarified lysate is loaded onto an anion exchange (AEX) column, and the bound mutant toxin is eluted using a sodium chloride gradient. Sodium citrate is added to the product pool from the AEX column and loaded on a hydrophobic interaction chromatography (HIC) column. The bound mutant toxin is eluted using a sodium citrate gradient. The HIC pool containing purified mutant toxin 20 polypeptide (SEQ ID NO: 6) is exchanged into the final buffer by diafiltration . The purified mutant toxin B is exchanged into the final drug substance intermediate buffer by diafiltration. After diafiltration, the retentate is filtered through a 0.2 micron filter prior to chemically inactivation to a final drug substance. The protein concentration is targeted to 1-3 mg/mL.

Example 20: Formaldehyde/Glycine Inactivation

After purification, the genetic mutant toxins A and B (SEQ ID NOs: 4 and 6, respectively) are inactivated for 48 hours at 25°C using 40 mM (1.2 mg/ml) of formaldehyde. The inactivation is carried out at pH 7.0 ± 0.5 in 10 mM phosphate, 150 mM sodium chloride buffer containing 40 mM (3 mg/ml) glycine. The inactivation period is set to exceed three times the period needed for reduction in the EC₅₀ in IMR90 cells to greater than 1000 μ g/mL. After 48 hours, the biological activity is reduced 7 to 8 \log_{10} relative to the native toxin. Following the 48 hour incubation, the inactivated mutant toxin is exchanged into the final drug substance buffer by diafiltration. For example, using a 100 kD regenerated cellulose acetate ultrafiltration cassette, the inactivated toxin is concentrated to 1-2 mg/mL and buffer-exchanged.

Example 21: N-(3-Dimethylaminopropyl)-N'-ethylcarbodiimide (EDC)/N-hydroxysuccinimide (NHS) Inactivation

After purification, the genetic mutant toxins (SEQ ID NO: 4 and SEQ ID NO: 6) are inactivated for 2 hours at 25°C using 0.5 mg EDC and 0.5 mg NHS per mg of

5 purified genetic mutant toxin A and B (approximately 2.6 mM and 4.4 mM respectively). The reaction is quenched by the addition of glycine to a final concentration of 100 mM and the reactions incubate for an additional 2 hours at 25°C. The inactivation is carried out at pH 7.0 ± 0.5 in 10 mM phosphate, 150 mM sodium chloride buffer. The inactivation period is set to exceed three times the period needed for reduction in the 10 EC₅₀ in IMR90 cells to greater than 1000 ug/mL. After 2 hours, the biological activity is reduced 7 to 8 log₁₀ relative to the native toxin. Following the 4 hour incubation, the inactivated mutant toxin is exchanged into the final drug substance buffer by diafiltration. For example, using a 100 kD regenerated cellulose acetate ultrafiltration cassette, the inactivated toxin is concentrated to 1-2 mg/mL and buffer-exchanged.

15 Unless otherwise stated, the following terms as used in the Examples section refer to a composition produced according to the present description in Example 21: “EDC/NHS-treated triple mutant toxin”; “EDC-inactivated mutant toxin”; “mutant toxin [A/B] drug substance”; “EI-mutant toxin”; “EDC/NHS-triple mutant toxin.” For example, the following terms are synonymous: “EDC/NHS-treated triple mutant toxin A”; “EDC- 20 inactivated mutant toxin A”; “mutant toxin A drug substance”; “EI-mutant toxin A”; “EDC/NHS-triple mutant toxin A.” As another example, the following terms are synonymous: “EDC/NHS-treated triple mutant toxin B”; “EDC-inactivated mutant toxin B”; “mutant toxin B drug substance”; “EI-mutant toxin B”; “EDC/NHS-triple mutant toxin B.”

25 The mutant toxin A drug substance and the mutant toxin B drug substance are each manufactured using a batch process, which includes (1) fermentation of a the toxin negative *C. difficile* strain (VPI 11186) containing a plasmid encoding the respective genetic triple mutant toxin polypeptide (in a medium including soy hydrolysate, yeast extract HY YEST™ 412 (Sheffield Bioscience), glucose, and thiamphenicol), (2) 30 purification of the genetic mutant toxin (the “drug substance intermediate”) from the cell-free lysate using ion exchange and hydrophobic interaction chromatographic procedures to at least greater than 95% purity, (3) chemical inactivation by treatment

with EDC/NHS followed by quenching/capping with glycine, and (4) exchange into the final buffer matrix.

Example 22: Studies Supporting Conditions of Inactivation and Formulation

To optimize the chemical inactivation of the genetic mutant toxins, a statistical design of experiment (DOE) was performed. Factors examined in the DOE included temperature, formaldehyde/glycine concentration, EDC/NHS concentration and time

5 (Table 9 and 10). To monitor loss of biological activity, EC₅₀ values in IMR90 cells were determined. In addition, cell morphology of IMR-90 cells various timepoints post-treatment were also observed. See Figure 9, showing morphology at 72 hours post treatment. To determine the effect on protein structure, epitope recognition was monitored using dot-blot analysis using a panel of monoclonal antibodies raised against

10 different domains of the toxin.

Table 9: Parameters Tested Formaldehyde/Glycine DOE

Parameters	Range tested
Time (days)	1 to 14
Temperature (°C)	4 to 37
Toxin concentration (mg/ml)	1 to 1.25
Formaldehyde concentration (mM)	2 to 80
Glycine concentration (mg/ml)	0 to 80

Table 10: Parameters Tested EDC/NHS DOE

Parameters	Range tested
Time (hours)	1 to 4
Temperature (°C)	25 to 35
Toxin concentration (mg/ml)	1 to 1.25
EDC (mg/mg toxin)	0.25 to 2.5
NHS (mg/mg toxin)	0 to 2.5

In the formaldehyde/glycine inactivation of *C. difficile* mutant toxins, final reaction conditions were chosen such that the desired level of reduction in cytotoxic activity (7 to

15 8 log₁₀) was achieved while maximizing epitope recognition. See Example 20 above.

In the EDC/NHS inactivation of *C. difficile* mutant toxins, final reaction conditions were chosen such that the desired level of reduction in cytotoxic activity (7 to 8 log₁₀) was achieved while maximizing epitope recognition. See Example 21 above.

In an alternative embodiment, the EDC-NHS reaction was quenched by addition of alanine, which sufficiently quenched the reaction. Use of alanine may result in a modification on the mutant toxin protein that is similar to the modification when the reaction is quenched by glycine. For example, quenching by adding alanine may result 5 in an alanine moiety on a side chain of a glutamic acid and/or aspartic acid residue of the mutant toxin. In another alternative embodiment, the EDC-NHS reaction was quenched by addition of glycine methyl ester, which sufficiently quenched the reaction.

Production of chemically inactive triple mutant *C. difficile* toxin A and toxin B under optimized conditions resulted in a further reduction of residual cytotoxicity to an 10 undetectable level (>1000 µg/mL - the highest concentration tested via the CPE assay), while retaining antigenicity as measured by their reactivity to the toxin-specific neutralizing antibodies. The results shown in Table 28 demonstrate a stepwise reduction in cytotoxicity from wt toxin through to EDC/NHS-treated triple mutant toxins. Immunofluorescence labelling confirmed that triple mutant toxins (SEQ ID NO: 4 and 6) 15 and mutant toxin drug substances exhibited comparable binding to the IMR-90 cells suggesting that the cytotoxicity loss was not due to reduced binding to the cells (data not shown). Compared to mutant toxin A drug substance, the mutant toxin B drug substance achieved higher fold-reduction in cytotoxicity, which is consistent with the observed ~600-fold higher potency of TcdB compared to TcdA.

Table 28. Cytotoxicity Summary

Toxin	Sample	EC ₅₀	Fold reduction in cytotoxicity
A	TcdA (SEQ ID NO: 1)	1.6 ng/mL	1
	Triple mutant toxin A (SEQ ID NO: 4)	12.5 µg/mL	7800
	Mutant toxin A Drug Substance	>1000 µg/mL	>625,000
B	TcdB (SEQ ID NO: 2)	2.5 pg/mL	1
	Triple mutant toxin B (SEQ ID NO: 6)	45 ng/mL	18,000
	Mutant toxin B Drug Substance	>1000 µg/mL	>400,000,000

Cytotoxicity assay results for mutant toxin B modified by EDC alone, or by EDC and sulfo-NHS were also assessed. See Table 29.

Table 29

Sample	Cytotoxicity	Comment
	EC ₅₀ , mg.mL ⁻¹ (CPE)	
TcdB TM (SEQ ID NO: 6), unmodified	0.03	
TM TcdB-EDC 1, no NHS	<0.97	Reacted with EDC alone
TM TcdB-EDC 2, no NHS	<0.97	Duplicate preparation
TM TcdB-EDC 3, sulfo-NHS (0.5x)	125	Reacted with EDC and sulfo-NHS
TM TcdB-EDC 4, sulfo-NHS (0.5x)	125	Duplicate preparation
TM TcdB-EDC 3, sulfo-NHS (1.0x)	250	Reacted with EDC and sulfo-NHS
TM TcdB-EDC 4, sulfo-NHS (2.0x)	750	Reacted with EDC and sulfo-NHS

Conditions: Triple mutant toxin B (“TM TcdB”)(SEQ ID NO: 6) was modified in the 5 weight ratios mutant toxin B:EDC:sulfo-NHS = 1: 0.5: 0.94. This ratio is the molar equivalent (corrected for higher MW of sulfo-NHS) to the standard EDC/NHS reaction as described in Example 21. To determine the affect of sulfo-NHS, the sulfo-NHS ratio was varied from 0.5x to 2x the standard ratio. Duplicate reactions were performed in 1 x PBS pH 7.0 at 25 °C, and were initiated by addition of EDC solution. After 2 hours, 10 reactions were quenched by the addition of 1 M glycine pH 7.0 (0.1 M final concentration) and incubated for a further 2 hours. Quenched reactions were desalted and mutant toxin B drug substance (“TM TcdB-EDC”) was concentrated using Vivaspin 20 devices, and sterile filtered into sterile vials and submitted for assessment in a cytotoxicity assay.

15 At the same molar ratio, sulfo-NHS reduced the EC₅₀ to about 250 ug/mL as compared to >1000 ug/mL for NHS. Even at twice the molar ratio, sulfo-NHS does not appear not as effective as NHS in decreasing cytotoxicity. See Table 30.

Table 30

Modification	reference Digest (TcdB EDC 004)	NHS control digest (TcdB EDC 001)	Sulfo-NHS Sample Digest
glycine adduct (+57 da)	49	29	35
beta-alanine (+71 da)	24	19	0
crosslinks (-18 da)	7	4	3
dehydroalanine (-34 da)	6	5	4
Unmodified	273	195	217

To determine the number and type of modifications, peptide mapping was performed on both EDC/NHS and EDC/sulfo-NHS inactivated triple mutant toxin B samples. Similar amounts of glycine adducts, crosslinks and dehydroalanine

5 modifications were observed in both samples. However in the sulfo-NHS sample, no beta-alanine was observed.

Wild-type toxin B (SEQ ID NO: 2) was inactivated using the standard protocol (see Example 21); toxin B:EDC:NHS 1:0.5:0.5, 25°C for 2 hours in 1 x PBS pH 7.0, then quench with 1 M glycine (0.1 M final concentration) and incubate for an additional 2
10 hours. The sample was desalted, concentrated and submitted for cytotoxicity assay. The EC₅₀ for this samples was <244 ng/mL.

Example 23: Reversion Studies

To determine if reversion occurs with either the formaldehyde/glycine or EDC/NHS inactivated *C. difficile* mutant toxins, samples of inactivated mutant toxins (1 mg/mL) were incubated at 25°C for five-six weeks. Aliquots were removed each week 5 and the EC₅₀ values in IMR90 cells were determined. One formaldehyde/glycine inactivated sample contained no formaldehyde and one sample contained 0.01% formaldehyde. The EC₅₀ was measured by the CPE assay.

Table 11: Results from Inactivated TcdA Reversion Study			
Time of Incubation (Days)	EC ₅₀ (IMR90 cell assay)		
	Formalin-inactivated		EDC/NHS
	No formaldehyde	0.01% formaldehyde	
0	1000 ug/ml	1000 ug/ml	1000 ug/ml
7	740 ug/mL	ND	1000 ug/ml
14	493 ug/mL	1000 ug/ml	1000 ug/ml
21	395 ug/mL	ND	1000 ug/ml
28	395 ug/mL	1000 ug/ml	1000 ug/ml
35	326 ug/M	ND	ND

10 At 25°C in the absence of residual formaldehyde, partial reversion is observed (Table 11). After five weeks, the cytotoxic activity increased approximately 3-fold. Although the cytotoxic activity increased, after five weeks there was still a 7 log₁₀ reduction relative to the native toxin. Reversion was completely prevented by inclusion 15 of formalin at a concentration of 0.010%. No reversion was observed in the EDC/NHS inactivated sample. Throughout the 6-week incubation, EC₅₀ values remained at the starting level of >1000 µg/mL for all four lots of both EDC/NHS-treated triple mutant toxin A (SEQ ID NO: 4) and EDC/NHS-treated triple mutant toxin B (SEQ ID NO: 6). In contrast, the EC₅₀ values of FI-treated triple mutant toxin A (SEQ ID NO: 4) and FI-treated triple mutant toxin B (SEQ ID NO: 6) were not stable and declined to 20 unacceptably low EC₅₀ values, indicating an increase in cytotoxicity or reversion of inactivation. See Table 11.

In addition to stably reducing the cytotoxicity to an undetectable level (>1000 µg/mL, as measured by the CPE assay), mutant toxins inactivated using EDC/NHS retained important epitopes that are targets of toxin-neutralizing mAbs. See Table 31. FI mutant toxins showed a loss of the same antigenic determinants.

5 **Table 31. EDC/NHS Inactivation Reduced Cytotoxicity of Genetic Mutant Toxins and Maintained Important Antigenic Determinants**

Sample	EC ₅₀	Reduction in cytotoxicity relative to wt toxin (log ₁₀) ^a	Max binding (R _{max}) ^b Neut mAb ^d		
			1 ^c	2	3
Triple mutant A (SEQ ID NO: 4)	12.5 µg/mL	4.5	100	100	100
FI- Triple mutant A	>1000 µg/mL	>6.4	55	59	53
EDC/NHS- Triple mutant A	>1000 µg/mL	>6.4	90	94	103
Triple mutant B (SEQ ID NO: 6)	69 ng/mL	4.3	100	100	100
FI-Triple mutant B	>1000 µg/mL	8.4	67	67	36
EDC/NHS-Triple mutant B	>1000 µg/mL	8.4	87	78	73

a cytotoxicity was measured using the CPE assay on IMR90 cells

b values determined by Biacore™ analysis using multiple neutralizing mAbs directed at various non-overlapping toxin epitopes

10 c values are averages of two experiments

d For the first three rows, the neut mAb “1,” “2,” “3” refer to mAbs A60-22, A80-29, and A65-33 for toxin A, respectively. For the bottom three rows, the neut mAb “1,” “2,” “3” refer to mAbs B8-26, B59-3, and B-56-15 for toxin B, respectively.

Example 24: Preclinical Immunogenicity Studies

Key preclinical objectives include testing compositions including *C. difficile* mutant toxins A and B in small animals and nonhuman primates (NHP). Mice and hamsters were immunized to determine, among other things, if the *C. difficile* compositions are capable of eliciting neutralizing antibodies against the mutant toxin A and B. The antigens were tested for induction of serum neutralization antibody responses following a series of immunizations in mice, hamsters, and cynomolgus macaques. The genetic and/or chemically-inactivated mutant toxins were formulated in either neutral buffer, aluminum phosphate buffer, or buffer containing ISCOMATRIX as an adjuvant in some embodiments. Neutralizing antibody responses were generally tested about two to four weeks after each boost or the final dose.

The toxin neutralization assay demonstrates the ability of an antiserum to neutralize the cytotoxic effect mediated by *C. difficile* TcdA or TcdB and is therefore able to measure the functional activity of antibodies that are present in a sample. A toxin neutralization assay was performed on a human lung fibroblast cell line, IMR-90, which is sensitive to both TcdA and TcdB. Briefly, a 96-well microtiter plate was seeded with IMR-90 cells serving as the target of toxin-mediated cytotoxicity. Each test serum sample was analyzed separately for the ability to neutralize TcdA and TcdB. Appropriate serial dilutions of test antisera were mixed with a fixed concentrations of TcdA or TcdB and incubated at 37°C for 90 minutes in a humidified incubator (37°C /5% CO₂) to allow for neutralization of the toxins to occur. For quality control, all plates included a Reference standard and controls which includes antitoxin antibodies of known titer. After 90 minutes, the toxin-antisera mixture was added to the IMR-90 cell monolayer and the plates were incubated for an additional 72 hours. Subsequently, CellTiter-Glo® substrate was added to the assay plate to determine the Adenosine Triphosphate (ATP) levels present in metabolically active cells and was measured as Relative Luminescence Units (RLU). A large ATP level indicates high cell viability, and levels are directly proportional to the amount of neutralization of the toxin by the antibody present in the sample. For preclinical data, the RLU data was plotted against the dilution value of the test antisera sample to generate a Four-Parameter Logistic (4-PL) regression response fit curve. The neutralization titers were expressed as the sample dilution value which exhibited 50% reduction in cytotoxicity.

Example 25: Mouse immunogenicity study: muC. difficile2010-06

The purpose of this study was to assess the immunogenicity of two forms of mutant *C. difficile* toxin B (SEQ ID NO: 6), each chemically-inactivated by different methods. In this study, the untreated mutant toxin B (SEQ ID NO: 6) (genetically 5 inactivated but not chemically inactivated) was used as a control, with and without adjuvant.

Groups of 10 mice were immunized intramuscularly with 10 µg of an immunogen according to Table 12.

Table 12. Testing chemically inactivated mutant toxin B (SEQ ID NO: 6) in mice

Group	Immunogen	Dose	No.	Route	Schedule
1	Formalin-Inactivated Mutant toxin B ^a in AlPO ₄ ^c	10 µg	10	IM	Prime wk 0, Boost wks 4, 8
2	Inactivated Mutant toxin B form 2 ^b in AlPO ₄ ^c	10 µg	10	IM	Prime wk 0, Boost wks 4, 8
3	Genetic-Inactivated Mutant toxin B unadjuvanted	10 µg	10	IM	Prime wk 0, Boost wks 4, 8,
4	Genetic-Inactivated Mutant toxin B in AlPO ₄ ^c	10 µg	10	IM	Prime wk 0, Boost wks 4, 8,

^a chemical inactivation = Formalin/glycine treated 10°C for 7 days
^b chemical inactivation = EDC/NHS treated, 30°C for 2 hours
^c aluminum ion concentration = 0.5 mg/mL

10 Results: There were no adverse events in the mice following each administration of the vaccine candidates. As illustrated in Figure 10, mice in each group developed significant robust anti-toxin B neutralizing antibodies after the third dose with the respective mutant toxins.

15 Based on the week 12 titers, it appears that in mice the EDC-inactivated mutant toxin B (Group 2) and the formalin-inactivated mutant toxins (Group 1) generated potent neutralizing responses.

In the absence of chemical inactivation, the genetic mutant toxin B (SEQ ID NO: 6) generated neutralizing responses after two doses (Groups 3-4, week 8), which were boosted after the third dose (Groups 3-4, week 12).

5

Example 26: Mouse immunogenicity study: muC. difficile2010-07:

The purpose of this study was to assess immunogenicity of chemically inactivated *C. difficile* mutant toxins A and B (SEQ ID NOs: 4 and 6, respectively), either alone or in combination. The immunogens for all groups were formulated with aluminum phosphate

5 as an adjuvant.

Groups of 5 mice were immunized intramuscularly with 10 µg of an immunogen according to Table 13.

Table 13. Testing Chemically Inactivated Genetic A and B mutant toxins (SEQ ID NOs: 4 and 6, respectively) in Mice

Group	Immunogen	Dose	No.	Group	Schedule
1	Formalin-Inactivated ^a Mutant toxin B (SEQ ID NO: 6) in AlPO ₄ ^c	10 µg	5	IM	Prime wk 0, Boost wks 4, 8, 12
2	EDC-Inactivated ^b Mutant toxin B (SEQ ID NO: 6) in AlPO ₄ ^c	10 µg	5	IM	Prime wk 0, Boost wks 4, 8, 12
3	Formalin-Inactivated Mutant toxin A (SEQ ID NO: 4) form 1 in AlPO ₄ ^c	10 µg	5	IM	Prime wk 0, Boost wks 4, 8, 12
4	EDC-Inactivated Mutant toxin A (SEQ ID NO: 4) in AlPO ₄ ^c	10 µg	5	IM	Prime wk 0, Boost wks 4, 8, 12
5	Formalin-Inactivated Mutant toxins A+B in AlPO ₄ ^c	10 µg each	5	IM	Prime wk 0, Boost wks 4, 8, 12

^a Formalin-treatment = formalin/glycine treated for 2 days at 25°C; mutant toxin was not cytotoxic and retained binding to all mutant toxin-specific monoclonal antibodies tested

^b EDC-treatment = EDC/NHS treated for 4hrs at 30°C; mutant toxin was not cytotoxic and retained binding to all mutant toxin-specific monoclonal antibodies tested

^c aluminum ion concentration = 0.5 mg/mL

Results: There were no adverse events in the mice following each administration of the vaccine candidates. As illustrated in Figure 11, after two doses of chemically inactivated genetic mutant toxins, the anti-toxin A neutralizing antibodies (Groups 3-5) were boosted to titers between 3 and $4\log_{10}$ while the anti-toxin B neutralizing antibodies (Groups 1-2, 5) remained low to undetectable, which is consistent with the data from the mouse study described above (Figure 10). Anti-toxin B neutralizing antibodies boosted to $2-3\log_{10}$ in groups 1, 2, and 5 following the third dose (week 12 titers) and reached their peak two weeks following the fourth dose (week 14 titers). The anti-toxin A neutralizing antibody titers in groups 3-5 increased slightly following the 5 third (week 12 titers) and fourth immunizations (week 14 titers).

10

Example 27: Hamster immunogenicity study: hamC. difficile2010-02:

The purpose of this study was to assess immunogenicity and protective potential of *C. difficile* triple mutant and chemically inactivated mutant toxins A and B in the Syrian golden hamster model. The Syrian golden hamster model represents the best

5 available challenge model for simulating human CDAD. The same batches of mutant toxins A and B used in mouse study muC. difficile2010-07 were used in this study. As a control, one group was given mutant toxins without aluminum-containing adjuvant.

Groups of 5 Syrian golden hamsters were immunized intramuscularly with 10 µg of an immunogen according to Table 14.

Table 14. Testing Chemically Inactivated Mutant Toxins A and B (SEQ ID NOs: 4 and 6, respectively) in Hamsters (hamC. difficile2010-02)

Group	Immunogen	Dose	No.	Route	Schedule
1	Formalin-Inactivated ^a Mutant toxins A+B (SEQ ID NOs: 4 and 6) in AlPO ₄ ^c	10 µg each	5	IM	Prime wk 0, Boost wks 4, 8, 12
2	Formalin-Inactivated Mutant toxins A+B (SEQ ID NOs: 4 and 6) in PBS (no adjuvant)	10 µg each	5	IM	Prime wk 0, Boost wks 4, 8, 12
3	EDC-Inactivated ^b Mutant toxins A+B (SEQ ID NOs: 4 and 6) in AlPO ₄ ^c	10 µg each	5	IM	Prime wk 0, Boost wks 4, 8, 12
4	List Biological toxoid in AlPO ₄ ^c	10 µg each	5	IM	Prime wk 0, Boost wks 4, 8, 12

^a Formalin-treatment = formalin/glycine treated for 2 days at 25°C; Mutant toxin was not cytotoxic and retained binding to all mutant toxin-specific monoclonal antibodies tested

^b EDC-treatment = EDC/NHS treated for 4hrs at 30°C; Mutant toxin was not cytotoxic and retained binding to all mutant toxin-specific monoclonal antibodies tested

^c aluminum ion concentration = 0.5 mg/mL

1. Animals: 15 Syrian golden hamsters, female, 6-8 weeks old/ 100-130g each.
2. Vaccination: IM, 0.05 ml each, according to above schedule. Toxoids will be provided by Process Development and will be formulated in AlPO₄ diluent by the Formulations Group. Group 2 will serve as a non-adjuvanted control group.
3. Bleed: All hamsters will be bled at weeks 0, 4, 8, and 12, just prior to each immunization.
4. Serum sample analysis: Neutralization assay

Results: There were no adverse events observed following immunization with the mutant toxins. As illustrated in Figure 12, after a single dose of mutant toxins, the anti-toxin A neutralizing responses were between 2-3 \log_{10} for the formalin-inactivated mutant toxins (Groups 1-2) and between 3-4 \log_{10} for the EDC-inactivated mutant toxins (Group 3). After the second dose, anti-toxin A antibodies boosted in all three groups. Anti-toxin A antibodies in all three groups did not appear to increase after the third dose. A similar result was observed after the fourth immunization, where an increase in titer was observed in the formalin-inactivated group that did not contain the aluminum adjuvant (Group 2).

10 The anti-toxin B neutralizing responses were undetectable in the formalin-inactivated mutant toxins groups (Groups 1-2) and were just over 2 \log_{10} for the EDC-inactivated mutant toxins (Group 3) after a single dose. After the second dose, anti-toxin B neutralizing antibody titers in the two formalin-inactivated groups (Groups 1-2) increased to 3-4 \log_{10} while those in the EDC-inactivated group (Group 3) increased to 15 4-5 \log_{10} . For all three groups, increases in anti-toxin B neutralizing antibody titers were observed after the third and/or fourth doses, with all groups reaching a peak titer at week 16 (after the last dose). See Figure 12.

In Figure 13, the level of neutralizing antibody responses against chemically inactivated genetic mutant toxins (Figure 12) was compared to those elicited by List 20 Biological Laboratories, Inc. (Campbell, California) (also referred herein as "List Bio" or "List Biologicals") toxoids (i.e., toxoids purchased from List Biological Laboratories were prepared by formalin inactivation of wild type toxins; control reagent used to establish the hamster challenge model).

As used herein, "FI" in figures and tables refers to formalin/glycine treatment of 25 the toxins, 2 days at 25°C, unless otherwise stated. As used herein, "EI" in figures and tables refers to EDC/NHS treatment for 4 hours at 30°C, unless otherwise stated. In Figure 13, 5 hamster animals were treated with the respective mutant toxin composition, whereas 11 hamster animals were treated with the toxoid purchased from List Biological.

The data in Figure 13 shows that, in hamsters administered according to Table 14, the respective neutralizing antibody titers against toxin A (Figure 13A) and toxin B (Figure 13B) induced by the immunogenic composition including EDC inactivated mutant toxins after two doses is higher than the respective neutralizing antibody titers 5 elicited by the List Biologicals toxoids.

Example 28: Hamster immunogenicity study: *C. difficile* ham2010-02 (continued)

To assess protective efficacy of the mutant toxins, immunized hamsters, along with one control group of non-immunized animals, were first given an oral dose of clindamycin antibiotic (30 mg/kg) to disrupt normal intestinal flora. After five days, the 5 hamsters were challenged with an oral dose of wild type *C. difficile* spores (630 strain, 100 cfu per animal). Animals were monitored daily for eleven days post-challenge for signs of CDAD, which in hamsters is known as wet tail. Using a system of clinical scoring a number of different parameters, animals determined to have severe CDAD were euthanized. The parameters included activity following stimulation, dehydration, 10 excrement, temperature, and weight, etc., which are known in the art.

At day 11, the study was terminated and all surviving animals were euthanized. Figure 14 shows the survival curves for each of the three immunized groups (Groups 1-3, according to Table 14) as compared to the non-immunized controls. As can be seen, the non-immunized animals all developed severe CDAD and required euthanasia 15 between days 1-3 post challenge (0% survival). Both groups administered with formalin-inactivated mutant toxin had 60% survival curves, with animals not requiring euthanasia until day 3 (Group 1) or day 4 (Group 2). The group administered with EDC-inactivated mutant toxin had an 80% survival curve, with 1 (out of 5) animal requiring euthanasia on day 7. Accordingly, the hamsters were protected from lethal challenge with *C. difficile* 20 spores.

Example 29: Hamster immunogenicity study: hamC. difficile2010-03: Immunogenicity of genetic and chemically-inactivated C. difficile mutant toxins

The purpose of this study was to assess immunogenicity of non-adjuvanted C. difficile triple mutant and chemically inactivated mutant toxins A and B (SEQ ID NOs: 4 and 6, respectively) in the Syrian golden hamster model. The same batches of mutant toxins A and B (SEQ ID NOs: 4 and 6, respectively) used in mouse study muC. difficile2010-07 were used in this study. As a control, one group (Group 1) was given a phosphate-buffered saline as placebo.

Groups of five or ten Syrian golden hamsters were immunized with an immunogen according to Table 15. Animals were given three doses. In addition, animals were dosed every two weeks.

Table 15: Experimental Design of Hamster Immunization and Challenge					
Group	Immunogen	Dose	No.	Route	Schedule
1	Placebo (PBS buffer)	NA	5	NA	
2	Mutant toxin A+B (SEQ ID NOs: 4 and 6, respectively); Formalin-inactivated	10 µg each	10	IM	Prime wk 0, Boost wks 2, 4
3	Mutant toxin A+B (SEQ ID NOs: 4 and 6, respectively); EDC-Inactivated	10 µg each	10	IM	Prime wk 0, Boost wks 2, 4
4	Mutant toxin A+B (SEQ ID NOs: 4 and 6, respectively); genetic	10 µg each	10	IM	Prime wk 0, Boost wks 2, 4

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Results: See Figure 15. No anti-toxin A or B antibodies were observed in the placebo control group. After one dose, anti-toxin A neutralizing antibodies were observed between $2-3 \log_{10}$ for the formalin-inactivated (Group 2) and genetic mutant toxin (Group 4) groups and between $3-4 \log_{10}$ for the EDC-inactivated group (Group 3). Anti-5 toxin A neutralizing antibodies increased in each of these groups (2-4) after the second immunization with the relevant mutant toxins (compare titers at week 2 to week 3 in Figure 15). After the third dose of mutant toxins (given at week 4), anti-toxin A neutralizing antibody titers in Groups 2-4 increased compared to their week 4 titers.

Anti-toxin B neutralizing antibodies were detectable after the second dose, 10 wherein the formalin-inactivated (Group 2) and EDC-inactivated (Group 3) anti-toxin B neutralizing antibodies increased to between $3-4 \log_{10}$ and to between $2-3 \log_{10}$ for the genetic triple mutant (Group 4). Following the third immunization (week 4), the anti-toxin B neutralizing antibody titers boosted to between $3-4 \log_{10}$ for the formalin-inactivated 15 mutant toxins (Group 2) and genetic mutant toxins (Group 4) and between $4-5 \log_{10}$ for the EDC-inactivated mutant toxins (Group 3).

For both anti-toxin A and anti-toxin B neutralizing antibodies, peak titers were observed at week 6 (post-dose 3) for all vaccinated groups (Groups 2-4).

Assessment of Immunogenic Compositions Adjuvanted with AlhydrogelTM/CpG or ISCOMATRIX

20 Hamsters immunized with an immunogenic composition including a chemically inactivated mutant toxin formulated with AlhydrogelTM/ISCOMATRIX, or AlhydrogelTM/CpG24555 (Alh/CpG) developed robust neutralizing antitoxin antisera. It was observed that peak antitoxin A and antitoxin B responses were 2-3-fold higher and statistically significant in groups immunized with mutant toxins formulated in Alh/CpG or 25 ISCOMATRIX when compared to vaccine formulated with AlhydrogelTM alone. See Table 32 showing 50% neutralization titers. Hamsters (n=10/group) were immunized IM at 0, 2, and 4 weeks with 10 μ g each mutant toxin A drug substance and mutant toxin B drug substance formulated with 100 μ g of AlhydrogelTM or 200 μ g of CpG 24555 + 100 μ g of AlhydrogelTM, or 10 U of ISCOMATRIX. Sera were collected at each time point and 30 analyzed in the toxin neutralization assay for functional antitoxin activity. Geometric mean titers are provided in Table 32. Asterisks (*) indicate statistical significance ($p<0.05$) when compared to titers in the AlhydrogelTM group.

Table 32: Immunogenicity of Adjuvanted Mutant Toxin Drug Substances in Hamsters

Antitoxin A		50% Neutralization Titer					
		Week 0	Week 1	Week 2	Week 3	Week 4	Week 6
Alhydrogel TM Titer:		10	26	88	7425	6128	15965
Alh/CpG Titer:		10	103	*688	*34572	*23028	*62203
ISCOMATRIX Titer:		10	27	*246	*12375	8566	*36244
Antitoxin B							
		Week 0	Week 1	Week 2	Week 3	Week 4	Week 6
Alhydrogel TM Titer:		10	15	10	218	1964	7703
Alh/CpG Titer:		10	10	18	*5550	*5212	*59232
ISCOMATRIX Titer:		10	12	12	*7412	*15311	*92927

Protective efficacy of the immunogenic composition including mutant toxin drug substances formulated with these adjuvants was tested. Hamsters were immunized and 5 were given oral clindamycin (30 mg/kg) on week 5 and challenged according to the method described above. One group of unimmunized hamsters (n=5) was included as a control. Increased efficacy was observed in hamsters immunized with mutant toxin drug substances adjuvanted with either Alh/CpG or ISCOMATRIX (100% survival) as compared to AlhydrogelTM alone (70% survival). Accordingly, the hamsters were 10 protected from lethal challenge with *C. difficile* spores.

Example 30: *Clostridium difficile* Vaccination in Cynomolgus Macaques

The purpose of this study was to test the immunogenicity of low and high doses of EDC-Inactivated and Formalin-Inactivated *C. difficile* mutant toxins in cynomolgus macaques. All mutant toxins were formulated in ISCOMATRIX® as an adjuvant except 5 for one group, which served as the unadjuvanted control (Group 5).

Table 16: Immunization of Cynomolgus Macaques

Group	Immunogen	Number	Dose	Route	Schedule
1	FI-Mutant toxins A+B (ISCOMATRIX)	5	10 µg each	IM	Prime wk 0, Boost wks 2, 4
2	FI-Mutant toxins A+B (ISCOMATRIX)	5	100 µg each	IM	Prime wk 0, Boost wks 2, 4
3	EI-Mutant toxins A+B (ISCOMATRIX)	5	10 µg each	IM	Prime wk 0, Boost wks 2, 4
4	EI-Mutant toxins A+B (ISCOMATRIX)	5	100 µg each	IM	Prime wk 0, Boost wks 2, 4
5	EI-Mutant toxins A+B (no adjuvant)	5	100 µg each	IM	Prime wk 0, Boost wks 2, 4

Animals: 25 cynomolgus macaques

The asterisk, “*”, in figure 16 refers to having only 4 cynos in the group for week 12, one cyno in the group was terminally bled week at 8

Vaccination: IM, 0.5mL per dose, at weeks 0, 2, and 4. Mutant toxin compositions were prepared as described above. The mutant toxin compositions were formulated in ISCOMATRIX, except Group 5 was formulated in buffer without adjuvant.

Bleed: Weeks -2, 0, 2, 3, 4, 6, 8, and 12. Euthanasia and terminal bleeds on animals with highest *C. difficile* titers at week 8.

Serum sample analysis: Protein ELISA and Neutralization assays

Results: Figure 16 shows the anti-toxin neutralizing antibody responses in these animals at weeks 0, 2, 3, 4, 6, 8, and 12. Anti-toxin A titers were between 2-3 \log_{10} for all five groups after a single dose (week 2 titers). These titers boosted after each subsequent dose for each group. In these animals, there was no drop in titer between 5 weeks 3 and 4. For all groups, the peak titers were between 4-5 \log_{10} . At all time points, the group without ISCOMATRIX adjuvant (Group 5) had the lowest titers, indicating the utility of ISCOMATRIX at boosting the immune responses. The no-adjuvant control group (Group 5) reached peak titers at week 12, as did the group immunized with the high dose of EDC-inactivated mutant toxins (Group 4); all other groups reached peak 10 titers at week 6, two weeks after the last dose. The titers in all groups boosted after the second dose (week 3 time point). As with the anti-toxin A responses, the anti-toxin B responses did not decrease from week 3 to week 4. After the third dose (week 6 time point), the anti-toxin B neutralizing antibody titers in all groups were between 3-4 \log_{10} , except in the low dose formalin-inactivated group (Group 1) and the high dose EDC- 15 inactivated group (Group 4), both of which had titers just $>4 \log_{10}$. The peak titers were observed at week 12 for all groups except the low dose EDC-inactivated group (Group 3), which had peak titers at week 8. All groups had peak titers $>4 \log_{10}$.

Example 31: Monoclonal Antibodies Production

Although toxins A and B share a lot of structural homology, the neutralizing activities of the antibodies were found to be toxin-specific. In this invention, several antibodies were identified that are specific to individual toxin, and directed to various epitopes and functional domains, and have high affinity and potent neutralizing activity toward native toxins. Antibodies were isolated from mice that were immunized with either a commercially available formalin inactivated (FI)-mutant toxin or recombinant holo-mutant toxin (SEQ ID NOs: 4 and 6) rendered non-toxic by introducing specific mutations in its catalytic site for producing toxin A and B mAb, respectively. Epitope mapping of the antibodies showed that the vast majority of the mAb against toxin A (49 out of 52) were directed to the non-catalytic C terminal domain of the toxin.

Monoclonals against toxin B were targeted to three domains of the protein. Out of a total of 17 toxin B specific mAb, 6 were specific to N-terminus (e.g., amino acids 1-543 of a wild-type *C. difficile* TcdB, such as 630), 6 to C-terminus (e.g., amino acids 1834-2366 of a wild-type *C. difficile* TcdB, such as 630) and 5 to mid-translocation domain (e.g., amino acids 799-1833 of a wild-type *C. difficile* TcdB, such as 630). The approach of using mutant *C. difficile* toxins (e.g., SEQ ID NO: 4 and 6) as immunizing antigens thus offers a key advantage of presenting most, if not all, antigenic epitopes as compared to the formalin inactivation process that tend to adversely affect the antigenic structure of the mutant toxin.

Example 32: Characterization of toxin A mAb, A3-25, which includes a variable light chain having the amino acid sequence of SEQ ID NO: 36 and a variable heavy chain having the amino acid sequence of SEQ ID NO: 37.

The mAb A3-25 was of particular interest since this antibody defied all attempts 5 to define its immunoglobulin (Ig) isotyping using the commonly available isotyping kits for IgG, IgM and IgA. Further analysis by western blot using Ig H-chain specific antisera showed that the A3-25 is of IgE isotype, a rare event in mAb production. This was further confirmed by the nucleotide sequencing of mRNA isolated from A3-25 10 hybridoma cells. The amino acid sequences deduced from the nucleotide sequences of the variable regions of H- and L-chain of A3-25 are shown in Figure 17.

In order to further evaluate the A3-25 mAb in animal model for *C difficile* infection and disease, its Ig isotype was changed to murine IgG1 by molecular grafting of the variable region of ϵ H chain onto the murine γ heavy chain according to the published methods.

15

Example 33: Neutralizing Ability and Epitope Mapping of Toxin Specific Antibodies

Further, in an effort to identify functional/neutralizing antibodies, all monoclonals were evaluated for the ability to neutralize wild type toxins in a standard cytopathic effect (CPE) assay or in a more stringent and quantitative assay based on measurement of ATP as cell viability indicator.

Out of a total of 52 toxin A specific antibodies, four mAb (A3-25, A65-33, A60-22 and A80-29 (Table 17 and Figure 18) exhibited varied levels of neutralizing activity. BiaCore competitive binding assay and hemagglutination inhibition (HI) assays were performed to map the antibody epitopes. Results indicated that these antibodies may be targeted to different epitopes of the toxin A protein (Table 17). To further identify the location of binding sites on the protein, the antibodies were individually evaluated in western blot or dot blot assays using toxin fragments of known sequences. All 4 neutralizing mAb were found to be directed to the C-terminus region of the toxin.

From a total of 17 toxin B specific antibodies, 9 were found to be neutralizing. Of the nine neutralizing mAb, six of them were directed to the N-terminus and the other three to the translocation domain of the B toxin (Table 18). Based on the Biacore competitive binding assay, the nine neutralizing monoclonal antibodies may be grouped into four epitope groups as shown in Figure 19.

TABLE 17: Characteristics of Selected Toxin A mAb

Epitope Group (Biacore)	mAb #	Neutralizing activity	Hemagglutination Inhibition	Binding Specificity	Ig Isotype
1	A3-25	+	-	C-terminal	IgE, κ
2	A65-33	+	-	C-terminal	IgG2a, κ
3	A80-29	+	+	C-terminal	IgG1, κ
ND	A60-22	+	+	C-terminal	IgG1, κ
4	A64-6	-	-	In progress	IgG1, κ
	A50-10	-	-	C-terminal	IgG1, κ
	A56-33	-	-	In progress	IgG1, κ
ND	A1	-	-	N-terminal	IgG1, κ

TABLE 18: Characteristics of Selected Toxin B mAb

Epitope Group (Biacore)	mAb #	Neutralizing activity	Binding Specificity	Ig isotype
1	B2-31	+	N-terminal	IgG1, κ
	B5-40			IgG1, κ
	B8-26			IgG1, κ
	B70-2			IgG1, κ
2	B6-30	+	N-terminal	IgG1, κ
	B9-30			IgG1, κ
3	B59-3	+	Translocation domain	IgG1, κ
	B60-2			IgG1, κ
4	B56-6	+	Translocation domain	IgG1, κ
	B58-4	-		IgG1, κ
5	B12-34	-	C-terminal	IgG1, κ
	B14-23			IgG1, κ
	B80-3			IgG1, κ
6	B66-29	-	C-terminal	IgG1, κ
7	B84-3	-	C-terminal	IgG1, κ

Example 34: Identification of Novel Toxin A Antibodies Combinations with Significantly Enhanced Neutralizing Activity:

The four toxin A mAb (A3-25, A65-33, A60-22 and A80-29) showed incomplete or partial neutralization of toxin A when tested individually in the ATP based 5 neutralization assay. The mAb A3-25 was the most potent antibody and the other three were less neutralizing with A80-29 barely above background (Figure 18). However, when A3-25 was combined with either one of the other three mAbs, a synergistic effect in neutralization was observed in all three combinations which was far greater than the sum total of neutralization of individual antibodies as shown in Figure 20A-C. In 10 addition, all three combinations exhibited complete neutralization capability normally observed with anti-toxin A polyclonal antibodies.

Example 35: Identification of Novel Toxin B Antibodies Combinations Showing Significantly Enhanced Neutralizing Activity:

We also observed synergistic neutralization with the Toxin B mAbs from the 15 different epitope groups identified by BiaCore analysis. Toxin B mAb B8-26, the most dominant mAb of group 1, was combined with multiple mAbs from group 3. The combinations were evaluated in a toxin B specific neutralization assay and the results are shown in Figure 21 and Table 19.

Table 19: Neutralization of Toxin B with mAbs

mAb	Neut titer	
	CPE	ATP
B8-26 alone	20,480	5,000
B59-3 alone	320	120
B60-2 alone	320	80
B8-26 + B59-3	655,360	~60,000
B8-26 + B60-2	327,680	nd
nd, not done		

20

The synergistic neutralizing effect was observed when B8-26 was combined with an epitope group 3 mAb (Figure 21B), but not any other mAb (data not shown).

Example 36: *In vitro* screening by mAb for Safe and Efficacious Mutant Toxin**Compositions:**

Genetic mutant toxins A and B of *C. difficile* (e.g., SEQ ID NO: 4 and 6) generated via genetic engineering showed residual cytotoxicity using an *in vitro* cytotoxicity assay.

5 Although we have achieved a ~4 log reduction in cytotoxicity for each mutant toxin C. *difficile* toxin (Table 20), further chemical inactivation of the mutant toxins, such as with formalin treatment was preferred. However, chemical inactivation treatments may be harsh and may adversely affect key antigenic epitopes of these toxins or mutant toxins.

Table 20: A Comparison of *In Vitro* Cytotoxicity of WT Toxin, Triple Mutant Toxin, and Formalin-Inactivated (FI, from List Biological) WT toxins (List Biological, commercial)

Tcd	Source/treatment	EC ₅₀ ng/mL	Fold Reduction in Cytotoxicity
TcdA			
Toxin A (SEQ ID NO: 1)	WT	0.92	1
Mutant toxin A (SEQ ID NO: 4)	Triple mutant	8600	9348
Toxoid A (FI)	Formalin treated, commercial	>20,000	>21,739
TcdB			
Toxin B (SEQ ID NO: 2)	WT	0.009	1
Mutant toxin B (SEQ ID NO: 6)	Triple mutant	74	8222
Toxoid B (FI)	Formalin treated, commercial	4300	477,778

For bioprocess optimization, a statistical design of experiment (DOE) was performed for the chemical inactivation of triple mutant Tcd A and B (1 mg/mL) using formalin and EDC/NHS treatment. To optimize formalin inactivation of triple mutant TcdA, we varied concentrations of formalin/glycine (20-40 mM), pH (6.5-7.5), and temperature 5 (25-40°C). For triple mutant TcdB, we varied the formalin/glycine concentration from 2 to 80 mM and the temperature and pH were 25 °C and 7.0 respectively. The incubation time for all formalin treatments was 24 hours. For the formalin inactivation, “40/40” in Tables 21 and 23 represents the concentration of formalin and glycine used in the reaction. For EDC/NHS treatment, we varied the concentrations of EDC/NHS from 0.25 10 to 2.5 mg/mg of triple mutant TcdA and from 0.125 to 2.5 mg/mg of triple mutant TcdB and incubated for four hours at 25 °C. At the end of the reactions, all samples were desalted in 10 mM phosphate, pH 7.0. After purification, the treated Tcds were analyzed for residual cytotoxicity and mAb recognition of epitopes by dot-blot analysis. The goal 15 was to identify treatment conditions that reduce cytotoxicity to the desired level (EC₅₀ > 1000 µg/mL) without negatively impacting epitopes recognized by a panel of neutralizing mAbs (++++ or +++). The treatment conditions (marked with a check mark “✓” in Tables 21-24) yielded potentially safe and efficacious immunogenic compositions that retained reactivity to at least four neutralizing mAbs while exhibiting 6-8 log₁₀ reduction in 20 cytotoxicity, relative to the respective wild-type toxin cytotoxicity. Select results are illustrated in Tables 21 to 24. Additional data from varying treatment conditions on the triple mutant toxins and the data from *in vitro* cytotoxicity and toxin neutralization assays are shown in Table 33 and Table 34. See also, for example, Examples 20 and 21 above, which provide further details regarding preferred crosslinking treatment conditions of the mutant toxins.

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Table 21: Cytotoxicity and Neutralizing mAb Reactivity of Formalin-inactivated Triple Mutant TcdA (SEQ ID NO: 4)

Chemical inactivation reaction conditions on Triple Mutant TcdA	CPE µg /mL	Reactivity with mAb (dot blot, non-denaturing conditions)					
		N-terminal Mab#6	Translocation Domain Mab# 102	C-terminal (neut)			
				A80-29	A3-25	A60-22	A65-33
25°C, pH 6.5, 20/20mM	250	++++	++++	++++	++++	++++	++++
25°C, pH 6.5, 40/40mM ✓	>1000	++++	++++	++++	++++	++++	++++
25°C, pH 7.5, 40/40mM ✓	>1000	++++	++++	++++	++++	++++	++++
40°C, pH 6.5, 40/40mM	>1000	++	++	++++	++++	++++	++++
40°C, pH 7.5, 40/40mM	>1000	++	++	++++	++++	++++	+++
None, Triple mutant toxin A	18.5-25	++++	++++	++++	++++	++++	++++
FI Toxoid A (List Biological)	ND	—	—	++	++	+++	+

Table 22: Cytotoxicity and Neutralizing mAb Reactivity of EDC-inactivated Triple Mutant TcdA (SEQ ID NO: 4)

Chemical inactivation reaction conditions on Triple Mutant TcdA	CPE µg /mL	Reactivity with mAb (dot blot, non-denaturing conditions)					
		N-terminal Mab#6	Translocation Domain Mab# 102	C-terminal (neut)			
				A80-29	A3-25	A60-22	A65-33
25°C, 0.25mg/mg, 4hr ✓	>1000	++++	++++	++++	++++	++++	++++
25°C, 0.5mg/mg, 4hr ✓	>1000	++++	++++	++++	++++	++++	++++
25°C, 1.25mg/mg, 4hr ✓	>1000	++	++++	++++	++	++++	++++
25°C, 2.5mg/mg, 4hr ✓	>1000	++	++++	++++	++	++++	+++
None, Triple mutant TcdA	18.5-25	++++	++++	++++	++++	++++	++++
FI Toxoid A (List Biological)	ND	—	—	++	++	+++	+

Table 23: Cytotoxicity and Neutralizing mAb Reactivity of Formalin-inactivated Triple Mutant TcdB (SEQ ID NO: 6)

Chemical inactivation reaction conditions on Triple Mutant TcdB	CPE (µg/mL)	mAb # (N-terminal aa 1 – 543)		mAb # (mid-/C-terminal aa 544 – 2366)	
		B8-26	B9-30	B56-6	B59-3
25°C, pH 7.0, 80/80mM, 24hr ✓	>1000	++++	++++	++++	+++
25°C, pH 7.0, 40/40mM, 24hr ✓	>1000	++++	++++	++++	++++
25°C, pH 7.0, 10/10mM, 24hr	15.6	++++	++++	++++	++++
25°C, pH 7.0, 2/2mM, 24hr	<0.98	++++	++++	++++	++++
None, Triple mutant TcdB	0.058	++++	++++	++++	++++
FI Toxoid B (List Biological)	ND	+++	+++	+++	++

Table 24: Cytotoxicity and Neutralizing mAb Reactivity of EDC-inactivated Triple Mutant TcdB (SEQ ID NO: 6)

Chemical inactivation reaction conditions on Triple Mutant TcdB	CPE (µg/mL)	mAb # (N-terminal aa 1 – 543)		mAb # (mid-/C-terminal aa 544 – 2366)	
		8-26	9-30	56-6	59-3
25°C, 0.125mg/mg, 4hr	3.9	++++	++++	++++	++++
25°C, 0.25mg/mg, 4hr	250	++++	++++	++++	++++
25°C, 0.5mg/mL, 4hr ✓	>1000	++++	++++	++++	++++
25°C, 1.25mg/mg, 4hr ✓	>1000	++++	+++	+++	+++
25°C, 2.5mg/mg, 4hr ✓	>1000	++++	+++	+++	+++
None, Triple mutant TcdB	0.058	++++	++++	++++	++++
FI Toxoid B (List Biological)	ND	+++	+++	+++	++

Table 33

		Cyto Assay (EC50)		Reactivity with mAb (dot blot, non-denaturing conditions)					
Sample #	Mutant toxin A (SEQ ID NO: 4) Sample ID	CPE; 24h μ g /mL	CPE, 72h μ g /mL	N-terminal Mab#6	Translocation Domain Mab# 102	C-terminal (neut)			
						80-29	3-25	60-22	65-33
1	L44166-157A	>1000	>1000	++++	++++	++++	+++	++++	++++
2	L44166-157B	>1000	>1000	+++	++++	++++	+++	++++	++++
3	L44166-157C	>1000	>1000	+++	+++	+++	+++	+++	+++
4	L44166-157D	>1000	>1000	+++	+++	++++	+++	++++	++++
5	L44905-160A	>1000	>1000	++	++	++++	++	++++	++++
6	L44166-166	>1000	>1000	++++	++++	++++	++++	++++	++++
7	L44905-170A	ND	>1000	+	+	++	++	++	+
8	L44897-61	>1000	ND	+++	++	++++	++++	++++	++++
9	L44897-63	>1000	ND	++++	++	++++	+++	++++	++++
10	L44897-72 Tube#1	250	ND	++++	++++	++++	++++	++++	++++
11	L44897-72 Tube#2	>1000	ND	++++	++++	++++	++++	++++	++++
12	L44897-72 Tube#3	>1000	ND	+++	+++	++++	++++	++++	++++
13	L44897-72 Tube#4	>1000	ND	+++	++++	++++	++++	++++	++++
14	L44897-72 Tube#5	>1000	ND	+++	++++	++++	++++	++++	++++
15	L44897-75 Tube#6	>1000	ND	+++	++++	++++	++++	++++	++++
16	L44897-75 Tube#7	>1000	ND	++++	++++	++++	++++	++++	++++
17	L44897-75 Tube#8	>1000	ND	++++	++++	++++	++++	++++	++++
18	L44897-75 Tube#9	>1000	ND	++	++	++++	++++	++++	++++
19	L44897-75 Tube#10	>1000	ND	++++	++++	++++	++++	++++	++++
20	L44897-75 Tube#11	>1000	ND	++	++	++++	++++	++++	+++
21	L44897-101 (pre-modification) TxA control	23.4	<7.8	++++	++++	++++	++++	++++	++++
22	L44897-101, 2 hr	187.5	155.9	+++	++++	++++	++++	++++	++++
23	L44897-101, 4 hr	375	380.3	+++	++++	++++	++++	++++	++++
24	L44897-101, 6 hr	500	429.6	+++	++++	++++	++++	++++	++++

25	L44897 102, 24 hr	>1000	>1000	++	++++	++++	++++	++++	++++	++++
26	L44897-103, 51 hr	>1000	>1000	+	+++	+++	++++	++++	++++	+++
27	L44897-104, 74 hr	>1000	>1000	—	+++	+++	+++	+++	+++	+++
28	L44897-105, 120 hr	>1000	>1000	—	++	++	+++	+++	++	++
29	L44980-004	>1000	>1000	++++	++++	++++	++++	++++	++++	++++
30	Reaction #1 Week 0, 25C	750 ug/mL	ND	ND	++	++	+++	+++	+++	++
31	Reaction #1 Week 1, 25C	375 ug/mL	ND	ND	+++	+++	+++	+++	+++	+++
32	Reaction #1 Week 2, 25C	375 ug/mL	ND	ND	+++	+++	+++	+++	+++	+++
33	Reaction #1 Week 3, 25C	375 ug/mL	ND	ND	+++	+++	+++	+++	+++	+++
34	Reaction #1 Week 4, 25C	250 ug/mL	ND	ND	+++	+++	+++	+++	+++	+++
35	Reaction #1 Week 3, 37C	93.8 ug/mL	ND	ND	++++	++++	++++	++++	++++	++++
36	Reaction #2 Week 0, 25C	375 ug/mL	ND	ND	+++	+++	+++	+++	+++	+++
37	Reaction #2 Week 1, 25C	375 ug/mL	ND	ND	++++	++++	++++	++++	++++	++++
38	Reaction #2 Week 2, 25C	750 ug/mL	ND	ND	++	++	++	+++	+++	++
39	Reaction #2 Week 3, 25C	250 ug/mL	ND	ND	+++	+++	+++	++++	++++	+++
40	Reaction #2 Week 4, 25C	250 ug/mL	ND	ND	+++	+++	+++	+++	++++	+++
41	Reaction #2 Week 3, 37C	187.5 ug/mL	ND	ND	+++	+++	++++	++++	++++	+++
42	TxA Control Week 3, 25C	18.8 ug/mL	ND	ND	++++	++++	++++	++++	++++	++++
43	TxA Control Week 3, 37C	25 ug/mL	ND	ND	++++	++++	++++	++++	++++	++++
44	L44897-116-6 29.5 hrs	> 2000 ug/mL	ND	ND	++	++	++	+++	+++	++
45	L44897-116-7 57.5 hrs	> 2000 ug/mL	ND	ND	++	++	++	+++	+++	++
46	L44897-116 -8 79.5 hrs	> 2000 ug/mL	ND	ND	+	+	+	+++	+++	+
47	L44897-116-9 123.5 hrs	> 2000 ug/mL	ND	ND	++	++	++	+++	+++	++
48	L44897-139	>1000	ND	++	++++	++++	++++	++++	++++	++++
49	L44166-204	>1000	ND	++++	++++	++++	++++	++++	++++	++++

Chemical crosslinking reaction conditions for the samples of triple mutant toxin A (SEQ ID NO: 4) referenced in Table 33

Samples 1-4 were modified with EDC/NHS. Conditions: 30°C, 20 mM MES/150 mM NaCl pH 6.5. Reactions were initiated by addition of EDC. After 2 hours reaction, 5 samples A, B, and C had 1 M glycine added to 50 mM glycine final concentration. Sample D had no glycine added. The reactions were set up with different weight ratios of Mutant toxin A (SEQ ID NO: 4):EDC:NHS as indicated below.

- 1 L44166-157A 1: 0.25: 0.25 w: w: w
- 2 L44166-157B 1: 1.25: 1.25
- 10 3 L44166-157C 1: 2.5: 2.5
- 4 L44166-157D 1: 2.5: 2.5

Sample 5 L44905-160A 80 mM HCHO, 80 mM glycine, 80mM NaPO₄ pH 7, 1mg/mL Mutant toxin A (SEQ ID NO: 4) Protein, 48 hrs reaction at 25 °C.

Sample 6 L44166-166 EDC/NHS modification of Mutant toxin A (SEQ ID NO: 4) 15 at 25 °C in 20 mM MES/150mM NaCl pH 6.5. Mutant toxin A (SEQ ID NO: 4):EDC:NHS = 1:0.5:0.5. Reaction initiated by addition of EDC. After 2 hours reaction, 1M glycine added to 0.1 M glycine final concentration and further 2 hour incubation. After this time, reaction buffer exchanged into 1 X PBS on SephadexTM G25.

Sample 7 L44905-170A 80 mM HCHO, 80 mM glycine, 80mM NaPO₄ pH 7, 20 1mg/mL Mutant toxin A (SEQ ID NO: 4) Protein, 48 hrs reaction at 35 C. This formalin reaction was directed at producing excessive crosslinking so that antigen binding would be severely diminished.

Sample 8 L44897-61 32 mM HCHO/80 mM glycine, 72 hrs reaction at 25 °C.

Sample 9 L44897-63 80 mM HCHO/80 mM glycine, 72 hrs reaction at 25 °C.

25

The following reactions all had 24 hrs reaction time.

Sample 10 L44897-72 Tube#1 25°C, 80 mM NaPi pH 6.5, 20 mM HCHO/20 mM glycine

Sample 11 L44897-72 Tube#2 25 °C, 80 mM NaPi pH 6.5, 40 mM HCHO/40 mM glycine

5 Sample 12 L44897-72 Tube#3 32.5 °C, 80 mM NaPi pH 7.0, 30 mM HCHO/30 mM glycine

Sample 13 L44897-72 Tube#4 32.5 °C, 80 mM NaPi pH 7.0, 30 mM HCHO/30 mM glycine

10 Sample 14 L44897-72 Tube#5 32.5 °C, 80 mM NaPi pH 7.0, 30 mM HCHO/30 mM glycine

Sample 15 L44897-75 Tube#6 25 °C, 80 mM NaPi pH 7.5, 20 mM HCHO/20 mM glycine

15 Sample 16 L44897-75 Tube#7 25 °C, 80 mM NaPi pH 7.5, 40 mM HCHO/40 mM glycine

Sample 17 L44897-75 Tube#8 40 °C, 80 mM NaPi pH 6.5, 20 mM HCHO/20 mM glycine

20 Sample 18 L44897-75 Tube#9 40 °C, 80 mM NaPi pH 6.5, 40 mM HCHO/40 mM glycine

Sample 19 L44897-75 Tube#10 40 °C, 80 mM NaPi pH 7.5, 20 mM HCHO/20 mM glycine

Sample 20 L44897-75 Tube#11 40 °C, 80 mM NaPi pH 7.5, 40 mM HCHO/40 mM glycine

The following 8 samples were reacted at 25 °C for the indicated times in 80 mM NaPi pH 7.0 containing 78 mM HCHO and 76 mM glycine

Sample 21 L44897-101 (pre-modification) TxA control time zero control sample, not modified or exposed to HCHO/glycine

5 Sample 22 L44897-101, 2 hr
 Sample 23 L44897-101, 4 hr
 Sample 24 L44897-101, 6 hr
 Sample 25 L44897 102, 24 hr
 Sample 26 L44897-103, 51 hr
10 Sample 27 L44897-104, 74 hr
 Sample 28 L44897-105, 120 hr

Sample 29 (L44980-004) was EDC/NHS modified Mutant toxin A (SEQ ID NO: 4) (triple mutant toxin A (SEQ ID NO: 4)-EDC). Reaction conditions are: 25 °C, buffer 15 was 20 mM MES/150 mM NaCl pH 6.6. Triple mutant toxin A (SEQ ID NO: 4):EDC:NHS = 1:0.5:0.5 w:w:w. Reaction initiated by addition of EDC. After 2 hours reaction, glycine added to 0.1 M final concentration and reacted further 2 hours at 25 C. Reaction terminated by desalting on Sephadex G25.

The following 12 samples and 2 controls were reversion experiments where samples were incubated at 25°C and 37 °C.

Reaction 1 = 25 °C, 80 mM NaPi pH 7.0, 40 mM HCHO only (no glycine), 24 hour reaction.

5 Reaction 2 = 25 °C, 80 mM NaPi pH 7.0, 40 mM HCHO/40 mM glycine, 24 hour reaction

	Sample	Reaction
10	30	Reaction #1 Week 0, 25°C
	31	Reaction #1 Week 1, 25°C
	32	Reaction #1 Week 2, 25°C
	33	Reaction #1 Week 3, 25°C
	34	Reaction #1 Week 4, 25°C
	35	Reaction #1 Week 3, 37°C
15	36	Reaction #2 Week 0, 25°C
	37	Reaction #2 Week 1, 25°C
	38	Reaction #2 Week 2, 25°C
	39	Reaction #2 Week 3, 25°C
	40	Reaction #2 Week 4, 25°C
	41	Reaction #2 Week 3, 37°C
20	42	TxA Control Week 3, 25°C
	43	TxA Control Week 3, 37°C

The next 4 samples were generated by reaction for the indicated times at 25 °C in 80 mM NaPi pH 7.0, 40 mM HCHO/40 mM glycine

25	44	L44897-116-6 29.5 hrs
	45	L44897-116-7 57.5 hrs
	46	L44897-116 -8 79.5 hrs
	47	L44897-116-9 123.5 hrs

30 Sample 48 L44897-139 48 hrs reaction at 25 °C, 80 mM NaPi pH 7.0, 40 mM HCHO/ 40 mM glycine.

Sample 49 L44166-204 EDC/NHS modification of Mutant toxin A (SEQ ID NO: 4). 25 C, buffer 1 x PBS pH7.0. Mutant toxin A (SEQ ID NO: 4):EDC:NHS = 1:0.5:0.5 w:w:w. 2 hours reaction with EDC/NHS, then 1 M glycine added to 0.1 M final concentration and further 2 hours reaction. Buffer exchanged on Sephadex G25 into 20 mM L-histidine/100 mM NaCl pH 6.5.

Table 34

	Cyto Assay (EC50)		Reactivity with neut mAb (dot blot, non-denaturing conditions)				Strong reactivities to all 4 mAbs
Mutant toxin B Sample ID	CPE; 24h	ATP, 72h	mAb # (N-terminal aa 1 – 543) 8-26 9-30	mAb # (mid-/C-terminal aa 544 – 2366) 56-6 59-3			
L44905-86-01 Triple mutant toxin B (SEQ ID NO: 6), Untreated Control	<0.1 µg /mL	<0.1µg/mL	++++	++++	++++	++++	✓
L44905-86-02 Triple mutant toxin B (SEQ ID NO: 6), Rxn1, 10°C, day1	≥100 µg/mL	2.2µg/mL	++++	++++	++++	++++	✓
L44905-86-03 Triple mutant toxin B (SEQ ID NO: 6), Rxn1, 25°C, day1	>100 µg/mL	>100 µg/mL	+++	++++	++	+++	✓*
L44905-86-04 Triple mutant toxin B (SEQ ID NO: 6), Rxn2, 10°C, day1	>100 µg/mL	5.2 µg/mL	++++	++++	++++	++++	✓
L44905-86-05 Triple mutant toxin B (SEQ ID NO: 6), Rxn2, 25°C, day1	>100 µg/mL	>100 µg/mL	++++	++++	++	+++	✓*

L44905-86-06 Triple mutant toxin B (SEQ ID NO: 6), Rxn3, 10°C, day1	>100 µg/mL	>100 µg/mL	++++	-	++++	+++	
L44905-86-07 Triple mutant toxin B (SEQ ID NO: 6), Rxn3, 25°C, day1	>100 µg/mL	>100 µg/mL	++++	-	++++	+++	
L44905-86-08 Triple mutant toxin B (SEQ ID NO: 6), Rxn1, 10°C, day5	>100 µg/mL	>100 µg/mL	++++	++++	++++	+++	✓
L44905-86-09 Triple mutant toxin B (SEQ ID NO: 6), Rxn1, 25°C, day5	>100 µg/mL	>100 µg/mL	++	++	-	+	
L44905-86-10 Triple mutant toxin B (SEQ ID NO: 6), Rxn2, 10°C, day5	>100 µg/mL	>100 µg/mL	++++	++++	++++	++++	✓
L44905-86-11 Triple mutant toxin B (SEQ ID NO: 6), Rxn2, 25°C, day5	>100 µg/mL	>100 µg/mL	++	++++	-	+	
L44905-86-12 Triple mutant toxin B (SEQ ID NO: 6), Rxn3, 10°C, day5	>100 µg/mL	>100 µg/mL	++++	-	++++	+++	
L44905-86-13 Triple mutant toxin B (SEQ ID NO: 6), Rxn3, 25°C, day5	>100 µg/mL	>100 µg/mL	++++	-	++++	++++	

L44905-86-14 Triple mutant toxin B (SEQ ID NO: 6), Rxn1, 10°C, day7	>100 µg/mL	>100 µg/mL	++++	++++	++++	++++	√
L44905-86-15 Triple mutant toxin B (SEQ ID NO: 6), Rxn1, 25°C, day7	>100 µg/mL	>100 µg/mL	+++	++++	-	+	
L44905-86-16 Triple mutant toxin B (SEQ ID NO: 6), Rxn2, 10°C, day7	>100 µg/mL	>100 µg/mL	+++	++++	+++	++++	√
L44905-86-17 Triple mutant toxin B (SEQ ID NO: 6), Rxn2, 25°C, day7	>100 µg/mL	>100 µg/mL	++	++	-	+	
L44905-86-18 Triple mutant toxin B (SEQ ID NO: 6), Rxn3, 10°C, day7	>100 µg/mL	>100 µg/mL	++++	-	++++	++++	
L44905-86-19 Triple mutant toxin B (SEQ ID NO: 6), Rxn3, 25°C, day7	>100 µg/mL	>100 µg/mL	+++	-	++	++	
L34346-30A	>100 µg/mL	>100 µg/mL	++++	++++	++++	++++	√
L34346-30B	>100 µg/mL	>100 µg/mL	+++	++++	++++	++++	√
Commercial, FI Toxoid B (List Biologicals)	ND	ND	++++	++++	++++	++++	√

Commercial, Control Toxin B wt (List Biologicals)	22.5 pg/ mL	7.8 pg/mL	+++	++	+++	+++	✓
Control, recombinant triple mutant toxin B (SEQ ID NO: 6)	78 ng/mL	72 ng/ml	+++	++	++++	+++	✓

Chemical crosslinking reaction conditions for the samples of mutant toxin B referenced in Table 34

Triple mutant toxin B (SEQ ID NO: 6) was chemically crosslinked and tested according

5 to the following reaction conditions. The L44905-86 samples were tested in an experiment involving three formalin reaction variations and two incubation temperatures. Each day, 6 samples were taken for a total of 18 samples. The first sample in the list is the untreated control (which makes 19 samples total). The untreated control included an untreated triple mutant toxin B polypeptide (SEQ ID NO: 6).

Reaction1 ("Rxn1") = 80 mM HCHO, 80 mM glycine, 80mM NaPO4 pH 7, 1mg/mL Triple mutant toxin B (SEQ ID NO: 6) Protein

Reaction2 ("Rxn2") = 80 mM HCHO, No glycine, 80mM NaPO4 pH 7, 1mg/mL Triple mutant toxin B (SEQ ID NO: 6) Protein

15 Reaction3 ("Rxn3") = 80 mM HCHO, No glycine, 80mM NaPO4 pH 7, 1mg/mL Triple mutant toxin B (SEQ ID NO: 6) Protein + Cyanoborohydride capping. Cyanoborohydride Capping involved 80mM CNBrH₄ added to desalted final reaction and incubated 24 hr at 36°C.

20 For Sample L34346-30A 0.5g EDC and NHS per gram of triple mutant toxin B (SEQ ID NO: 6), 4 hours at 30°C,in 20 mM MES, 150mM NaCl, pH 6.5.

For Sample L34346-30B 0.5g EDC and NHS per gram of triple mutant toxin B (SEQ ID NO: 6), 2 hours at 30°C followed by addition of glycine (final concentration of g/L) and incubated another 2 hours at 30°C, in 20 mM MES, 150mM NaCl, pH 6.5. The 25 only difference between the two reactions for L34346-30A and L34346-30B is the addition of glycine to reaction L34346-30B.

Example 37: Antibodies Induced by Immunogenic Compositions are Capable of Neutralizing Toxins from Various *C. difficile* Strains

To assess whether antibodies induced by the immunogenic compositions including the mutant toxin drug substances can neutralize a broad spectrum of diverse 5 toxin sequences, strains representing diverse ribotypes and toxinotypes were sequenced to identify the extent of genetic diversity among the various strains compared to the mutant toxin drug substances. Culture supernatants containing secreted toxins from the various strains were then tested in an *in vitro* neutralization assay using sera from immunized hamsters to determine the coverage of the 10 immunogenic composition and to determine the ability of the immunogenic composition to protect against diverse toxins from circulating clinical strains.

Both HT-29 cells (colon carcinoma cell line) and IMR-90 cells were used to test the neutralization of toxins expressed from CDC strains. HT-29 cells are more sensitive to TcdA; the EC₅₀ of the purified TcdA in these cells is 100 pg/mL as compared to 3.3 15 ng/mL for TcdB. On the other hand IMR-90 cells are more sensitive to TcdB; the EC₅₀ of the purified TcdB in these cells ranges between 9-30 pg/mL as compared to 0.92-1.5 ng/mL for TcdA. The assay specificity for both TcdA and TcdB in these cell lines was confirmed by using both polyclonal and monoclonal toxin-specific antibodies. For assay normalization, culture filtrates of the 24 CDC isolates were tested at a 20 concentration four times their respective EC₅₀ value. Three of the strains had toxin levels that were too low for testing in the neutralization assay.

Twenty-four strains representing diverse ribotypes/ toxinotypes covering greater than 95% of the circulating strains of *C. difficile* in the USA and Canada were obtained from the CDC. Among these isolates were strains representing ribotypes 027, 001 and 25 078, three epidemic strains of CDAD in the United States, Canada and UK. Strains 2004013 and 2004118 represented ribotype 027; strain 2004111 represented ribotype 001 and strains 2005088, 2005325 and 2007816 represented ribotype 078. To identify the extent of genetic diversity between the disease-causing clinical isolates and the 630 strain, the toxin genes (*tcdA* and *tcdB*) from these clinical strains were fully sequenced. 30 See Table 35. The amino acid sequences of the toxins were aligned using ClustalW in the Megalign™ program (DNASTAR® Lasergene®) and analyzed for sequence identity. For *tcdA*, genomic alignment analysis showed that all of the clinical isolates and strain 630 shared overall about 98-100% amino acid sequence identity. The C-terminal

portion of the *tcdA* gene was slightly more divergent. The same analysis was performed for the *tcdB* gene which exhibited greater sequence divergence. Notably strains 2007838/NAP7/126 and 2007858/NAP1/unk5 displayed the most divergent patterns from the 630 strain in the N terminal (79-100%) and the C terminal domains

5 (88-100%; data not shown).

A hamster serum pool (HS) was collected from the Syrian golden hamsters that were immunized with an immunogen including mutant TcdA (SEQ ID NO: 4) and mutant TcdB (SEQ ID NO: 6), wherein the mutant toxins were inactivated with EDC, according to, for example, Example 29, Table 15, described above, and formulated with aluminum phosphate. The results in Table 35 show that at least toxin B from the respective culture supernatants were neutralized, in an *in vitro* neutralization assay, by sera from the immunized hamsters.

10

Table 35. Description of *C. difficile* strains from CDC and Ability of Immune Hamster Sera to Neutralize Various Toxins

Strain	PFGE Type	Ribotype	Neutralized by Hamster Sera
2005088	NAP7	78	yes
2007816	NAP7-related	78	yes
2005325	NAP7	78	yes
2004013	NAP1	27	yes
2007886	NAP1		yes
2008222	NAP4	77	yes
2004206	NAP4	154	yes
2005283	NAP5	Unk3	Not tested ^b
2009141	NAP2		yes
2007838	NAP7	126	yes
2004111	NAP2	1	yes
2007070	NAP10	70	yes
2006017	NAP12	15	yes
2009078	NAP11	106	Not tested ^b
2007217	NAP8	126	yes
2006376	NAP9	17	yes
2007302	NAP11	Unk2	yes
2004118	NAP1	27	yes
2005022	NAP3	53	yes
2009292	NAP1		yes
2004205	NAP6	2	yes
2007858	NAP1	Unk5	yes
2009087	NAP11	106	Not tested ^b
2005359	NAP1-related		yes

b Toxin levels were too low to perform the neutralization assay.

Figure 23 depicts the results of the neutralization assay using toxin preparations from various *C. difficile* strains on IMR-90 cells. The data show TcdB neutralizing antibodies in the hamster antisera were capable of neutralizing toxins from all 21 isolates tested, including hypervirulent strains and a TcdA-negative, TcdB-positive 5 strain. At least 16 different strains of *C. difficile* were obtained from the CDC (Atlanta, GA)(previously described) and were cultured in *C. difficile* culture media under suitable conditions as known in the art and as described above. Culture supernatants containing the secreted toxins were analyzed to determine their cytotoxicity (EC₅₀) on IMR-90 monolayers and subsequently tested in a standard *in vitro* neutralization assay 10 at 4 times the EC₅₀ using various dilutions of sera from hamsters immunized with mutant toxin A drug substance and mutant toxin B drug substance, formulated with aluminium phosphate. Crude toxin obtained from culture supernatants of each strain and purified toxin (commercial toxin obtained from List Biologicals)(not purified from respective supernatants) were tested for cytotoxicity to IMR-90 cells using the *in vitro* 15 cytotoxicity assay described above.

In Figures 23A-K, the graphs show results from *in vitro* cytotoxicity tests (previously described) in which the ATP levels (RLUs) are plotted against increasing concentrations of: *C. difficile* culture media and the hamster serum pool (■); crude toxin and the hamster serum pool (●); purified toxin and the hamster serum pool (▲); crude 20 toxin (▼), control; and purified toxin (◆), control. The toxins from the respective strains were added to the cells at 4xEC₅₀ values.

As shown in Figures 23A-K, the hamsters that received the described immunogen surprisingly developed neutralizing antibodies that exhibited neutralizing activity against toxins from at least the following 16 different CDC strains of *C. difficile*, 25 in comparison to the respective toxin only control: 2007886 (Figure 23A); 2006017 (Figure 23B); 2007070 (Figure 23C); 2007302 (Figure 23D); 2007838 (Figure 23E); 2007886 (Figure 23F); 2009292 (Figure 23G); 2004013 (Figure 23H); 2009141 (Figure 23I); 2005022 (Figure 23J); 2006376 (Figure 23K). See also Table 35 for additional *C. difficile* strains from which toxins were tested and were neutralized by the immunogenic 30 composition including a mutant toxin A drug substance and mutant toxin B drug substance, formulated in aluminum phosphate.

In another study, culture supernatants containing secreted toxins from the various *C. difficile* strains (obtained from the CDC and from Leeds Hospital, UK) were tested in the *in vitro* neutralization assay using sera from hamsters that were administered with mutant toxin A drug substance and mutant toxin B drug substance, 5 formulated with AlhydrogelTM. See Table 36 for the experimental design. The results are shown in Table 37 and Table 38.

Table 36: Experimental design

Assay Control	In assay using HT-29 cells: Rabbit anti-serum (Anti-Toxin A polyclonal Fitzgerald Industries, #70-CR65) and Reference Toxin A (wild-type toxin A from List Biologicals) In assay using IMR-90 cells: Rabbit anti-serum (Anti-Toxin B polyclonal Meridian Life Science, #B01246R) and Reference Toxin B (wild-type toxin B from List Biologicals)
Sample Controls	In assay using HT-29 cells: HS serum + Reference Toxin A
	In assay using IMR-90 cells: HS serum + Reference Toxin B
	HS serum+630 wt toxin
	HS serum+ Culture media of IMR-90 or HT-29 cell line
	HS serum+ culture supernatant of VPI11186
Test Sample	HS + respective <i>C. difficile</i> culture supernatant
Source of Hamster antiserum (HS)	Animals administered with mutant toxin A drug substance and mutant toxin B drug substance formulated with Alhydrogel TM

Table 37: Immunogenic Composition-induced Antibodies Neutralized Toxin A and Toxin B from Various Wild-type *C. difficile* Strains from the CDC, including Hypervirulent strains

Cdiff Strain	PFGE Type	Ribotype	Toxinotype	Other Typing Method	Neutralized by HS (IMR-90, Toxin B)	Neutralized by HS (HT-29, Toxin A)
2004111	NAP2	1	0	Respective toxin sequence has 100% Homology to toxin from Strain 630	Yes	Yes
2009141	NAP2		0		Yes	Yes
2006017	NAP12	15	0		Yes	Yes
2007302	NAP11	Unk2	0		Yes	Yes
2009087	NAP11	106	0		Yes	Yes
2005022	NAP3	53	0		Yes	Yes
2005283	NAP5	Unk3	0		Yes	Yes
2009078	NAP5	53	0		Yes	Yes
2004206	NAP4	154	0		Yes	Yes
2008222	NAP4	77	0		Yes	Yes
2004205	NAP6	2	0		Yes	Yes
2007070	NAP10	70	0		Yes	Yes
2006376	NAP9	17	VIII	txnA-/txnB+	Yes	N/A
2007816	NAP7-related	78	V	Increasing prevalence in US and Europe	Yes	Yes
2007838	NAP7	126			Yes	Yes
2005088	NAP7	78			Yes	Yes
2005325	NAP7	78			Yes	Yes
2007217	NAP8	126			Yes	Yes
2004013	NAP1	27	III	Hypervirulent NAP1/027/III	Yes	Yes
2004118	NAP1	27			Yes	Yes
2009292	NAP1				Yes	Yes
2005359	NAP1-related				Yes	Yes
2007858	NAP1	Unk5	IX/XXIII	Other	Yes	Yes
2007886	NAP1		IX/XXIII		Yes	Yes

Table 38: Immunogenic Composition-induced Antibodies Neutralized Toxin A and Toxin B from Various Wild-type *C. difficile* Strains from Europe, including Hypervirulent strains

Cdiff Strain	PFGE Type	Other Typing Method	Toxin type	Neutralized by HS (IMR-90, Toxin B)	Neutralized by HS (HT-29, Toxin A)	
001	NAP2	Toxinotype 0 Strains	0	Yes	Yes	
002	NAP6			Yes	Yes	
012 (004)	NAPCR1			Yes	Yes	
014	UK			Yes	Yes	
015	NAP12			Yes	Yes	
020	NAP4			Yes	Yes	
029	UK			Yes	Yes	
046	UK			Yes	Yes	
053	NAP5			Yes	Yes	
059	UK			Yes	Yes	
077	UK			Yes	Yes	
078	UK			Yes	Yes	
081	UK			Yes	Yes	
087	UK			Yes	Yes	
095	UK	txnA-/txnB+	VIII	Yes	Yes	
106	UK			Yes	Yes	
117	UK			Yes	Yes	
017	NAP9	Hypervirulent	III	Yes	NA	
027	NAP1			Yes	Yes	
075	UK			Yes	Yes	
003	NAP10	Other	I	Yes	Yes	
023	UK		IV	Yes	Yes	
070	UK		XIII	Yes	Yes	
126	UK		UK	Yes	Yes	
131	UK		UK	In Progress	Yes	
Wild-type <i>C. difficile</i> strains obtained from Leeds Hospital, UK.						
"UK" = unknown status						
NA, not applicable; strain does not make toxin A; was not tested in Toxin A neutralization assay						

Example 38: Peptide mapping of EDC/NHS triple mutant toxins

To characterize the EDC/NHS inactivated triple mutant toxins, peptide mapping experiments were performed on four lots of EDC/NHS-treated triple mutant toxin A (SEQ ID NO: 4) and four lots of EDC/NHS-treated triple mutant B (SEQ ID NO: 6). After 5 digesting the mutant toxins with trypsin, the resulting peptide fragments were separated using reverse-phase HPLC. Mass spectral analysis was used to identify modifications that occur as a result of the inactivation process. For both mutant toxin A drug substance and mutant toxin B drug substance, greater than 95% of the theoretical tryptic peptides were identified. Crosslinks and glycine adducts (glycine was used as 10 the capping agent) were identified. In both mutant toxin A drug substance and mutant toxin B drug substance, beta-alanine adducts were also observed. Without being bound by mechanism or theory, the beta-alanine adducts appear to result from the reaction of three moles of NHS with one mole of EDC which forms NHS activated beta-alanine. This molecule can then react with lysine groups to form beta-alanine adducts (+70 Da). 15 In the EDC/NHS-treated triple mutant toxin B samples, low levels (0.07 moles/mole protein) of dehydroalanine (-34 Da) were also observed. Dehydroalanine is a result of de-sulfonation of a cysteine residue. The same type and degree of modification was observed in all four batches of each mutant toxin, indicating that the process produces a consistent product. Peptide mapping (at greater than 95% sequence coverage) 20 confirms that modifications are present. A summary of the modifications are shown in Table 39. See also Figures 24-25. In addition, the size and charge heterogeneity of the triple mutant toxin A drug substance and of the triple mutant toxin B drug substance increased, as compared to the size and charge heterogeneity of the respective triple mutant toxin A and triple mutant toxin B in the absence of chemical inactivation. As a 25 result, the size-exclusion chromatography (SEC) and anion-exchange chromatography (AEX) profiles had relatively broad peaks (data not shown).

Table 39. Summary of Modifications Observed in Mutant Toxin Drug Substances

Modification	# of Modified Residues	Total # of Residues	Degree of Modification	Moles modified/ mole protein
Mutant toxin A drug substance				
Crosslink	2	313 Asp/Glu	16-40%	0.6
Glycine moiety	8	313 Asp/Glu	10-53%	2.2
Beta Alanine moiety	19	233 Lys	10-60%	4.7
Mutant toxin B drug substance				
Crosslink	3	390 Asp/Glu	11-63%	0.8
Glycine moiety	23	390 Asp/Glu	10-31%	3.9
Beta Alanine moiety	10	156 Lys	12-42%	2.6
dehydroalanine	2	8 Cys	1.0-3.5%	.07

The degree of modification is calculated by dividing the HPLC area of modified peptide by the HPLC area of the native + modified peptide.

Example 39: Drug Product Production

The *C. difficile* immunogenic composition (drug product) contains two active pharmaceutical ingredients (mutant toxin A drug substance and mutant toxin B drug substance). An exemplary drug product is a lyophilized formulation containing 10 mM Tris buffer pH 7.4, 4.5% (w/w) trehalose dihydrate, and 0.01% (w/v) polysorbate 80, including each of a mutant toxin A drug substance and a mutant toxin B drug substance. See Table 40. The immunogenic composition is prepared for injection by resuspending the lyophilized vaccine either with diluent or with diluent containing Alhydrogel.TM The placebo will include a sterile normal saline solution for injection (0.9% sodium chloride).

10 **Table 40**

Component	Selected
Formulation dosage form	Lyophilized
Antigen dose per 0.5 mL	25, 50, 100 µg of each EDC/NHS-treated triple mutant toxin A (SEQ ID NO: 4) and EDC/NHS-treated triple mutant toxin B (SEQ ID NO: 6)
pH	7.4 ± 0.5
Buffer	10 mM Tris
Stabilizer/Bulking agent	4.5% Trehalose dihydrate (3 – 6%)
Surfactant	0.01% Polysorbate 80 (0.005 – 0.015%)
Container closures	2 mL 13 mm Type 1 flint glass Vial, Blowback, West - Flurotec

Buffer preparation

Water for injection (WFI) is added to a compounding vessel. While mixing, the excipients are added and dissolved until into solution. The pH is measured. If required, 15 the pH is adjusted to 7.4 ± 0.1 with HCl. The solution is diluted to the final weight with WFI then filtered using a 0.22 µm Millipore ExpressTM SHC XL150 filter. A pre-filtration bioburden reduction sample is taken prior to filtration. The filtered buffer is sampled for osmolality and pH.

Formulation preparation

The thawed mutant toxin Drug Substances are pooled in the formulation vessel based on the precalculated amounts in the following order of operation: 50% of the target dilution buffer volume to achieve 0.6 mg/mL is added to the vessel first, followed

5 by addition of mutant toxin A drug substance and mixed for 5 minutes at 100 rpm. Mutant toxin B drug substance is then added to the vessel and the solution is further diluted to 0.6 mg/mL dilution point and then mixed for another 5 minutes at 100 rpm. A sample is removed and tested for total mutant toxin concentration. The solution is

10 diluted to 100 percent volume based on the in-process mutant toxin concentration value then mixed for 15 minutes at 100 rpm. The formulated drug product is sampled for pH and bioburden pre-filtration. The formulated drug product is then filtered using a Millipore ExpressTM SHC XL150 for overnight storage, or brought to the filling line for

15 sterile filtration.

The formulated bulk is brought to the filling area, sampled for bioburden, and

15 then sterile filtered with two in-series Millipore ExpressTM SHC XL150 filters. The formulated bulk is filled into depyrogenated glass vials at a target fill volume of 0.73 mL. The filled vials are partially stoppered and then loaded into the freeze dryer. The lyophilization cycle is executed as shown in Table 41. At the completion of cycle, the lyophilization chamber is back-filled with nitrogen to 0.8 atm and then the stoppers are

20 fully seated. The chamber is unloaded and the vials are capped using flip-off seals.

Table 41. *C. difficile* Drug Product Lyophilization Cycle Set Points

Step	Temperature (°C)	Ramp (minutes)	Soak (minutes)	Pressure
Loading	5°C	N/A	60	-
Freezing 1	-50°C	183	60	-
Annealing	-10°C	133	180	
Freezing 2	-45°C	117	90	
Vacuum Initiation	-45°C	-	60	50
Primary Drying	-30°C	75	3420	50
Secondary Drying	30°C	300	600	50
Storage	5°C	50	-	50

Drug product stability data is summarized in Table 42. The data suggest that the drug product is physically and chemically stable during storage at 2-8°C for at least 3 months or at least 1 month at 25° or 40°C. Under both storage conditions, the level of impurities detected by size exclusion chromatography (SEC) did not change, nor were there 5 changes in *in vitro* antigenicity through the latest timepoints tested.

Table 42: Stability of Lyophilized Drug Product^a

	Drug Product Formulation 200 µg/mL mutant toxin A drug substance, 200 µg/mL mutant toxin B drug substance, 4.5% Trehalose dihydrate, 0.01% Polysorbate 80, 10mM Tris buffer pH 7.4			
Test	t=0	1 Month@25°C	1 Month@40°C	3 months @2-8°C
Appearance before Reconstitution.	White cake essentially free from visible foreign particulate matter	White cake essentially free from visible foreign particulate matter	White cake essentially free from visible foreign particulate matter	White cake essentially free from visible foreign particulate matter
Appearance after Reconstitution.	Clear colorless solution	Clear colorless solution	Clear colorless solution	Clear colorless solution
pH	7.5	7.6	7.6	7.5
Strength by AEX (µg/mL)	<ul style="list-style-type: none"> mutant toxin A drug substance 212 mutant toxin B drug substance 235 	<ul style="list-style-type: none"> Mutant toxin A drug substance 193 mutant toxin B drug substance 223 	<ul style="list-style-type: none"> Mutant toxin A drug substance 191 Mutant toxin B drug substance 222 	<ul style="list-style-type: none"> Mutant toxin A drug substance 193 mutant toxin B drug substance 230
Impurity by SEC	< 2.5%	2.8%	2.8%	2.9%
Characterization by SEC	HMMS: 29.6% Monomer: 68.0%	HMMS: 30.2% Monomer: 67.1%	HMMS: 30.2% Monomer: 67.1%	HMMS: 28.5% Monomer: 68.7%
Moisture	0.5	NA	NA	NA

a Lyophilized DP is reconstituted with 60 mM NaCl diluent for these tests.

Example 40: Vaccine diluents

For saline, 60 mM NaCl is used as a diluent for the lyophilized drug product without any adjuvant to ensure an isotonic solution upon reconstitution.

AlhydrogelTMAlhydrogelTM85 2% (Brenntag) is a commercially available Good

5 Manufacturing Practice (GMP) grade product composed of octahedral crystalline sheets of aluminum hydroxide. An exemplary AlhydrogelTM diluent formulation is shown in Table 43. The exemplary formulation may be used in combination with the drug product described above.

Component	Selected
Formulation dosage form	Liquid Suspension
Adjuvant dose per 0.5 mL	0.5 mg Al
pH	6.5 ± 0.5
Buffer	10 mM His
Salt	60 mM NaCl
Container closures	2 mL 13 mm Type 1 Flint Glass Vial, Blowback, West - Flurotec

10 Studies with the AlhydrogelTM adjuvant show 100% binding of mutant toxin A drug substance and mutant toxin B drug substance to 1 mg Al/mL AlhydrogelTM from pH 6.0 to 7.5. Maximum binding of both drug substances was seen at the highest protein concentration tested (300 µg/mL each).

15 The binding of the proteins to AlhydrogelTM was also tested with the lyophilized drug product formulation containing 200 µg/mL of each drug substance and AlhydrogelTM ranging from 0.25 to 1.5 mg/ml. The drug product was reconstituted with diluents containing the varying concentrations of AlhydrogelTM and the percent of each mutant toxin bound was measured. All tested concentrations of AlhydrogelTM demonstrated 100% binding of the antigens.

20 The binding kinetics of the proteins to AlhydrogelTM at the target dose of mutant toxin A drug substance and mutant toxin B drug substance (200 µg/mL each) were also assessed. The results show that 100% of the mutant toxin drug substances were bound to AlhydrogelTM throughout the 24-hour RT time course.

CpG 24555 and AlhydrogelTM: CpG 24555 is a synthetic 21-mer oligodeoxynucleotide (ODN) having a sequence 5-TCG TCG TTTTC GGT GCT TTT-3 (SEQ ID NO: 48). An exemplary formulation for a combination of CpG 24555 and AlhydrogelTM diluents is shown in Table 44. The exemplary formulation may be used in combination with the drug product described above.

Table 44: Formulation Rationale for CpG/AlhydrogelTM Diluent

Component	Selected
Formulation dosage form	Liquid Suspension
Adjuvant dose per 0.5 mL	0.5 mg Al and 1mg cpG
pH	6.5 ± 0.5
Buffer	10 mM His
Salt	60 mM NaCl
Container closures	2 mL 13 mm Type 1 Flint Glass Vial, Blowback, West - Flurotec

ISCOMATRIX®: The ISCOMATRIX® adjuvant is a saponin-based adjuvant known in the art. An exemplary formulation for the ISCOMATRIX® adjuvant formulation is shown in Table 45. The exemplary formulation may be used in combination with the drug product described above.

Table 45: Formulation Rationale for ISCOMATRIX® Diluent

Component	Selected
Formulation dosage form	Liquid Suspension
Adjuvant dose per 0.5 mL	45units
pH	6.2 ± 0.5
Buffer	10 mM phosphate
Salt	60 mM NaCl
Container closures	2 mL 13 mm Type 1 Flint Glass Vial, Blowback, West - Flurotec

**Example 41: Immunogenicity of Mutant Toxin Drug Substance Compositions
Adjuvanted with AlhydrogelTM in NHP Model and Preclinical Proof of Concept**

The immunogenicity of mutant toxin A drug substance and mutant toxin B drug substance compositions adjuvanted with AlhydrogelTM in NHPs was assessed, specifically

5 cynomolgus macaques. NHPs immunized at two-week intervals (weeks 0, 2, 4) with 10 μ g of each mutant toxin A drug substance and mutant toxin B drug substance compositions (formulated with AlhydrogelTM) per dose, developed robust neutralizing antitoxin responses. See Table 46. Both antitoxin A and antitoxin B neutralizing responses reached a protective range after the third immunization and remained within 10 or above the protective range at least through week 33 (last timepoint studied).

Cynomolgus macaques (n=8) were immunized IM at 0, 2 and 4 weeks with 10 μ g each of mutant toxin A drug substance and mutant toxin B drug substance formulated in 250 μ g of AlhydrogelTM. Sera was collected at each time point and analyzed in the toxin neutralization assay for functional antitoxin activity. GMTs are provided in Table 46.

15 The protective titer range provided in the table depicts the neutralizing antibody titer range which correlates to significant reduction in recurrence of *C. difficile* infection in the MerckTM monoclonal antibody therapy trial.

Table 46: Immunogenicity of Mutant Toxin A Drug Substance and Mutant Toxin B Drug Substance (Formulated in 250 μ g AlhydrogelTM) in Cynomolgus Monkeys (50% Neutralization Titer)

Antitoxin A (Merck/Medarex protective range: 666- 6,667 for antitoxin A)													
Week:	Wk 0	Wk 1	Wk 2	Wk 3	Wk 4	Wk 5	Wk 6	Wk 8	Wk 12	Wk 25	Wk 29	Wk 33	
Titer:	15	19	129	382	336	246	306	217	159	152	154	217	8

Antitoxin B (Merck/Medarex protective range: 222- 2,222 for antitoxin B)													
Titer:	10	10	10	10	20	311	410	446	676	163	297	351	0

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Correlation of Human Protective Antibody Titers from Merck mAb Therapy Trial to Titers Induced by Pfizer's Vaccine Candidate in NHPs

The Phase 2 efficacy study with Merck/Medarex mAbs (Lowy et al., *N Engl J Med.* 2010 Jan 21;362(3):197-205) seemed to demonstrate a correlation between the 5 level of neutralizing antitoxin mAbs in the serum and the prevention of recurrence of CDAD. After administration of the toxin-specific mAbs to humans, serum antibody levels in human recipients in the range of 10 to 100 µg/mL appear to protect against recurrences (70% reduction in the recurrence of CDAD).

10 Immunogenic compositions including the mutant toxin drug substances were tested to gauge whether the immunogenic compositions are capable of inducing a potentially efficacious neutralizing antibody responses in humans by comparing published data from the Merck/Medarex Phase 2 study to the levels of antibody induced by the immunogenic compositions in the NHP model. This was accomplished by utilizing previously published characteristics of the Merck/Medarex mAbs to convert the 15 range of these mAbs in the serum obtained from subjects that displayed no sign of recurrences (10-100 µg/mL) into 50% neutralization titers and comparing these titers ("protective titer range") to the titers observed in the preclinical models described herein. As shown in Table 46, the immunogenic compositions including the mutant toxin A drug 20 substance and mutant toxin B drug substance adjuvanted with AlhydrogelTM generated immune responses in NHPs that reached the "protective range" after the third dose and have remained within or above this range through week 33. The level of toxin-neutralizing antibodies induced in NHPs by the inventive *C. difficile* immunogenic composition is comparable to the serum antibody levels in the Merck/Medarex trial subjects who appeared to be protected from recurrences of CDAD.

25

Example 42: Immunogenicity of Mutant Toxin Drug Substance Compositions Adjuvanted with ISCOMATRIX or AlhydrogelTM/CpG 24555 (Alh/CpG) in NHP Model

In NHPs, both ISCOMATRIX and Alh/CpG statistically significantly enhanced antitoxin A and B neutralization titers when compared to vaccine administered with

5 AlhydrogelTM alone (Table 47). Antitoxin responses above background were elicited at earlier time points by vaccine administered with either Alh/CpG or ISCOMATRIX (week 2-4) as compared to AlhydrogelTM alone (week 4-6), which may have an important effect on protection from recurrence of CDAD in humans. Compared to AlhydrogelTM, the immunogenic composition adjuvanted with Alh/CpG or with ISCOMATRIX generated

10 antitoxin neutralization titers that reached the protective range (see also Example 41) more swiftly and that have remained within or above this range through week 33.

As shown in Table 47, Cynomolgus macaques were immunized IM at weeks 0, 2, and 4 with 10 μ g each of mutant toxin A drug substance and mutant toxin B drug substance formulated in 250 μ g of AlhydrogelTM (n=8), or 500 μ g of CpG + 250 μ g of

15 AlhydrogelTM (n=10), or 45 U of ISCOMATRIX (n=10). Sera were collected at each time point and analyzed in the toxin neutralization assay described above for functional antitoxin activity. GMTs are listed in the tables. Asterisks (*) indicate statistical significance (p<0.05) when compared to titers in the AlhydrogelTM group. The protective titer range represents the neutralizing antibody titer range which correlates to significant

20 reduction in recurrence of *C. difficile* infection according to the Merck/Medarex mAb therapy trial.

Table 47: Immunogenicity of Adjuvanted Mutant Toxin Drug Substances in NHPs (50% Neutralization Titer)							
Antitoxin A (Merck/Medarex protective range: 666- 6,667 for antitoxin A)							
Week:	Wk 0	Wk 2	Wk 4	Wk 6	Wk 12	Wk 25	Wk 33
Alhydrogel™ Titer:	15	129	336	3069	1599	1520	2178
Alhydrogel™ + CpG Titer:	17	*1004	*2162	*15989	*7179	*5049	*7023
ISCOMATRIX Titer:	25	*1283	*3835	*19511	*12904	*6992	*7971
Antitoxin B (Merck/Medarex protective range: 222- 2,222 for antitoxin B)							
Alhydrogel™ Titer:	10	10	20	410	676	1631	3510
Alhydrogel™ + CpG Titer:	10	13	*136	*2163	*5076	*9057	*27971
ISCOMATRIX Titer:	10	10	*269	*5325	*9161	*19479	*25119

The dose of mutant toxin A drug substance and mutant toxin B drug substance administered, in the presence of ISCOMATRIX or Alh/CpG adjuvants, on neutralizing 5 antitoxin antibody titers generated in NHPs was also evaluated. In one study, NHPs were administered a low (10 μ g) or a high (100 μ g) dose of each mutant toxin drug substance formulated in ISCOMATRIX. Responses were compared at each time point after immunization. As shown in Table 48, antitoxin neutralization titers were robust in both treatment groups. The antitoxin A titers were nearly equivalent at most time points 10 between the low dose and high dose groups, while there was a trend for the antitoxin B titers to be higher in the high dose group.

Table 48: Neutralizing Antitoxin Titers in NHPs Following Immunization with Either 10 μ g or 100 μ g of Each of Mutant Toxin Drug Substance and Mutant Toxin Drug Substance Administered with ISCOMATRIX (50% Neutralization Titer)

Antitoxin A (Merck/Medarex protective range: 666- 6,667 for antitoxin A)

Week	Wk 0	Wk 2	Wk 3	Wk 4	Wk 6	Wk 8	Wk 12
10μg Titer:	11	585	3522	4519	19280	10225	12084
100 μg Titer:	11	400	1212	2512	9944	10283	18337
Antitoxin B (Merck/Medarex protective range: 222- 2,222 for antitoxin B)							
10μg Titer:	10	10	112	266	3710	2666	7060
100 μg Titer:	10	10	303	469	6016	4743	20683

As shown in Table 48, Cynomolgus macaques (n=5) were immunized IM at weeks 0, 2, and 4 with 10 μ g or 100 μ g each of mutant toxin A drug substance and 5 mutant toxin B drug substance formulated with 45U of ISCOMATRIX. Sera were collected at each time point and analyzed in the toxin neutralization assay for functional antitoxin activity. GMTs are listed in the table. The protective titer range represents the neutralizing antibody titer range which correlates to significant reduction in recurrence of *C. difficile* infection in the Merck/Medarex mAb therapy trial.

In an effort to enhance the kinetics of antitoxin B responses, NHPs were immunized with a constant dose of mutant toxin A drug substance (10 µg) that was mixed with an increasing dose of mutant toxin B drug substance (10, 50, or 100 µg) in the presence of ISCOMATRIX or Alh/CpG adjuvants. Regardless of adjuvant, there
5 was a trend for groups that received higher doses of mutant toxin B drug substance (either 50 or 100 µg) to induce higher antitoxin B neutralizing responses in comparison to the 10 µg dose of mutant toxin B drug (Table 50, marked by * to indicate statistically significant increases). This trend was observed at most time points after the final
10 immunization. However, in some cases, antitoxin A neutralizing responses showed a statistically significant decrease (marked by ^ in Table 49) when the amount of mutant toxin B was increased.

As shown in Table 49 and Table 50, NHPs (10 per group) were immunized IM at weeks 0, 2, and 4 with different ratios of mutant toxin A drug substance and mutant toxin B drug substance (10µg of mutant toxin A drug substance plus either 10, 50, or
15 100 µg of mutant toxin B drug substance; designated 10A:10B, 10A:50B and 10A:100B, respectively, in Table 49 and Table 50), formulated with ISCOMATRIX (45U per dose) or with Alh/CpG/ (250 µg / 500 µg per dose). Table 49 shows Antitoxin A titers. Table 50 shows Antitoxin B titers. GMTs are listed in the tables. The protective titer range represents the neutralizing antibody titer range which correlates to significant
20 reduction in recurrence of *C. difficile* infection in the Merck mAb therapy trial. The symbol ^, represents statistically significant decrease in neutralizing titers (p<0.05) compared to the 10A: 10B group. The asterisk symbol, *, represents statistically significant increase in neutralizing titers (p<0.05) compared to the 10A: 10B group.

Table 49: Neutralizing Antitoxin Titers in NHPs Following Immunization with 10µg Mutant Toxin A Drug Substance Combined with 10, 50, or 100 µg Mutant Toxin B Drug Substance using ISCOMATRIX or Alh/CpG as Adjuvants (50% Neutralization Titer)

Antitoxin A (Merck/Medarex protective range: 666- 6,667 for antitoxin A)

ISCOMATRIX

Week:	Wk 0	Wk 2	Wk 4	Wk 6	Wk 12	Wk 25	Wk 33
10A:10B Titer:	25	1283	3835	19511	12904	6992	7971
10A:50B Titer:	29	906	2917	16126	^7756	^4208	5965
10A:100B Titer:	20	982	2310	^5034	^5469	^4007	3780

Antitoxin A (Merck/Medarex protective range: 666- 6,667 for antitoxin A)

Alh/CpG

10A:10B Titer:	17	1004	2162	15989	7179	5049	7023
10A:50B Titer:	20	460	1728	16600	6693	6173	8074
10A:100B Titer:	27	^415	1595	13601	6465	5039	6153

Table 50: Neutralizing Antitoxin Titers in NHPs Following Immunization with 10µg Mutant Toxin A Drug Substance Combined with 10, 50, or 100 µg Mutant Toxin B Drug Substance using ISCOMATRIX or Alh/CpG as Adjuvants (50% Neutralization Titer)

Antitoxin B (Merck/Medarex protective range: 222- 2,222 for antitoxin B)

Week:	Wk 0	Wk 2	Wk 4	Wk 6	Wk 12	Wk 25	Wk 33
ISCOMATRIX	10	10	269	5325	9161	19479	25119
Titer:							
Titer:	13	*20	*604	4861	10801	20186	*57565
Titer:	10	*23	*862	*10658	10639	*33725	*56073

Antitoxin B (Merck/Medarex protective range: 222- 2,222 for antitoxin B)

Alh/CpG	10	13	136	2163	5076	9057	27971
Titer:							
Titer:	10	15	*450	*5542	*9843	15112	50316
Titer:	11	17	*775	*13533	*11708	*17487	26600

Example 43: Five-Week Repeat-Dose IM Toxicity Study with an Immunogenic Composition in Cynomolgus Monkeys, with a 4-Week Recovery Period

The 5-week IM repeat-dose toxicity study with PF-06425095 (an immunogenic composition including triple mutant toxin A drug substance and triple mutant toxin B

5 drug substance in a combination with adjuvants aluminum hydroxide and CpG 24555) in Cynomolgus monkeys was conducted to assess the potential toxicity and immunogenicity of *C. difficile* triple mutant toxin A drug substance and triple mutant toxin B drug substance in a combination with the adjuvants aluminum hydroxide and CpG 24555 (PF-06425095). PF-06425095 at 0.2 or 0.4 mg/dose triple mutant toxin A
10 drug substance and triple mutant toxin B drug substance (low- and high-dose immunogenic composition groups, respectively), 0.5 mg aluminum as aluminum hydroxide, and 1 mg CpG 24555 and the adjuvant combination alone (aluminum hydroxide + CpG 24555; PF-06376915) were administered IM to cynomolgus monkeys (6/sex/group) as a prime dose followed by 3 booster doses (Days 1, 8, 22, and 36). A
15 separate group of animals (6/sex) received 0.9% isotonic saline at an approximate pH of 7.0. The immunogenic composition vehicle was composed of 10 mM Tris buffer at pH 7.4, 4.5% trehalose dihydrate, and 0.1% polysorbate 80. The adjuvant control vehicle was composed of 10 mM histidine buffer with 60 nM NaCl at pH 6.5. The total dose volume was 0.5 mL per injection. All doses were administered into the left and/or
20 right quadriceps muscle. Selected animals underwent a 4-week dose-free observation period to assess for reversibility of any effects observed during the dosing phase of the study.

There were no adverse findings in this study. PF-06425095 was well-tolerated and produced only local inflammatory reaction without evidence of systemic toxicity.

25 During the dosing phase, dose-dependent increases from pretest in fibrinogen (23.1% to 2.3x) on Days 4 and 38 and C-reactive protein on Days 4 (2.1x to 27.5x) and 38 (2.3x to 101.5x), and globulin (11.1% to 24.1%) on Day 36 and/or 38, were seen in immunogenic composition-treated groups and were consistent with the expected inflammatory response to administration of an adjuvanted immunogenic composition.

The increases in fibrinogen and C-reactive protein noted on Day 4 had partially recovered by Day 8 with increases in fibrinogen (25.6% to 65.5%) and C-reactive protein (4.5x and 5.6x) in the high-dose immunogenic composition group only.

Increases in interleukin (IL)-6 were observed in the low- and high-dose immunogenic 5 composition groups on Day 1, Hour 3 (8.3x to 127.2x individual values Day 1, Hour 0, dose responsive) and Day 36, Hour 3 (9.4x to 39.5x individual values Day 36, Hour 0). There were no changes observed in the other cytokines (IL-10, IL-12, Interferon-Inducible Protein (IP-10), and Tumor Necrosis Factor α (TNF- α). Increases in these acute phase proteins and cytokine were part of the expected normal physiologic 10 response to the administration of foreign antigen. There were no PF 06425095-related or adjuvant-related alterations in these clinical pathology parameters in the recovery phase (cytokines were not evaluated during the recovery phase). In addition, there were localized changes at the injection sites, which were of similar incidence and severity in the adjuvant control group and the low- and high-dose immunogenic 15 composition groups; hence, they were not directly related to PF-06425095. During the dosing phase, the changes included minimal to moderate chronic-active inflammation that was characterized by separation of muscle fibers by infiltrates of macrophages, which often contained basophilic granular material (interpreted as aluminum-containing adjuvant), lymphocytes, plasma cells, neutrophils, eosinophils, necrotic debris, and 20 edema. The basophilic granular material was also present extracellularly within these foci of chronic-active inflammation. At the end of the recovery phase, there was minimal to moderate chronic inflammation and mononuclear cell infiltrate, and minimal fibrosis. These injection site findings represent a local inflammatory response to the adjuvant. Other microscopic changes included minimal to moderate increased lymphoid cellularity 25 in the iliac (draining) lymph node and minimal increased cellularity in germinal centers in the spleen that were noted during the dosing phase in the adjuvant control group and the low- and high-dose immunogenic composition groups. At the end of the recovery phase, these microscopic findings were of lower severity. These effects represent an immunologic response to antigenic stimulation, and were a pharmacologic response to 30 the adjuvant or PF-06425095. There was no test article-related increase in anti-DNA antibodies.

Based on absence of adverse findings, the no observed adverse effect level (NOAEL) in this study is the high-dose immunogenic composition group (0.4 mg of triple mutant toxin A drug substance and triple mutant toxin B drug substance/dose as PF-06425095) administered as two 0.5 mL injections for four doses.

5

Example 44: Efficacy of Seropositive NHP Sera Passively Transferred to Hamsters

Groups of 5 Syrian golden hamsters were administered an oral dose of clindamycin antibiotic (30 mg/kg) to disrupt normal intestinal flora. After five days, the 5 hamsters were challenged with an oral dose of wild type *C. difficile* spores (630 strain, 100 cfu per animal), and administered intraperitoneally (IP) with NHP sera according to Table 51. Without being bound by mechanism or theory, disease symptoms following challenge with the spores typically manifest beginning about 30-48 hours post-challenge.

10 The NHP sera that were administered to the hamsters were pooled from NHP serum samples exhibiting the highest titer (anti-toxin A sera and anti-toxin B sera) following three immunizations with mutant toxin A drug substance and mutant toxin B drug substance (10:10, 10:50, and 10:100 A:B ratios), formulated with ISCOMATRIX (see Example 42, Table 49, and Table 50). The NHP sera were collected from 15 timepoints at weeks 5, 6, and 8 (immunizations occurred at weeks 0, 2, and 4), as described in Example 42. Results are shown in Tables 52-54 below. The symbol “+” indicates a Geometric mean (GM) in () that does not include animal #3, non-responder. “*TB” represents terminal bleed, the day the animal was euthanized, which is not the same for all animals.

20

Table 51: Experimental design

Group	Administered composition	No. animals	Route	Schedule
1 “1 dose”	Seropositive NHP sera (unconcentrated)	5	IP	Challenge Day 0 Dose day 0 Bleed days 0, 1, 2, TB on day11
2 “2 dose”	Seropositive NHP sera (unconcentrated)	5	IP	Challenge Day 0 Dose days 0, 1 Bleed days 0, 1, 2, TB on day11

Table 52: Anti-toxin A Neutralization Titers in Hamster Sera Following 1 or 2 IP doses of NHP Sera (50% Neutralization Titer in RLU)

	Day	Hamster 1	Hamster 2	Hamster 3	Hamster 4	Hamster 5	GM	SE
1 dose	D0	50	50	50	50	50	50	0
	D1	2877	4008	2617	4917	1872	3081	538
	D2	1983	3009	2750	2902	1117	2214	357
		3239		155				
	TB* (d4)		537 (d9)	(d11)	977 (d9)	972 (d2)	762	538
2 dose	D0	50	50	50	50	50	50	0
	D1	1154	2819	50	429	1174	606 (1131) ⁺	475
	D2	4119	4674	1899	545		2113	862
		1236	1267	1493		1877		
	TB* (d9)		(d8)	(d4)	50 (d11)	(d9)	738	306
Input NHP sera = 41976								

Table 53: Anti-toxin B Neutralization Titers in Hamster Sera Following 1 or 2 IP doses of NHP Sera (50% Neutralization Titer in RLU)

	Day	Hamster 1	Hamster 2	Hamster 3	Hamster 4	Hamster 5	GM	SE
1 dose	D0	50	50	50	50	50	50	0
	D1	1846	4254	1347	5178	406	1859	904
	D2	992	1795	2585	2459	1145	1669	327
	TB* (d4)	1744	50		265	544		317
2 dose	D0	50	50	50	50	50	50	0
	D1	1189	2229	50	550	3920	778 (1546) ⁺	687
	D2	2288	2706	1452	287		1268	477
	TB* (d9)	301	694	682	50	1334	394	217
Input NHP sera = 23633								

Table 54: Percentage of hamsters protected from severe CDAD following 1 or 2 IP doses of NHP sera

Days post-infection	0	2	4	6	8	10	11
1 dose NHP Sera	100%	80%	60%	60%	60%	20%	20%
2 dose NHP Sera	100%	100%	80%	80%	60%	20%	20%
Placebo	100%	75%	50%	25%	0%	n/a	n/a

In another study, Syrian golden hamsters were administered an oral dose of clindamycin antibiotic (30 mg/kg) to disrupt normal intestinal flora. After five days, the hamsters were challenged with an oral dose of wild type *C. difficile* spores (630 strain, 100 cfu per animal), and administered intraperitoneally (IP) NHP sera according to 5 Table 55. Without being bound by mechanism or theory, disease symptoms following challenge with the spores typically manifest beginning about 30-48 hours post-challenge.

The NHP sera that were administered to the hamsters were pooled from samples collected from NHPs following three immunizations with mutant toxin A drug substance 10 and mutant toxin B drug substance (10:10, 10:50, and 10:100 A:B ratios), formulated with AlhydrogelTM and CpG 24555 (see Example 42, Table 49, and Table 50). The NHP sera were collected from timepoints at weeks 5, 6, 8, and 12 as described in Examine 42 (NHPs were immunized on weeks 0, 2, and 4). Results are shown in Tables 56-59 below. Sera from the hamsters were further investigated to determine inhibitory 15 concentration (IC₅₀) value, which were determined using the toxin neutralization assay described above. The level of toxin-neutralizing antibodies induced in hamsters by the inventive *C. difficile* immunogenic composition is comparable to the serum antibody levels in the Merck/Medarex trial subjects who appeared to be protected from recurrences of CDAD.

20

Table 55: Experimental Design				
Group	Administered Composition	No.	Route	Schedule
1	Seropositive NHP sera	5	IP	Challenge D0 Dose D0, 1, 3, 5, 7
2	Seropositive NHP sera	5	IP	no challenge Dose D0, 1, 3, 5, 7
3	Seropositive NHP sera	10	IP	Challenge D0 Dose D0, 1, 3, 5, 7
4	Placebo	5	IM	Challenge D0

Table 56: Anti-toxin A Neutralization Titers^a in Hamster Sera Following 1 or 2 IP doses of NHP Sera (50% Neutralization Titer in RLU)

Day	Challenged (Groups 1 and 3)	Not Challenged (Group 2)	p Value
0	11	12	0.5933
1	380	720	0.034*
3	666	1220	0.0256*
5	864	1367	0.0391*
7	564	1688	0.0411*
11	263	1281	0.001*

Input NHP sera pool = 9680

a, titers expressed as geometric means for each group (n=15 at day 0 for “challenged” group, n=5 for “not challenged” group)

Merck/Medarex protective range: 666- 6,667 for antitoxin A

The asterisk “*” indicates a significant difference.

Table 57: Anti-toxin B Neutralization Titers^a in Hamster Sera Following 1 or 2 IP doses of NHP Sera (50% Neutralization Titer in RLU)

Day	Challenged (Groups 1 and 3)	Not Challenged (Group 2)	p Value
0	10	10	0.3343
1	465	828	0.0579
3	765	1400	0.0273*
5	941	1734	0.0226*
7	611	1877	0.0498*
11	194	1436	0.0047*

Input NHP sera pool = 19631

a, titers expressed as geometric means for each group (n=15 at day 0 for “challenged” group, n=5 for “not challenged” group)

Merck/Medarex protective range: 222- 2,222 for antitoxin B

The asterisk “*” indicates a significant difference.

Table 58: Percentage of hamsters protected from severe CDAD following IP dose of NHP sera

Days post-infection	0	2	4	6	8	10	11
Groups 1 and 3	100%	73%	53%	53%	47%	33%	33%
Placebo (Group 2)	100%	50%	0%				

Table 59: IC₅₀ values from Toxin-specific 50% Neutralization Titers

		IC ₅₀ of Anti Toxin A								IC ₅₀ of Anti Toxin B					
		Day of Post Dose								Day of Post Dose					
Challenged	Animal ID	0	1	3	5	7	11	Animal ID	0	1	3	5	7	11	
	1-1	10	50	338	died	D4		1-1	10	50	254	died			
	1-2	10	614	579	777	605	192	1-2	10	720	659	896	475	157	
	1-3	10	710	1035	845	548	D10	1-3	10	867	1017	988	694		
	1-4	10	850	588	942	1116	296	1-4	10	1158	555	1158	1806	250	
	1-5	10	780	895*				1-5	10	910	687*				
	3-1	10	647	D2				3-1	10	598	D2				
	3-2	10	331	D2				3-2	10	290	D2				
	3-3	10	660	1273	849	692	640	3-3	10	717	1623	870	791	574	
	3-4	10	536	493	1102	1314	D9	3-4	10	618	598	977	1478	D9	
	3-5	10	817	807	774	1077	187	3-5	10	772	1260	850	913	243	
	3-6	10	117	649	803	50	186	3-6	10	1038	773	883	50	50	
	3-7	10	50	D2				3-7	10	50	D2				
	3-8	10	149	659	650*			3-8	10	121	1010	517*			
	3-9	30	797	1170*				3-9	10	1008	1720*				
	3-10	10	792	D2				3-10	10	835	D2				
	GeoMean	11	380	666	864	564	263	GeoMean	10	465	765	941	611	194	
Not Challenged	Std Error	1	78	86	41	163	88	Std Error	0	94	125	38	224	88	
	2-1	10	697	1634	1597	2219	1709	2-1	10	890	1777	1910	3229	1355	
	2-2	10	779	1207	1322	1755	1327	2-2	10	939	1378	1564	1897	1379	
	2-3	10	581	669	722	1401	1118	2-3	10	828	837	865	1484	1404	
	2-4	26	856	1540	1875	1830	1826	2-4	10	748	1780	2939	1880	2650	

	2-5	10	715	1331	1668	1374	744	2-5	10	752	1475	2064	1364	880
	GeoMean	12	720	1220	1367	1688	1281	GeoMean	10	828	1400	1734	1877	1436
	Std Error	3	46	169	199	156	197	Std Error	0	38	173	338	332	296

*= deceased on that day

Example 45: Characterization of Mutant Toxin Drug Substances

The primary structure of triple mutant toxin A is shown in SEQ ID NO: 4. The NH₂-terminal Met residue at position 1 of SEQ ID NO: 4 is originated from the initiation codon of SEQ ID NO: 12 and is absent in isolated protein (e.g., see SEQ ID NO: 84).

5 Accordingly, in Example 12 to Example 45, “SEQ ID NO: 4” refers to SEQ ID NO: 4 wherein the initial methionine (at position 1) is absent. Both purified triple mutant toxin A (SEQ ID NO: 4) (Drug Substance Intermediate - Lot L44993-132) and EDC/NHS treated triple mutant toxin A (SEQ ID NO: 4) (“mutant toxin A Drug Substance” - Lot L44898-012) displayed a single NH₂-terminal sequence starting at SLISKEELIKLAYSI
10 (positions 2-16 of SEQ ID NO: 4).

The primary structure of triple mutant toxin B is shown in SEQ ID NO: 6. The NH₂-terminal Met residue at position 1 of SEQ ID NO: 6 is originating from the initiation codon and is absent in isolated protein (e.g., see SEQ ID NO: 86). Accordingly, in Example 12 to Example 45, “SEQ ID NO: 6” refers to SEQ ID NO: 6 wherein the initial 15 methionine (at position 1) is absent. Both purified triple mutant toxin B (SEQ ID NO: 6) (Drug Substance Intermediate - Lot 010) and EDC/NHS treated triple mutant toxin B (SEQ ID NO: 6) (“mutant toxin B Drug Substance” - Lot L44906-153) displayed a single NH₂-terminal sequence starting at SLVNRKQLEKMANVR (positions 2-16 of SEQ ID NO: 6).

20 Circular dichroism (CD) spectroscopy was used to assess secondary and tertiary structure of triple mutant A (SEQ ID NO: 4) and mutant toxin A drug substance. CD spectroscopy was also used to assess secondary and tertiary structure of the triple mutant toxin B (SEQ ID NO: 6) and the mutant toxin B drug substance. CD spectroscopy was also used to assess potential effects of pH on structure. The effect of 25 EDC treatment on triple mutant toxin A was analyzed by comparing CD data obtained for mutant toxin A drug substance to the data obtained for triple mutant toxin A. The effects of EDC treatment on triple mutant toxin B (SEQ ID NO: 6) were analyzed by comparing CD data obtained for mutant toxin B drug substance to the data obtained for triple mutant toxin B.

Mutant toxin A drug substance far-UV CD data were obtained at various pH. Spectra recorded at pH 5.0-7.0 are indicative of high proportion of α -helices in the secondary structure, suggesting that polypeptide backbone of the protein adopts well-defined conformation dominated by α -helices.

5 Near-UV CD spectra of mutant toxin A drug substance were also obtained. Strong negative ellipticity between 260 and 300 nm is an indication that aromatic side chains are in the unique rigid environment, i.e. mutant toxin A drug substance possesses tertiary structure. In fact, characteristic features arising from individual types of aromatic side chains can be distinguished within the spectrum: shoulder at \sim 290 nm
10 and largest negative peak at \sim 283 nm are due to absorbance of the polarized light by ordered tryptophan side chains, negative peak at 276 nm is from the tyrosine side chains, and minor shoulders at 262 and 268 nm are indicative of the phenylalanine residues participating in tertiary contacts. Far- and near-UV results provide evidence that mutant toxin A drug substance retains compactly folded structure at physiological
15 pH. Nearly identical far- and near-UV CD spectra observed at pH 5.0 - 7.0 indicate that no detectable structural changes are taking place within this pH range. CD data could not be collected at pH 3.0 and 4.0, since the protein was insoluble at these pH points. In comparing far- and near-UV CD spectra of mutant toxin A drug substance with those of the triple mutant toxin A, spectra of both proteins are essentially identical under all of
20 the experimental conditions studied, indicating that EDC treatment had no detectable effects on secondary and tertiary structure of the triple mutant toxin A. This finding is in agreement with the gel-filtration and analytical ultracentrifugation results, which show no detectable changes in Stokes radii and sedimentation/frictional coefficients, respectively.

25

Mutant toxin A drug substance (as well as triple mutant toxin A) contains 25 tryptophan residues that are spread throughout the primary sequence and can serve as convenient intrinsic fluorescence probes. Fluorescence emission spectra of mutant toxin A drug substance between 300 and 400 nm as a function of temperature were 5 obtained. At 6.8°C mutant toxin A drug substance shows characteristic tryptophan fluorescence emission spectrum upon excitation at 280 nm. Fluorescence emission maximum is observed at ~335 nm, indicating that tryptophan residues are in non-polar environment, typical of protein interiors rather than of polar aqueous environments. The fluorescence emission spectra results, together with the results of the CD experiments 10 presented in this report, confirm that mutant toxin A drug substance retains compact folded structure.

Fluorescence of the extrinsic probe 8-anilino-1-naphthalene sulfonic acid (ANS) was used to characterize possible conformational changes in mutant toxin A drug substance and triple mutant toxin A upon changes in pH. As can be seen from the 15 results, there is essentially no increase in ANS fluorescence intensity when either mutant toxin A drug substance or triple mutant toxin A are titrated with the probe at pH 7.0, suggesting that no hydrophobic surfaces are exposed on the proteins under these conditions. Shifting pH to 2.6 leads to a dramatic increase in ANS fluorescence quantum yield upon increase in probe's concentration, until fluorescence quantum yield 20 reaches apparent saturation. This increase in ANS fluorescence quantum yield indicates that at low pH (2.6), both mutant toxin A drug substance and triple mutant toxin A undergo pH-induced conformational change that exposes hydrophobic surfaces. Such conformational changes indicate that EDC-induced modification and inactivation 25 of triple mutant toxin A did not restrict conformational plasticity of mutant toxin A drug substance (DS).

Effect of EDC treatment on hydrodynamic properties of triple mutant toxin A was evaluated using size-exclusion chromatography on a G4000 SWXL column. Mutant toxin A drug substance and triple mutant toxin A were injected onto the G4000 SWXL column equilibrated at pH 7.0, 6.0, and 5.0. The data indicate that no differences in the 30 Stoke's radius of mutant toxin A drug substance and triple mutant toxin A can be detected using size exclusion chromatography. Therefore, EDC treatment has not dramatically affected hydrodynamic properties and, correspondingly, overall molecular shape of the triple mutant toxin A.

Further analysis of triple mutant toxin A and mutant toxin A drug substance was performed using multi-angle laser light scattering (MALLS) technique. Treatment of triple mutant toxin A with EDC resulted in generation of heterogeneous mixture composed of various multimeric and monomeric species. Such heterogeneity reflects 5 introduction of a large number of EDC-induced inter- and intra-molecular covalent bonds between carboxyls and primary amines of the protein.

Obtained data provide physical and chemical characteristics of triple mutant toxin A and mutant toxin A drug substance (triple mutant toxin A treated with EDC) and describe the key features of their primary, secondary, and tertiary structure. Generated 10 data demonstrate that treatment of triple mutant toxin A with EDC resulted in covalent modification of its polypeptide chain but did not affect secondary and tertiary structures of the protein. Treatment with EDC leads to intra- and intermolecular cross-linking. The biochemical and biophysical parameters obtained for mutant toxin A drug substance (as well as triple mutant toxin A) are presented in Table 60.

15

Table 60: Major Biochemical and Biophysical Parameters Obtained for Triple Mutant Toxin A (SEQ ID NO: 4) and Mutant Toxin A Drug Substance

Parameter	Triple Mutant toxin A (SEQ ID NO: 4)	Mutant Toxin A Drug Substance
Number of amino acid residues	2709	2709
N-terminal sequence	SLISKEELIKLAYS (positions 2-16 of SEQ ID NO: 4)	SLISKEELIKLAYS (positions 2-16 of SEQ ID NO: 4)
Mol mass (from AA sequence)	308 kDa	308 kDa
Mol mass (from SEC-MALLS)	299 kDa	300 kDa and 718 -1139 kDa
Extinction coefficient at 280 nm	1.292 or 1.275 (mg/ml) ⁻¹ cm ⁻¹	1.292 or 1.275 275 (mg/ml) ⁻¹ cm ⁻¹
Theoretical pl	5.57	ND
Partial specific mol volume at 20 °C	0.735 cm ³ /g	0.735 cm ³ /g
Anhydrous volume monomer	3.8x10 ⁻¹⁹ cm ³	3.8x10 ⁻¹⁹ cm ³
Sedimentation coefficient/monomer	9.2S	9.2S
Frictional coefficient ratio (f/f ₀)	1.69	1.69
Stokes radius /monomer	78.4±1.1	77.9
Fluorescence max (λ _{ex} = 280 nm)	334-335 nm	334-335 nm
Near-UV CD spectrum minima	284 nm and 278 nm	284 nm and 278 nm
Mean res ellipticity at 284 & 278 nm	-138±7 & -130±7	-138±8 & 131±10
Mean res ellipticity at 222 nm	-8989±277	-7950±230
DSC unfolding transitions maxima (PBS, pH 7.4)	47.3 °C and 53.6 °C	47.9±0.2 °C and 54.1±0.2 °C

Mutant toxin B drug substance far-UV CD data were obtained at various pH. Spectra recorded at pH 5.0-7.0 are indicative of high proportion of α -helices in the secondary structure, suggesting that polypeptide backbone of the protein adopts well-defined conformation dominated by α -helices.

5 Near-UV CD spectra of mutant toxin B drug substance were also obtained. Strong negative ellipticity between 260 and 300 nm is an indication that aromatic side chains are in the unique rigid environment, i.e. mutant toxin B drug substance possesses tertiary structure. In fact, characteristic features arising from individual types of aromatic side chains can be distinguished within the spectrum: shoulder at \sim 290 nm
10 and largest negative peak at \sim 283 nm are due to absorbance of the polarized light by ordered tryptophan side chains, negative peak at 276 nm is from the tyrosine side chains, and minor shoulders at 262 and 268 nm are indicative of the phenylalanine residues participating in tertiary contacts. Far- and near-UV CD spectra provide evidence that mutant toxin B drug substance retains compactly folded structure at
15 physiological pH. Very similar far- and near-UV CD spectra observed at pH 5.0 - 7.0 indicate that no detectable secondary or tertiary structural changes are taking place within this pH range. CD data could not be collected at pH 3.0 and 4.0, since the protein was insoluble at these pH points.

In comparing far- and near-UV CD spectra of mutant toxin B drug substance with
20 those of the triple mutant toxin B, spectra of both proteins are very similar between pH 5.0 and 7.0, indicating that EDC treatment had no detectable effects on secondary and tertiary structure of the protein.

Triple mutant toxin B contains 16 tryptophan residues that are spread throughout the primary sequence and can serve as convenient intrinsic fluorescence probes.
25 Fluorescence emission spectra of mutant toxin B drug substance between 300 and 400 nm as a function of temperature were obtained. At 7°C mutant toxin B drug substance shows characteristic tryptophan fluorescence emission spectrum upon excitation at 280 nm. Fluorescence emission maximum is observed at \sim 335 nm, indicating that tryptophan residues are in non-polar environment, typical of protein interiors rather than
30 of polar aqueous environments. This result, together with the results of the CD experiments (see above), confirm that mutant toxin B drug substance retains compact folded structure.

Fluorescence of the extrinsic probe 8-anilino-1-naphthalene sulfonic acid (ANS) was used to characterize possible conformational changes in mutant toxin B drug substance and triple mutant toxin B upon changes in pH. As can be seen from the results, there is essentially no increase in ANS fluorescence intensity when either 5 mutant toxin B drug substance or triple mutant toxin B are titrated with the probe at pH 7.0, suggesting that no hydrophobic surfaces are exposed on the proteins under these conditions. Shifting pH to 2.6 leads to a dramatic increase in ANS fluorescence quantum yield upon increase in probe's concentration in the presence of mutant toxin B drug substance, until fluorescence quantum yield reaches apparent saturation. This 10 increase in ANS fluorescence quantum yield indicates that at low pH (2.6), mutant toxin B drug substance undergoes pH-induced conformational change that exposes hydrophobic surfaces. Such conformational changes indicate that EDC-induced modification and inactivation of triple mutant toxin B did not restrict conformational plasticity of mutant toxin B drug substance (DS).

15 Effect of EDC treatment on hydrodynamic properties of triple mutant toxin B was evaluated using size-exclusion chromatography on a G4000 SWXL column. mutant toxin B drug substance and triple mutant toxin B were injected onto the G4000 SWXL column equilibrated at pH 7.0, 6.0, 5.0. The data indicate that no differences in the Stoke's radius of mutant toxin B drug substance and triple mutant toxin B can be 20 detected using size-exclusion chromatography, therefore EDC treatment has not dramatically affected hydrodynamic properties and, correspondingly, overall molecular shape of the protein.

25 Further analysis of triple mutant toxin B and mutant toxin B drug substance was performed using multi-angle laser light scattering (MALLS) technique. Treatment of triple mutant toxin B with EDC resulted in generation of more heterogeneous mixture that is composed of various multimeric and monomeric species. Such heterogeneity reflects introduction of a large number of EDC-induced inter- and intra-molecular covalent bonds between carboxyls and primary amines of the protein.

Obtained data provide physical and chemical characteristics of triple mutant toxin B and mutant toxin B drug substance (triple mutant toxin B treated with EDC) and describe the key features of their primary, secondary, and tertiary structure. Generated data demonstrate that treatment of triple mutant toxin B with EDC resulted in covalent 5 modification of its polypeptide chain but did not affect secondary and tertiary structures of the protein. Treatment with EDC leads to intra- and intermolecular cross-linking. The major biochemical and biophysical parameters obtained for mutant toxin B drug substance (as well as triple mutant toxin B) are presented in Table 61.

Table 61: Major Biochemical and Biophysical Parameters Obtained for Triple Mutant Toxin B (SEQ ID NO: 6) and Mutant Toxin B Drug Substance

Parameter	Triple mutant toxin B (SEQ ID NO: 6)	Mutant Toxin B Drug Substance
Number of amino acid residues	2365	2365
N-terminal sequence	SLVNRKQLEKMANVR (positions 2-16 of SEQ ID NO: 6)	SLVNRKQLEKMANVR (positions 2-16 of SEQ ID NO: 6)
Mol mass (from AA sequence)	269.5 kDa	269.5 kDa
Mol mass (from SEC-MALLS)	255 kDa and ~1,754 kDa	264, 268, 706, and 2,211 kDa
Extinction coefficient at 280 nm	1.067 (mg/ml) ⁻¹ cm ⁻¹	1.067 (mg/ml) ⁻¹ cm ⁻¹
Theoretical pl	4.29	ND
Partial specific mol volume at 20 °C	0.734 cm ³ /g	0.734 cm ³ /g
Anhydrous volume monomer	3.3x10 ⁻¹⁹ cm ³	3.3x10 ⁻¹⁹ cm ³
Sedimentation coefficient/monomer	9.1±0.2S	9.4S
Frictional coefficient ratio (f/f ₀)	1.58±0.03	1.53
Stokes radius /monomer	76.2	76.2
Fluorescence max (λ _{ex} = 280 nm)	335 nm	335 nm
Near-UV CD negative bands	290, 283, 276, 268, 262 nm	290, 283, 276, 268, 262 nm
Far-UV CD negative bands	208 and 222 nm	208 and 222 nm
DSC unfolding transition midpoints T _{m1} and T _{m2} (PBS, pH 7.0)	48.8±0.0 °C and 52.0±0.1 °C	48.2±0.3 °C and 54.3±0.2 °C

Example 46: Perfusion Fermentation for Toxoid B (triple mutant)

Toxoid B (triple mutant) seed cultures: inoculated each 1L bottle containing 400 mL of medium with 1 ml seed, incubate at 37°C, stationary overnight (~15 hrs). The final OD₆₀₀ should be 3.0 – 4.0. Working vol: 3L (2.7L medium + 300 mL inoculum). Each fermenter had 1

5 Rushton impeller and a tube sparger. Initial conditions: Temperature: 37 °C, N2 flow: ~0.5 vvm, sparged. Controllers: pH controlled at 7.0 with 5N NaOH. Foam controlled by automatic addition of PPG-2000, with 0.25 mL/L added to the fermenter medium before sterilization.

Perfusion culture of *C. difficile* was performed using a stack of 2 SARTOCON Slice Cassettes, 0.2 µm pore size of HYDROSART filter material, 0.1 square meter surface area per

10 cassette. The 3L was added to the fermentors in the 10 L glass fermentors. The perfusion started when the OD reaches the target ~4 for 2 hour intervals of increasing speed as 0.75 L/hr, 1.5 L/hr, 2.25 L/hr, 3 L/hr for example 1, Figure 28. The perfusion will be started with fermentation medium when the OD reaches the target ~4 for 2 hour intervals of increasing speed as 0.75 L/hr, 1.5 L/hr, 3 L/hr, and 6 L/hr for example 2, Figure 29. At the start signal for 15 perfusion, the recirculation pump was started up at the desired speed at 1.3 L/min crossflow.

Example 1, Figure 28: Final OD 50 and toxoid B titer 243 mg/L were obtained.

Example 2, Figure 29: Final OD 59 and toxoid B titer 306 mg/L were obtained.

The invention also provides the following embodiments as defined in the clauses below:

Clause 1. An isolated polypeptide comprising SEQ ID NO: 183.

Clause 2. An isolated polypeptide comprising SEQ ID NO: 184.

Clause 3. An immunogenic composition comprising an isolated polypeptide

5 comprising SEQ ID NO: 183.

Clause 4. An immunogenic composition comprising an isolated polypeptide comprising SEQ ID NO: 184.

Clause 5. A culture medium comprising soy hydrolysate, yeast extract, and glucose, and wherein the medium comprises a *Clostridium difficile* bacterium derived from VPI

10 11186, wherein the bacterium lacks an endogenous polynucleotide encoding a toxin, wherein the bacterium comprises a polynucleotide encoding a mutant *C. difficile* toxin, wherein the bacterium further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter.

Clause 6. A method of culturing *Clostridium difficile* comprising culturing *C. difficile* in

15 a medium, wherein the medium comprises soy hydrolysate, yeast extract, and glucose, and wherein the medium comprises a *Clostridium difficile* bacterium derived from VPI 11186, wherein the bacterium lacks an endogenous polynucleotide encoding a toxin, wherein the bacterium comprises a polynucleotide encoding a mutant *C. difficile* toxin, wherein the bacterium further comprises a *Clostridium sporogenes* ferredoxin (fdx)

20 promoter.

Clause 7. A method of producing a *Clostridium difficile* toxin comprising culturing *C. difficile* in a medium under suitable conditions to produce a toxin, and isolating the toxin from the medium; wherein the medium comprises soy hydrolysate, yeast extract, and glucose, and wherein the medium comprises a *Clostridium difficile* bacterium derived

25 from VPI 11186, wherein the bacterium lacks an endogenous polynucleotide encoding a toxin, wherein the bacterium comprises a polynucleotide encoding a mutant *C. difficile* toxin, wherein the bacterium further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter.

CLAIMS

1. A culture medium comprising soy hydrolysate, yeast extract, and glucose as a carbon source, wherein the medium comprises a recombinant *Clostridium difficile* cell, wherein the cell lacks an endogenous polynucleotide encoding *C. difficile* toxin A, wherein the cell comprises a polynucleotide which encodes a polypeptide comprising the amino acid sequence set forth in any one of SEQ ID NOs: 1-8, 15, 17, 19, 21, 23, 25, 28-35, 82-761, and wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide.
2. The medium according to claim 1, further comprising polyethylene glycol 2000.
3. The medium according to claim 1, further comprising thiamphenicol.
4. The medium according to claim 1, wherein the cell is a recombinant *C. difficile* VPI 11186.
5. A method of culturing a *Clostridium difficile* cell comprising culturing the *C. difficile* cell in a medium, wherein the medium comprises soy hydrolysate, yeast extract, and glucose as a carbon source, and wherein the *Clostridium difficile* cell is a recombinant *C. difficile* VPI 11186, wherein the cell lacks an endogenous polynucleotide encoding *C. difficile* toxin A, wherein the cell comprises a polynucleotide encoding a polypeptide comprising the amino acid sequence set forth in any one of SEQ ID NOs: 1-8, 15, 17, 19, 21, 23, 25, 28-35, 82-761, wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide.
6. A method of producing a polypeptide comprising the amino acid sequence set forth in any one of SEQ ID NOs: 1-8, 15, 17, 19, 21, 23, 25, 28-35, 82-761, the method comprising culturing a *Clostridium difficile* cell in a medium under suitable conditions to produce the polypeptide, and isolating the polypeptide from the medium; wherein the medium comprises soy hydrolysate, yeast extract, and glucose as a carbon source, and wherein the *C. difficile* cell is a recombinant *C. difficile* VPI 11186, wherein the cell lacks an endogenous polynucleotide encoding *C. difficile* toxin A, wherein the cell comprises a polynucleotide encoding the polypeptide, wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide.

7. A method of producing a mutant *Clostridium difficile* toxin, the method comprising culturing a recombinant *C. difficile* cell in a medium under suitable conditions to produce a polypeptide comprising the amino acid sequence set forth in any one of SEQ ID NOS: 3, 4, 7, 28, 29, 32, 34, 36, 83, 84, 183, 196-212, 223-245, 250-253, 255-263, 276-323, 374-421, 472-567, 664-711, 5, 6, 8, 30, 31, 33, 35, 82, 85, 86, 184, 213-222, 246-249, 264-269, 324-373, 422-471, 568-663, 712-761, and 185-195, and isolating the polypeptide from the medium; wherein the medium comprises soy hydrolysate, yeast extract, and glucose as a carbon source, wherein the cell comprises a polynucleotide which encodes the polypeptide, wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide; and wherein the method further comprises contacting the polypeptide with a crosslinker comprising 1 -ethyl-3-(3- dimethylaminopropyl) carbodiimide) (EDC), thereby producing the mutant *C. difficile* toxin.
8. A method of producing a mutant *Clostridium difficile* toxin, the method comprising culturing a recombinant *C. difficile* cell in a medium under suitable conditions to produce a polypeptide comprising the amino acid sequence set forth in any one of SEQ ID NOS: 3, 4, 7, 28, 29, 32, 34, 36, 83, 84, 183, 196-212, 223-245, 250-253, 255-263, 276-323, 374-421, 472-567, 664-711, 5, 6, 8, 30, 31, 33, 35, 82, 85, 86, 184, 213-222, 246-249, 264-269, 324-373, 422-471, 568-663, 712-761, and 185-195, and isolating the polypeptide from the medium; wherein the medium comprises soy hydrolysate, yeast extract, and glucose as a carbon source, wherein the cell comprises a polynucleotide which encodes the polypeptide, wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide; and wherein the method further comprises contacting the polypeptide with a crosslinker comprising N- Hydroxysuccinimide (NHS), thereby producing the mutant *C. difficile* toxin.
9. The method according to any one of claims 5-8, wherein the yeast extract comprises HY-
YEAST.
10. The method according to any one of claims 5-9, wherein the medium further comprises polyethylene glycol 2000.

11. The method according to any one of claims 5-10, wherein the medium further comprises thiamphenicol.
12. The method according to any one of claims 7-11, wherein the cell is a recombinant *C. difficile* VPI 11186 (ATCC 700057TM).
13. The method according to any one of claims 7-11, wherein the cell is a recombinant *C. difficile* 1351 (ATCC 43593TM).
14. The method according to any one of claims 7-11, wherein the cell is a recombinant *C. difficile* 3232 (ATCC BAA-1801TM).
15. The method according to any one of claims 7-11, wherein the cell is a recombinant *C. difficile* 7322 (ATCC 43601TM).
16. The method according to any one of claims 7-11, wherein the cell is a recombinant *C. difficile* 5036 (ATCC 43603TM).
17. The method according to any one of claims 7-11, wherein the cell is a recombinant *C. difficile* 4811 (ATCC 43602TM).
18. The medium according to claim 1, wherein the cell is a recombinant *C. difficile* 1351 (ATCC 43593TM).
19. The medium according to claim 1, wherein the cell is a recombinant *C. difficile* 3232 (ATCC BAA-1801TM).
20. The medium according to claim 1, wherein the cell is a recombinant *C. difficile* 7322 (ATCC 43601TM).
21. The medium according to claim 1, wherein the cell is a recombinant *C. difficile* 5036 (ATCC 43603TM).
22. The medium according to claim 1, wherein the cell is a recombinant *C. difficile* 4811 (ATCC 43602TM).

23. A culture medium comprising soy hydrolysate, yeast extract, and glucose as a carbon source, wherein the medium comprises a recombinant *Clostridium difficile* cell, wherein the cell lacks an endogenous polynucleotide encoding *C. difficile* toxin A, wherein the cell comprises a polynucleotide which encodes a polypeptide comprising the amino acid sequence set forth in SEQ ID NO: 4, and wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide.
24. The medium according to claim 23, further comprising polyethylene glycol 2000.
25. The medium according to any one of claims 23-24, further comprising thiamphenicol.
26. The medium according to any one of claims 23-25, wherein the cell is a recombinant *C. difficile* VPI 11186.
27. The medium according to any one of claims 23-25, wherein the cell is a recombinant *C. difficile* 1351 (ATCC 43593TM).
28. The medium according to any one of claims 23-25, wherein the cell is a recombinant *C. difficile* 3232 (ATCC BAA-1801TM).
29. The medium according to any one of claims 23-25, wherein the cell is a recombinant *C. difficile* 7322 (ATCC 43601TM).
30. The medium according to any one of claims 23-25, wherein the cell is a recombinant *C. difficile* 5036 (ATCC 43603TM).
31. The medium according to any one of claims 23-25, wherein the cell is a recombinant *C. difficile* 4811 (ATCC 43602TM).
32. A culture medium comprising soy hydrolysate, yeast extract, and glucose as a carbon source, wherein the medium comprises a recombinant *Clostridium difficile* cell, wherein the cell lacks an endogenous polynucleotide encoding *C. difficile* toxin A, wherein the cell comprises a polynucleotide which encodes a polypeptide comprising the amino acid sequence set forth in SEQ ID NO: 6, and wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide.

33. The medium according to claim 32, further comprising polyethylene glycol 2000.
34. The medium according to any one of claims 32-33, further comprising thiamphenicol.
35. The medium according to any one of claims 32-34, wherein the cell is a recombinant *C. difficile* VPI 11186.
36. The medium according to any one of claims 32-34, wherein the cell is a recombinant *C. difficile* 1351 (ATCC 43593TM).
37. The medium according to any one of claims 32-34, wherein the cell is a recombinant *C. difficile* 3232 (ATCC BAA-1801TM).
38. The medium according to any one of claims 32-34, wherein the cell is a recombinant *C. difficile* 7322 (ATCC 43601TM).
39. The medium according to any one of claims 32-34, wherein the cell is a recombinant *C. difficile* 5036 (ATCC 43603TM).
40. The medium according to any one of claims 32-34, wherein the cell is a recombinant *C. difficile* 4811 (ATCC 43602TM).
41. A method of culturing a *Clostridium difficile* cell comprising culturing the *C. difficile* cell in a medium, wherein the medium comprises soy hydrolysate, yeast extract, and glucose as a carbon source, and wherein the *Clostridium difficile* cell is a recombinant *C. difficile* VPI 11186, wherein the cell lacks an endogenous polynucleotide encoding *C. difficile* toxin A, wherein the cell comprises a polynucleotide encoding a polypeptide comprising the amino acid sequence set forth in SEQ ID NO: 4, wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide.
42. A method of producing a polypeptide comprising the amino acid sequence set forth in SEQ ID NO: 4, the method comprising culturing a *Clostridium difficile* cell in a medium under suitable conditions to produce the polypeptide, and isolating the polypeptide from the medium; wherein the medium comprises soy hydrolysate, yeast extract, and glucose as a carbon source, and wherein the *C. difficile* cell is a recombinant *C. difficile* VPI 11186,

wherein the cell lacks an endogenous polynucleotide encoding *C. difficile* toxin A, wherein the cell comprises a polynucleotide encoding the polypeptide, wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide.

43. A method of producing a mutant *Clostridium difficile* toxin, the method comprising culturing a recombinant *C. difficile* cell in a medium under suitable conditions to produce a polypeptide comprising the amino acid sequence set forth in SEQ ID NO: 4, and isolating the polypeptide from the medium; wherein the medium comprises soy hydrolysate, yeast extract, and glucose as a carbon source, wherein the cell comprises a polynucleotide which encodes the polypeptide, wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide; and wherein the method further comprises contacting the polypeptide with 1 -ethyl-3-(3-dimethylaminopropyl) carbodiimide) (EDC), thereby producing the mutant *C. difficile* toxin.
44. A method of producing a mutant *Clostridium difficile* toxin, the method comprising culturing a recombinant *C. difficile* cell in a medium under suitable conditions to produce a polypeptide comprising the amino acid sequence set forth in SEQ ID NO: 4, and isolating the polypeptide from the medium; wherein the medium comprises soy hydrolysate, yeast extract, and glucose as a carbon source, wherein the cell comprises a polynucleotide which encodes the polypeptide, wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide; and wherein the method further comprises contacting the polypeptide with N- Hydroxysuccinimide (NHS), thereby producing the mutant *C. difficile* toxin.
45. The method according to any one of claims 41-44, wherein the yeast extract comprises HY-YEST.
46. The method according to any one of claims 41-45, wherein the medium further comprises polyethylene glycol 2000.

47. The method according to any one of claims 41-46, wherein the medium further comprises thiamphenicol.
48. The method according to any one of claims 43-47, wherein the cell is a recombinant *C. difficile* VPI 11186 (ATCC 700057TM).
49. The method according to any one of claims 43-47, wherein the cell is a recombinant *C. difficile* 1351 (ATCC 43593TM).
50. The method according to any one of claims 43-47, wherein the cell is a recombinant *C. difficile* 3232 (ATCC BAA-1801TM).
51. The method according to any one of claims 43-47, wherein the cell is a recombinant *C. difficile* 7322 (ATCC 43601TM).
52. The method according to any one of claims 43-47, wherein the cell is a recombinant *C. difficile* 5036 (ATCC 43603TM).
53. The method according to any one of claims 43-47, wherein the cell is a recombinant *C. difficile* 4811 (ATCC 43602TM).
54. A method of culturing a *Clostridium difficile* cell comprising culturing the *C. difficile* cell in a medium, wherein the medium comprises soy hydrolysate, yeast extract, and glucose as a carbon source, and wherein the *Clostridium difficile* cell is a recombinant *C. difficile* VPI 11186, wherein the cell lacks an endogenous polynucleotide encoding *C. difficile* toxin A, wherein the cell comprises a polynucleotide encoding a polypeptide comprising the amino acid sequence set forth in SEQ ID NO: 6, wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide.
55. A method of producing a polypeptide comprising the amino acid sequence set forth in SEQ ID NO: 6, the method comprising culturing a *Clostridium difficile* cell in a medium under suitable conditions to produce the polypeptide, and isolating the polypeptide from the medium; wherein the medium comprises soy hydrolysate, yeast extract, and glucose as a carbon source, and wherein the *C. difficile* cell is a recombinant *C. difficile* VPI 11186, wherein the cell lacks an endogenous polynucleotide encoding *C. difficile* toxin A, wherein

the cell comprises a polynucleotide encoding the polypeptide, wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide.

56. A method of producing a mutant *Clostridium difficile* toxin, the method comprising culturing a recombinant *C. difficile* cell in a medium under suitable conditions to produce a polypeptide comprising the amino acid sequence set forth in SEQ ID NO: 6, and isolating the polypeptide from the medium; wherein the medium comprises soy hydrolysate, yeast extract, and glucose as a carbon source, wherein the cell comprises a polynucleotide which encodes the polypeptide, wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide; and wherein the method further comprises contacting the polypeptide with 1 -ethyl-3-(3-dimethylaminopropyl) carbodiimide) (EDC), thereby producing the mutant *C. difficile* toxin.
57. A method of producing a mutant *Clostridium difficile* toxin, the method comprising culturing a recombinant *C. difficile* cell in a medium under suitable conditions to produce a polypeptide comprising the amino acid sequence set forth in SEQ ID NO: 6, and isolating the polypeptide from the medium; wherein the medium comprises soy hydrolysate, yeast extract, and glucose as a carbon source, wherein the cell comprises a polynucleotide which encodes the polypeptide, wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide; and wherein the method further comprises contacting the polypeptide with N- Hydroxysuccinimide (NHS), thereby producing the mutant *C. difficile* toxin.
58. The method according to any one of claims 54-57, wherein the yeast extract comprises HY-
YEAST.
59. The method according to any one of claims 54-58, wherein the medium further comprises polyethylene glycol 2000.
60. The method according to any one of claims 54-59, wherein the medium further comprises thiamphenicol.

61. The method according to any one of claims 54-60, wherein the cell is a recombinant *C. difficile* VPI 11186 (ATCC 700057TM).
62. The method according to any one of claims 54-60, wherein the cell is a recombinant *C. difficile* 1351 (ATCC 43593TM).
63. The method according to any one of claims 54-60, wherein the cell is a recombinant *C. difficile* 3232 (ATCC BAA-1801TM).
64. The method according to any one of claims 54-60, wherein the cell is a recombinant *C. difficile* 7322 (ATCC 43601TM).
65. The method according to any one of claims 54-60, wherein the cell is a recombinant *C. difficile* 5036 (ATCC 43603TM).
66. The method according to any one of claims 54-60, wherein the cell is a recombinant *C. difficile* 4811 (ATCC 43602TM).
67. The method according to claim 7 or 8, wherein the crosslinker comprises a combination of 1 -ethyl-3-(3- dimethylaminopropyl) carbodiimide) (EDC) and N- Hydroxysuccinimide (NHS).
68. A method of producing a mutant *Clostridium difficile* toxin, the method comprising culturing a recombinant *C. difficile* cell in a medium under suitable conditions to produce a polypeptide comprising the amino acid sequence set forth in any one of SEQ ID NOs: 1, 15, 17, 19, 87, 88-109, 134-140, 148-157, 168-172, 178, and 180, and isolating the polypeptide from the medium; wherein the medium comprises soy hydrolysate, yeast extract, and glucose as a carbon source, wherein the cell comprises a polynucleotide which encodes the polypeptide, wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide; and wherein the method further comprises contacting the polypeptide with a crosslinker comprising 1 -ethyl-3-(3- dimethylaminopropyl) carbodiimide) (EDC), thereby producing the mutant *C. difficile* toxin.

69. A method of producing a mutant *Clostridium difficile* toxin, the method comprising culturing a recombinant *C. difficile* cell in a medium under suitable conditions to produce a polypeptide comprising the amino acid sequence set forth in any one of SEQ ID NOs: 2, 21, 23, 25, 110-133, 141-167, 173-179, 181, and 182, and isolating the polypeptide from the medium; wherein the medium comprises soy hydrolysate, yeast extract, and glucose as a carbon source, wherein the cell comprises a polynucleotide which encodes the polypeptide, wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide; and wherein the method further comprises contacting the polypeptide with a crosslinker comprising 1 -ethyl-3-(3-dimethylaminopropyl) carbodiimide) (EDC), thereby producing the mutant *C. difficile* toxin.
70. A method of producing a mutant *Clostridium difficile* toxin, the method comprising culturing a recombinant *C. difficile* cell in a medium under suitable conditions to produce a polypeptide comprising the amino acid sequence set forth in any one of SEQ ID NOs: 1, 15, 17, 19, 87, 88-109, 134-140, 148-157, 168-172, 178, and 180, and isolating the polypeptide from the medium; wherein the medium comprises soy hydrolysate, yeast extract, and glucose as a carbon source, wherein the cell comprises a polynucleotide which encodes the polypeptide, wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide; and wherein the method further comprises contacting the polypeptide with a crosslinker comprising N-Hydroxysuccinimide (NHS), thereby producing the mutant *C. difficile* toxin.
71. A method of producing a mutant *Clostridium difficile* toxin, the method comprising culturing a recombinant *C. difficile* cell in a medium under suitable conditions to produce a polypeptide comprising the amino acid sequence set forth in any one of SEQ ID NOs: 2, 21, 23, 25, 110-133, 141-167, 173-179, 181, and 182, and isolating the polypeptide from the medium; wherein the medium comprises soy hydrolysate, yeast extract, and glucose as a carbon source, wherein the cell comprises a polynucleotide which encodes the polypeptide, wherein the cell further comprises a *Clostridium sporogenes* ferredoxin (fdx) promoter that drives expression of the polypeptide; and wherein the method further

comprises contacting the polypeptide with a crosslinker comprising N-Hydroxysuccinimide (NHS), thereby producing the mutant *C. difficile* toxin.

72. The method according to any one of claims 68-71, wherein the crosslinker comprises a combination of 1 -ethyl-3-(3- dimethylaminopropyl) carbodiimide) (EDC) and N-Hydroxysuccinimide (NHS).

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FIG. 1A

MSLISKEELIKLAYSIRPRENEYKTILTNLDEYNKLTNNNNENKYLQLKK 50
 MSLISKEELIKLAYSIRPRENEYKTILTNLDEYNKLTNNNNENKYLQLKK 50
 MSLISKEELIKLAYSIRPRENEYKTILTNLDEYNKLTNNNNENKYLQLKK 50
 MSLISKEELIKLAYSIRPRENEYKTILTNLDEYNKLTNNNNENKYLQLKK 50
 MSLISKEELIKLAYSIRPRENEYKTILTNLDEYNKLTNNNNENKYLQLKK 50

LNESIDVFMNKYKTSSRNRALSNLKKDILKEVILIKNSNTSPVEKNLHFV 100
 LNESIDVFMNKYKTSSRNRALSNLKKDILKEVILIKNSNTSPVEKNLHFV 100
 LNESIDVFMNKYKTSSRNRALSNLKKDILKEVILIKNSNTSPVEKNLHFV 100
 LNESIDVFMNKYKNSSRNRALSNLKKDILKEVILIKNSNTSPVEKNLHFV 100
 LNESIDVFMNKYKNSSRNRALSNLKKDILKEVILIKNSNTSPVEKNLHFV 100

WIGGEVSDIALEYIKQWADINAEYNIKLWYDSEAFLVNTLKKAIIVESSTT 150
 WIGGEVSDIALEYIKQWADINAEYNIKLWYDSEAFLVNTLKKAIIVESSTT 150
 WIGGEVSDIALEYIKQWADINAEYNIKLWYDSEAFLVNTLKKAIIVESSTT 150
 WIGGEVSDIALEYIKQWADINAEYNIKLWYDSEAFLVNTLKKAIIVESSTT 150
 WIGGEVSDIALEYIKQWADINAEYNIKLWYDSEAFLVNTLKKAIIVESSTT 150

EALQLLEEEIIONQPQFDNMKFYKKRMEFIYDRQKRFINYYKSQINKPTVPT 200
 EALQLLEEEIIONQPQFDNMKFYKKRMEFIYDRQKRFINYYKSQINKPTVPT 200
 EALQLLEEEIIONQPQFDNMKFYKKRMEFIYDRQKRFINYYKSQINKPTVPT 200
 EALQLLEEEIIONQPQFDNMKFYKKRMEFIYDRQKRFINYYKSQINKPTVPT 200
 EALQLLEEEIIONQPQFDNMKFYKKRMEFIYDRQKRFINYYKSQINKPTVPT 200

IDDIIKSHLVSEYNRDETVLESYRTNSLRKINSNHGIDIRANSLFTEQEL 250
 IDDIIKSHLVSEYNRDETVLESYRTNSLRKINSNHGIDIRANSLFTEQEL 250
 IDDIIKSHLVSEYNRDETVLESYRTNSLRKINSNHGIDIRANSLFTEQEL 250
 IDDIIKSHLVSEYNRDETLLESYRTNSLRKINSNHGIDIRANSLFTEQEL 250
 IDDIIKSHLVSEYNRDETLLESYRTNSLRKINSNHGIDIRANSLFTEQEL 250

LNIYSQELLNRGNLAAASDIVRLLALKNFGGVYLDVDMLPGIHSDFLFTI 300
 LNIYSQELLNRGNLAAASDIVRLLALKNFGGVYLDVDMLPGIHSDFLFTI 300
 LNIYSQELLNRGNLAAASDIVRLLALKNFGGVYLDVDMLPGIHSDFLFTI 300
 LNIYSQELLNRGNLAAASDIVRLLALKNFGGVYLDVDMLPGIHSDFLFTI 300
 LNIYSQELLNRGNLAAASDIVRLLALKNFGGVYLDVDMLPGIHSDFLFTI 300

SRPSSIGLDRWEMIKLEAIMKYKKYINNYTSENFDKLDQQLKDNFKLIIE 350
 SRPSSIGLDRWEMIKLEAIMKYKKYINNYTSENFDKLDQQLKDNFKLIIE 350
 SRPSSIGLDRWEMIKLEAIMKYKKYINNYTSENFDKLDQQLKDNFKLIIE 350
 PRPSSIGLDRWEMIKLEAIMKYKKYINNYTSENFDKLDQQLKDNFKLIIE 350
 PRPSSIGLDRWEMIKLEAIMKYKKYINNYTSENFDKLDQQLKDNFKLIIE 350

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FIG. 1B

SKSEKSEIFSKLENLNVSDEIKIAFALGSVINQALISKQGSYLTNLVIE 400
 SKSEKSEIFSKLENLNVSDEIKIAFALGSVINQALISKQGSYLTNLVIE 400
 SKSEKSEIFSKLENLNVSDEIKIAFALGSVINQALISKQGSYLTNLVIE 400
 SKSEKSEIFSKLENLNVSDEIKIAFALGSVINQALISKQGSYLTNLVIE 400
 SKSEKSEIFSKLENLNVSDEIKIAFALGSVINQALISKQGSYLTNLVIE 400

QVKNRYQFLNQHLPNAIESDNNFTDTKIFHDSLFLNSATAENSMFLTKIA 450
 QVKNRYQFLNQHLPNAIESDNNFTDTKIFHDSLFLNSATAENSMFLTKIA 450
 QVKNRYQFLNQHLPNAIESDNNFTDTKIFHDSLFLNSATAENSMFLTKIA 450
 QVKNRYQFLNQHLPNAIESDNNFTDTKIFHDSLFLNSATAENSMFLTKIA 450
 QVKNRYQFLNQHLPNAIESDNNFTDTKIFHDSLFLNSATAENSMFLTKIA 450

PYLQVGMPEARSTISLSGPAYASAYYDFINLQENTIEKTLKASDLIEF 500
 PYLQVGMPEARSTISLSGPAYASAYYDFINLQENTIEKTLKASDLIEF 500
 PYLQVGMPEARSTISLSGPAYASAYYDFINLQENTIEKTLKASDLIEF 500
 PYLQVGMPEARSTISLSGPAYASAYYDFINLQENTIEKTLKASDLIEF 500
 PYLQVGMPEARSTISLSGPAYASAYYDFINLQENTIEKTLKASDLIEF 500

KFPENNLSQLTEQEINSLWSFDQASAKYQFEKYVRDYTGGSISEDNGVDF 550
 KFPENNLSQLTEQEINSLWSFDQASAKYQFEKYVRDYTGGSISEDNGVDF 550
 KFPENNLSQLTEQEINSLWSFDQASAKYQFEKYVRDYTGGSISEDNGVDF 550
 KFPENNLSQLTEQEINSLWSFDQASAKYQFEKYVRDYTGGSISEDNGVDF 550
 KFPENNLSQLTEQEINSLWSFDQASAKYQFEKYVRDYTGGSISEDNGVDF 550

NKNTALDKNYLLNNKIPSNVVEAGSKNYVHYIIQLQGDDISYEATCNLF 600
 NKNTALDKNYLLNNKIPSNVVEAGSKNYVHYIIQLQGDDISYEATCNLF 600
 NKNTALDKNYLLNNKIPSNVVEAGSKNYVHYIIQLQGDDISYEATCNLF 600
 NKNTALDKNYLLNNKIPSNVVEAGSKNYVHYIIQLQGDDISYEATCNLF 600
 NKNTALDKNYLLNNKIPSNVVEAGSKNYVHYIIQLQGDDISYEATCNLF 600

SKNPKNSIIIQRNMNESAKSYFLSDDGESILELNKYRIPERLKNKEVKV 650
 SKNPKNSIIIQRNMNESAKSYFLSDDGESILELNKYRIPERLKNKEVKV 650
 SKNPKNSIIIQRNMNESAKSYFLSDDGESILELNKYRIPERLKNKEVKV 650
 SKNPKNSIIIQRNMNESAKSYFLSDDGESILELNKYRIPERLKNKEVKV 650
 SKNPKNSIIIQRNMNESAKSYFLSDDGESILELNKYRIPERLKNKEVKV 650

TFIGHGKDEFNTSEFARLSVDSLNEISSFLDTIKLDISPKNVEVNLLGC 700
 TFIGHGKDEFNTSEFARLSVDSLNEISSFLDTIKLDISPKNVEVNLLGA 700
 TFIGHGKDEFNTSEFARLSVDSLNEISSFLDTIKLDISPKNVEVNLLGC 700
 TFIGHGKDEFNTSEFARLSVDSLNEISSFLDTIKLDISPKNVEVNLLGC 700
 TFIGHGKDEFNTSEFARLSVDSLNEISSFLDTIKLDISPKNVEVNLLGC 700

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FIG. 1C

NMFSYDFNVEETYPGKLLSIMDKITSTLPDVNKNSITIGANQYEVRI 750
 NMFSYDFNVEETYPGKLLSIMDKITSTLPDVNKNSITIGANQYEVRI 750
 NMFSYDFNVEETYPGKLLSIMDKITSTLPDVNKNSITIGANQYEVRI 750
 NMFSYDFNVEETYPGKLLSIMDKITSTLPDVNKDSITIGANQYEVRI 750
 NMFSYDFNVEETYPGKLLSIMDKITSTLPDVNKDSITIGANQYEVRI 750

EGRKELLAHSGKWINKEEAIMSDLSSKEYIFFDSIDNKLAKSKNIPGLA 800
 EGRKELLAHSGKWINKEEAIMSDLSSKEYIFFDSIDNKLAKSKNIPGLA 800
 EGRKELLAHSGKWINKEEAIMSDLSSKEYIFFDSIDNKLAKSKNIPGLA 800
 EGRKELLAHSGKWINKEEAIMSDLSSKEYIFFDSIDNKLAKSKNIPGLA 800
 EGRKELLAHSGKWINKEEAIMSDLSSKEYIFFDSIDNKLAKSKNIPGLA 800

SISEDIKTLLLDASVPDTKFILNNLKLNISSIGDYIYYEKLEPVKNII 850
 SISEDIKTLLLDASVPDTKFILNNLKLNISSIGDYIYYEKLEPVKNII 850
 SISEDIKTLLLDASVPDTKFILNNLKLNISSIGDYIYYEKLEPVKNII 850
 SISEDIKTLLLDASVPDTKFILNNLKLNISSIGDYIYYEKLEPVKNII 850
 SISEDIKTLLLDASVPDTKFILNNLKLNISSIGDYIYYEKLEPVKNII 850

HNSIDDLIDEFNLLLENVSDELYELKKLNNLDEKYLISFEDISKNNSTYSV 900
 HNSIDDLIDEFNLLLENVSDELYELKKLNNLDEKYLISFEDISKNNSTYSV 900
 HNSIDDLIDEFNLLLENVSDELYELKKLNNLDEKYLISFEDISKNNSTYSV 900
 HNSIDDLIDEFNLLLENVSDELYELKKLNNLDEKYLISFEDISKNNSTYSV 900
 HNSIDDLIDEFNLLLENVSDELYELKKLNNLDEKYLISFEDISKNNSTYSV 900

RFINKSNGESVYVETEKEIFSKYSEHITKEISTIKNSIITDVNGNLLDNI 950
 RFINKSNGESVYVETEKEIFSKYSEHITKEISTIKNSIITDVNGNLLDNI 950
 RFINKSNGESVYVETEKEIFSKYSEHITKEISTIKNSIITDVNGNLLDNI 950
 RFINKSNGESVYVETEKEIFSKYSEHITKEISTIKNSIITDVNGNLLDNI 950
 RFINKSNGESVYVETEKEIFSKYSEHITKEISTIKNSIITDVNGNLLDNI 950

QLDHTSQVNTLNAAFFIQSLIDYSSNKDVNLSTSVKVQLYAQLFSTGL 1000
 QLDHTSQVNTLNAAFFIQSLIDYSSNKDVNLSTSVKVQLYAQLFSTGL 1000
 QLDHTSQVNTLNAAFFIQSLIDYSSNKDVNLSTSVKVQLYAQLFSTGL 1000
 QLDHTSQVNTLNAAFFIQSLIDYSSNKDVNLSTSVKVQLYAQLFSTGL 1000
 QLDHTSQVNTLNAAFFIQSLIDYSSNKDVNLSTSVKVQLYAQLFSTGL 1000

NTIYDSIQLVNLISNAVNDTINVLPITIEGIPIVSTILDGINLGAAIKEL 1050
 NTIYDSIQLVNLISNAVNDTINVLPITIEGIPIVSTILDGINLGAAIKEL 1050
 NTIYDSIQLVNLISNAVNDTINVLPITIEGIPIVSTILDGINLGAAIKEL 1050
 NTIYDSIQLVNLISNAVNDTINVLPITIEGIPIVSTILDGINLGAAIKEL 1050
 NTIYDSIQLVNLISNAVNDTINVLPITIEGIPIVSTILDGINLGAAIKEL 1050

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FIG. 1D

LDEHDPLLKELEAKVGVLAINMSLSIAATVASIVGIGAEVTIFLLPIAG 1100
 LDEHDPLLKELEAKVGVLAINMSLSIAATVASIVGIGAEVTIFLLPIAG 1100
 LDEHDPLLKELEAKVGVLAINMSLSIAATVASIVGIGAEVTIFLLPIAG 1100
 LDEHDPLLKELEAKVGVLAINMSLSIAATVASIVGIGAEVTIFLLPIAG 1100
 LDEHDPLLKELEAKVGVLAINMSLSIAATVASIVGIGAEVTIFLLPIAG 1100

ISAGIPSLVNNELILHDKATSVVNYFNHLSESKKYGPLKTEDDKILVPID 1150
 ISAGIPSLVNNELILHDKATSVVNYFNHLSESKKYGPLKTEDDKILVPID 1150
 ISAGIPSLVNNELILHDKATSVVNYFNHLSESKKYGPLKTEDDKILVPID 1150
 ISAGIPSLVNNELILHDKATSVVNYFNHLSESKYGPLKTEDDKILVPID 1150
 ISAGIPSLVNNELILHDKATSVVNYFNHLSESKYGPLKTEDDKILVPID 1150

DLVISEIDFNNNSIKLGTNCNILAMEGGSGHTVTGNIDHFFSSPSISSHIP 1200
 DLVISEIDFNNNSIKLGTNCNILAMEGGSGHTVTGNIDHFFSSPSISSHIP 1200
 DLVISEIDFNNNSIKLGTNCNILAMEGGSGHTVTGNIDHFFSSPSISSHIP 1200
 DLVISEIDFNNNSIKLGTNCNILAMEGGSGHTVTGNIDHFFSSPSISSHIP 1200
 DLVISEIDFNNNSIKLGTNCNILAMEGGSGHTVTGNIDHFFSSPSISSHIP 1200

SLSIYSAIGIETENLDFSKKIMMLPNAPSrvFWWETGAVPGLRSLENDGT 1250
 SLSIYSAIGIETENLDFSKKIMMLPNAPSrvFWWETGAVPGLRSLENDGT 1250
 SLSIYSAIGIETENLDFSKKIMMLPNAPSrvFWWETGAVPGLRSLENDGT 1250
 SLSVYSAIGIKTENLDFSKKIMMLPNAPSrvFWWETGAVPGLRSLENNGT 1250
 SLSVYSAIGIKTENLDFSKKIMMLPNAPSrvFWWETGAVPGLRSLENNGT 1250
 ;**;*****

RLLDSIRDLYPGKFYWRFYAFFDYAITTLKPVYEDTNIKIKLDKDTRNFI 1300
 RLLDSIRDLYPGKFYWRFYAFFDYAITTLKPVYEDTNIKIKLDKDTRNFI 1300
 RLLDSIRDLYPGKFYWRFYAFFDYAITTLKPVYEDTNIKIKLDKDTRNFI 1300
 KLLDSIRDLYPGKFYWRFYAFFDYAITTLKPVYEDNTKIKLDKDTRNFI 1300
 KLLDSIRDLYPGKFYWRFYAFFDYAITTLKPVYEDNTKIKLDKDTRNFI 1300
 :*****

MPTITTNEIRNKLSYSFDGAGGTYSLLLSSYPISTNINLSKDDLWIFNID 1350
 MPTITTNEIRNKLSYSFDGAGGTYSLLLSSYPISTNINLSKDDLWIFNID 1350
 MPTITTNEIRNKLSYSFDGAGGTYSLLLSSYPISTNINLSKDDLWIFNID 1350
 MPTITTDEIRNKLSYSFDGAGGTYSLLLSSYPISMNINLSKDDLWIFNID 1350
 MPTITTDEIRNKLSYSFDGAGGTYSLLLSSYPISMNINLSKDDLWIFNID 1350
 *****;*****

NEVREISIENGTIKKGKLIKDVLSKIDINKNKLIIIGNQTIDFSGDIDNKD 1400
 NEVREISIENGTIKKGKLIKDVLSKIDINKNKLIIIGNQTIDFSGDIDNKD 1400
 NEVREISIENGTIKKGKLIKDVLSKIDINKNKLIIIGNQTIDFSGDIDNKD 1400
 NEVREISIENGTIKKGNLIEDVLSKIDINKNKLIIIGNQTIDFSGDIDNKD 1400
 NEVREISIENGTIKKGNLIEDVLSKIDINKNKLIIIGNQTIDFSGDIDNKD 1400
 *****;*****

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FIG. 1E

RYIFLTCELDDKISLIETNLVAKSYSLLLGDKNYLISNLSNIIEKINT 1450
 RYIFLTCELDDKISLIETNLVAKSYSLLLGDKNYLISNLSNIIEKINT 1450
 RYIFLTCELDDKISLIETNLVAKSYSLLLGDKNYLISNLSNTIEKINT 1450
 RYIFLTCELDDKISLIETNLVAKSYSLLLGDKNYLISNLSNTIEKINT 1450
 RYIFLTCELDDKISLIETNLVAKSYSLLLGDKNYLISNLSNTIEKINT 1450

LGLDSKNIAINYTDESNNKYFGAISKTSQKSIIHYKKDSKNILEFYNDST 1500
 LGLDSKNIAINYTDESNNKYFGAISKTSQKSIIHYKKDSKNILEFYNDST 1500
 LGLDSKNIAINYTDESNNKYFGAISKTSQKSIIHYKKDSKNILEFYNDST 1500
 LGLDSKNIAINYTDESNNKYFGAISKTSQKSIIHYKKDSKNILEFYNGST 1500
 LGLDSKNIAINYTDESNNKYFGAISKTSQKSIIHYKKDSKNILEFYNGST 1500

LEFNSKDFIAEDINVFMKDDINTITGKYVVDNNNTDKSIDFSISLVSKNQV 1550
 LEFNSKDFIAEDINVFMKDDINTITGKYVVDNNNTDKSIDFSISLVSKNQV 1550
 LEFNSKDFIAEDINVFMKDDINTITGKYVVDNNNTDKSIDFSISLVSKNQV 1550
 LEFNSKDFIAEDINVFMKDDINTITGKYVVDNNNTDKSIDFSISLVSKNQV 1550

KVNGLYLNESVYSSYLDVFVKNSDGHHNTSNFMNLFLDNISFWKLFGFENI 1600
 KVNGLYLNESVYSSYLDVFVKNSDGHHNTSNFMNLFLDNISFWKLFGFENI 1600
 KVNGLYLNESVYSSYLDVFVKNSDGHHNTSNFMNLFLDNISFWKLFGFENI 1600
 KVNGLYLNESVYSSYLDVFVKNSDGHHNTSNFMNLFLNNISFWKLFGFENI 1600
 KVNGLYLNESVYSSYLDVFVKNSDGHHNTSNFMNLFLNNISFWKLFGFENI 1600

NFVIDKYFTLVGKTNLGYVEFICDNNKNIDIFYFGEWKTSSSKSTIFSGNG 1650
 NFVIDKYFTLVGKTNLGYVEFICDNNKNIDIFYFGEWKTSSSKSTIFSGNG 1650
 NFVIDKYFTLVGKTNLGYVEFICDNNKNIDIFYFGEWKTSSSKSTIFSGNG 1650
 NFVIDKYFTLVGKTNLGYVEFICDNNKNIDIFYFGEWKTSSSKSTIFSGNG 1650
 NFVIDKYFTLVGKTNLGYVEFICDNNKNIDIFYFGEWKTSSSKSTIFSGNG 1650

RNVVVEPIYNPDTGEDISTSLSDFSYEPLYGIDRYINKVLIAPDLYTSLIN 1700
 RNVVVEPIYNPDTGEDISTSLSDFSYEPLYGIDRYINKVLIAPDLYTSLIN 1700
 RNVVVEPIYNPDTGEDISTSLSDFSYEPLYGIDRYINKVLIAPDLYTSLIN 1700
 RNVVVEPIYNPDTGEDISTSLSDFSYEPLYGIDRYINKVLIAPDLYTSLIN 1700
 RNVVVEPIYNPDTGEDISTSLSDFSYEPLYGIDRYINKVLIAPDLYTSLIN 1700

INTNYYSEYYPEIIIVLPNTFHKKVNIINLDSSSFYKWSTEGSDFILVR 1750
 INTNYYSEYYPEIIIVLPNTFHKKVNIINLDSSSFYKWSTEGSDFILVR 1750
 INTNYYSEYYPEIIIVLPNTFHKKVNIINLDSSSFYKWSTEGSDFILVR 1750
 INTNYYSEYYPEIIIVLPNTFHKKVNIINLDSSSFYKWSTEGSDFILVR 1750
 INTNYYSEYYPEIIIVLPNTFHKKVNIINLDSSSFYKWSTEGSDFILVR 1750

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FIG. 1F

YLEESNKKILQKIRIKGILSNTQSFNKMISDFKDIKKLSLGYIMSNFKSF 1800
 YLEESNKKILQKIRIKGILSNTQSFNKMISDFKDIKKLSLGYIMSNFKSF 1800
 YLEESNKKILQKIRIKGILSNTQSFNKMISDFKDIKKLSLGYIMSNFKSF 1800
 YLEESNKKILQKIRIKGILSNTQSFNKMISDFKDIKKLSLGYIMSNFKSF 1800
 YLEESNKKILQKIRIKGILSNTQSFNKMISDFKDIKKLSLGYIMSNFKSF 1800

NSENELDRDHLGFKIIDNKTYYYDEDSKLVKGLININNSLFYFDPIEFNL 1850
 NSENELDRDHLGFKIIDNKTYYYDEDSKLVKGLININNSLFYFDPIEFNL 1850
 NSENELDRDHLGFKIIDNKTYYYDEDSKLVKGLININNSLFYFDPIEFNL 1850
 NSENELDRDHLGFKIIDNKTYYYDEDSKLVKGLININNSLFYFDPIESNL 1850
 NSENELDRDHLGFKIIDNKTYYYDEDSKLVKGLININNSLFYFDPIESNL 1850

VTGWQTINGKKYYFDINTGAALISYKIIINGKHFYFNNNDGVMQLGVFKGPD 1900
 VTGWQTINGKKYYFDINTGAALISYKIIINGKHFYFNNNDGVMQLGVFKGPD 1900
 VTGWQTINGKKYYFDINTGAALTSYKIIINGKHFYFNNNDGVMQLGVFKGPD 1900
 VTGWQTINGKKYYFDINTGAASTSYKIIINGKHFYFNNNGVMQLGVFKGPD 1900
 VTGWQTINGKKYYFDINTGAASTSYKIIINGKHFYFNNNGVMQLGVFKGPD 1900

GFEYFAPANTQNNNIEGQAIIVYQSKFLTLNGKKYYFDNDSKAVTGWRIN 1950
 GFEYFAPANTQNNNIEGQAIIVYQSKFLTLNGKKYYFDNDSKAVTGWRIN 1950
 GFEYFAPANTQNNNIEGQAIIVYQSKFLTLNGKKYYFDNNNSKAVTGWRIN 1950
 GFEYFAPANTQNNNIEGQAIIVYQSKFLTLNGKKYYFDNDNSKAVTGWRIN 1950
 GFEYFAPANTQNNNIEGQAIIVYQSKFLTLNGKKYYFDNDNSKAVTGWRIN 1950

NEKYYFNPNNAIAAVGLQVIDNNKYYFNPDATIISKGWQTVNGSRYYFDT 2000
 NEKYYFNPNNAIAAVGLQVIDNNKYYFNPDATIISKGWQTVNGSRYYFDT 2000
 NEKYYFNPNNAIAAVGLQVIDNNKYYFNPDATIISKGWQTVNGSRYYFDT 2000
 NEKYYFNPNNAIAAVGLQVIDNNKYYFNPDATIISKGWQTVNGSRYYFDT 2000
 NEKYYFNPNNAIAAVGLQVIDNNKYYFNPDATIISKGWQTVNGSRYYFDT 2000

DTAIAFNGYKTIDGKHFYFDSDCVVKIGVFSTSNGFEYFAPANTYNNIE 2050
 DTAIAFNGYKTIDGKHFYFDSDCVVKIGVFSTSNGFEYFAPANTYNNIE 2050
 DTAIAFNGYKTIDGKHFYFDSDCVVKIGVFSTSNGFEYFAPANTYNNIE 2050
 DTAIAFNGYKTIDGKHFYFDSDCVVKIGVFSGSNGFEYFAPANTYNNIE 2050
 DTAIAFNGYKTIDGKHFYFDSDCVVKIGVFSGSNGFEYFAPANTYNNIE 2050

GQAIVYQSKFLTLNGKKYYFDNNNSKAVTGWQTIDSKYYFNTNTAEATG 2100
 GQAIVYQSKFLTLNGKKYYFDNNNSKAVTGWQTIDSKYYFNTNTAEATG 2100
 GQAIVYQSKFLTLNGKKYYFDNNNSKAVTGQLTIDSKYYFNTNTAEATG 2100
 GQAIVYQSKFLTLNGKKYYFDNNNSKAVTGWQTIDSKYYFNTNTAEATG 2100
 GQAIVYQSKFLTLNGKKYYFDNNNSKAVTGWQTIDSKYYFNTNTAEATG 2100

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FIG. 1G

WQTIDGKKYYFNTNTAEEATGWQTIDGKKYYFNTNTAIASTGYTIINGKH 2150
 WQTIDGKKYYFNTNTAEEATGWQTIDGKKYYFNTNTAIASTGYTIINGKH 2150
 WQTIDGKKYYFNTNTAEEATGWQTIDGKKYYFNTNTAIASTGYTIINGKH 2150
 WQTIDGKKYYFNTNTAEEATGWQTIDGKKYYFNTNTSIASTGYTIINGKY 2150
 WQTIDGKKYYFNTNTAEEATGWQTIDGKKYYFNTNTSIASTGYTIINGKY 2150
 *****:*****:*****:*****:*****:*****:*****:*****:*****:

FYFNTDGIMQIGVFKGPNGFEYFAPANTDANNIEGQAILYQNEFLTLNGK 2200
 FYFNTDGIMQIGVFKGPNGFEYFAPANTDANNIEGQAILYQNEFLTLNGK 2200
 FYFNTDGIMQIGVFKGPNGFEYFAPANTDANNIEGQAILYQNEFLTLNGK 2200
 FYFNTDGIMQIGVFKVPNGFEYFAPANTHNNNIEGQAILYQNKFLTLNGK 2200
 FYFNTDGIMQIGVFKVPNGFEYFAPANTHNNNIEGQAILYQNKFLTLNGK 2200
 *****:*****:*****:*****:*****:*****:*****:*****:

KYYFGSDSKAVTGWIINNKYYFNPNNAAIAAIHLCTINNDKYYFSYDGI 2250
 KYYFGSDSKAVTGWIINNKYYFNPNNAAIAAIHLCTINNDKYYFSYDGI 2250
 KYYFGSDSKAVTGWIINNKYYFNPNNAAIAAIHLCTINNDKYYFSYDGI 2250
 KYYFGSDSKAIGWQTIDGKKYYFNPNNAAIAATHLCTINNDKYYFSYDGI 2250
 KYYFGSDSKAIGWQTIDGKKYYFNPNNAAIAATHLCTINNDKYYFSYDGI 2250
 *****:*****:*****:*****:*****:*****:*****:*****:

LQNGYITIERNNFYFDANNESKMTGVFKGPNGFEYFAPANTHNNNIEGQ 2300
 LQNGYITIERNNFYFDANNESKMTGVFKGPNGFEYFAPANTHNNNIEGQ 2300
 LQNGYITIERNNFYFDANNESKMTGVFKGPNGFEYFAPANTHNNNIEGQ 2300
 LQNGYITIERNNFYFDANNESKMTGVFKGPNGFEYFAPANTHNNNIEGQ 2300
 LQNGYITIERNNFYFDANNESKMTGVFKGPNGFEYFAPANTHNNNIEGQ 2300
 *****:*****:*****:*****:*****:*****:*****:*****:

AIVYQNKFLTLNGKKYYFDNDSSKAVTGWQTIDGKKYYFNLNTAEEATGWQ 2350
 AIVYQNKFLTLNGKKYYFDNDSSKAVTGWQTIDGKKYYFNLNTAEEATGWQ 2350
 AIVYQNKFLTLNGKKYYFDNDSSKAVTGWQTIDGKKYYFNLNTAEEATGWQ 2350
 AIVYQNKFLTLNGKKYYFDNDSSKAVTGWQTIDSKKYYFNLNTAVAVTGWQ 2350
 AIVYQNKFLTLNGKKYYFDNDSSKAVTGWQTIDSKKYYFNLNTAVAVTGWQ 2350
 *****:*****:*****:*****:*****:*****:*****:*****:

TIDGKKYYFNLNTAEEATGWQTIDGKKYYFNTNTFIASTGYSINGKHY 2400
 TIDGKKYYFNLNTAEEATGWQTIDGKKYYFNTNTFIASTGYSINGKHY 2400
 TIDGKKYYFNLNTAEEATGWQTIDGKKYYFNTNTFIASTGYSINGKHY 2400
 TIDGEKYYFNLNTAEEATGWQTIDGKRYYYFNTNTYIASTGYTIINGKHY 2400
 TIDGEKYYFNLNTAEEATGWQTIDGKRYYYFNTNTYIASTGYTIINGKHY 2400
 *****:*****:*****:*****:*****:*****:*****:

FNTDGIMQIGVFKGPNGFEYFAPANTHNNNIEGQAILYQNKFLTLNGKKY 2450
 FNTDGIMQIGVFKGPNGFEYFAPANTHNNNIEGQAILYQNKFLTLNGKKY 2450
 FNTDGIMQIGVFKGPNGFEYFAPANTDANNIEGQAILYQNKFLTLNGKKY 2450
 FNTDGIMQIGVFKGPNGFEYFAPANTHNNNIEGQAILYQNKFLTLNGKKY 2450
 FNTDGIMQIGVFKGPNGFEYFAPANTHNNNIEGQAILYQNKFLTLNGKKY 2450
 *****:*****:*****:*****:*****:*****:*****:

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FIG. 1H

YFGSDSKAVTGLRTIDGKKYYFNTNTAVAVTGWQTINGKKYYFNTNTSIA 2500
 YFGSDSKAVTGLRTIDGKKYYFNTNTAVAVTGWQTINGKKYYFNTNTSIA 2500
 YFGSDSKAVTGLRTIDGKKYYFNTNTAVAVTGWQTINGKKYYFNTNTSIA 2500
 YFGSDSKAVTGLRTIDGKKYYFNTNTAVAVTGWQTINGKKYYFNTNTYIA 2500
 YFGSDSKAVTGLRTIDGKKYYFNTNTAVAVTGWQTINGKKYYFNTNTYIA 2500

STGYTIISGKHFYFNTDGIMQIGVFKGPDGFYFAPANTDANNIEGQAIR 2550
 STGYTIISGKHFYFNTDGIMQIGVFKGPDGFYFAPANTDANNIEGQAIR 2550
 STGYTIISGKHFYFNTDGIMQIGVFKGPDGFYFAPANTDANNIEGQAIR 2550
 STGYTIISGKHFYFNTDGIMQIGVFKGPDGFYFAPANTDANNIEGQAIR 2550
 STGYTIISGKHFYFNTDGIMQIGVFKGPDGFYFAPANTDANNIEGQAIR 2550

YQNRFLYLHDNIYYFGNNSKAATGWVTIDGNRYYFEPNTAMGANGYKTID 2600
 YQNRFLYLHDNIYYFGNNSKAATGWVTIDGNRYYFEPNTAMGANGYKTID 2600
 YQNRFLYLHDNIYYFGNNSKAATGWVTIDGNRYYFEPNTAMGANGYKTID 2600
 YQNRFLYLHDNIYYFGNDSKAATGWATIDGNRYYFEPNTAMGANGYKTID 2600
 YQNRFLYLHDNIYYFGNDSKAATGWATIDGNRYYFEPNTAMGANGYKTID 2600

NKNFYFRNGLPQIGVFKGSNGFEYFAPANTDANNIEGQAIRYQNRLFHLL 2650
 NKNFYFRNGLPQIGVFKGSNGFEYFAPANTDANNIEGQAIRYQNRLFHLL 2650
 NKNFYFRNGLPQIGVFKGSNGFEYFAPANTDANNIEGQAIRYQNRLFHLL 2650
 NKNFYFRNGLPQIGVFKGPNGFEYFAPANTDANNIDGQAIRYQNRLFHLL 2650
 NKNFYFRNGLPQIGVFKGPNGFEYFAPANTDANNIDGQAIRYQNRLFHLL 2650

GKIYYFGNNSKAVTGWQTINGKVVYFMPDTAMAAAGGLFEIDGVIYFFGV 2700
 GKIYYFGNNSKAVTGWQTINGKVVYFMPDTAMAAAGGLFEIDGVIYFFGV 2700
 GKIYYFGNNSKAVTGWQTINGKVVYFMPDTAMAAAGGLFEIDGVIYFFGV 2700
 GKIYYFGNNSKAVTGWQTINSKVYYFMPDTAMAAAGGLFEIDGVIYFFGV 2700
 GKIYYFGNNSKAVTGWQTINSKVYYFMPDTAMAAAGGLFEIDGVIYFFGV 2700

DGVKAPGIYG 2710 (SEQ ID NO: 1, 630)
 DGVKAPGIYG 2710 (SEQ ID NO: 4)
 DGVKAPGIYG 2710 (SEQ ID NO: 19, VPI10463)
 DGVKAPGIYG 2710 (SEQ ID NO: 15, R20291)
 DGVKAPGIYG 2710 (SEQ ID NO: 17, CD196)

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Fig. 2A

MSLVNRKQLEKMANVRFRQEDEVAILDALEEYHNMSENTVVEKYLKLKDINSLTDIYI 60
 MSLVNRKQLEKMANVRFRQEDEVAILDALEEYHNMSENTVVEKYLKLKDINSLTDIYI 60
 MSLVNRKQLEKMANVRFRQEDEVAILDALEEYHNMSENTVVEKYLKLKDINSLTDIYI 60
 MSLVNRKQLEKMANVRFRQEDEVAILDALEEYHNMSENTVVEKYLKLKDINSLTDIYI 60
 MSLVNRKQLEKMANVRFRQEDEVAILDALEEYHNMSENTVVEKYLKLKDINSLTDIYI 60

DTYKKSGRNKALKFKEYLVTEVLELKNNNLTPVEKNLHFVWIGGQINDTAINYINQWKD 120
 DTYKKSGRNKALKFKEYLVTEVLELKNNNLTPVEKNLHFVWIGGQINDTAINYINQWKD 120
 DTYKKSGRNKALKFKEYLVTEVLELKNNNLTPVEKNLHFVWIGGQINDTAINYINQWKD 120
 DTYKKSGRNKALKFKEYLVTEVLELKNNNLTPVEKNLHFVWIGGQINDTAINYINQWKD 120
 DTYKKSGRNKALKFKEYLVTEVLELKNNNLTPVEKNLHFVWIGGQINDTAINYINQWKD 120

VNSDYNVNVFYDSNAFLINTLKKTVVESAINDTLESFRENLNDPRFDYNKFFRKRMETIY 180
 VNSDYNVNVFYDSNAFLINTLKKTVVESAINDTLESFRENLNDPRFDYNKFFRKRMETIY 180
 VNSDYNVNVFYDSNAFLINTLKKTVVESAINDTLESFRENLNDPRFDYNKFFRKRMETIY 180
 VNSDYNVNVFYDSNAFLINTLKKTIVESATNDTLESFRENLNDPRFDYNKFYRKRMETIY 180
 VNSDYNVNVFYDSNAFLINTLKKTIVESATNDTLESFRENLNDPRFDYNKFYRKRMETIY 180

DKQKNFINYYKAQREENPELIIDDIVKTYLSNEYSKEIDELNTYIEESLNKITQNSGNDV 240
 DKQKNFINYYKAQREENPELIIDDIVKTYLSNEYSKEIDELNTYIEESLNKITQNSGNDV 240
 DKQKNFINYYKAQREENPELIIDDIVKTYLSNEYSKEIDELNTYIEESLNKITQNSGNDV 240
 DKQKNFINYYKTQREENPDLIIDDIVKIYLSNEYSKIDELNSYIEESLNKTENSGNDV 240
 DKQKNFINYYKTQREENPDLIIDDIVKIYLSNEYSKIDELNSYIEESLNKTENSGNDV 240

RNFEEFKNGESFNLYEQELVERWNLAAASDILRISALKEIGGYLDVDMLRGIQPDLFES 300
 RNFEEFKNGESFNLYEQELVERWNLAAASDILRISALKEIGGYLDVDMLRGIQPDLFES 300
 RNFEEFKNGESFNLYEQELVERWNLAAASDILRISALKEIGGYLAVAMILPGIQPDLFES 300
 RNFEEFKGGESFKLYEQELVERWNLAAASDILRISALKEVGGYLDVDMLRGIQPDLFES 300
 RNFEEFKGGESFKLYEQELVERWNLAAASDILRISALKEVGGYLDVDMLRGIQPDLFES 300

IEKPSSVTVDWEMTKLEAIMKYKEYIPEYTSEHFMDLDEEVQSSFESVLASKSDKSEIF 360
 IEKPSSVTVDWEMTKLEAIMKYKEYIPEYTSEHFMDLDEEVQSSFESVLASKSDKSEIF 360
 IEKPSSVTVDWEMTKLEAIMKYKEYIPEYTSEHFMDLDEEVQSSFESVLASKSDKSEIF 360
 IEKPSSVTVDWEMVKLEAIMKYKEYIPEYTSEHFMDLDEEVQSSFESVLASKSDKSEIF 360
 IEKPSSVTVDWEMVKLEAIMKYKEYIPEYTSEHFMDLDEEVQSSFESVLASKSDKSEIF 360

SSLGDMEASPLEVKIAFNSKGIIINQGLISVKDSYCSNLIVKQIENRYKILNNSLNPAISE 420
 SSLGDMEASPLEVKIAFNSKGIIINQGLISVKDSYCSNLIVKQIENRYKILNNSLNPAISE 420
 SSLGDMEASPLEVKIAFNSKGIIINQGLISVKDSYCSNLIVKQIENRYKILNNSLNPAISE 420
 SSLGDMEASPLEVKIAFNSKGIIINQGLISVKDSYCSNLIVKQIENRYKILNNSLNPAISE 420
 SSLGDMEASPLEVKIAFNSKGIIINQGLISVKDSYCSNLIVKQIENRYKILNNSLNPAISE 420

DNDFTTTNTFIDSIMAEANADNGRFMELGKYLRVGFFPDVKTTINLSGPEAYAAAYQD 480
 DNDFTTTNTFIDSIMAEANADNGRFMELGKYLRVGFFPDVKTTINLSGPEAYAAAYQD 480
 DNDFTTTNTFIDSIMAEANADNGRFMELGKYLRVGFFPDVKTTINLSGPEAYAAAYQD 480

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FIG. 2B

DNDFNTTNAFIDSIMAEANADNGRFMMELGKYLRVGFDPDVKTINLSGPEAYAAAYQD 480
 DNDFNTTNAFIDSIMAEANADNGRFMMELGKYLRVGFDPDVKTINLSGPEAYAAAYQD 480
 *****;*****;

LLMFKEGSMNIHLIEADLRNFEISKTNISQSTEQEMASLWSFDDARAKAQFEEYKRNYFE 540
 LLMFKEGSMNIHLIEADLRNFEISKTNISQSTEQEMASLWSFDDARAKAQFEEYKRNYFE 540
 LLMFKEGSMNIHLIEADLRNFEISKTNISQSTEQEMASLWSFDDARAKAQFEEYKRNYFE 540
 LLMFKEGSMNIHLIEADLRNFEISKTNISQSTEQEMASLWSFDDARAKAQFEEYKKNYFE 540
 LLMFKEGSMNIHLIEADLRNFEISKTNISQSTEQEMASLWSFDDARAKAQFEEYKKNYFE 540
 *****;

GSLGEDDNLDFSQNIVVDKEYLLEKISSLARSSERGYIHYIVQLQGDKISYEACNLFAK 600
 GSLGEDDNLDFSQNIVVDKEYLLEKISSLARSSERGYIHYIVQLQGDKISYEACNLFAK 600
 GSLGEDDNLDFSQNIVVDKEYLLEKISSLARSSERGYIHYIVQLQGDKISYEACNLFAK 600
 GSLGEDDNLDFSQNIVVDKEYLLEKISSLARSSERGYIHYIVQLQGDKISYEACNLFAK 600
 GSLGEDDNLDFSQNIVVDKEYLLEKISSLARSSERGYIHYIVQLQGDKISYEACNLFAK 600
 *****;

TPYDSVLFQKNIEDSEIAYYYNPGDGEIQEIDKYKIPSIISDRPKIKLTFIGHGKDEFNT 660
 TPYDSVLFQKNIEDSEIAYYYNPGDGEIQEIDKYKIPSIISDRPKIKLTFIGHGKDEFNT 660
 TPYDSVLFQKNIEDSEIAYYYNPGDGEIQEIDKYKIPSIISDRPKIKLTFIGHGKDEFNT 660
 TPYDSVLFQKNIEDSEIAYYYNPGDGEIQEIDKYKIPSIISDRPKIKLTFIGHGKDEFNT 660
 TPYDSVLFQKNIEDSEIAYYYNPGDGEIQEIDKYKIPSIISDRPKIKLTFIGHGKDEFNT 660
 *****;

DIFAGFDVDSLSTEIEAAIDLAKEDISPKSIEINLLGCNMFSYSINVEETYPGKLLLKV 720
 DIFAGFDVDSLSTEIEAAIDLAKEDISPKSIEINLLGCNMFSYSINVEETYPGKLLLKV 720
 DIFAGFDVDSLSTEIEAAIDLAKEDISPKSIEINLLGCNMFSYSINVEETYPGKLLLKV 720
 DIFAGLDVDSLSTEIETAAIDLAKEDISPKSIEINLLGCNMFSYSVNVEETYPGKLLLKV 720
 DIFAGLDVDSLSTEIETAAIDLAKEDISPKSIEINLLGCNMFSYSVNVEETYPGKLLLKV 720
 *****;

DKISELMPsisQDSIIVSANQYEV RinSEGRRELLDHSGEWINKEESIIKDISSKEYISF 780
 DKISELMPsisQDSIIVSANQYEV RinSEGRRELLDHSGEWINKEESIIKDISSKEYISF 780
 DKISELMPsisQDSIIVSANQYEV RinSEGRRELLDHSGEWINKEESIIKDISSKEYISF 780
 DKVSELMPsisQDSIIVSANQYEV RinSEGRRELLDHSGEWINKEESIIKDISSKEYISF 780
 DKVSELMPsisQDSIIVSANQYEV RinSEGRRELLDHSGEWINKEESIIKDISSKEYISF 780
 **;

NPKENKITVSKKNLPELSTLLQEI RNNNSNSDIELEEKVMLTECEINVISNIDTQIVEER 840
 NPKENKITVSKKNLPELSTLLQEI RNNNSNSDIELEEKVMLTECEINVISNIDTQIVEER 840
 NPKENKITVSKKNLPELSTLLQEI RNNNSNSDIELEEKVMLTECEINVISNIDTQIVEER 840
 NPKENKIIVKSKKNLPELSTLLQEI RNNNSNSDIELEEKVMLAECEINVISNIDTQVVEGR 840
 NPKENKIIVKSKKNLPELSTLLQEI RNNNSNSDIELEEKVMLAECEINVISNIDTQVVEGR 840
 *****;

IEEAKNLTSDSINYIKDEFKLIESISDALCDLKQQNELEDSHFISFEDISSETDEGFSIRF 900
 IEEAKNLTSDSINYIKDEFKLIESISDALCDLKQQNELEDSHFISFEDISSETDEGFSIRF 900
 IEEAKNLTSDSINYIKDEFKLIESISDALCDLKQQNELEDSHFISFEDISSETDEGFSIRF 900
 IEEAKSLTSDSINYIKNEFKLIESISDALYDLKQQNELEESHFISFEDILETDEGFSIRF 900
 IEEAKSLTSDSINYIKNEFKLIESISDALYDLKQQNELEESHFISFEDILETDEGFSIRF 900
 *****;

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FIG. 2C

INKETGESIFVETEKTIFSEYANHITEEISKIKGTIFDTVNGKLVKKVNLDTHEVNTLN 960
 INKETGESIFVETEKTIFSEYANHITEEISKIKGTIFDTVNGKLVKKVNLDTHEVNTLN 960
 INKETGESIFVETEKTIFSEYANHITEEISKIKGTIFDTVNGKLVKKVNLDTHEVNTLN 960
 IDKETGESIFVETEKAIFSEYANHITEEISKIKGTIFDTVNGKLVKKVNLDATHEVNTLN 960
 IDKETGESIFVETEKAIFSEYANHITEEISKIKGTIFDTVNGKLVKKVNLDATHEVNTLN 960
 *;*****:*****:*****:*****:*****:*****

AAFFIQSLIEYNSSKESLSNLNVAMKVQVYAQLFSTGLNTITDAAKVVELVSTALDETID 1020
 AAFFIQSLIEYNSSKESLSNLNVAMKVQVYAQLFSTGLNTITDAAKVVELVSTALDETID 1020
 AAFFIQSLIEYNSSKESLSNLNVAMKVQVYAQLFSTGLNTITDAAKVVELVSTALDETID 1020
 AAFFIQSLIEYNSSKESLSNLNVAMKVQVYAQLFSTGLNTITDAAKVVELVSTALDETID 1020
 AAFFIQSLIEYNSSKESLSNLNVAMKVQVYAQLFSTGLNTITDAAKVVELVSTALDETID 1020

LLPTLSEGLPTIATIIDGVSLGAAIKELSETSDPLLQEIETAKIGIMAVNLTTATTAIIT 1080
 LLPTLSEGLPTIATIIDGVSLGAAIKELSETSDPLLQEIETAKIGIMAVNLTTATTAIIT 1080
 LLPTLSEGLPTIATIIDGVSLGAAIKELSETSDPLLQEIETAKIGIMAVNLTTATTAIIT 1080
 LLPTLSEGLPVIATIIDGVSLGAAIKELSETSDPLLQEIETAKIGIMAVNLTAATTAIIT 1080
 LLPTLSEGLPVIATIIDGVSLGAAIKELSETSDPLLQEIETAKIGIMAVNLTAATTAIIT 1080

SSLGIAASGFSILLVPLAGISAGIPSLVNNELVLRDKATKVVDFKHSVSLVETEGVFTLLD 1140
 SSLGIAASGFSILLVPLAGISAGIPSLVNNELVLRDKATKVVDFKHSVSLVETEGVFTLLD 1140
 SSLGIAASGFSILLVPLAGISAGIPSLVNNELVLRDKATKVVDFKHSVSLVETEGVFTLLD 1140
 SSLGIAASGFSILLVPLAGISAGIPSLVNNELILRDKATKVVDFFSHISLAESEGAFSLD 1140
 SSLGIAASGFSILLVPLAGISAGIPSLVNNELILRDKATKVVDFFSHISLAESEGAFSLD 1140

DKIMMPQDDLVISEIDFNNNSIVLGKCEIWRMEGGSGHTVTDDIDHFFSAPSITYREPHL 1200
 DKIMMPQDDLVISEIDFNNNSIVLGKCEIWRMEGGSGHTVTDDIDHFFSAPSITYREPHL 1200
 DKIMMPQDDLVISEIDFNNNSIVLGKCEIWRMEGGSGHTVTDDIDHFFSAPSITYREPHL 1200
 DKIMMPQDDLVISEIDFNNNSITLGKCEIWRMEGGSGHTVTDDIDHFFSAPSITYREPHL 1200
 DKIMMPQDDLVISEIDFNNNSITLGKCEIWRMEGGSGHTVTDDIDHFFSAPSITYREPHL 1200

SIYDVLEVQKEELDLSKDLMVLPNAPNRVFAWETGWTPLRSLENDGTKLLDRIRDNYEG 1260
 SIYDVLEVQKEELDLSKDLMVLPNAPNRVFAWETGWTPLRSLENDGTKLLDRIRDNYEG 1260
 SIYDVLEVQKEELDLSKDLMVLPNAPNRVFAWETGWTPLRSLENDGTKLLDRIRDNYEG 1260
 SIYDVLEVQKEELDLSKDLMVLPNAPNRVFAWETGWTPLRSLENDGTKLLDRIRDNYEG 1260
 SIYDVLEVQKEELDLSKDLMVLPNAPNRVFAWETGWTPLRSLENDGTKLLDRIRDNYEG 1260

EFYWRYFAFIADALITTLKPRYEDTNIRINLDSNTRSFIGVPIITTEYIREKLSYSFYGSG 1320
 EFYWRYFAFIADALITTLKPRYEDTNIRINLDSNTRSFIGVPIITTEYIREKLSYSFYGSG 1320
 EFYWRYFAFIADALITTLKPRYEDTNIRINLDSNTRSFIGVPIITTEYIREKLSYSFYGSG 1320
 EFYWRYFAFIADALITTLKPRYEDTNIRINLDSNTRSFIGVPIITTEYIREKLSYSFYGSG 1320
 EFYWRYFAFIADALITTLKPRYEDTNIRINLDSNTRSFIGVPIITTEYIREKLSYSFYGSG 1320

GTYALSLSQYNMGINIELSESVDWIIDVDNVVRDVTIESDKIKKGDLIEGILSTLSIEEN 1380
 GTYALSLSQYNMGINIELSESVDWIIDVDNVVRDVTIESDKIKKGDLIEGILSTLSIEEN 1380
 GTYALSLSQYNMGINIELSESVDWIIDVDNVVRDVTIESDKIKKGDLIEGILSTLSIEEN 1380

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FIG. 2D

GTYALSLSQYNMNIELNENDTWVIDVDNVVRDVTIESDKIKKGDLIENILSKLSIEDN 1380
GTYALSLSQYNMNIELNENDTWVIDVDNVVRDVTIESDKIKKGDLIENILSKLSIEDN 1380

NHIQQKIDYIGFNSELQKNI PYSFVDSEKENG FINGSTKEGLFVSEL PDVVLISKVYMD 1500
NHIQQKIDYIGFNSELQKNI PYSFVDSEKENG FINGSTKEGLFVSEL PDVVLISKVYMD 1500
NHIQQKIDYIGFNSELQKNI PYSFVDSEKENG FINGSTKEGLFVSEL PDVVLISKVYMD 1500
NSVQQKIDYIGLNSELQKNI PYSFMDDKGKENGFINCSTKEGLFVSEL SDVVLISKVYMD 1500
NSVQQKIDYIGLNSELQKNI PYSFMDDKGKENGFINCSTKEGLFVSEL SDVVLISKVYMD 1500
* . * * * * * * * . * * * * * * * * . * * * * * * * * . * * * * * * * * . * * * * * * * *

FEFICDENDNIQPYFIKFNTLETNYTLYVGNRQN MIVEPNYDLD DSGDISS TVINFSQKY 1680
FEFICDKDNNIQPYFIKFNTLETKYTLYVGNRQN MIVEPNYDLD DSGDISS TVINFSQKY 1680
FEFICDKDNNIQPYFIKFNTLETKYTLYVGNRQN MIVEPNYDLD DSGDISS TVINFSQKY 1680

LYGIDSCVNKVVISPNIYTDEINITPVYETNNNTYPEEVIVLDANYINEKINVNINDLSIRY 1740
LYGIDSCVNKVVISPNIYTDEINITPVYETNNNTYPEEVIVLDANYINEKINVNINDLSIRY 1740
LYGIDSCVNKVVISPNIYTDEINITPVYETNNNTYPEEVIVLDANYINEKINVNINDLSIRY 1740
LYGIDSCVNKVIISPNIYTDEINITPIYEANNTYPEEVIVLDTNYISEKINININDLSIRY 1740
LYGIDSCVNKVIISPNIYTDEINITPIYEANNTYPEEVIVLDTNYISEKINININDLSIRY 1740

VWSNDGND FILMST SEENKV SQVK IRFVNFKD TLANKL SFNFS DKQDVPVSEI ILSFT 1800
VWSNDGND FILMST SEENKV SQVK IRFVNFKD TLANKL SFNFS DKQDVPVSEI ILSFT 1800
VWSNDGND FILMST SEENKV SQVK IRFVNFKD TLANKL SFNFS DKQDVPVSEI ILSFT 1800
VWSNDGSD FILMST DEENKV SQVK IRFTNFKGNTISDKISFNFS DKQDVSINKVISTFT 1800
VWSNDGSD FILMST DEENKV SQVK IRFTNFKGNTISDKISFNFS DKQDVSINKVISTFT 1800

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FIG. 2E

PSYYEDGLIGYDLGLVSLYNEKEFYINNFGMMVSGLIYINDSLYYFKPPVNNLITGFVTVG 1860
 PSYYEDGLIGYDLGLVSLYNEKEFYINNFGMMVSGLIYINDSLYYFKPPVNNLITGFVTVG 1860
 PSYYEDGLIGYDLGLVSLYNEKEFYINNFGMMVSGLIYINDSLYYFKPPVNNLITGFVTVG 1860
 PSYYVEGLLNLYDLGLISLYNEKEFYINNFGMMVSGLVYINDSLYYFKPPIKNLITGFTTIG 1860
 PSYYVEGLLNLYDLGLISLYNEKEFYINNFGMMVSGLVYINDSLYYFKPPIKNLITGFTTIG 1860
 **** ;*:*****:*****:*****:*****:*****:*****:*****:*****:*****:*

DDKYYFNPINGGAASIGETIIDDKNYYFNQSGVLQTGVFSTEDGFKYFAPANTLDENLEG 1920
 DDKYYFNPINGGAASIGETIIDDKNYYFNQSGVLQTGVFSTEDGFKYFAPANTLDENLEG 1920
 DDKYYFNPINGGAASIGETIIDDKNYYFNQSGVLQTGVFSTEDGFKYFAPANTLDENLEG 1920
 DDKYYFNPDNGGAASVGETIIDGKNEYFSQNGVLQTGVFSTEDGFKYFAPADTLDENLEG 1920
 DDKYYFNPDNGGAASVGETIIDGKNEYFSQNGVLQTGVFSTEDGFKYFAPADTLDENLEG 1920
 *****:*****:*****:*****:*****:*****:*****:*****:*****:*****:*****

EAIDFTGKLIIDENIYYFDDNYRGAIVEWKELDGEHYFSPETGKAFKGLNQIGDYKYYFN 1980
 EAIDFTGKLIIDENIYYFDDNYRGAIVEWKELDGEHYFSPETGKAFKGLNQIGDYKYYFN 1980
 EAIDFTGKLIIDENIYYFDDNYRGAIVEWKELDGEHYFSPETGKAFKGLNQIGDYKYYFN 1980
 EAIDFTGKLTIIDENVYYFGDNYRAAIEWQTLDEVYYFSTDTGRAFKGLNQIGDDKFYFN 1980
 EAIDFTGKLTIIDENVYYFGDNYRAAIEWQTLDEVYYFSTDTGRAFKGLNQIGDDKFYFN 1980
 *****:*****:*****:*****:*****:*****:*****:*****:*****:*****:*****

SDGVMQKGFVSINDNKHYFDDSGVMKVGYTEIDGKHFYFAENGEMQIGVFNTEDEGFKYFA 2040
 SDGVMQKGFVSINDNKHYFDDSGVMKVGYTEIDGKHFYFAENGEMQIGVFNTEDEGFKYFA 2040
 SDGVMQKGFVSINDNKHYFDDSGVMKVGYTEIDGKHFYFAENGEMQIGVFNTEDEGFKYFA 2040
 SDGIMQKGFVNINDKTFYFDDSGVMKSGYTEIDGKYFYFAENGEMQIGVFNTEDEGFKYFA 2040
 SDGIMQKGFVNINDKTFYFDDSGVMKSGYTEIDGKYFYFAENGEMQIGVFNTEDEGFKYFA 2040
 :**:*****:*****:*****:*****:*****:*****:*****:*****:*****

HHNEDLGNEEGEEISYSGLNFKIYYFDDSFATAVVGWKDLEDGSKYYFDEDTAEAYIG 2100
 HNNEDLGNEEGEEISYSGLNFKIYYFDDSFATAVVGWKDLEDGSKYYFDEDTAEAYIG 2100
 HNNEDLGNEEGEEISYSGLNFKIYYFDDSFATAVVGWKDLEDGSKYYFDEDTAEAYIG 2100
 HHDEDLGNEEGEALSYSGLNFKIYYFDDSFATAVVGWKDLEDGSKYYFDEDTAEAYIG 2100
 HHDEDLGNEEGEALSYSGLNFKIYYFDDSFATAVVGWKDLEDGSKYYFDEDTAEAYIG 2100
 :***:*****:*****:*****:*****:*****:*****:*****:*****:*****

LSLINDGQYYFNDGIMQVGFVTINDKVFYFSDSGIIESGVQNIDDNYFYIDNGIVQIG 2160
 LSLINDGQYYFNDGIMQVGFVTINDKVFYFSDSGIIESGVQNIDDNYFYIDNGIVQIG 2160
 LSLINDGQYYFNDGIMQVGFVTINDKVFYFSDSGIIESGVQNIDDNYFYIDNGIVQIG 2160
 ISIINDGKYYFNDGIMQIGFVTINNEVFYFSDSGIVESGMQNIDDNYFYIDENGLVQIG 2160
 ISIINDGKYYFNDGIMQIGFVTINNEVFYFSDSGIVESGMQNIDDNYFYIDENGLVQIG 2160
 :*:*****:*****:*****:*****:*****:*****:*****:*****:*****:*****

VFDTSDEGYKYFAPANTVNDNIYQAVEYSGLVRVGEDVYYFGETYTIETGWIYDMENESD 2220
 VFDTSDEGYKYFAPANTVNDNIYQAVEYSGLVRVGEDVYYFGETYTIETGWIYDMENESD 2220
 VFDTSDEGYKYFAPANTVNDNIYQAVEYSGLVRVGEDVYYFGETYTIETGWIYDMENESD 2220
 VFDTSDEGYKYFAPANTVNDNIYQAVEYSGLVRVGEDVYYFGETYTIETGWIYDMENESD 2220
 VFDTSDEGYKYFAPANTVNDNIYQAVEYSGLVRVGEDVYYFGETYTIETGWIYDMENESD 2220
 *****:*****:*****:*****:*****:*****:*****:*****:*****:*****:*****

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Fig 2F

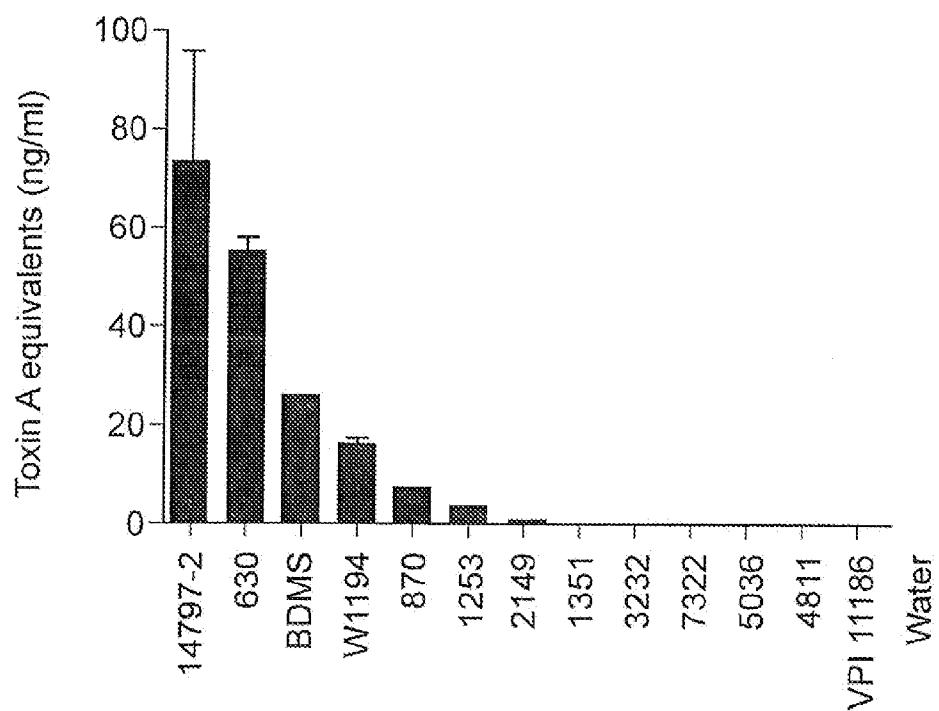
KYYFNPETKKACKGINLIDDIKYYFDEKGIMRTGLISFENNYYFNENGEMQFGYINIED 2280
 KYYFNPETKKACKGINLIDDIKYYFDEKGIMRTGLISFENNYYFNENGEMQFGYINIED 2280
 KYYFNPETKKACKGINLIDDIKYYFDEKGIMRTGLISFENNYYFNENGEMQFGYINIED 2280
 KYYFDPETKKAYKGINVDDIKYYFDENGIMRTGLITFEDNHYYFNEDGIMQYGYLNIED 2280
 KYYFDPETKKAYKGINVDDIKYYFDENGIMRTGLITFEDNHYYFNEDGIMQYGYLNIED 2280
 ****:***** * *;*****:*****:*****:*****:*****:*****:*****:*****

KMFYFGEDGVMQIGVFNTPDGFKYFAHQNTLDENFEGESINYTGWLDLDEKRYYFTDEYI 2340
 KMFYFGEDGVMQIGVFNTPDGFKYFAHQNTLDENFEGESINYTGWLDLDEKRYYFTDEYI 2340
 KMFYFGEDGVMQIGVFNTPDGFKYFAHQNTLDENFEGESINYTGWLDLDEKRYYFTDEYI 2340
 KTFYFSEDGIMQIGVFNTPDGFKYFAHQNTLDENFEGESINYTGWLDLDEKRYYFTDEYI 2340
 KTFYFSEDGIMQIGVFNTPDGFKYFAHQNTLDENFEGESINYTGWLDLDEKRYYFTDEYI 2340
 * ***.***:*****:*****:*****:*****:*****:*****:*****:*****

AATGSVIIDGEEYYFDPDTAQLVISE 2366 (SEQ ID NO: 2, 630)
 AATGSVIIDGEEYYFDPDTAQLVISE 2366 (SEQ ID NO: 25, VPI10463)
 AATGSVIIDGEEYYFDPDTAQLVISE 2366 (SEQ ID NO: 6)
 AATGSVIIDGEEYYFDPDTAQLVISE 2366 (SEQ ID NO: 21, R20291)
 AATGSVIIDGEEYYFDPDTAQLVISE 2366 (SEQ ID NO: 23, CD196)

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FIG. 3



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FIG. 4A

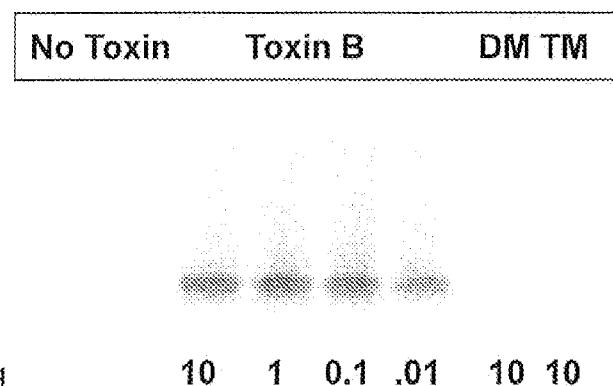
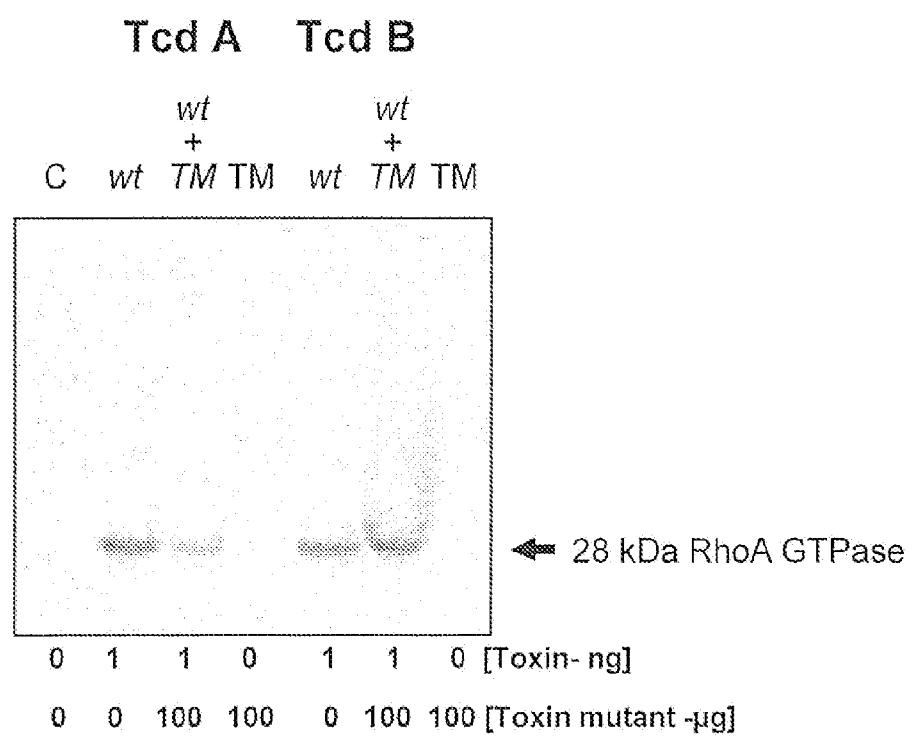
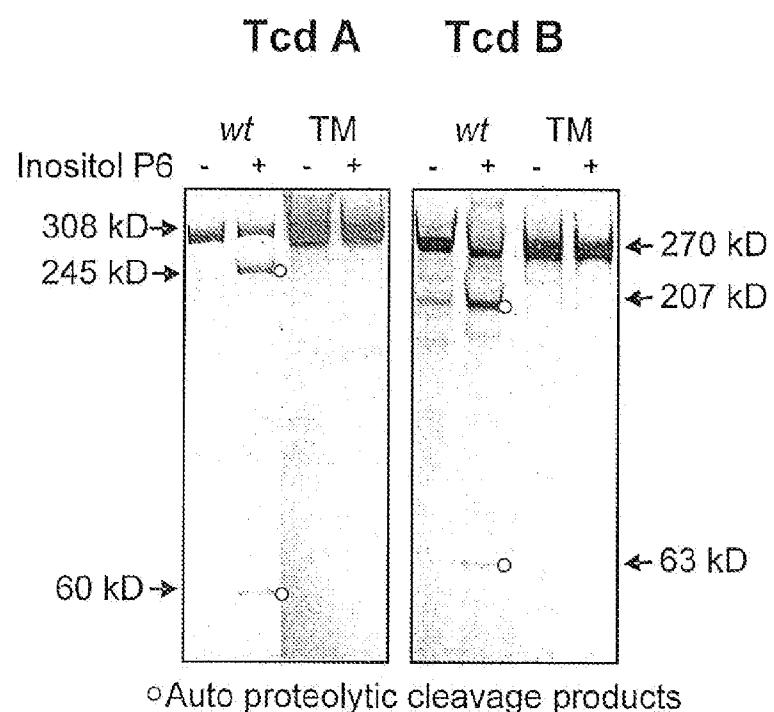


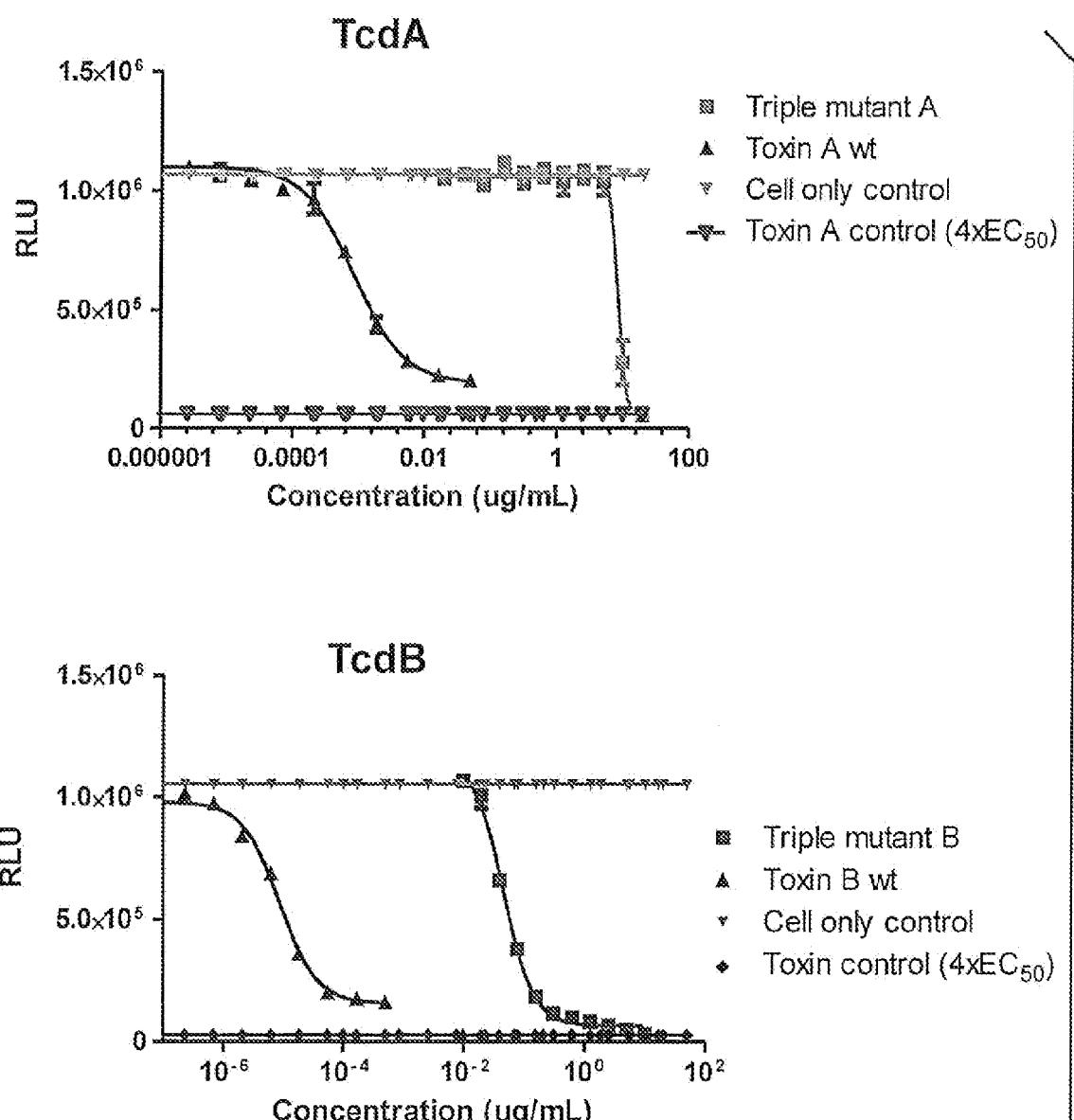
FIG. 4B



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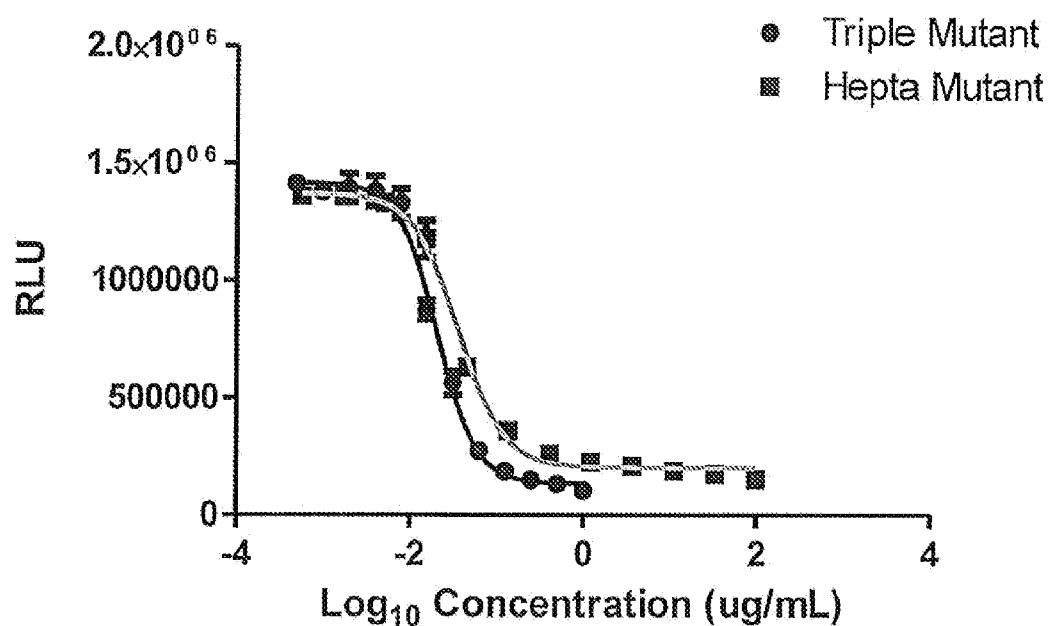
FIG. 5

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FIG. 6

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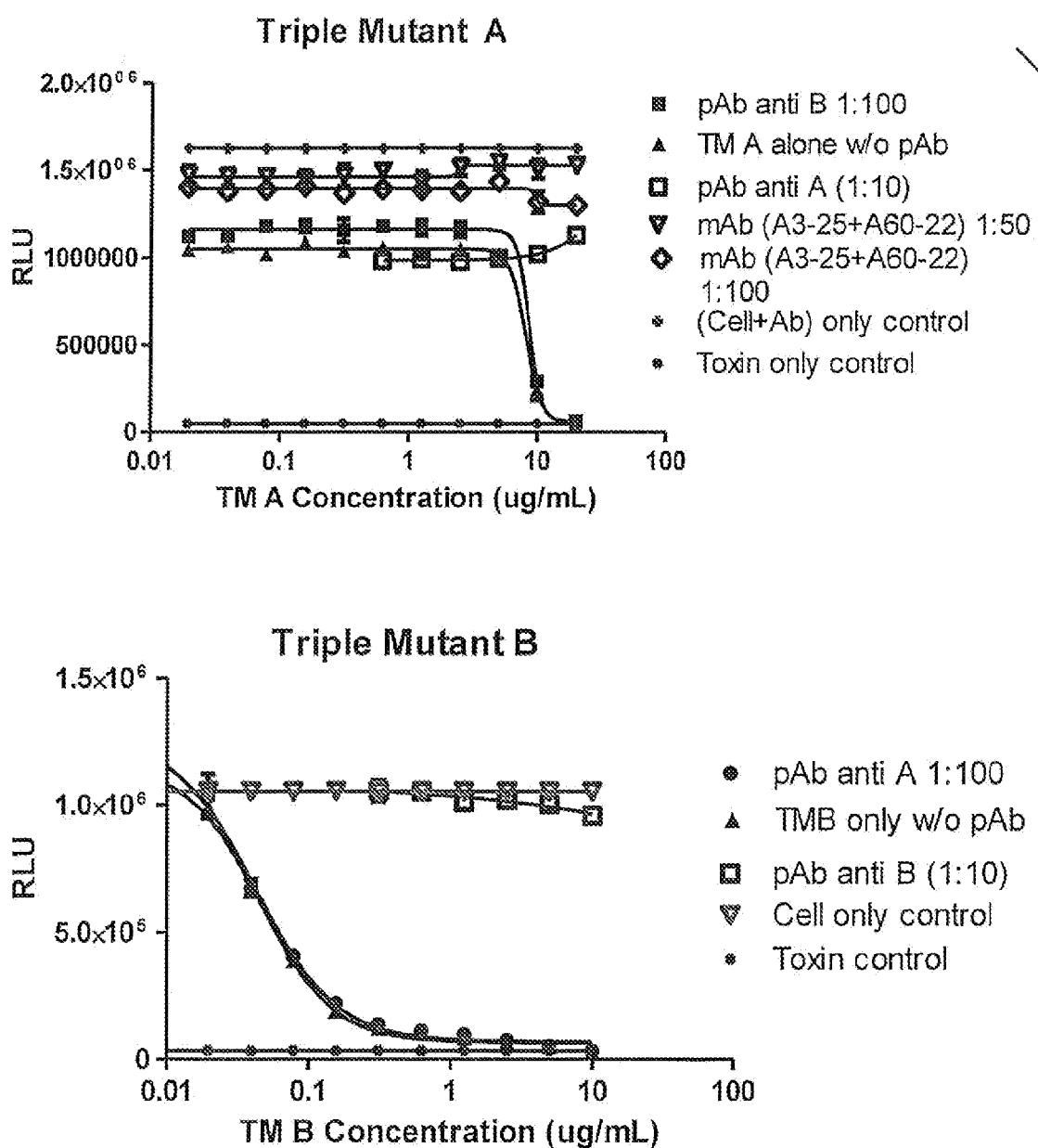
FIG. 7



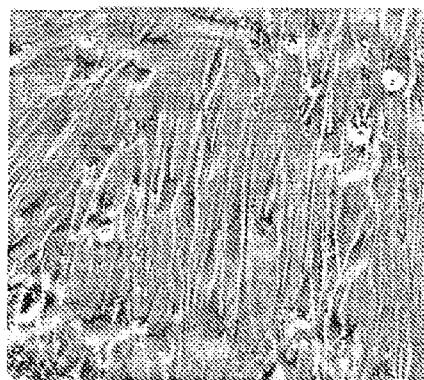
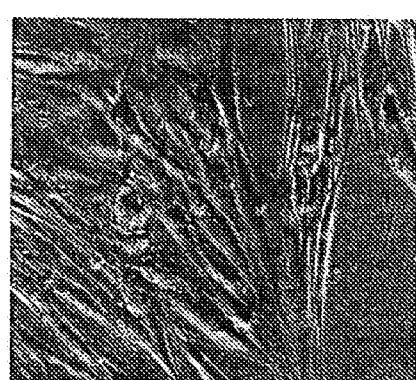
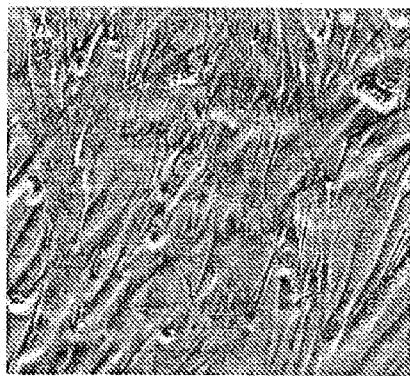
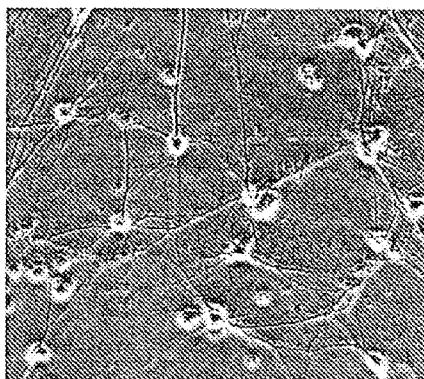
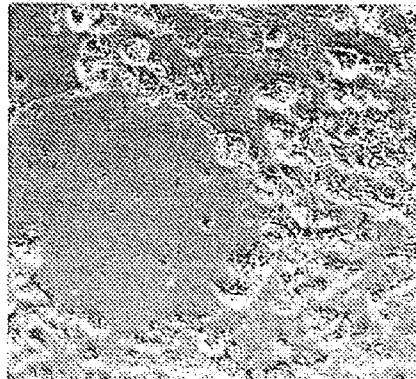
	Triple Mutant	Hepta Mutant
EC ₅₀	0.02078	0.03590

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FIG. 8

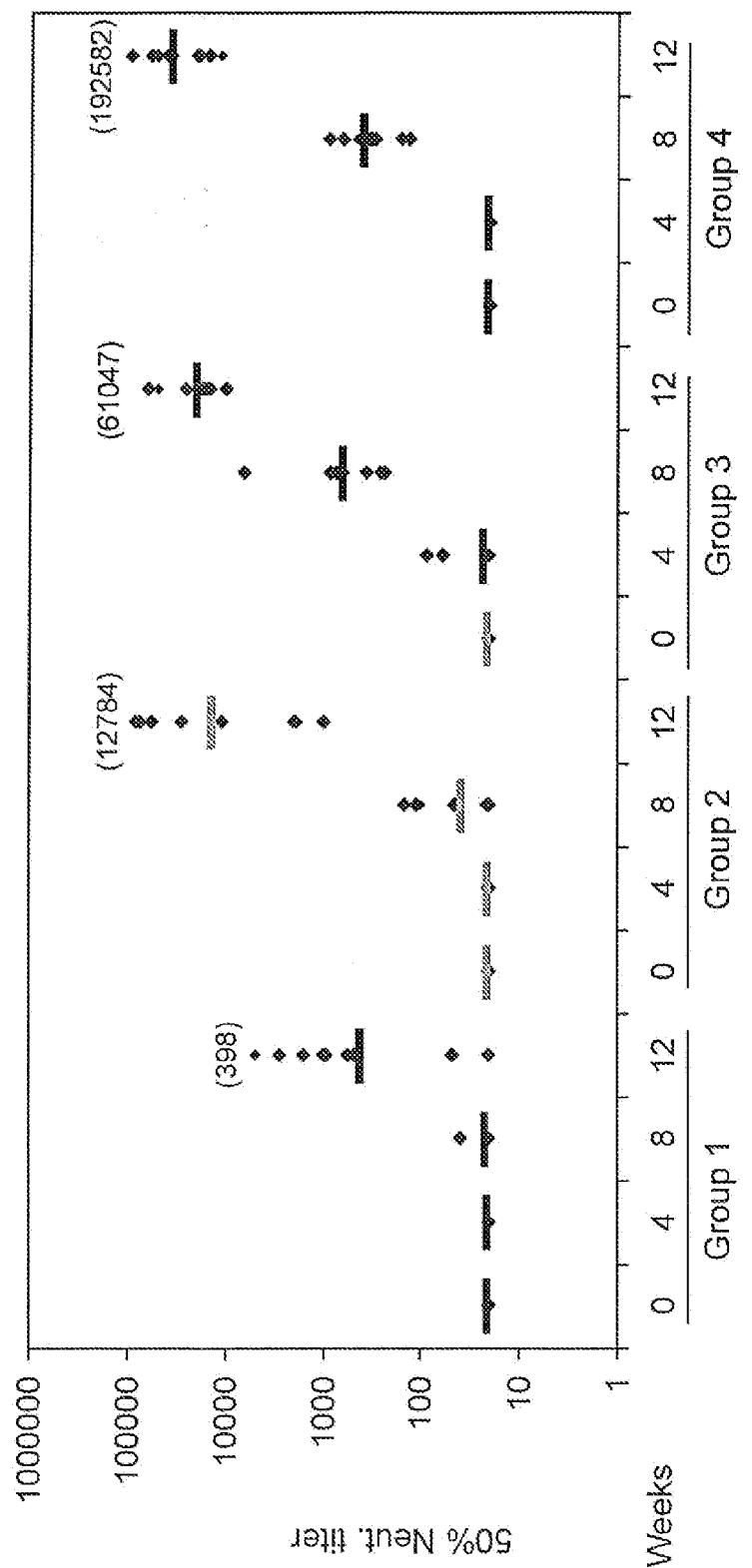


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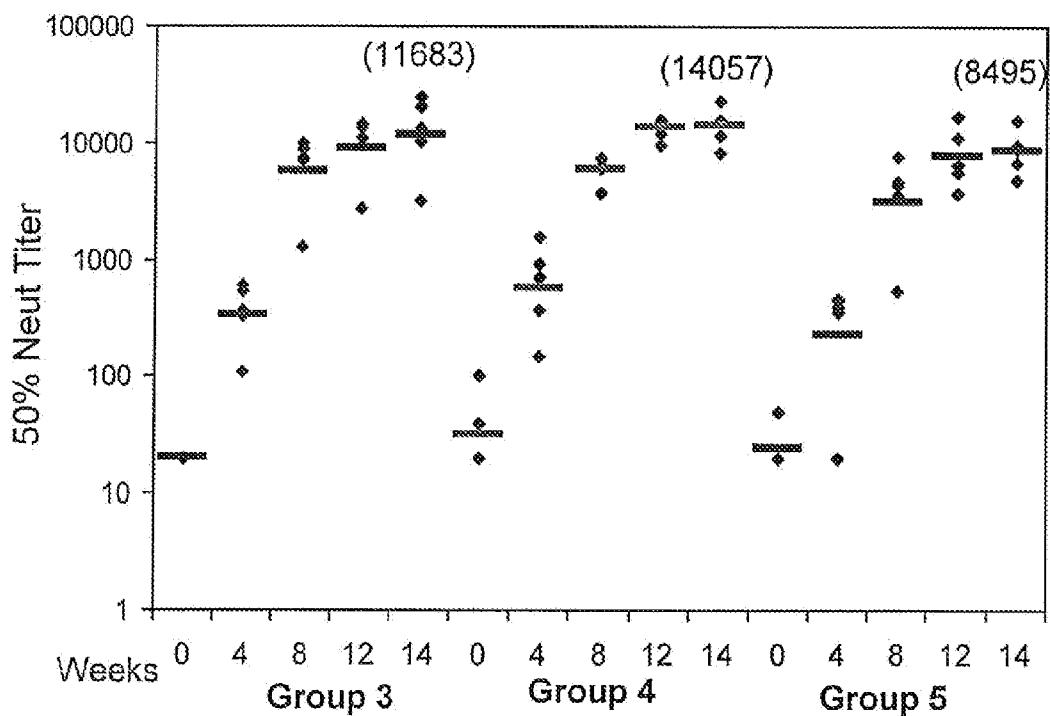
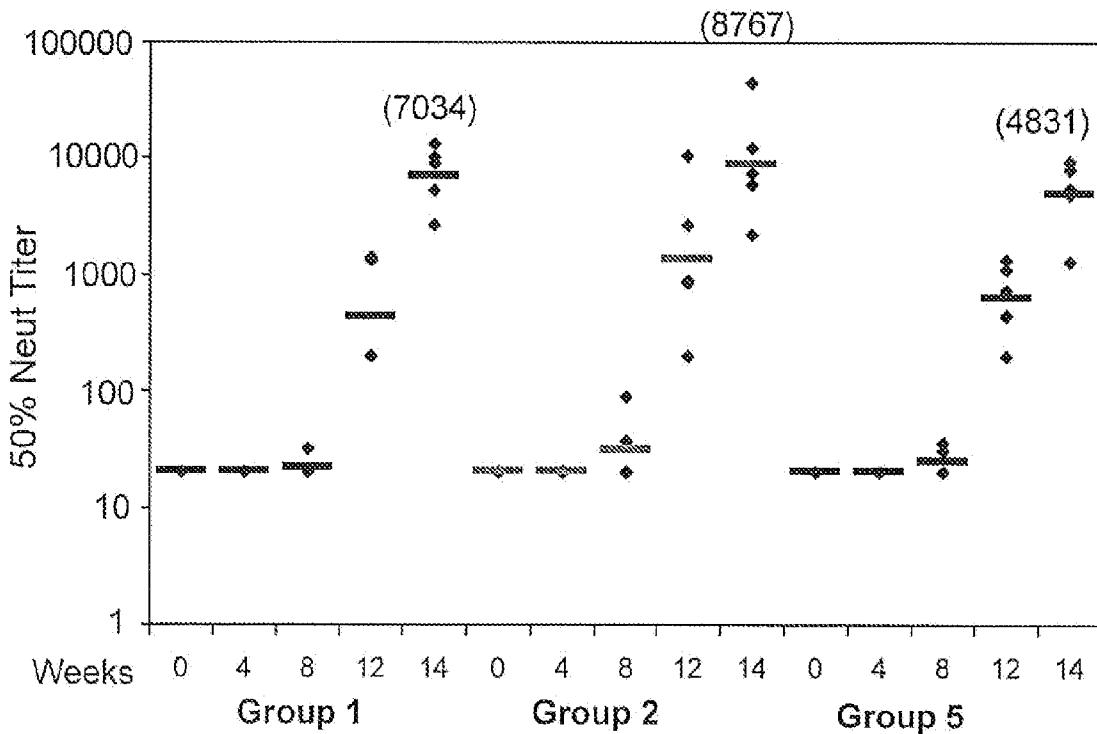
FIG. 9**A****B****C****D****E**

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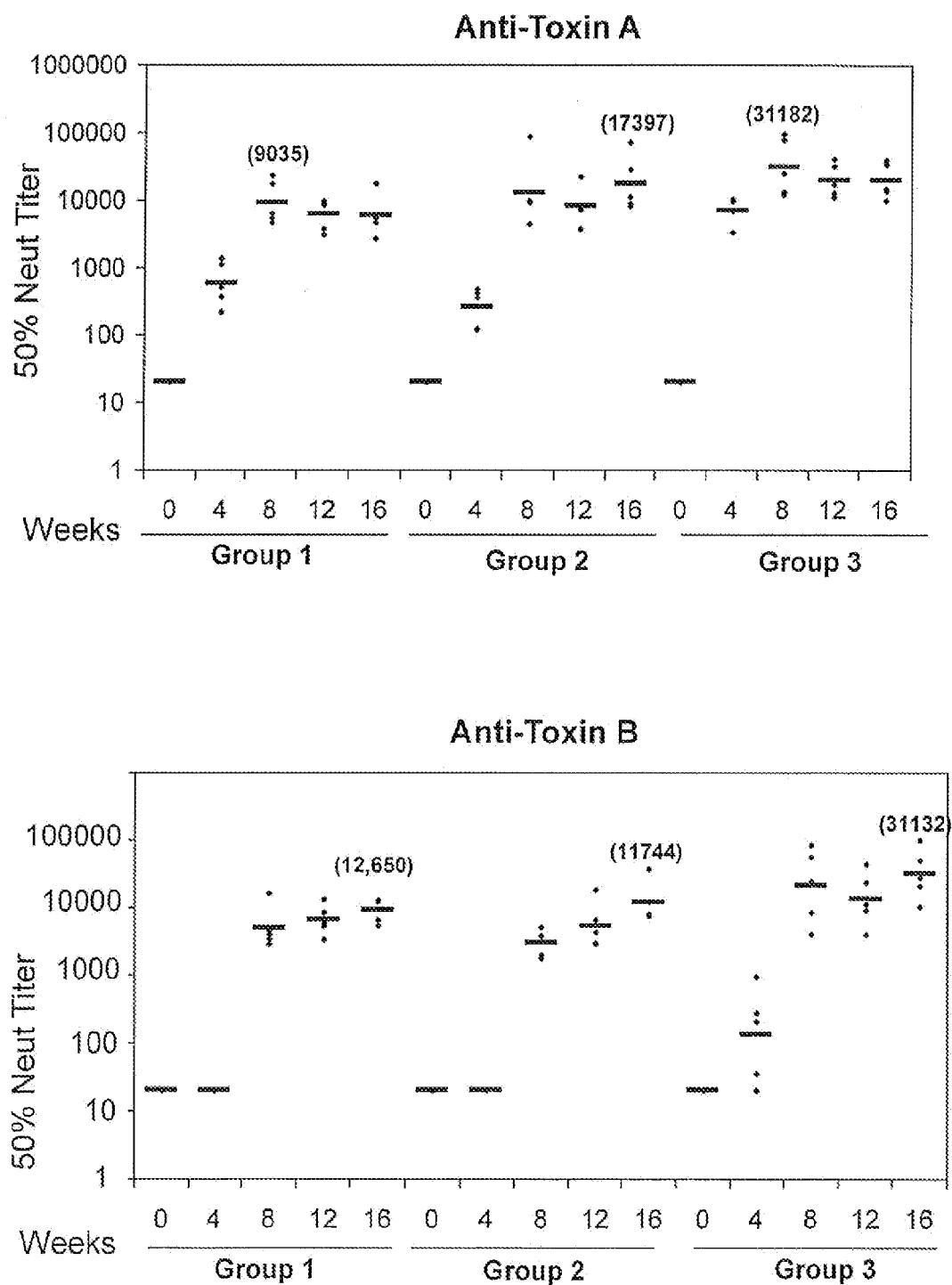
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FIG. 11A**FIG. 11B**

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FIG. 12

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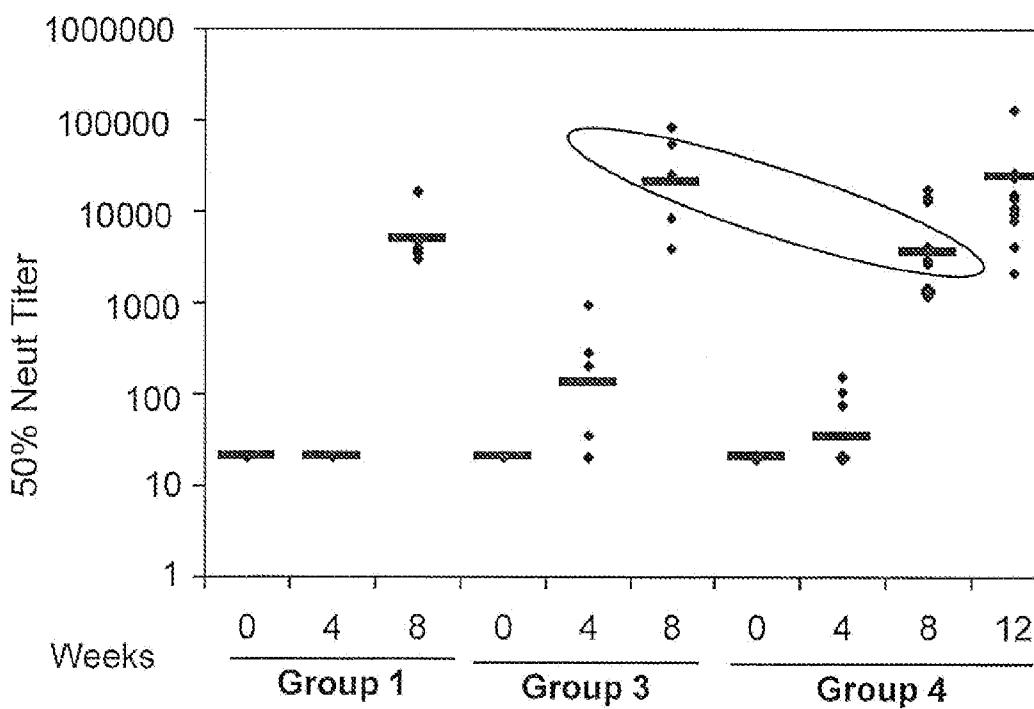
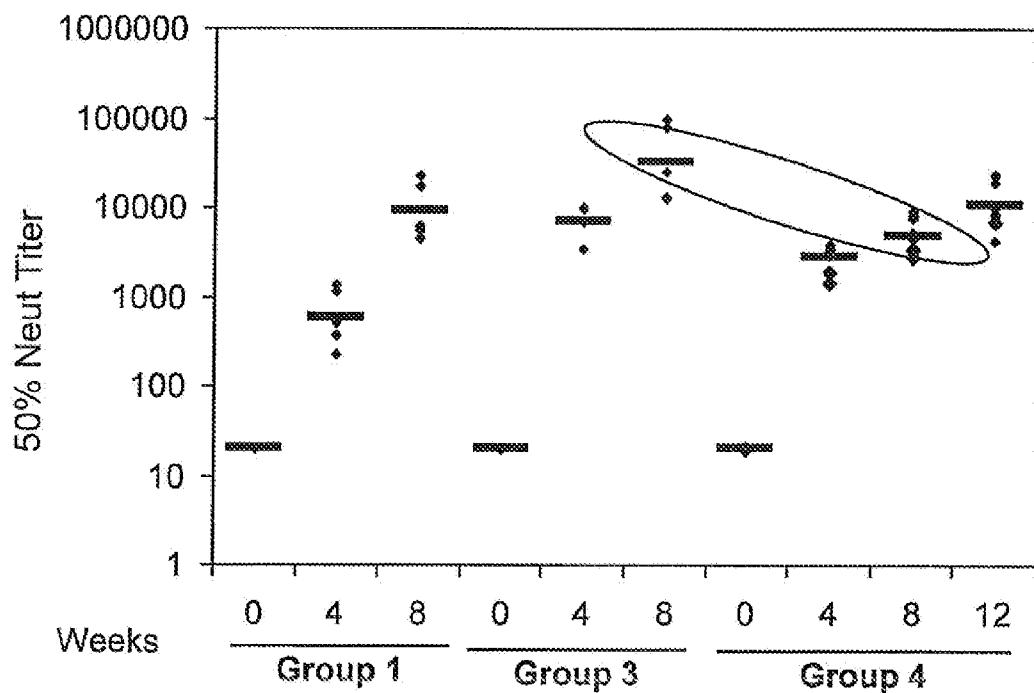
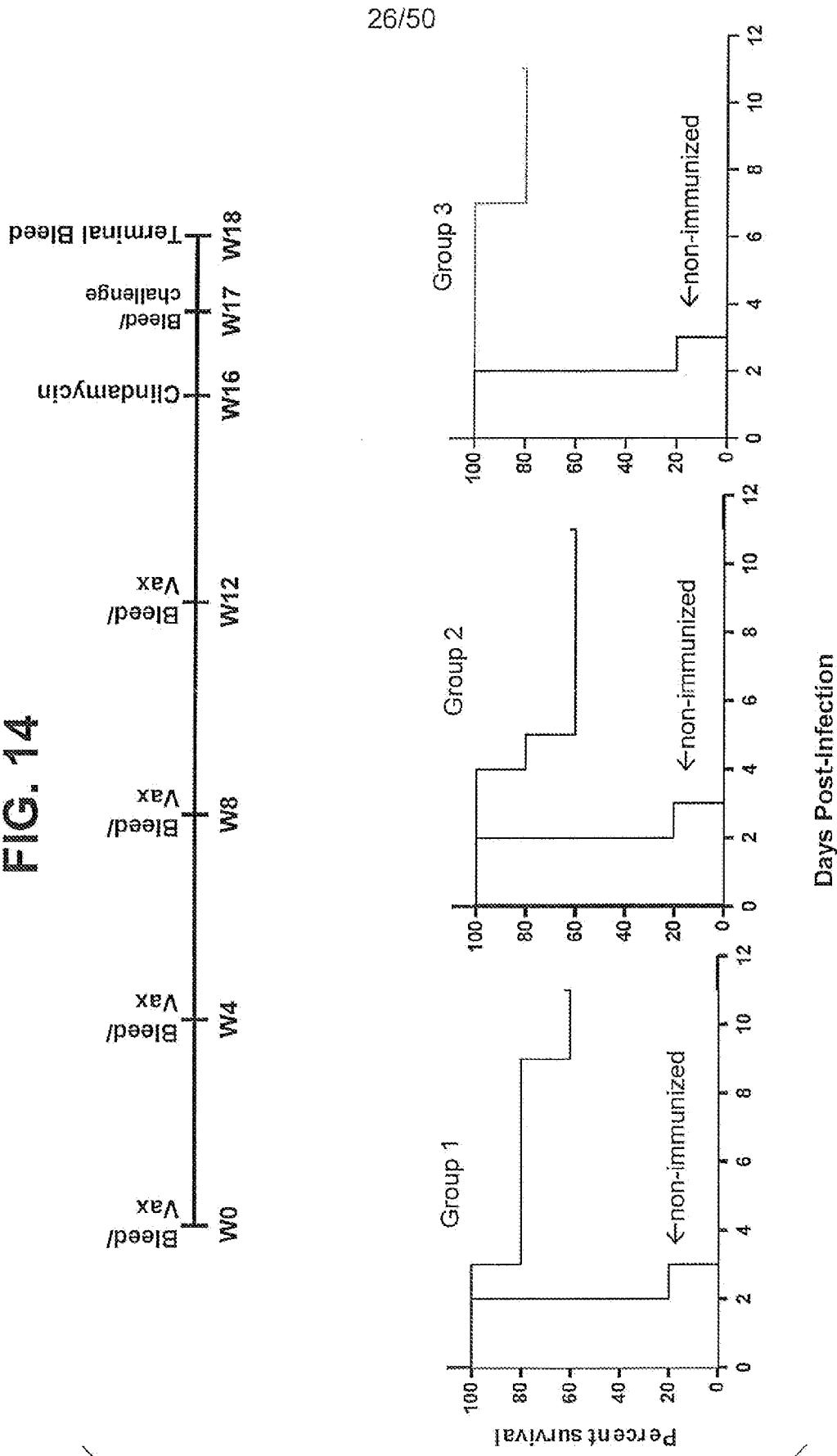
FIG. 13

FIG. 14



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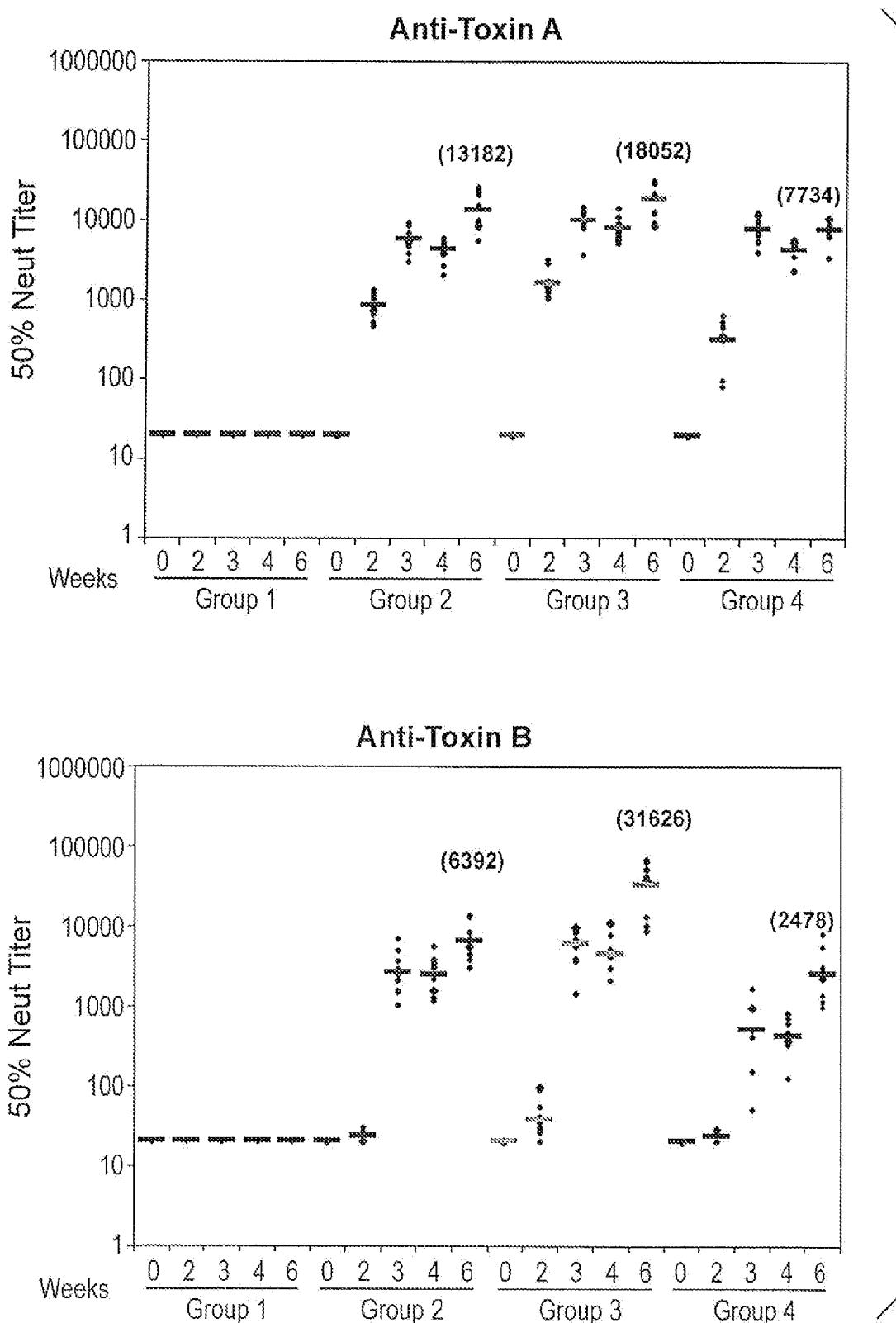
FIG. 15

FIG. 16A

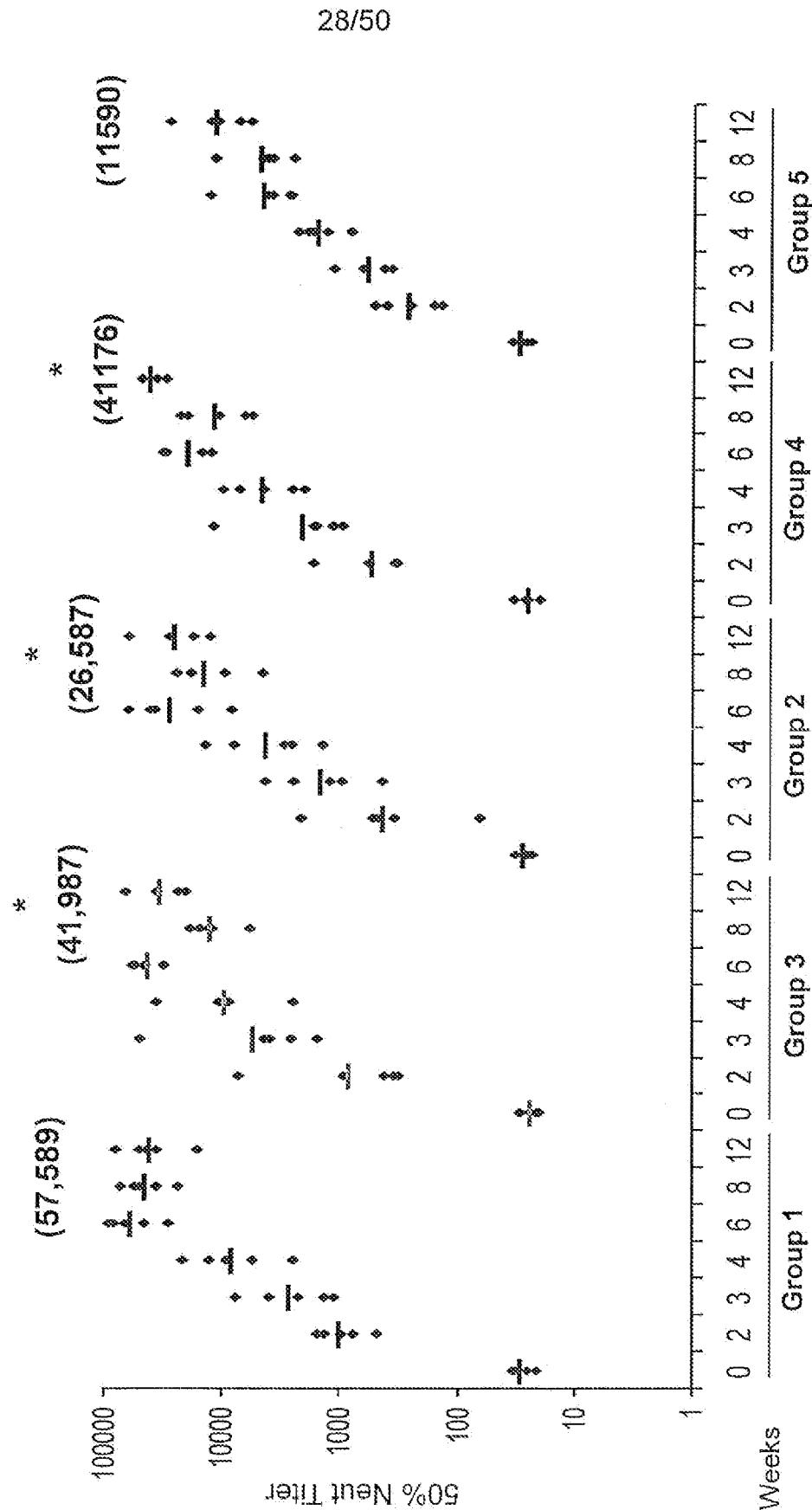
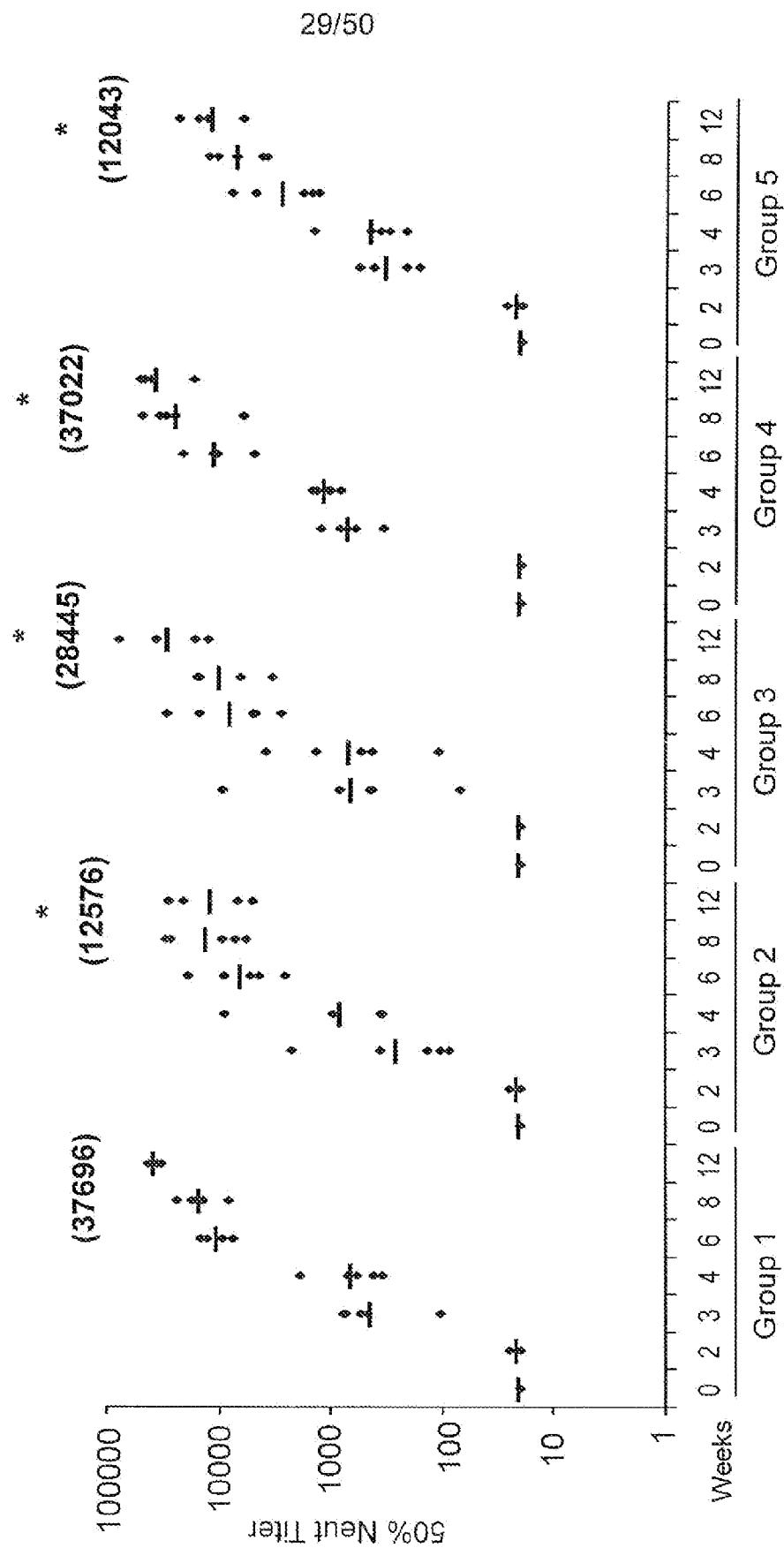


FIG. 16B

Anti-Toxin B



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FIG. 17

Variable Light Chain (muK)

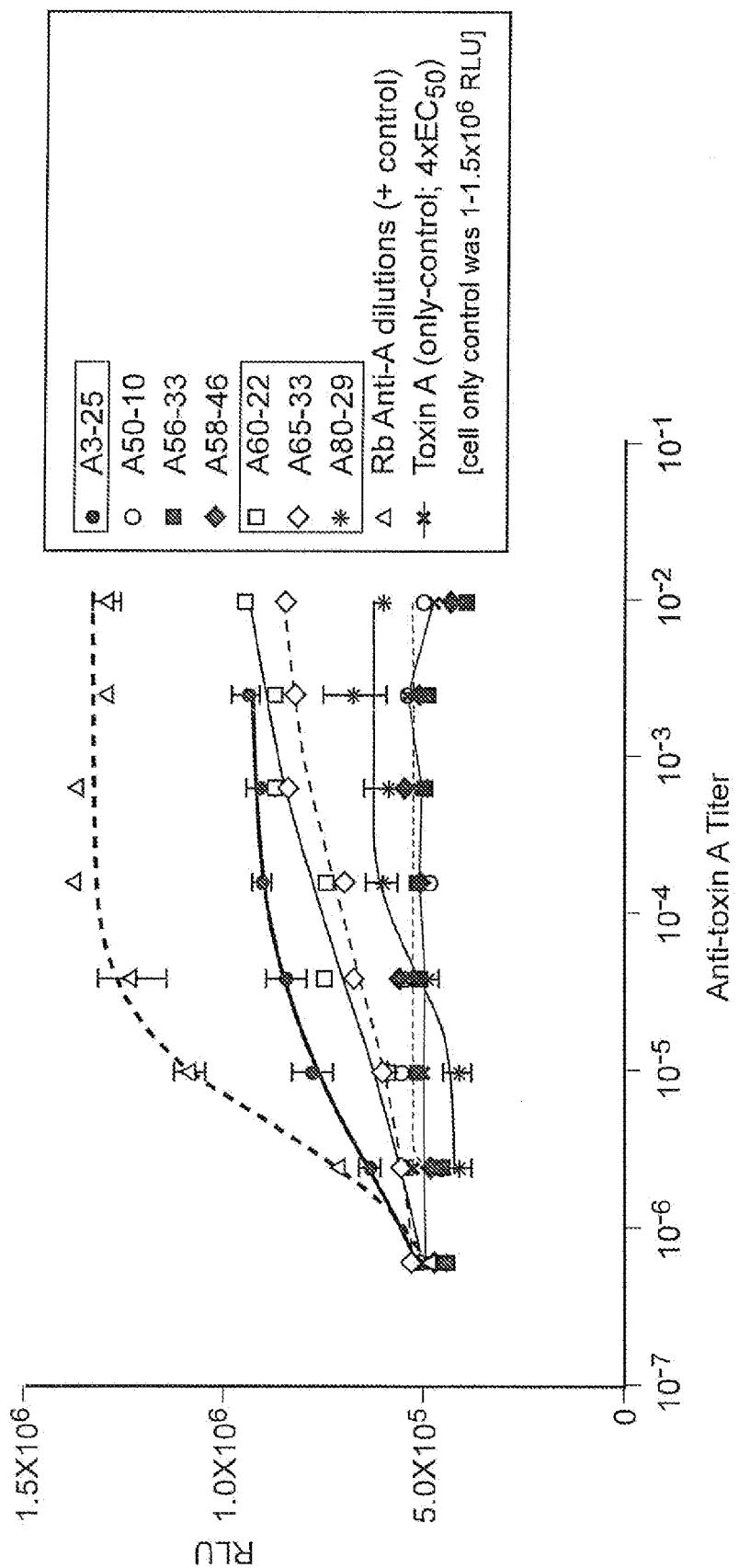
MKLPVRLLVLMFWIPGSSS DVVMTHTPLSLPVSLGDQASMSCRS
SQSLIHNGNTYLHWYLQKPGQSPKLLISKVSNRFSGVPDRFSG
SGSGTDFTLKISRVEAEDLGVYFCSQTTYFPYTFGGGTREIKRAD
AAPTVSIFPPSS (SEQ ID NO: 36)

Variable Heavy Chain (mlgE)

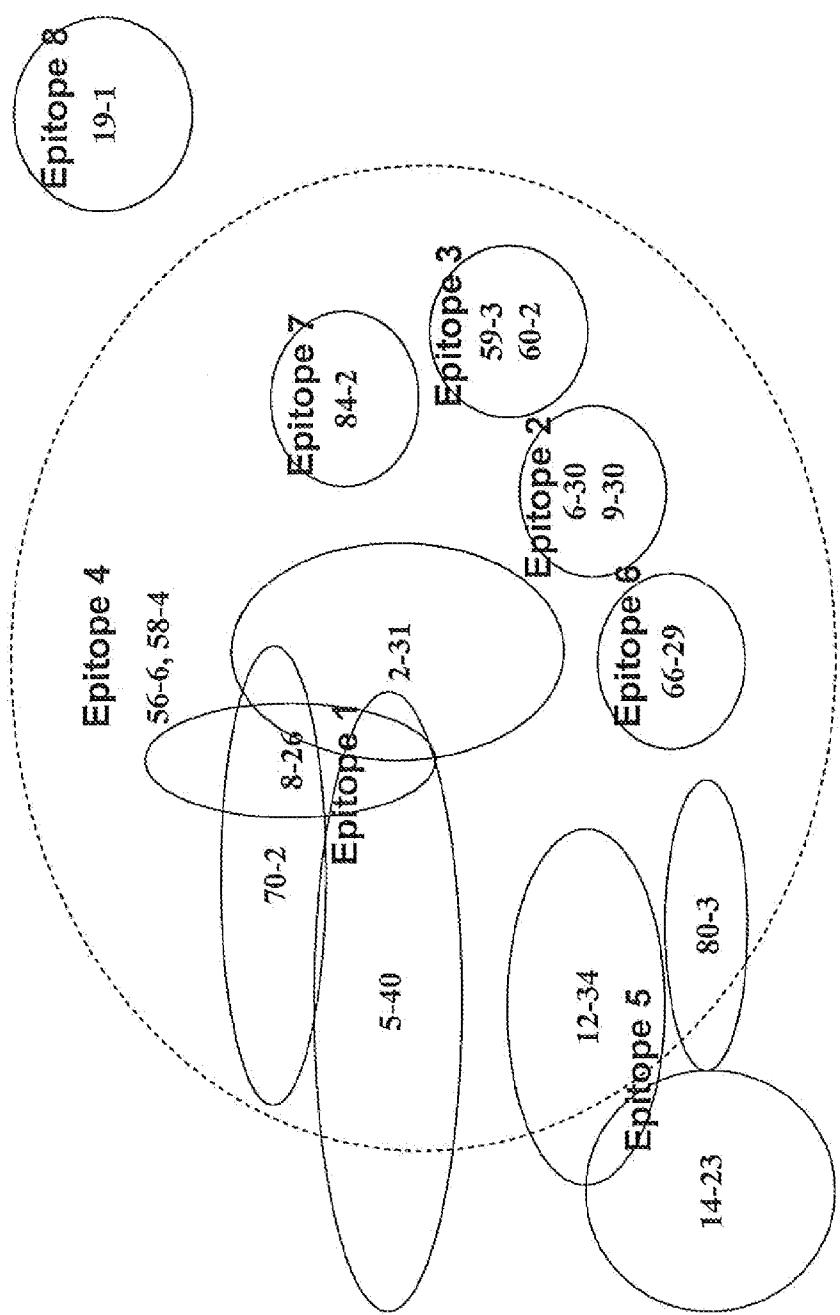
MYLGLNCVFIVFLLKGVQSEVNLEESGGGLVQPGGSMKLSCVAS
GFTFTNYWMNWVRQSPEKGLEWIAEIRLKSHNYATHFAESVKG
RFTISRDDSKSAVSLQMTNLTPEDTGIFYCTWDYYGNPAFVYWG
QGTLTVSAASIRNPQLYPLKPCKGTAMTLGCLVKDYFPGPVT
VTWYSDSLNMSTVN (SEQ ID NO: 37)

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FIG. 18

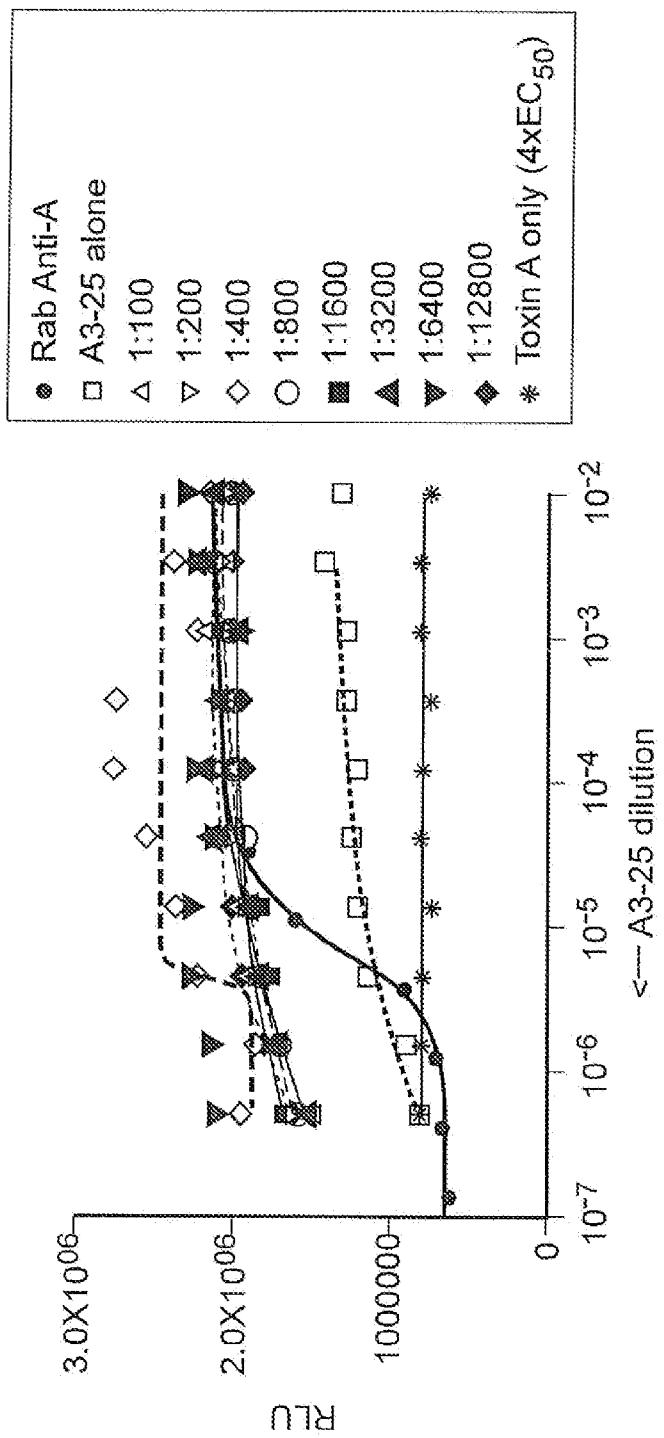


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FIG. 19

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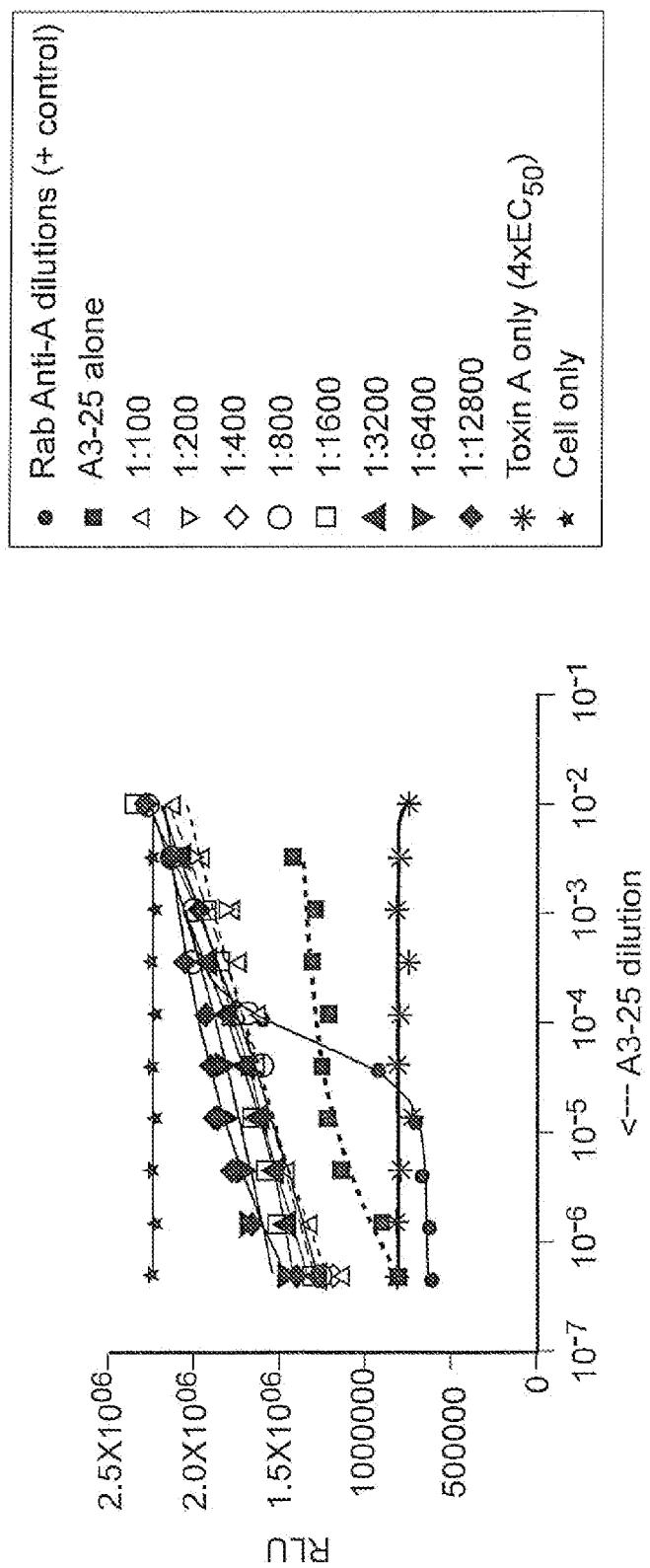
FIG. 20A



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FIG. 20B

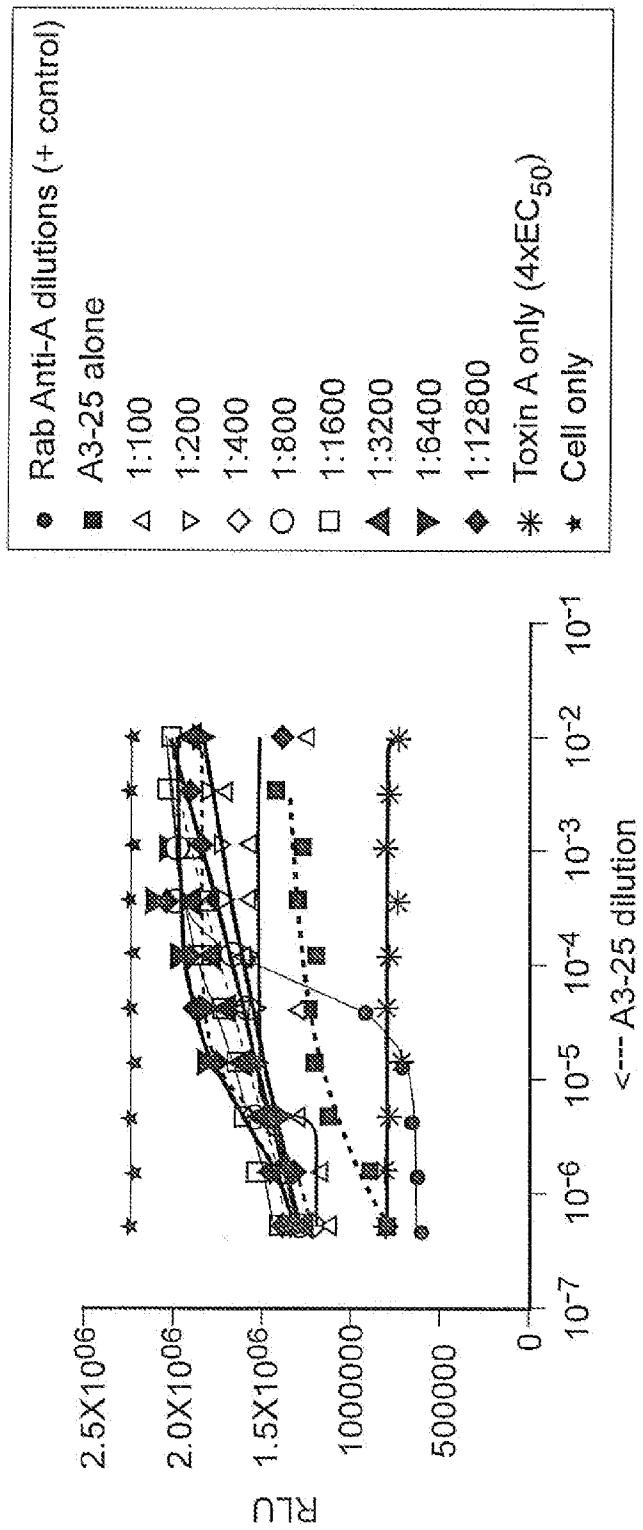
Toxin A Neutralization by mAb A3-25 plus A65-33 Dilutions



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FIG. 20C

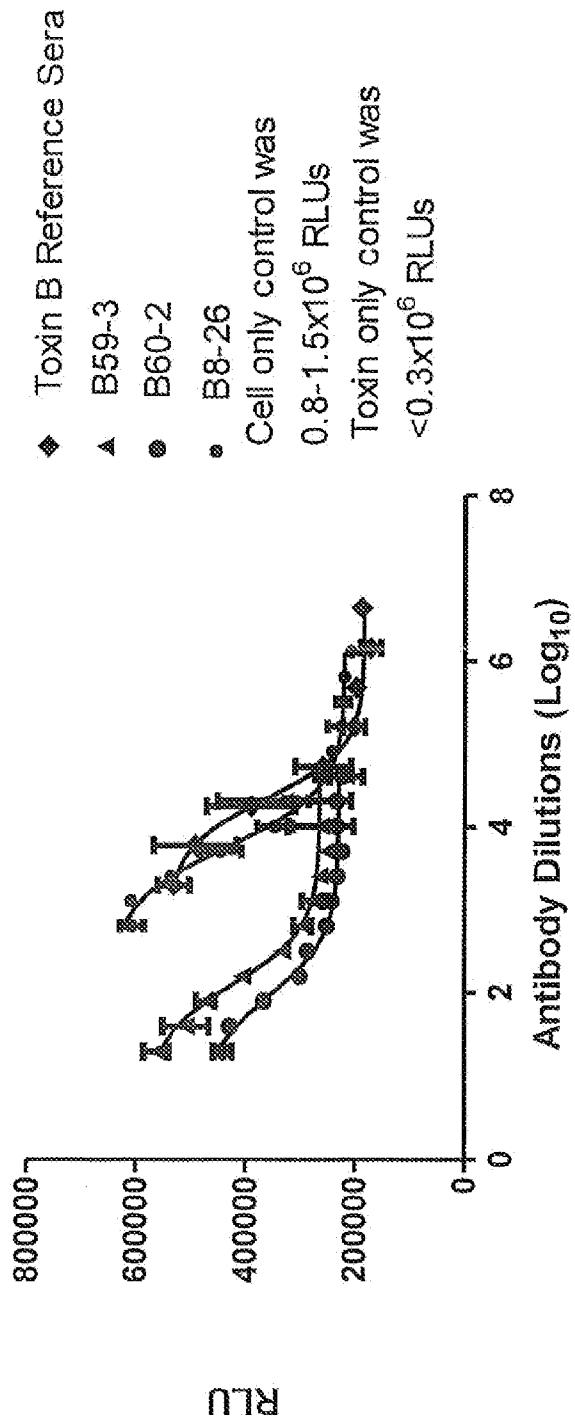
Toxin A Neutralization by mAb A3-25 plus A80-29 Dilutions



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FIG. 21A

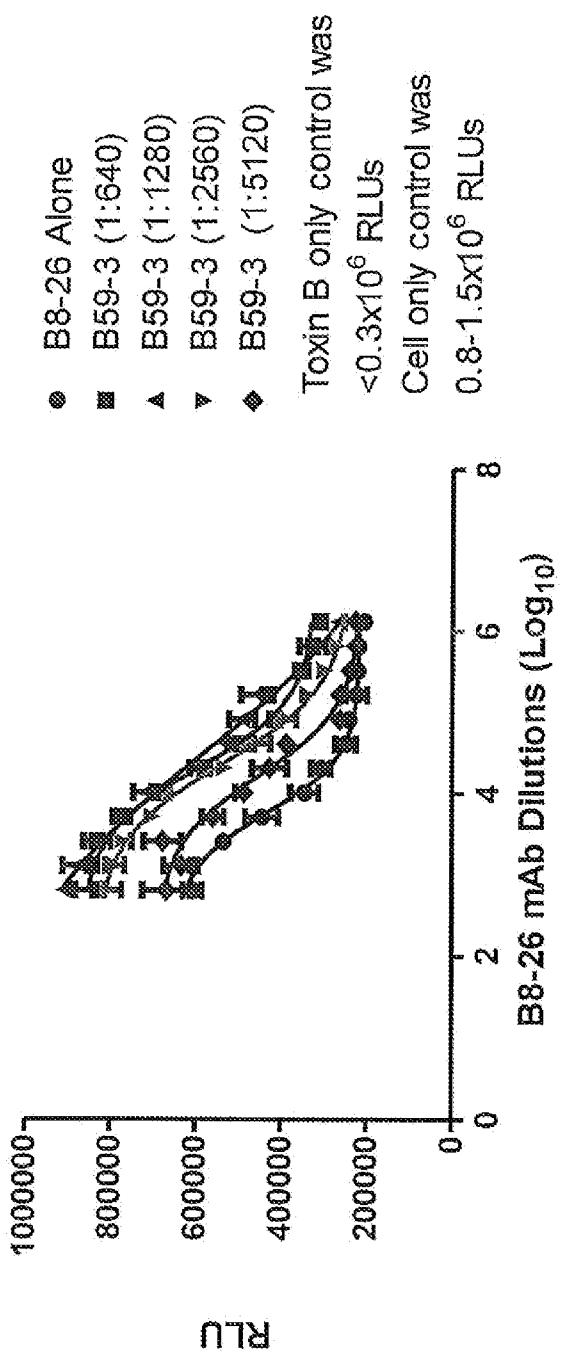
Toxin B Neutralization by Reference Sera and Different mAbs



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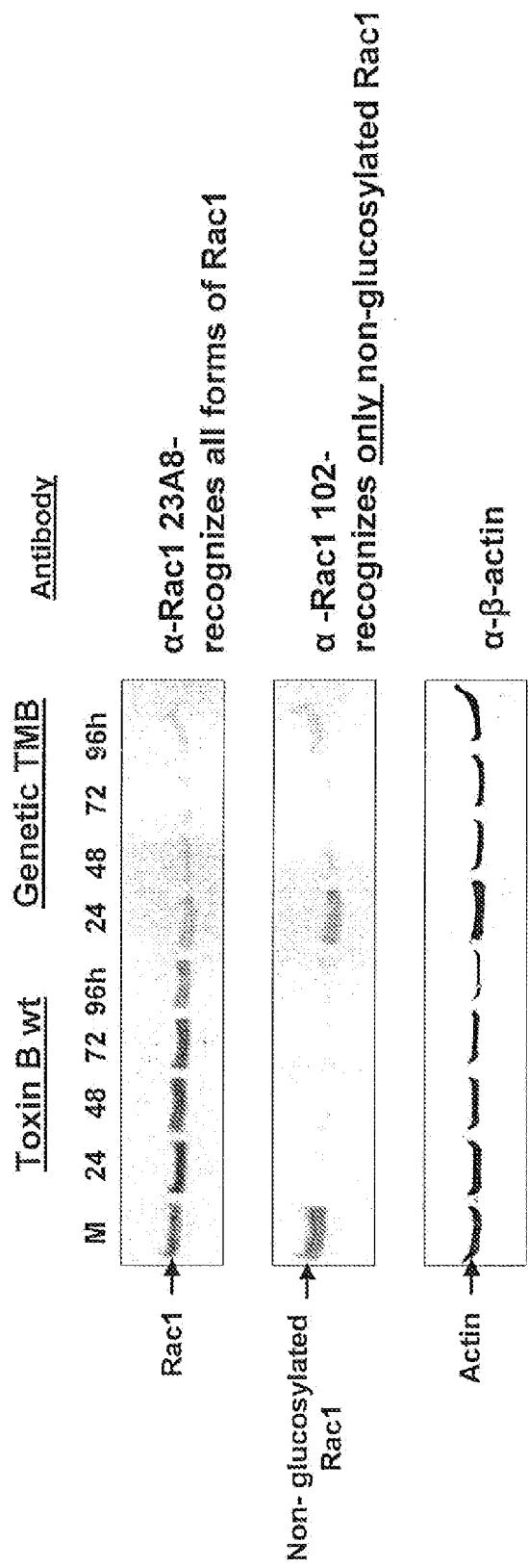
FIG. 21B

Toxin B Neutralization by B8-26 mAb plus B59-3 Dilutions



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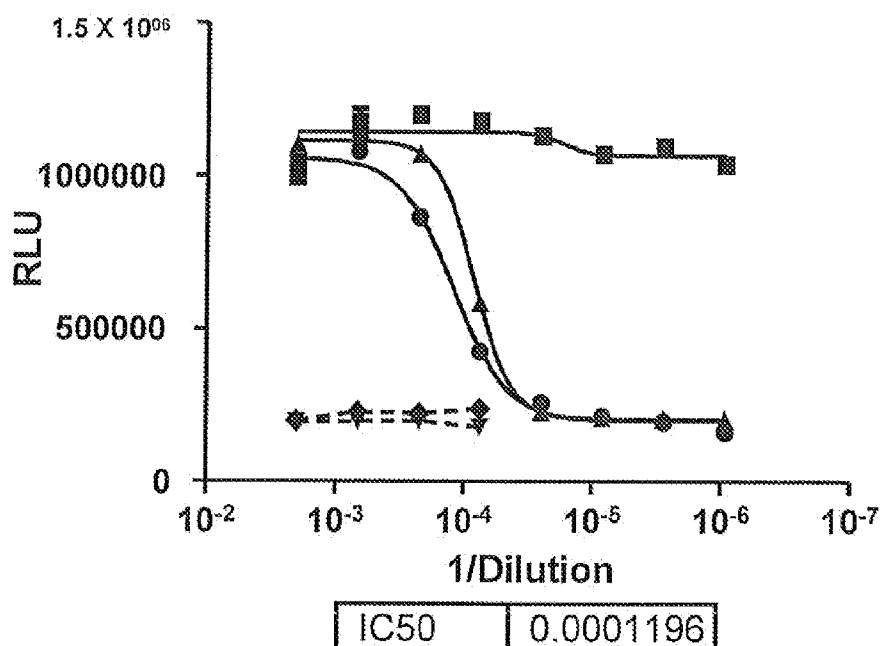
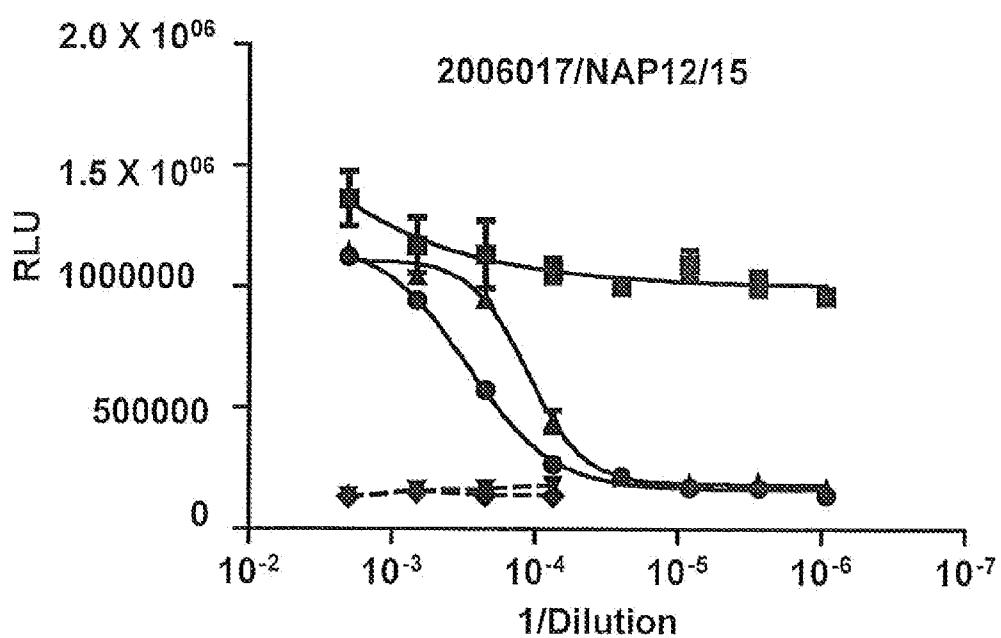
FIG. 22



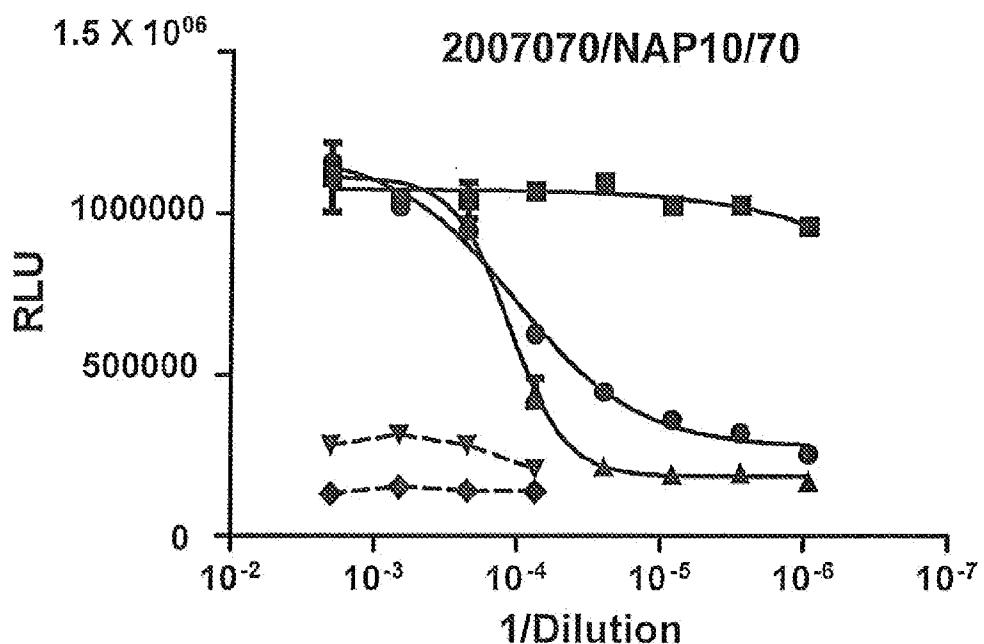
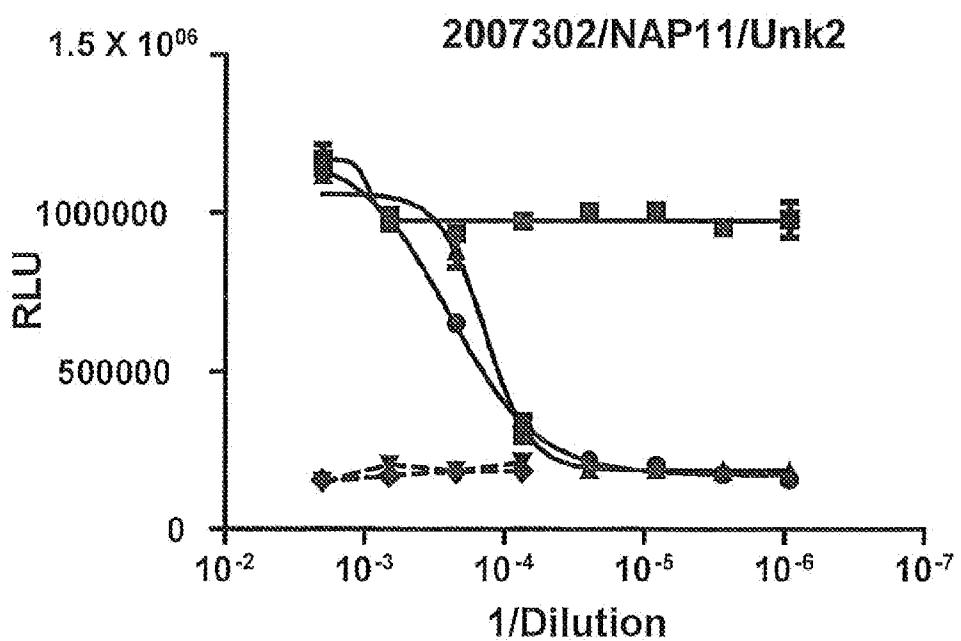
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FIG. 23A

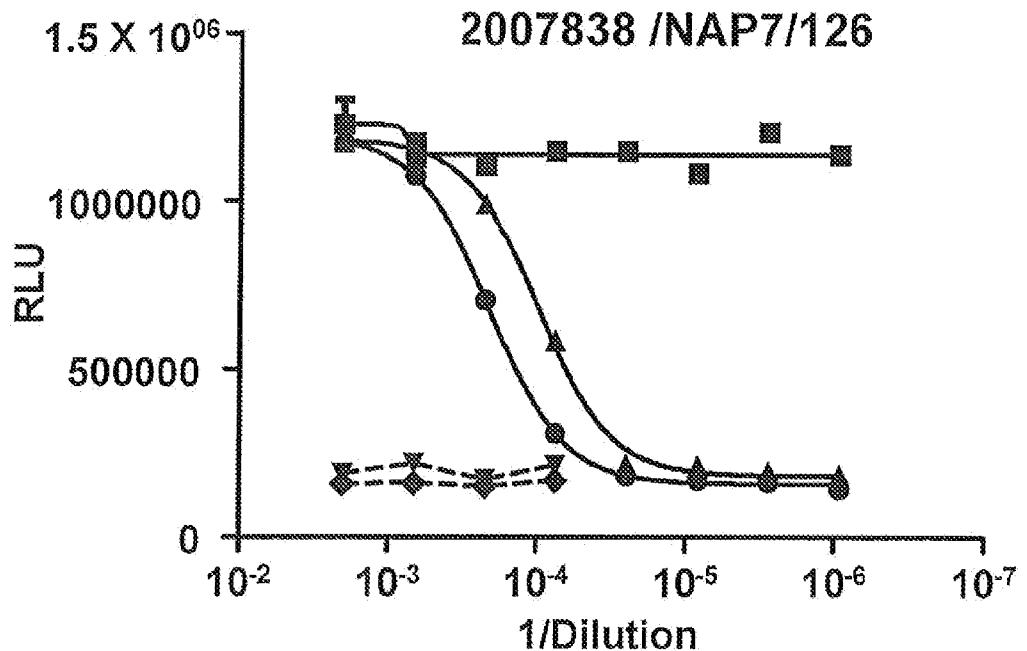
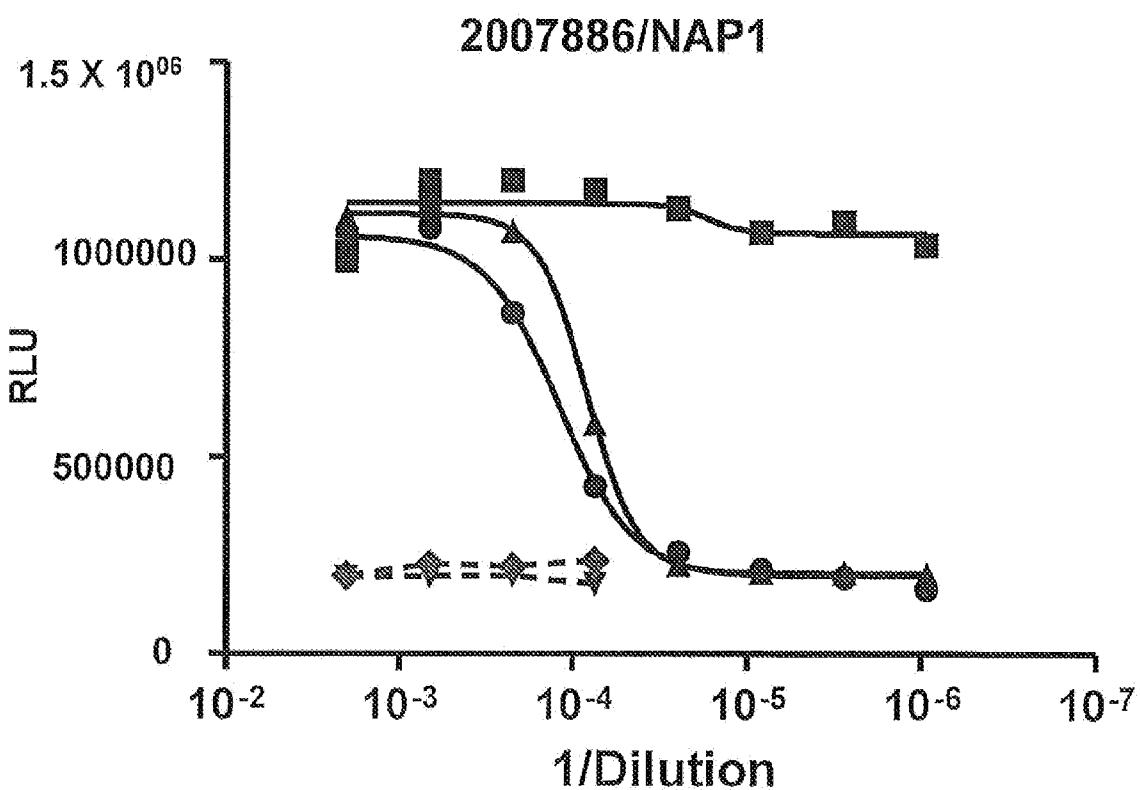
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**FIG. 23B**

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FIG. 23C**FIG. 23D**

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FIG. 23E**FIG. 23F**

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FIG. 23G

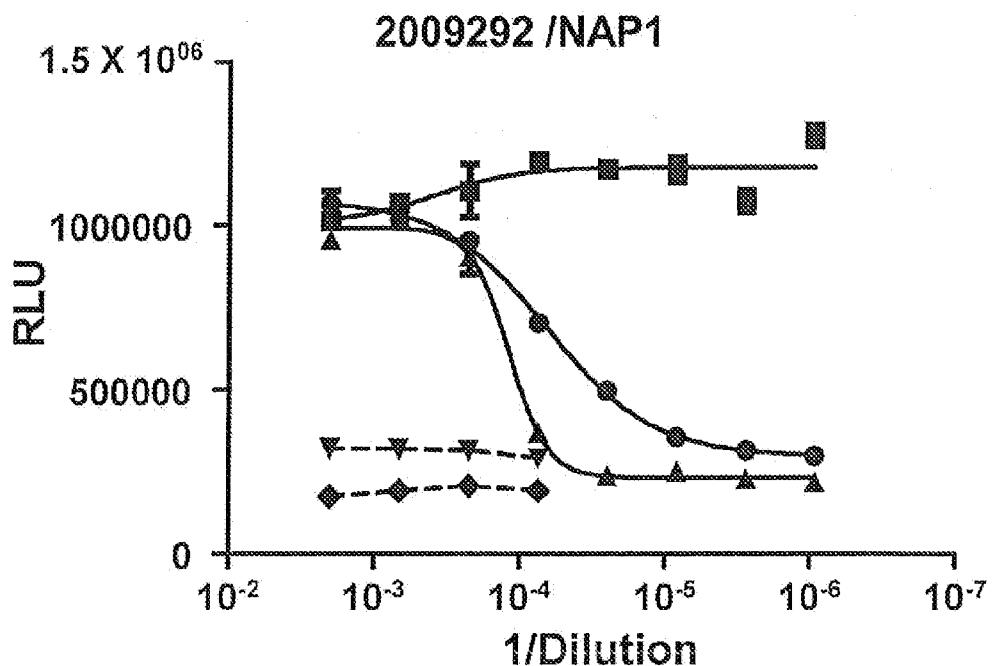
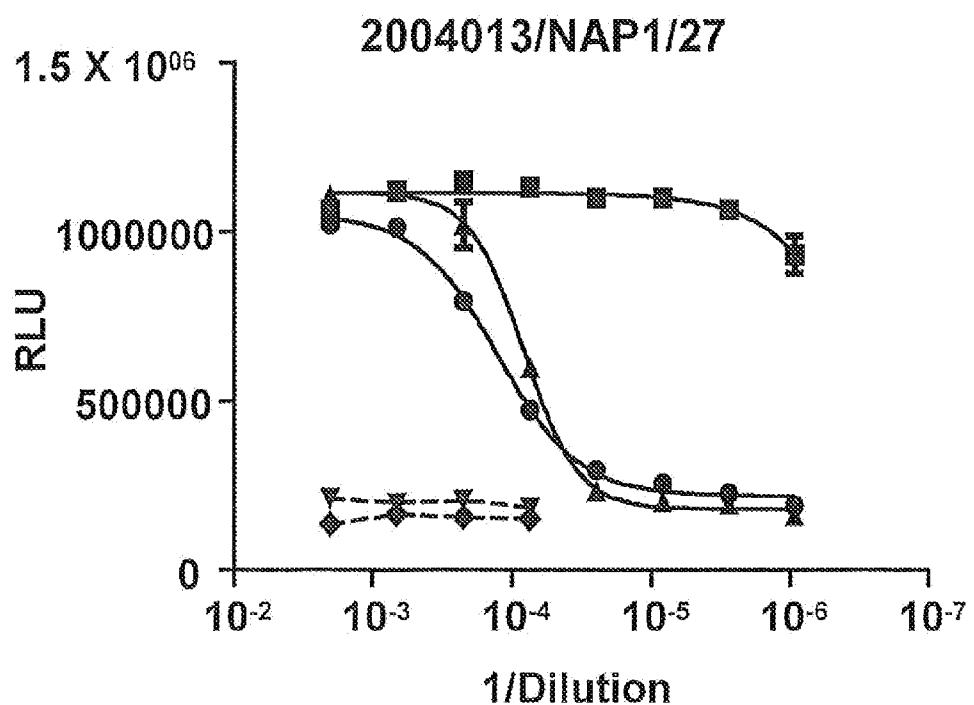
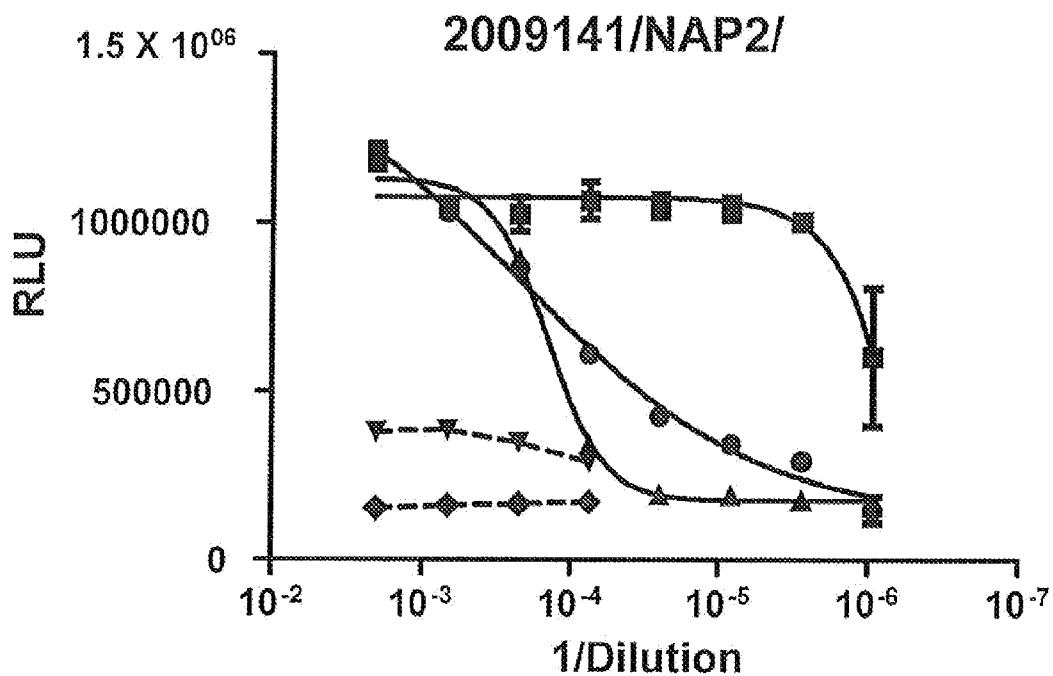
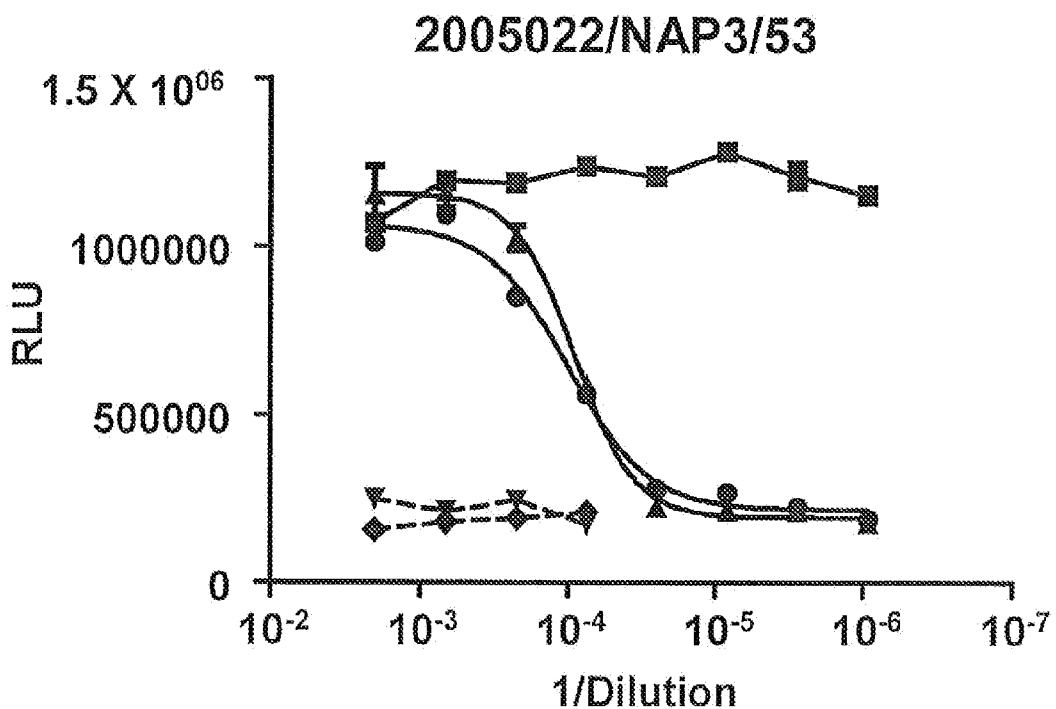


FIG. 23H

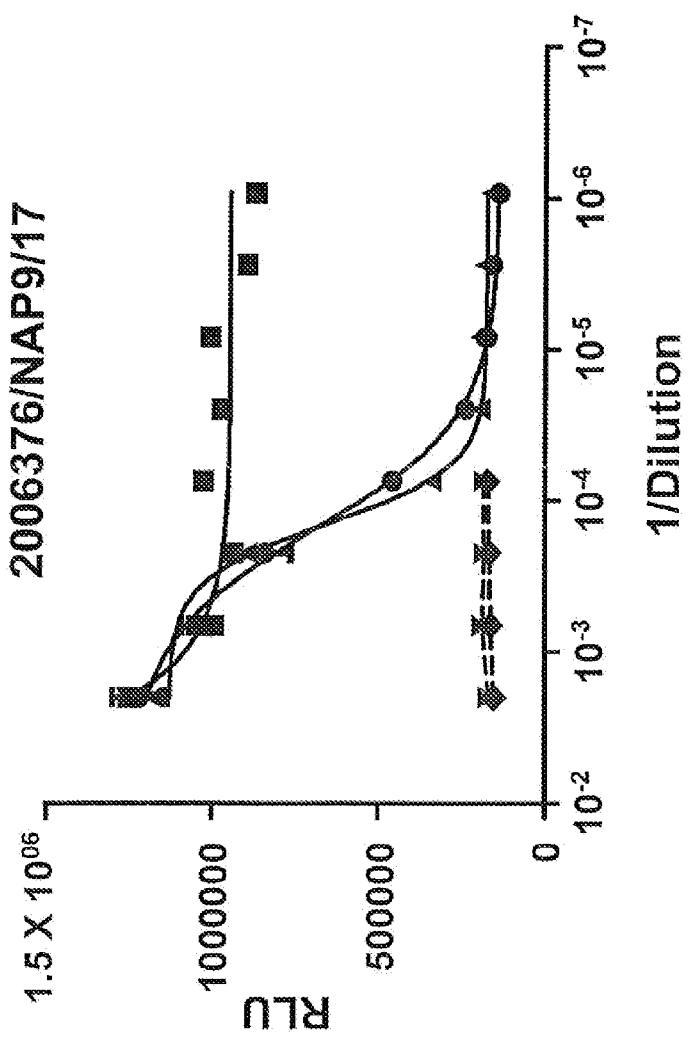


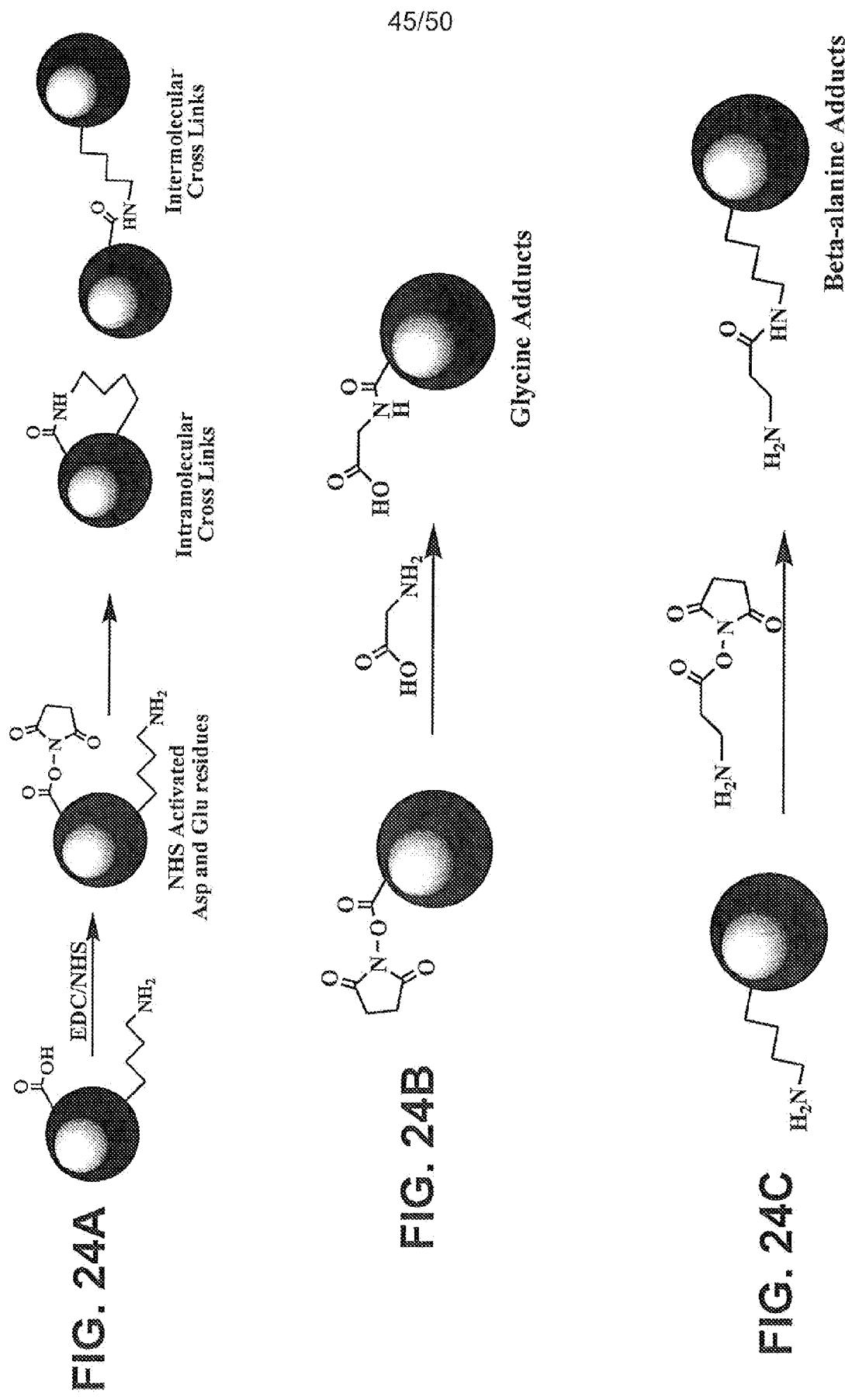
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FIG. 23 I**FIG. 23J**

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FIG. 23K





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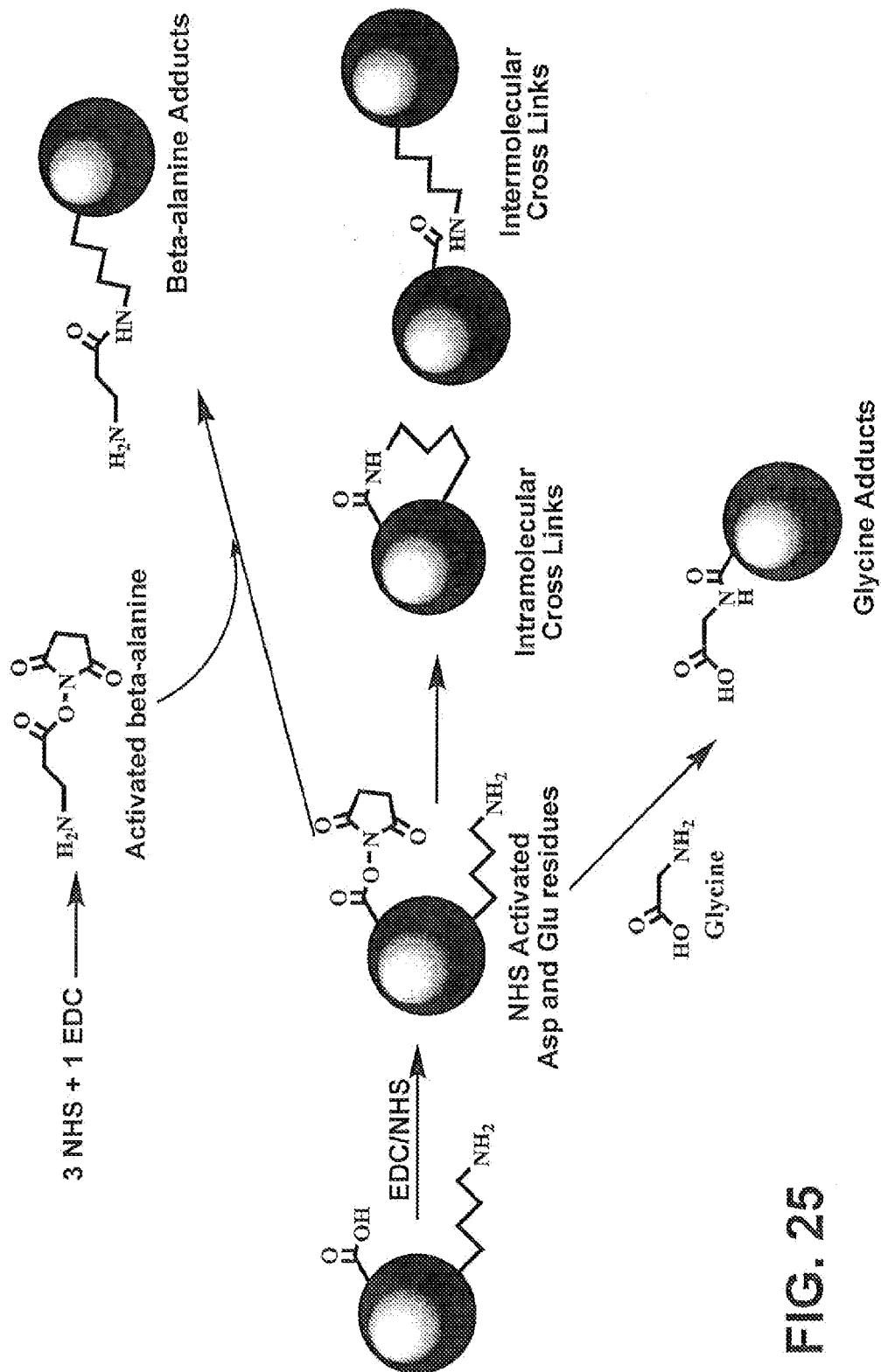
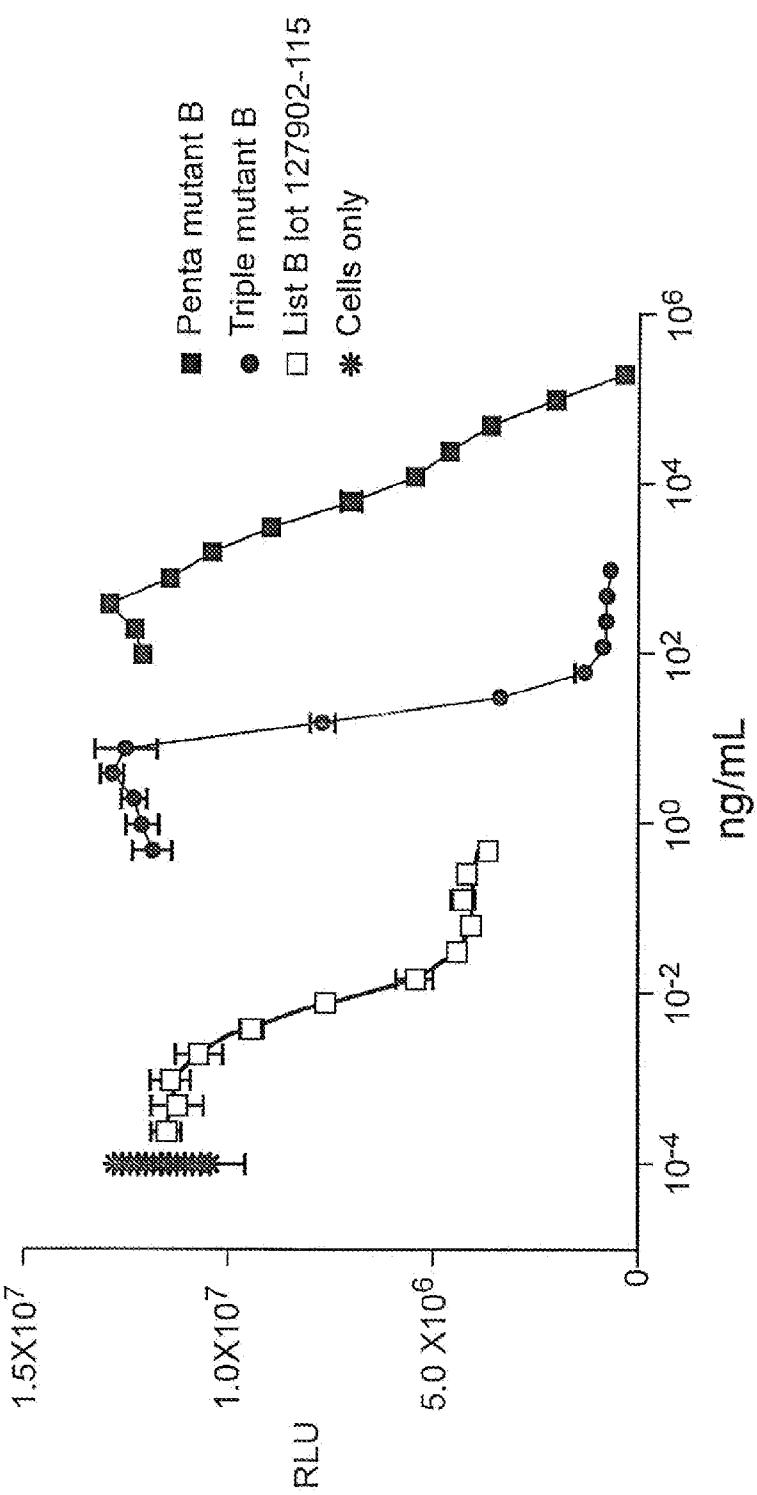


FIG. 25

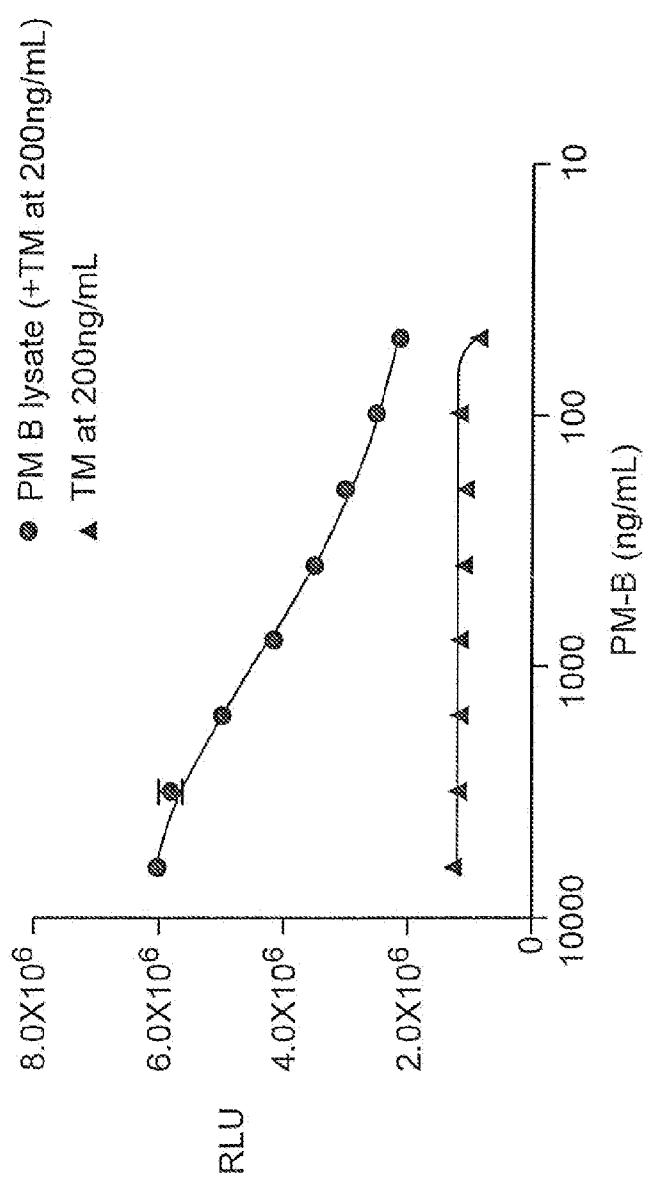
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FIG. 26



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FIG. 27



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FIG. 28

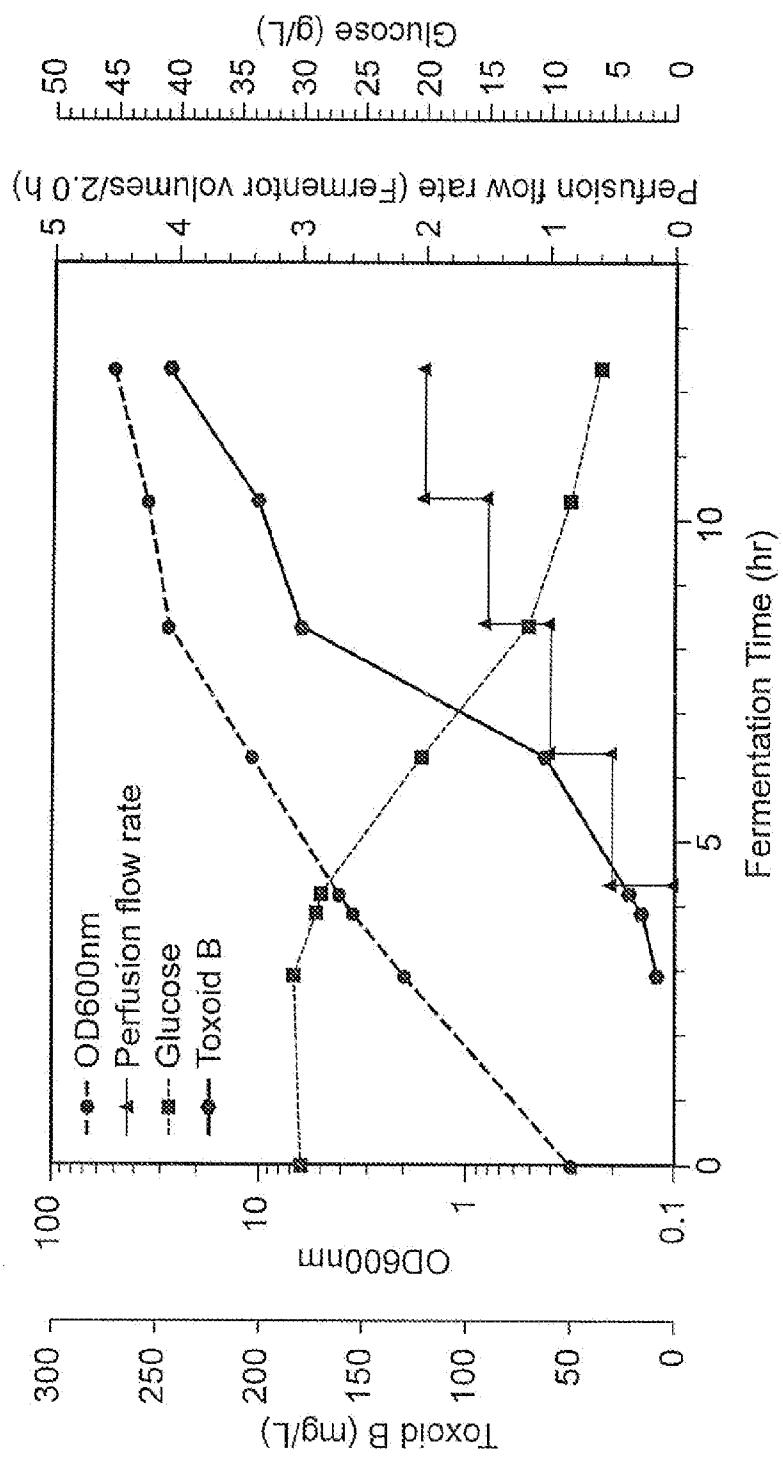


FIG. 29

