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(54) STABLE FORMULATIONS OF NEISSERIA MENINGITIDIS RLP2086 ANTIGENS

STABILE FORMULIERUNGEN VON NEISSERIA-MENINGITIDIS-RLP2086-ANTIGENEN

FORMULATIONS STABLES DES ANTIGÈNES RLP2086 DE NEISSERIA MENINGITIDIS

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- **MASIGNANI V. ET AL.: "Vaccination against Neisseria meningitidis using three variants of the lipoprotein GNA1870", THE JOURNAL OF EXPERIMENTAL MEDICINE, ROCKEFELLER UNIVERSITY PRESS, US, vol. 197, no. 6, 17 March 2003 (2003-03-17), pages 789 - 799, XP002286107, ISSN: 0022-1007, DOI: 10.1084/JEM.20021911**

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- HOUGHTEN R.A.: "GENERAL METHOD FOR THE RAPID SOLID-PHASE SYNTHESIS OF LARGE NUMBERS OF PEPTIDES: SPECIFICITY OF ANTIGEN-ANTIBODY INTERACTION AT THE LEVEL OF INDIVIDUAL AMINO ACIDS", PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF USA, NATIONAL ACADEMY OF SCIENCE, WASHINGTON, DC; US, vol. 82, 1 August 1995 (1995-08-01), pages 5131 - 5135, XP000611717, ISSN: 0027-8424, DOI: 10.1073/PNAS.82.15.5131

DescriptionFIELD OF THE INVENTION

5 **[0001]** The present invention relates to a method for stabilizing the potency of a LP2086 (fHBP) Subfamily A polypeptide and a LP2086 (fHBP) Subfamily B polypeptide in an immunogenic composition.

Background of the Invention

10 **[0002]** rLP2086 is a recombinant 28-kDa lipoprotein that induces cross-reactive bacterial antibodies against a number of *Neisseria meningitidis* strains. Based on deduced amino acid sequence homology, two different subfamilies of rLP2086 were identified, A and B. These two subfamilies were used in the formulation of the MnB-rLP2086 vaccine samples containing 20, 60, 120, and 200 µg/mL each in 10 mM Histidine (pH 6.0), 150 mM NaCl, and 0.5 mg/mL aluminum with varying levels of Polysorbate 80 (PS-80). Polysorbate 80, also known, as TWEEN 80, is a nonionic surfactant and emulsifier derived from sorbitol, and is frequently used in pharmaceutical formulations as an emulsifier, solubilizer and stabilizer. The presence of Polysorbate 80 in the MnB rLP2086 immunogenic composition is believed to prevent aggregation during formulation, processing, filtration, filling and shipping, reduce filter membrane absorption, and reduce tubing absorption.

20 Summary of the Invention

[0003] The invention relates to a method for stabilizing the potency of a LP2086 (fHBP) Subfamily A polypeptide and a LP2086 (fHBP) Subfamily B polypeptide in an immunogenic composition, by storing the LP2086 (fHBP) Subfamily A polypeptide and the LP2086 (fHBP) Subfamily B polypeptide in a buffer with between about 0.1 mg/ml and about 1 mg/ml aluminum and a molar ratio of detergent to protein of between about 0.5:1 and about 10:1, wherein the immunogenic composition does not include further active immunogens. In some embodiments, the present invention provides a stable immunogenic composition, wherein the potency of a LP2086 Subfamily B polypeptide is maintained for at least about 1-12 months, about 6-18 months, about 12-24 months, about 24-36 months, or about 36-48 months. The immunogenic composition further comprises a LP2086 Subfamily A polypeptide.

30 **[0004]** The immunogenic composition further comprises a detergent. In some embodiments, the molar ratio of the detergent to protein is between about 0.5:1 and about 10:1; between about 1:1 and about 5:1; or between about 1.4:1 and 4.2:1. In some embodiments, the molar ratio of the detergent to protein is about 2.8:1. In some embodiments the amount of detergent is sufficient to reduce polypeptide binding to silicon in a container, such as a syringe or a vial. In some embodiments, the detergent is a non-ionic detergent, such as a polysorbate detergent. In some embodiments, the detergent is Polysorbate-80.

35 **[0005]** The immunogenic composition further comprises aluminum. In some embodiments, the immunogenic composition comprises aluminum as aluminum phosphate, aluminum hydroxide, aluminum sulfate, or alum. The concentration of aluminum is between about 0.1 mg/mL and 1.0 mg/mL. In some embodiments, the concentration of aluminum is about 0.5 mg/mL.

40 **[0006]** In some embodiments, the immunogenic composition further comprises histidine. In some embodiments, the concentration of histidine is between about 2mM and about 20mM or between about 5mM and about 15mM. In some embodiments, the concentration of histidine is about 10mM. In some embodiments, the pH of the histidine is between about 5.0 and about 8.0 or between about 5.8 and about 6.0. In some embodiments, the concentration of histidine is 10mM, pH 6.0.

45 **[0007]** In some embodiments, the immunogenic composition further comprises succinate. In some embodiments, the concentration of succinate is between about 2mM and about 10mM or between about 3mM and about 7mM. In some embodiments, the concentration of succinate is about 5mM. In some embodiments, the pH of the succinate is between about 5.0 and about 8.0 or between about 5.8 and about 6.0. In some embodiments, the concentration of succinate is 5mM, pH 6.0.

50 **[0008]** In some embodiments, the immunogenic composition is lyophilized. In some embodiments, the lyophilized composition is resuspended in a buffer comprising aluminum. In some embodiments, the aluminum is present as aluminum phosphate, aluminum hydroxide, aluminum sulfate, or alum.

55 **[0009]** In some embodiments, the immunogenic composition comprises about a 2.8:1 molar ratio of Polysorbate 80 to protein, 0.5 mg/mL aluminum as AlPO₄, 10mM histidine pH 6.0, and 150 mM NaCl. In some embodiments, the immunogenic composition consists essentially of 200ug/mL LP2086 (fHBP) Subfamily B polypeptide, about a 2.8:1 molar ratio of Polysorbate 80 to protein, 0.5 mg/mL aluminum as AlPO₄, 10mM histidine pH 6.0, and 150 mM NaCl. In some embodiments, the immunogenic composition consists essentially of 200ug/mL rLP2086 (fHBP) Subfamily A polypeptide, 200ug/mL LP2086 (fHBP) Subfamily B polypeptide, about a 2.8:1 molar ratio of Polysorbate 80 to protein, 0.5 mg/mL

of aluminum as AlPO₄, 10mM histidine pH 6.0, and 150 mM NaCl.

[0010] In some embodiments, the immunogenic composition comprises about a 2.8:1 molar ratio of Polysorbate 80 to protein, 0.5 mg/mL aluminum as AlPO₄, 5mM succinate pH 6.0, and 150 mM NaCl. In some embodiments, the immunogenic composition consists essentially of 200ug/mL LP2086 (fHBP) Subfamily B polypeptide, about a 2.8:1 molar ratio of Polysorbate 80 to protein, 0.5 mg/mL aluminum as AlPO₄, 5mM succinate pH 6.0, and 150 mM NaCl. In some embodiments, the immunogenic composition consists essentially of 200ug/mL rLP2086 (fHBP) Subfamily A polypeptide, 200ug/mL LP2086 (fHBP) Subfamily B polypeptide, about a 2.8:1 molar ratio of Polysorbate 80 to protein, 0.5 mg/mL of aluminum as AlPO₄, 5mM succinate pH 6.0, and 150 mM NaCl.

[0011] In another aspect, the invention provides for a method of stabilizing the potency of a LP2086 Subfamily A polypeptide and a LP2086 Subfamily B polypeptide in an immunogenic composition by storing the LP2086 Subfamily A polypeptide and the LP2086 Subfamily B polypeptide in a buffer with between about 0.1mg/mL and about 10mg/mL aluminum and a molar ratio of detergent to protein between about 0.5:1 and 10:1. In some embodiments, the molar ratio of detergent to protein is between about 1:1 and about 5:1; or between about 1.4:1 and about 4.2:1. In some embodiments, the molar ratio of detergent to protein is about 2.8:1. In some embodiments the amount of detergent is sufficient to reduce polypeptide binding to silicon in a container, such as a syringe or a vial. In some embodiments, the detergent is a non-ionic detergent, such as a polysorbate detergent. In some embodiments, the detergent is Polysorbate-80.

[0012] In some embodiments, the aluminum is present as aluminum phosphate, aluminum hydroxide, aluminum sulfate, or alum. In some embodiments, the concentration of aluminum is about 0.5 mg/mL.

[0013] In some embodiments, the buffer further comprises histidine. In some embodiments, the concentration of histidine is between about 2mM and about 20mM or between about 5mM and about 15mM. In some embodiments, the concentration of histidine is about 10mM. In some embodiments, the pH of the histidine is between about 5.0 and about 8.0 or between about 5.8 and about 6.0. In some embodiments, the concentration of histidine is 10mM, pH 6.0.

[0014] In some embodiments, the buffer further comprises succinate. In some embodiments, the concentration of succinate is between about 2mM and about 10mM or between about 3mM and about 7mM. In some embodiments, the concentration of succinate is about 5mM. In some embodiments, the pH of the succinate is between about 5.0 and about 8.0 or between about 5.8 and about 6.0. In some embodiments, the concentration of succinate is 10mM, pH 6.0.

[0015] In some embodiments, the immunogenic composition is lyophilized. In some embodiments, the lyophilized composition is resuspended in a buffer comprising aluminum. In some embodiments, the aluminum is present as aluminum phosphate, aluminum hydroxide, aluminum sulfate, or alum.

[0016] In some embodiments, the buffer consists essentially of about a 2.8:1 molar ratio of Polysorbate 80 to protein, 0.5 mg/mL aluminum as AlPO₄, 10mM histidine pH 6.0, and 150 mM NaCl. In some embodiments, the immunogenic composition consists essentially of 200ug/mL LP2086 (fHBP) Subfamily A polypeptide, 200ug/mL LP2086 (fHBP) Subfamily B polypeptide, about a 2.8:1 molar ratio of Polysorbate 80 to protein, 0.5 mg/mL aluminum as AlPO₄, 10mM histidine pH 6.0, and 150 mM NaCl.

[0017] In some embodiments, the buffer consists essentially of about a 2.8:1 molar ratio of Polysorbate 80 to protein, 0.5 mg/mL aluminum as AlPO₄, 5mM succinate pH 6.0, and 150 mM NaCl. In some embodiments, the immunogenic composition consists essentially of 200ug/mL LP2086 (fHBP) Subfamily A polypeptide, 200ug/mL LP2086 (fHBP) Subfamily B polypeptide, about a 2.8:1 molar ratio of Polysorbate 80 to protein, 0.5 mg/mL aluminum as AlPO₄, 5mM succinate pH 6.0, and 150 mM NaCl.

Brief Description of the Drawings

[0018]

- Figure 1: Stability of Subfamily B in Formulations with Various Polysorbate 80 Concentrations.
- Figure 2: Accelerated Stability of Subfamily B with Various Polysorbate 80 Concentrations
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- Figure 5: Potency Results for 200 μg/mL with Different Molar Ratios
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- Figure 23: Potency and Bound Molar Ratio Results for Subfamily A
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- Figure 25: Binding of Subfamily A with AlPO₄ in Succinate and Histidine Buffers
- Figure 26: Binding of Subfamily B with AlPO₄ in Succinate and Histidine Buffers
- Figure 27: Comparison of Binding in Succinate, Histidine, and Phosphate buffer
- Figure 28: pH-Dependent Binding of Subfamily A with AlPO₄
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Detailed Description of the Invention

20 **[0019]** Unless defined otherwise, all technical and scientific terms used herein have the same meaning as those commonly understood by one of ordinary skill in the art to which this invention belongs. Although methods and materials similar or equivalent to those described herein can be used in the practice or testing of the present invention, suitable methods and materials are described below. The materials, methods and examples are illustrative only, and are not intended to be limiting. All publications, patents and other documents mentioned herein are incorporated by reference in their entirety.

25 **[0020]** Throughout this specification, the word "comprise" or variations such as "comprises" or "comprising" will be understood to imply the inclusion of a stated integer or groups of integers but not the exclusion of any other integer or group of integers.

Definitions

30 **[0021]** As used herein, the singular forms "a," "an," and "the" include plural references unless the context clearly dictates otherwise. Thus, *e.g.*, references to "the method" includes one or more methods, and/or steps of the type described herein and/or which will become apparent to one of ordinary skill in the art upon reading this disclosure and so forth.

35 **[0022]** As used herein, the plural forms include singular references unless the context clearly dictates otherwise. Thus, *e.g.*, references to "the methods" includes one or more methods, and/or steps of the type described herein and/or which will become apparent to one of ordinary skill in the art upon reading this disclosure and so forth.

40 **[0023]** As used herein, "about" means within a statistically meaningful range of a value such as a stated concentration range, time frame, molecular weight, temperature or pH. Such a range can be within an order of magnitude, typically within 20%, more typically still within 10%, and even more typically within 5% of a given value or range. The allowable variation encompassed by the term "about" will depend upon the particular system under study, and can be readily appreciated by one of ordinary skill in the art. Whenever a range is recited within this application, every whole number integer within the range is also contemplated as an embodiment of the invention.

45 **[0024]** The term "adjuvant" refers to a compound or mixture that enhances the immune response to an antigen as further described and exemplified herein. Non-limiting examples of adjuvants that can be used in the vaccine of the present invention include the RIBI adjuvant system (Ribi Inc., Hamilton, Mont.), alum, mineral gels such as aluminum hydroxide gel, oil-in-water emulsions, water-in-oil emulsions such as, *e.g.*, Freund's complete and incomplete adjuvants, Block copolymer (CytRx, Atlanta Ga.), QS-21 (Cambridge Biotech Inc., Cambridge Mass.), SAF-M (Chiron, Emeryville Calif.), AMPHIGEN[®] adjuvant, saponin, Quil A or other saponin fraction, monophosphoryl lipid A, and Avridine lipid-amine adjuvant.

50 **[0025]** The term "binding of aluminum to protein" refers to the percentage of protein molecules in a composition that are bound to aluminum. The binding of aluminum to protein can be determined using methods disclosed herein or known in the art.

55 **[0026]** The term "effective immunogenic amount" as used herein refers to an amount of a polypeptide or composition comprising a polypeptide which is effective in eliciting an immune response in a vertebrate host. For example, an effective immunogenic amount of a rLP2086 protein of this invention is an amount that is effective in eliciting an immune response in a vertebrate host. The particular "effective immunogenic dosage or amount" will depend upon the age, weight and medical condition of the host, as well as on the method of administration. Suitable doses are readily determined by

persons skilled in the art.

[0027] The term "molar ratio" as used herein refers to the ratio of the number of moles of two different elements in a composition. In some embodiments, the molar ratio is the ratio of moles of detergent to moles of protein. In some embodiments, the molar ratio is the ratio of moles of Polysorbate 80 to moles of protein. Based on the protein and Polysorbate 80 concentrations, the Molar Ratio is calculated using the following equation:

$$\text{Molar Ratio} = \frac{\% \text{PS80}}{\text{mg/mL Protein}} \times 216$$

For example, a composition comprising 0.01% Polysorbate 80 and 200 μ g has a detergent-to-protein molar ratio of 10.8:1 [(0.01/0.2) \times 216]. A ratio of 3 moles Polysorbate 80 to 2 moles of protein would be expressed as a molar ratio of PS80 to protein of 3:2. Further, if a molar ratio is recited as a single number, it refers to a ratio of that single number to 1. For example, Polysorbate 80-to-protein ratios of 0.5, 2, and 10 refer to ratios of 0.5:1, 2:1 and 10:1, respectively. As used herein, the terms "detergent to protein" molar ratio and "Polysorbate 80 to protein" molar ratio refer in general to the molar ratio of detergent (or Polysorbate 80) to protein antigens, particularly P2086 antigens. Based on the teachings disclosed herein, one of skill in the art would be able to determine how to calculate molar ratios for other detergents and the optimal molar ratio for formulations with other detergents. As used herein, a "low" molar ratio refers in general to a molar ratio of the detergent to protein antigen in the immunogenic composition that is less than a "high" molar ratio. A "high" molar ratio refers in general to a molar ratio of the detergent to protein antigen in the immunogenic composition that is greater than a "low" molar ratio. In some embodiments, a "high molar ratio" of detergent to protein refers to a molar ratio greater than 10:1. In some embodiments, a "low molar ratio" of detergent to protein refer to a molar ratio between 0.5:1 and 10:1.

[0028] The term "ORF2086" as used herein refers to Open Reading Frame 2086 from a *Neisseria* species bacteria. *Neisseria* ORF2086, the proteins encoded therefrom, fragments of those proteins, and immunogenic compositions comprising those proteins are known in the art and are described, e.g., in U.S. Patent Application Publication Nos. US 20060257413 and US 20090202593, each of which is hereby incorporated by reference in its entirety. The term "P2086" generally refers to the protein encoded by ORF2086. The P2086 proteins of the invention may be lipidated or non-lipidated. "LP2086" and "P2086" typically refer to lipidated and non-lipidated forms of a 2086 protein, respectively. The P2086 protein of the invention may be recombinant. "rLP2086" and "rP2086" typically refer to lipidated and non-lipidated forms of a recombinant 2086 protein, respectively.

[0029] The term "pharmaceutically acceptable carrier" as used herein is intended to include any and all solvents, dispersion media, coatings, antibacterial and antifungal agents, isotonic and absorption delaying agents, and the like, compatible with administration to humans or other vertebrate hosts. Typically, a pharmaceutically acceptable carrier is a carrier approved by a regulatory agency of a Federal, a state government, or other regulatory agency, or listed in the U.S. Pharmacopeia or other generally recognized pharmacopeia for use in animals, including humans as well as non-human mammals. The term "carrier" refers to a diluent, adjuvant, excipient, or vehicle with which the pharmaceutical composition is administered. Such pharmaceutical carriers can be sterile liquids, such as water and oils, including those of petroleum, animal, vegetable or synthetic origin. Water, saline solutions and aqueous dextrose and glycerol solutions can be employed as liquid carriers, particularly for injectable solutions. Suitable pharmaceutical excipients include starch, glucose, lactose, sucrose, gelatin, malt, rice, flour, chalk, silica gel, sodium stearate, glycerol monostearate, talc, sodium chloride, dried skim milk, glycerol, propylene, glycol, water, ethanol and the like. The composition, if desired, can also contain minor amounts of wetting, bulking, emulsifying agents, or pH buffering agents. These compositions can take the form of solutions, suspensions, emulsion, sustained release formulations and the like. Examples of suitable pharmaceutical carriers are described in "Remington's Pharmaceutical Sciences" by E. W. Martin. The formulation should suit the mode of administration. The appropriate carrier will be evident to those skilled in the art and will depend in large part upon the route of administration.

[0030] The term "potency" refers to an antigen's ability to raise an immunogenic response. In some embodiments, potency is measured by an epitopes ability to bind to an antibody. Potency may be lost or reduced over time due to loss of antigen or epitope integrity or a change in antigen or epitope conformation. Potency may be lost or reduced due to factors including, but not limited to, light, temperature, freeze/thaw cycles, agitation, and pH. Potency can be measured by the methods disclosed herein and by assays known in the art. Such potency determination assays include, but are not limited to, animal vaccination models, serum bactericidal assays (SBA), flow cytometry, and in vitro potency assays. The preferred methods for determining potency are SBA and in vitro potency assays. A more preferred method for determining potency is SBA. In some embodiments, potency can be determined using at least one monoclonal antibody directed against at least one epitope that is involved in immune response. In some embodiments, potency of a test sample is compared against potency of a reference standard. In some embodiments, the reference standard is the test sample at T₀. In some embodiments, the reference standard is an immunogenic composition without a detergent. In

some embodiments, the reference standard is an immunogenic composition with a detergent-to-protein molar ratio higher than 10:1.

[0031] A "protective" immune response refers to the ability of an immunogenic composition to elicit an immune response, either humoral or cell mediated, which serves to protect the subject from an infection. The protection provided need not be absolute, i.e., the infection need not be totally prevented or eradicated, if there is a statistically significant improvement compared with a control population of subjects, e.g. infected animals not administered the vaccine or immunogenic composition. Protection may be limited to mitigating the severity or rapidity of onset of symptoms of the infection. In general, a "protective immune response" would include the induction of an increase in antibody levels specific for a particular antigen in at least 50% of subjects, including some level of measurable functional antibody responses to each antigen. In particular situations, a "protective immune response" could include the induction of a two fold increase in antibody levels or a four fold increase in antibody levels specific for a particular antigen in at least 50% of subjects, including some level of measurable functional antibody responses to each antigen. In certain embodiments, opsonising antibodies correlate with a protective immune response. Thus, protective immune response may be assayed by measuring the percent decrease in the bacterial count in an opsonophagocytosis assay, for instance those described below. In some embodiments, there is a decrease in bacterial count of at least 10%, 25%, 50%, 65%, 75%, 80%, 85%, 90%, 95% or more, as compared to the bacterial count in the absence of the immunogenic composition.

[0032] The terms "protein", "polypeptide" and "peptide" refer to a polymer of amino acid residues and are not limited to a minimum length of the product. Thus, peptides, oligopeptides, dimers, multimers, and the like, are included within the definition. Both full-length proteins and fragments thereof are encompassed by the definition. The terms also include modifications, such as deletions, additions and substitutions (generally conservative in nature, but which may be non-conservative), to a native sequence, preferably such that the protein maintains the ability to elicit an immunological response within an animal to which the protein is administered. Also included are post-expression modifications, e.g. glycosylation, acetylation, lipidation, phosphorylation and the like.

[0033] The term "recombinant" as used herein refers to any protein, polypeptide, or cell expressing a gene of interest that is produced by genetic engineering methods. The term "recombinant" as used with respect to a protein or polypeptide, means a polypeptide produced by expression of a recombinant polynucleotide. The proteins of the present invention may be isolated from a natural source or produced by genetic engineering methods. "Recombinant," as used herein, further describes a nucleic acid molecule, which, by virtue of its origin or manipulation, is not associated with all or a portion of the polynucleotide with which it is associated in nature. The term "recombinant" as used with respect to a host cell means a host cell which includes a recombinant polynucleotide.

[0034] The terms "stable" and "stability" refer the ability of an antigen to remain immunogenic over a period of time. Stability may be measured in potency over time. The terms "stable" and "stability" further refer to the physical, chemical, and conformational stability of the immunogenic composition. Instability of a protein composition may be caused by chemical degradation or aggregation of the protein molecules to form higher order polymers, by dissociation of the heterodimers into monomers, deglycosylation, modification of glycosylation, or any other structural modification that reduces at least one biological activity of the protein composition included in the present invention. Stability may be assessed by methods well-known in the art, including measurement of a sample's light scattering, apparent attenuation of light (absorbance, or optical density), size (e.g. by size exclusion chromatography), in vitro or in vivo biological activity and/or properties by differential scanning calorimetry (DSC). Other methods for assessing stability are known in the art and can also be used according to the present invention.

[0035] In some embodiments, an antigen in a stable formulation of the invention may maintain at least 50%, 60%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98%, 99% or 100% potency, as compared to a reference standard, for at least 1 month, 2 months, 3 months, 4 months, 5 months, 6 months, 9 months, 12 months, 18 months, 24 months, 30 months, 36 months, 42 months, 48 months, 54 months, or 60 months. In some embodiments, an antigen in a stable formulation of the invention may maintain at least 50% potency, as compared to a reference standard, for at least 1 year, 2 years, 3 years, 4 years or 5 years. The terms "stable" and "stability" also refer to the ability of an antigen to maintain epitopes or immunoreactivity over a period of time. For example, an antigen in a stable formulation of the invention may maintain at least 50%, 60%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98%, 99% or 100% of its epitopes or immunoreactivity, as compared to a reference standard, for at least 1 month, 2 months, 3 months, 4 months, 5 months, 6 months, 9 months, 12 months, 18 months, 24 months, 30 months, 36 months, 42 months, 48 months, 54 months, or 60 months. In some embodiments, stability is measured with respect to an environmental condition. Non-limiting examples of environmental conditions include light, temperature, freeze/thaw cycles, agitation, and pH. One of skill in the art would be able to determine the presence of antigenic epitopes or immunoreactivity using the methods disclosed herein or other methods known in the art. See, e.g., McNeil et al. Vaccine, 27: 3417-3421 (2009). In some embodiments, the stability of an antigen is measured from the date of its formulation. In some embodiments, the stability of an antigen is measured from the date of a change in its storage conditions. Non-limiting examples of changes in storage conditions include changing from frozen to refrigerated, changing from frozen to room temperature, changing from refrigerated to room temperature, changing from refrigerated to frozen, changing from room temperature to frozen,

changing from room temperature to refrigerated, changing from light to dark, or introduction of agitation.

[0036] The terms "stablizer" refers to a compound that binds to an antigen and maintains the epitopes or immunoreactivity of the antigen over a period of time. Stabilizers are known in the art. Examples of stabilizers include multivalent cations, for example, calcium or aluminum.

[0037] The term "subject" refers to a mammal, bird, fish, reptile, or any other animal. The term "subject" also includes humans. The term "subject" also includes household pets. Non-limiting examples of household pets include: dogs, cats, pigs, rabbits, rats, mice, gerbils, hamsters, guinea pigs, ferrets, birds, snakes, lizards, fish, turtles, and frogs. The term "subject" also includes livestock animals. Non-limiting examples of livestock animals include: alpaca, bison, camel, cattle, deer, pigs, horses, llamas, mules, donkeys, sheep, goats, rabbits, reindeer, yak, chickens, geese, and turkeys.

[0038] The terms "vaccine" or "vaccine composition", which are used interchangeably, refer to pharmaceutical compositions comprising at least one immunogenic composition that induces an immune response in a subject.

General Description

[0039] The present invention arises out of the novel discovery that rLP2086 Subfamily B antigens, but not rLP2086 Subfamily A antigens, lose potency over time in a bivalent vaccine formulation and are thus unstable. By varying components in the bivalent formulation, it was determined that high molar ratios of detergent to protein in the bivalent vaccine formulation resulted in rLP2086 Subfamily B antigen specific instability. Reducing the molar ratio of detergent to protein in bivalent and monovalent formulations resulted in increased stability, as determined by maintenance of potency over time, of rLP2086 Subfamily B antigen without affecting the stability of rLP2086 Subfamily A antigen. This result is surprising because lipoproteins are typically purified and stored using high detergent concentrations to prevent aggregation of their hydrophobic lipid moieties.

[0040] Further studies revealed that low molar ratio formulations resulted in aggregation of rLP2086 Subfamily A and B antigens upon agitation of the low molar ratio immunogenic compositions. Increasing aluminum concentration in low molar ratio compositions, however, prevented aggregation of rLP2086 Subfamily A and B antigens, even with agitation. Moreover, rLP2086 Subfamily A antigens are more sensitive to the effects of low detergent molar ratios in the absence of aluminum.

Immunogenic Compositions

[0041] Immunogenic compositions that include a protein encoded by a nucleotide sequence from *Neisseria meningitidis* ORF2086 are known in the art. Exemplary immunogenic compositions include those described in US patent application publication numbers US 20060257413 and US 20090202593, which are incorporated herein by reference in their entirety. Such immunogenic compositions described therein include a protein exhibiting bactericidal activity identified as ORF2086 protein, immunogenic portions thereof, and/or biological equivalents thereof. The ORF2086 protein refers to a protein encoded by open reading frame 2086 of *Neisseria* species.

[0042] The protein may be a recombinant protein or an isolated protein from native *Neisseria* species. For example, *Neisseria* ORF2086 proteins may be isolated from bacterial strains, such as those of *Neisseria* species, including strains of *Neisseria meningitidis* (serogroups A, B, C, D, W-135, X, Y, Z, and 29E), *Neisseria gonorrhoeae*, and *Neisseria lactamica*, as well as immunogenic portions and/or biological equivalents of said proteins.

[0043] The ORF2086 proteins include 2086 Subfamily A proteins and Subfamily B proteins, immunogenic portions thereof, and/or biological equivalents thereof. The ORF2086 proteins or equivalents thereof, etc. may be lipidated or non-lipidated. Preferably, the *Neisseria* ORF2086 protein is lipidated.

[0044] In an one embodiment, the immunogenic composition includes an isolated protein having at least 95% amino acid sequence identity to a protein encoded by a nucleotide sequence from *Neisseria* ORF2086.

[0045] In one embodiment, the immunogenic composition includes an isolated protein having at least 95% amino acid sequence identity to a Subfamily A protein encoded by a nucleotide sequence from *Neisseria* ORF2086. Preferably, the immunogenic composition includes an isolated Subfamily A protein encoded by a nucleotide sequence from *Neisseria* ORF2086.

[0046] In another embodiment, the immunogenic composition includes an isolated protein having at least 95% amino acid sequence identity to a Subfamily B protein encoded by a nucleotide sequence from *Neisseria* ORF2086. Preferably, the immunogenic composition includes an isolated Subfamily B protein encoded by a nucleotide sequence from *Neisseria* ORF2086. In some embodiments, the ORF2086 Subfamily B protein is a B01 variant.

[0047] In yet another embodiment, the immunogenic composition includes an isolated protein having at least 95% amino acid sequence identity to a Subfamily A protein encoded by a nucleotide sequence from *Neisseria* ORF2086, and an isolated protein having at least 95% amino acid sequence identity to a Subfamily B protein encoded by a nucleotide sequence from *Neisseria* ORF2086. Preferably, the immunogenic composition includes an isolated Subfamily A protein encoded by a nucleotide sequence from *Neisseria* ORF2086 and an isolated Subfamily B protein encoded by a nucleotide

sequence from *Neisseria* ORF2086.

[0048] In one embodiment, the immunogenic composition includes a 1:1 ratio of a Subfamily A protein to a Subfamily B protein.

[0049] The immunogenic composition may include a protein encoded by a nucleotide sequence from *Neisseria* ORF2086, polynucleotides, or equivalents thereof as the sole active immunogen in the immunogenic composition.

[0050] The present invention also contemplates multi-immunization regimens wherein any composition useful against a pathogen may be combined therein or therewith the compositions of the present invention. For example, without limitation, a patient may be administered the immunogenic composition of the present invention and another immunological composition for immunizing against human papillomavirus virus (HPV), such as the HPV vaccine GARDASIL[®], as part of a multi-immunization regimen. Persons of skill in the art would be readily able to select immunogenic compositions for use in conjunction with the immunogenic compositions of the present invention for the purposes of developing and implementing multi-immunization regimens.

[0051] Immunogenic compositions of the invention preferably include a pharmaceutically acceptable carrier. Suitable pharmaceutically acceptable carriers and/or diluents include any and all conventional solvents, dispersion media, fillers, solid carriers, aqueous solutions, coatings, antibacterial and antifungal agents, isotonic and absorption delaying agents, and the like. Suitable pharmaceutically acceptable carriers include, for example, one or more of water, saline, phosphate buffered saline, dextrose, glycerol, ethanol and the like, as well as combinations thereof.

[0052] Pharmaceutically acceptable carriers may further include minor amounts of auxiliary substances such as wetting or emulsifying agents, preservatives or buffers, which enhance the shelf life or effectiveness of the antibody. The preparation and use of pharmaceutically acceptable carriers is well known in the art. Except insofar as any conventional media or agent is incompatible with the active ingredient, use thereof in the immunogenic compositions of the present invention is contemplated.

[0053] Immunogenic compositions can be administered parenterally, e.g., by injection, either subcutaneously or intramuscularly, as well as orally or intranasally. Methods for intramuscular immunization are described by Wolff et al. *Biotechniques*;11(4):474-85, (1991). and by Sedegah et al. *PNAS* Vol. 91, pp. 9866-9870, (1994). Other modes of administration employ oral formulations, pulmonary formulations, suppositories, and transdermal applications, for example, without limitation. Oral formulations, for example, include such normally employed excipients as, for example, pharmaceutical grades of mannitol, lactose, starch, magnesium stearate, sodium saccharine, cellulose, magnesium carbonate, and the like, without limitation. Preferably, the immunogenic composition is administered intramuscularly.

[0054] The immunogenic compositions of the invention can include one or more adjuvants. Exemplary adjuvants include, but are not limited to aluminum hydroxide; aluminum phosphate; STIMULON[™] QS-21 (Aquila Biopharmaceuticals, Inc., Framingham, Mass.); MPL[™] (3-O-deacylated monophosphoryl lipid A; Corixa, Hamilton, Mont.), 529 (an amino alkyl glucosamine phosphate compound, Corixa, Hamilton, Mont.), IL-12 (Genetics Institute, Cambridge, Mass.); GM-CSF (Immunex Corp., Seattle, Wash.); N-acetyl-muramyl-L-theronyl-D-isoglutamine (thr-MDP); N-acetyl-nor-muramyl-L-alanyl-D-isoglutamine (CGP 11637, referred to as nor-MDP); N-acetylmuramyl-L-alanyl-D-isoglutaminyl-L-alanine-2-(1'-2'-dipalmitoyl-sn-glycero-3-hydroxyphosphoryloxy-ethylamine) (CGP 19835A, referred to as MTP-PE); and cholera toxin. In certain preferred embodiments, the adjuvant is QS-21.

[0055] Additional exemplary adjuvants include non-toxic derivatives of cholera toxin, including its A subunit, and/or conjugates or genetically engineered fusions of the *N. meningitidis* polypeptide with cholera toxin or its B subunit ("CTB"), procholeraenoid, fungal polysaccharides, including schizophyllan, muramyl dipeptide, muramyl dipeptide ("MDP") derivatives, phorbol esters, the heat labile toxin of *E. coli*, block polymers or saponins.

[0056] Aluminum phosphate has been used as the adjuvant in a phase 1 clinical trial to a concentration 0.125 mg/dose, much lower than the limit of 0.85 mg/ dose specified by the US Code of Federal Regulations [610.15(a)]. Aluminum-containing adjuvants are widely used in humans to potentiate the immune response of antigens when administered intramuscularly or subcutaneously.

[0057] In certain preferred embodiments, the proteins of this invention are used in an immunogenic composition for oral administration which includes a mucosal adjuvant and used for the treatment or prevention of *N. meningitidis* infection in a human host. The mucosal adjuvant can be a cholera toxin; however, preferably, mucosal adjuvants other than cholera toxin which may be used in accordance with the present invention include non-toxic derivatives of a cholera holotoxin, wherein the A subunit is mutagenized, chemically modified cholera toxin, or related proteins produced by modification of the cholera toxin amino acid sequence. For a specific cholera toxin which may be particularly useful in preparing immunogenic compositions of this invention, see the mutant cholera holotoxin E29H, as disclosed in Published International Application WO 00/18434, which is hereby incorporated herein by reference in its entirety. These may be added to, or conjugated with, the polypeptides of this invention. The same techniques can be applied to other molecules with mucosal adjuvant or delivery properties such as *Escherichia coli* heat labile toxin (LT). Other compounds with mucosal adjuvant or delivery activity may be used such as bile; polycations such as DEAE-dextran and polyomithine; detergents such as sodium dodecyl benzene sulphate; lipid-conjugated materials; antibiotics such as streptomycin; vitamin A; and other compounds that alter the structural or functional integrity of mucosal surfaces. Other mucosally

active compounds include derivatives of microbial structures such as MDP; acridine and cimetidine. STIMULON™ QS-21, MPL, and IL-12, as described above, may also be used.

[0058] The immunogenic compositions of this invention may be delivered in the form of ISCOMS (immune stimulating complexes), 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. The proteins of this invention may also be incorporated into oily emulsions.

[0059] An amount (i.e., dose) of immunogenic composition that is administered to the patient can be determined in accordance with standard techniques known to those of ordinary skill in the art, taking into consideration such factors as the particular antigen, the adjuvant (if present), the age, sex, weight, species, condition of the particular patient, and the route of administration.

[0060] For example, a dosage for an adolescent human patient may include at least 0.1 μg, 1 μg, 10 μg, or 50 μg of a *Neisseria* ORF2086 protein, and at most 80 μg, 100 μg, 150 μg, or 200 μg of a *Neisseria* ORF2086 protein. Any minimum value and any maximum value may be combined to define a suitable range.

In vitro potency assay

[0061] Potency is determined by quantitation of functional epitopes in Subfamily A and Subfamily B proteins in an immunogenic composition using conformation-specific monoclonal antibodies against a rLP2086 reference material. Potency is determined by quantitative measure of the functional epitopes in Subfamily A or Subfamily B rLP2086 proteins that will elicit immune response in vivo to generate bactericidal antibodies. Quantitative technology is used for potency assay with selected monoclonal antibodies (mAbs). Two functional monoclonal antibodies that are conformational and non-overlapping are selected for each Subfamily rLP2086 protein in the immunogenic compositions. Between the two purified monoclonal antibodies, the first antibody is conjugated to a first tag, wherein the first tag is used to capture of rLP2086 protein molecule. In some embodiments, the first tag is biotin, glutathione-S transferase (GST), a 6xHis tag, or beads (e.g. carboxylated polystyrene beads or paramagnetic beads). In some embodiments the first tag is captured with streptavidin beads, a streptavidin column, nickel beads, a nickel column, centrifugation or with a magnetic field. The second antibody is conjugated to a second tag, wherein the second tag is quantifiable. In some embodiments, the second tag is biotin, horseradish peroxidase (HRP), a fluorophore or a radiolabel. In some embodiments, the second tag is detected with streptavidin conjugated to a fluorophore or HRP, by electrochemiluminescence, detection of fluorescence, or detection of radioactivity. Only the proteins that exhibit both epitopes recognized by the two mAbs in each immunogenic composition will be measured. Changes in any one or both epitopes of the protein will be reflected. The potency of the sample is reported relative to the potency of the reference material.

[0062] The disclosure encompasses a method for determining the potency of a 2086 protein. The method comprises the steps of (1) incubating a first monoclonal Ab and a second mAb with an immunogenic composition comprising a 2086 protein, wherein the first mAb is conjugated to a first tag that is used to capture the mAb and the second mAb is conjugated to a second tag that is detectable and wherein the first and second mAbs are directed to different conformational epitopes on a 2086 reference protein; (2) capturing the first mAb-bound 2086 protein using the first tag; and (3) detecting and quantifying the amount of captured second mAb-bound 2086 protein using the second tag. In some embodiments, the 2086 protein is a Subfamily A protein. In some embodiments, the 2086 protein is a Subfamily B protein. In some embodiments, the 2086 protein is lipidated. In some embodiments, the 2086 protein is non-lipidated. In some embodiments, the 2086 protein is recombinant. In some embodiments, the first tag is biotin, a 6xHis tag, or beads (e.g. carboxylated polystyrene beads or paramagnetic beads). In some embodiments, the first tag is captured with streptavidin beads, a streptavidin column, glutathione beads, glutathione column, nickel beads, a nickel column, centrifugation or with a magnetic field. In some embodiments, the second tag is biotin, HRP, a fluorophore or a radiolabel. In some embodiments, the second tag is detected with streptavidin conjugated to a fluorophore or HRP, by electrochemiluminescence, detection of fluorescence, or detection of radioactivity. In some embodiments, the immunogenic composition comprises multiple 2086 protein variants.

Stability of rLP2086 Subfamily A and B Antigen Potency

[0063] In some embodiments, the invention provides an immunogenic composition for stabilizing rLP2086 Subfamily A and/or rLP2086 Subfamily B antigens over time comprising a buffer with a high stabilizer concentration and a low detergent-to-protein molar ratio.

[0064] In some embodiments, the detergent-to-protein molar ratio in the immunogenic composition is between about 0.5 and about 10. In some embodiments, the detergent-to-protein molar ratio in the immunogenic composition is between about 1 and about 5. In some embodiments, the detergent-to-protein molar ratio in the immunogenic composition is between about 1.4 and about 4.2. In some embodiments, the detergent-to-protein molar ratio in the immunogenic composition is about 0.5, about 0.6, about 0.7, about 0.8, about 0.9, about 1.0, about 1.1, about 1.2, about 1.3, about

1.4, about 1.5, about 1.6, about 1.7, about 1.8, about 1.9, about 2.0, about 2.1, about 2.2, about 2.3, about 2.4, about 2.5, about 2.6, about 2.7, about 2.8, about 2.9, about 3.0, about 3.1, about 3.2, about 3.3, about 3.4, about 3.5, about 3.6, about 3.7, about 3.8, about 3.9, about 4.0, about 4.1, about 4.2, about 4.3, about 4.4, about 4.5, about 4.6, about 4.7, about 4.8, about 4.9, about 5.0, about 5.5, about 6.0, about 6.5, about 7.0, about 7.5, about 8.0, about 8.5, about 9.0, about 9.5, or about 10. In some embodiments, the detergent is a non-ionic detergent. In some embodiments, the detergent is a polysorbate detergent. In some embodiments, the detergent is Polysorbate 80.

[0065] The multivalent cation is aluminum. In some embodiments, the aluminum is present as one or more of AlPO_4 , $\text{Al}(\text{OH})_3$, $\text{Al}_2(\text{SO}_4)_3$ and alum. In some embodiments, the immunogenic composition comprises between about 0.1 mg/mL and about 1 mg/mL; between about 0.25 mg/mL and about 0.75 mg/mL, or between about 0.4mg/mL and about 0.6 mg/mL aluminum. In some embodiments, the immunogenic composition comprises about 0.1 mg/mL, about 0.15 mg/mL; about 0.2 mg/mL, about 0.25 mg/mL, about 0.3 mg/mL, about 0.35 mg/mL, about 0.4mg/mL, about 0.45 mg/mL, about 0.5 mg/mL, about 0.55 mg/mL, about 0.6 mg/mL, about 0.65 mg/mL, about 0.7 mg/mL, about 0.75 mg/mL, about 0.8 mg/mL, about 0.85 mg/mL, 0.9 mg/mL, about 0.95 mg/mL, or about 1 mg/mL aluminum. In some embodiments, there is at least 90%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99% or 100% binding of aluminum to protein.

[0066] In some embodiments, the immunogenic composition further comprises a buffer comprising histidine. In some embodiments, the concentration of histidine is between about 2mM and about 20mM; between about 5mM and about 15mM, or between about 8mM and 12mM. In some embodiments, the concentration of histidine is about 2mM, about 3mM, about 4mM, about 5mM, about 6mM, about 7mM, about 8mM, about 9mM, about 10mM, about 11mM, about 12mM, about 13mM, about 14mM, about 15mM, about 16mM, about 17mM, about 18mM, about 19mM or about 20mM.

[0067] In some embodiments, the immunogenic composition further comprises a buffer comprising succinate. In some embodiments, the concentration of succinate is between about 2mM and about 20mM; between about 2mM and about 10mM, or between about 3mM and 7mM. In some embodiments, the concentration of succinate is about 2mM, about 3mM, about 4mM, about 5mM, about 6mM, about 7mM, about 8mM, about 9mM, about 10mM, about 11mM, about 12mM, about 13mM, about 14mM, about 15mM, about 16mM, about 17mM, about 18mM, about 19mM or about 20mM.

[0068] In some embodiments, the immunogenic composition has a pH of between about 5.0 and about 8.0; between about 5.5 and about 7.0; or between about 5.8 and about 6.0. In some embodiments, the immunogenic composition has a pH of about 5.0, about 5.1, about 5.2, about 5.3, about 5.4, about 5.5, about 5.6, about 5.7, about 5.8, about 5.9, about 6.0, about 6.1, about 6.2, about 6.3, about 6.4, or about 6.5.

[0069] In some embodiments, the formulation of MnB rLP2086 Subfamily A and B protein antigens is 10 mM histidine-buffered saline, pH 6.0, containing 0.5 mg/mL aluminum as aluminum phosphate and Polysorbate 80:protein molar ratio of 2.8.

[0070] In some embodiments, the formulation of MnB rLP2086 Subfamily B protein antigen immunogenic composition is 5 mM succinate-buffered saline, pH 6.0, containing 0.5 mg/mL aluminum as aluminum phosphate and Polysorbate 80:protein molar ratio of 2.8.

[0071] In some embodiments, the invention provides a method of stabilizing rLP2086 Subfamily A and/or rLP2086 Subfamily B antigens over time comprising storing the antigens in a buffer with a high stabilizer concentration and a low detergent-to-protein molar ratio.

[0072] In some embodiments, the detergent-to-protein molar ratio less than 10:1. In some embodiments, the detergent-to-protein molar ratio in the buffer is between about 0.5 and about 10. In some embodiments, the detergent-to-protein molar ratio in the buffer is between about 1 and about 5. In some embodiments, the detergent-to-protein molar ratio in the buffer is between about 1.4 and about 4.2. In some embodiments, the detergent-to-protein molar ratio in the buffer is about 0.5, about 0.6, about 0.7, about 0.8, about 0.9, about 1.0, about 1.1, about 1.2, about 1.3, about 1.4, about 1.5, about 1.6, about 1.7, about 1.8, about 1.9, about 2.0, about 2.1, about 2.2, about 2.3, about 2.4, about 2.5, about 2.6, about 2.7, about 2.8, about 2.9, about 3.0, about 3.1, about 3.2, about 3.3, about 3.4, about 3.5, about 3.6, about 3.7, about 3.8, about 3.9, about 4.0, about 4.1, about 4.2, about 4.3, about 4.4, about 4.5, about 4.6, about 4.7, about 4.8, about 4.9, about 5.0, about 5.5, about 6.0, about 6.5, about 7.0, about 7.5, about 8.0, about 8.5, about 9.0, about 9.5, or about 10. In some embodiments, the detergent is a non-ionic detergent. In some embodiments, the detergent is a polysorbate detergent. In some embodiments, the detergent is Polysorbate 80.

[0073] The stabilizer in the buffer is aluminum. In some embodiments, the aluminum is present as one or more of AlPO_4 , $\text{Al}(\text{OH})_3$, $\text{Al}_2(\text{SO}_4)_3$ and alum. In some embodiments, the stabilizer in the buffer is between about 0.1 mg/mL and about 1 mg/mL; between about 0.25 mg/mL and about 0.75 mg/mL, or between about 0.4mg/mL and about 0.6 mg/mL aluminum. In some embodiments, the stabilizer in the buffer is about 0.1 mg/mL, about 0.15 mg/mL; about 0.2 mg/mL, about 0.25 mg/mL, about 0.3 mg/mL, about 0.35 mg/mL, about 0.4mg/mL, about 0.45 mg/mL, about 0.5 mg/mL, about 0.55 mg/mL, about 0.6 mg/mL, about 0.65 mg/mL, about 0.7 mg/mL, about 0.75 mg/mL, about 0.8 mg/mL, about 0.85 mg/mL, 0.9 mg/mL, about 0.95 mg/mL, or about 1 mg/mL aluminum. In some embodiments, there is at least 90%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99% or 100% binding of aluminum to protein.

[0074] In some embodiments, the buffer further comprises histidine. In some embodiments, the concentration of histidine is between about 2mM and about 20mM; between about 5mM and about 15mM, or between about 8mM and

12mM. In some embodiments, the concentration of histidine is about 2mM, about 3mM, about 4mM, about 5mM, about 6mM, about 7mM, about 8mM, about 9mM, about 10mM, about 11mM, about 12mM, about 13mM, about 14mM, about 15mM, about 16mM, about 17mM, about 18mM, about 19mM or about 20mM.

5 **[0075]** In some embodiments, the buffer further comprises succinate. In some embodiments, the concentration of succinate is between about 2mM and about 20mM; between about 2mM and about 10mM, or between about 3mM and 7mM. In some embodiments, the concentration of succinate is about 2mM, about 3mM, about 4mM, about 5mM, about 6mM, about 7mM, about 8mM, about 9mM, about 10mM, about 11mM, about 12mM, about 13mM, about 14mM, about 15mM, about 16mM, about 17mM, about 18mM, about 19mM or about 20mM.

10 **[0076]** In some embodiments, the buffer has a pH of between about 5.0 and about 8.0; between about 5.5 and about 7.0; or between about 5.8 and about 6.0. In some embodiments, the buffer has a pH of about 5.0, about 5.1, about 5.2, about 5.3, about 5.4, about 5.5, about 5.6, about 5.7, about 5.8, about 5.9, about 6.0, about 6.1, about 6.2, about 6.3, about 6.4, or about 6.5.

15 **[0077]** In some embodiments, the buffer that the MnB rLP2086 Subfamily A and B protein antigens are stored in is 10 mM histidine-buffered saline, pH 6.0, containing 0.5 mg/mL aluminum as aluminum phosphate and Polysorbate 80:protein molar ratio of 2.8.

20 **[0078]** In some embodiments, the buffer that the MnB rLP2086 Subfamily A and B protein antigens are stored in is 5 mM succinate-buffered saline, pH 6.0, containing 0.5 mg/mL aluminum as aluminum phosphate and Polysorbate 80:protein molar ratio of 2.8.

[0079] In order that this invention may be better understood, the following examples are set forth.

25 EXAMPLES

Example 1: Experimental Procedures

25 Determination of Aluminum Binding

30 **[0080]** A composition comprising aluminum and at least one protein antigen was centrifuged such that the aluminum was pelleted. Centrifugation of aluminum absorbed proteins is known in the art. See e.g., Egan et al., Vaccine, Vol. 27(24): 3175-3180 (2009). Aluminum-bound protein was also pelleted, while non-aluminum-bound protein remained in the supernatant. Total protein in the supernatant and pellet were determined by Lowry Assay. The percentage bound protein was calculated by dividing the total protein in the supernatant by the total protein added to the composition and multiplying by 100%. Similarly, the percentage unbound protein was calculated by dividing the total protein in the supernatant by the total protein added to the composition and multiplying by 100%.

35 **[0081]** For compositions comprising both Subfamily A and Subfamily B antigens, the individual Subfamily A and B protein concentrations in the supernatant were determined by ion-exchange chromatography. The separation and elution of Subfamily A and B proteins was carried out using a strong anion column and a high salt concentration eluent. Both Subfamily A and B proteins were detected and quantified using a fluorescence detector set at Excitation = 280 nm and Emission = 310 nm. Subfamily A and Subfamily B proteins elute at distinct retention times and were quantified using a standard curve generated against a rLP2086 protein reference material. The percentage unbound protein was calculated by dividing the total protein in the supernatant by the total protein added to the composition and multiplying by 100%. The percentage bound protein was calculated by subtracting the percentage unbound protein from 100%.

In Vitro Potency Assay

45 **[0082]** The rLP2086 potency assay is a homogeneous capture assay or sandwich assay that relies on two functional monoclonal antibodies that recognize conformational and non-overlapping epitopes on a single protein molecule of the rLP2086 drug substance. One purified monoclonal antibody serves as the capture antibody (mAb) and is chemically conjugated to carboxylated polystyrene beads which have a unique color-coded identifier. The second antibody is biotinylated and serves as a detection antibody that is subsequently bound by streptavidin conjugated to the fluorophore R-phycoerythrin (SA-PE). The fluidics of a Bio-Plex detection instrument quantifies individual microspheres and their associated SA-PE signal. A fluorescence signal from the R-phycoerythrin associated with the microsphere will be detected only by ternary complex formation between bead-conjugated antibody, antigen, and detection antibody and will be proportional to the number of functional epitopes in the rLP2086 samples. A change in one or both epitopes resulting in a loss of fluorescence relative to a reference standard will indicate a loss in potency.

Reagents

[0083]

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- Monoclonal antibody conjugated microspheres (conjugated to Luminex MicroPlex Microsphere bead region #12 or to bead region #66).
- Biotinylated monoclonal antibody.
- rLP2086 reference materials, Subfamilies A and B, 2 mg/ml. Store at -70 °C.
- rLP2086 Subfamily A and B bivalent control
- Streptavidin, R-phycoerythrin conjugated, lyophilized

Buffers

[0084]

- 10 mM Histidine, 150 mM NaCl, pH 6.0
- 5% w/v polysorbate 80 (PS-80) in 0.85% w/v saline.
- Matrix Buffer (10 mM Histidine, 0.02% polysorbate 80, 150 mM NaCl, pH 6.0).
- Assay Buffer (PBS, pH 7.4 with 0.1% BSA, 0.02% polysorbate 80, 0.1% azide).
- 100x Streptavidin, R-phycoerythrin-conjugated (SA-PE) - Open vial of lyophilized streptavidin, R-phycoerythrin and add 1 mL of distilled water. Vortexed until completely dissolved.

Procedure

[0085] 200 µL of Subfamily A protein and 200 µL of Subfamily B protein were added to 600 µL of Matrix Buffer for a concentration of 400 µg/ml of each subfamily. A standard curve of eight concentrations (3333-1.5 ng/mL) was generated by diluting the stock solution in Assay Buffer.

[0086] 200 µL of the bivalent control was added to 800 µL of Matrix Buffer for a concentration of 400 µg/mL of each subfamily. The 400 µg/mL stock to make 100, 50, and 12.5 ng/mL working concentrations diluted in Assay Buffer. 100 and 12.5 ng/mL represented high and low controls (CH) and (CL) respectively.

[0087] Test samples were diluted in Matrix Buffer to a concentration of 400 µg/mL. 100, 50, and 12.5 ng/mL working solutions were prepared from the 400 µg/mL stock.

[0088] A homogenous assay mixture using a conjugated bead concentration of 2×10^5 beads/mL and a detection antibody concentration of 30 µg/mL in Assay Buffer was prepared. A sample plate was prepared by adding 0.4 mL of standard, control, sample or blank to a 2 mL 96-well deep well plate. The filters of a 96-well MultiScreen_{HTS}-BV filter plate were pre-wet by adding 100 µL of Assay Buffer, which was then drawn through the filter by vacuum suction. 25 µL of the prepared homogenous assay mixture was added to 96-well plate. 25 µL of each standard, control, sample or blank solution was added to each well of the 96-well filter plate. The plates were incubated at room temperature for one hour with shaking.

[0089] After the antigen-antibody incubation buffer was removed by vacuum aspiration through the filter. The filter of each well was washed three times with 100 µL of Assay Buffer followed by vacuum aspiration. After the final wash, 50 µL of 1x SA-PE was added to each well. The plate was incubated 10 minutes at room temperature with shaking on a titer in the dark.

[0090] Following the SA-PE incubation, 75 µL of Assay Buffer was added to each well of the plate for a total volume of 125 µL. The plate was immediately read on a Bio-Plex 200 System.

Serum bactericidal assay

[0091] New Zealand White female rabbits, 2.5-3.0 kg, obtained from Charles River Canada (St. Constant, QC, Canada), were prescreened by whole cell ELISA to identify those with low reactivity against two different meningococcal strains (one from each P2086 subfamily). The rabbits, in general, had very low backgrounds, and those with the lowest values were selected for use. The rabbits were vaccinated intramuscularly at weeks 0, 4, and 9 with either monovalent rLP2086-A05, monovalent rLP2086-B01 or a bivalent r LP2086-A05 + B01 vaccine. Each dose contained 100µg of protein for the monovalent vaccines and 100µg of each protein for the bivalent vaccine, formulated in 10mM histidine buffer pH 6.0, 150mM NaCl, 0.02% Polysorbate 80 and 250µg AlPO₄. The vaccine was injected intramuscularly into the right hind leg (0.5 ml/dose). As a control, one group of rabbits was vaccinated with the formulation buffer alone. Pre-immune (week 0) and immune (week 10) serum samples were obtained for analyses. All animal protocols adhered to the established Institutional Animal Care and Use Committee guidelines.

[0092] Serum bactericidal antibodies in rabbits immunized with rLP2086 vaccine were determined using SBAs with human complement. Rabbit immune sera were heat-inactivated to remove intrinsic complement activity and subsequently serially diluted 1:2 in Dulbecco's PBS with Ca²⁺ and Mg²⁺ (D-PBS) in a 96-well microtiter plate to test for serum bactericidal activity against *N. meningitidis* strains. Bacteria used in the assay were grown in GC media supplemented

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with Kellogg's supplement (GCK) and monitored by optical density at 650 nm. Bacteria were harvested for use in the assay at a final OD₆₅₀ of 0.50-0.55, diluted in D-PBS and 1000-3000 CFU were added to the assay mixture with 20% human complement.

5 [0093] Human serum with no detectable bactericidal activity was used as the exogenous complement source. Complement sources were tested for suitability against each individual test strain. A complement source was used only if the number of bacteria surviving in controls without added immune sera was >75%. Ten unique complement sources were required to perform the SBAs described in this study.

10 [0094] After a 30 min incubation at 37°C with 5% CO₂, D-PBS was added to the reaction mixture and aliquots transferred to microfilter plates filled with 50% GCK media. The microfilter plates were filtered, incubated overnight at 37°C with 5% CO₂ and microcolonies were stained and quantified. The serum bactericidal titers were defined as the interpolated reciprocal serum dilution that yielded a 50% reduction in CFU compared to the CFU in control wells without immune sera. The SBA titer is defined as the reciprocal of the interpolated dilution of test serum that causes a 50% reduction in bacterial counts after a 30min incubation at 37°C. Susceptibility to killing with P2086 immune sera was established if there was a 4-fold or greater rise in SBA titer for P2086 immune sera compared to the corresponding pre-immune sera. 15 The limit of detection was a titer of 8 for the rabbit sera. Sera that were negative against the assay strain at the starting dilution were assigned a titer of one half the limit of detection for the assay (i.e. 4 for the rabbit).

Flow Cytometry

20 [0095] MnB cells were grown to an OD₆₅₀ of 0.45-0.55 and subsequently fixed in 1% (v/v) paraformaldehyde in 1 × PBS for 10min. One hundred microliters/well of bacteria were plated into 96-well U-bottom polystyrene plates, spun down and washed once in 1% (w/v) BSA in 1 × PBS. Anti-LP2086 monoclonal antibodies were added to the bacterial pellets, resuspended and incubated on ice for 30min. After two washes in 1% BSA/PBS, biotinylated goat anti-mouse 25 IgG (subclasses 1 + 2a + 2b + 3) (Jackson ImmunoResearch) was added to the cell pellets, resuspended and incubated on ice for 30min. The cells were washed twice and resuspended in streptavidin-PE (BD Biosciences) and incubated on ice for 30min. After two washes in 1% BSA/PBS, the cell pellets were resuspended in 1% paraformaldehyde. Mouse IgG was included as a negative control. Twenty thousand (20,000) events per well were acquired on a BD LSR II flow cytometer and analyzed using FlowJo v7 software (Treestar, Ashland, Oregon). The mean fluorescence intensity (MFI) of the PE channel was determined for each sample after gating on bacterial cells in the logarithmic FSC versus SSC 30 dot plot. An MFI was considered positive if the MFI was three times that of the control mouse IgG MFI.

Example 2: Polysorbate 80 Binding to rLP2086 Proteins

35 [0096] To understand the stability of Polysorbate 80 binding to each rLP2086 protein A and B, a rLP2086 formulated sample with 200 µg/mL Subfamily A with aluminum (Al) and another rLP2086 sample formulated with 200 µg/mL Subfamily B both stored at 2-8°C and 25°C were tested 5 months later for their protein and Polysorbate 80 content. A placebo (buffer + Al without protein) was also analyzed. The Polysorbate 80 distribution in the Placebo is shown in Figure 14, while the Polysorbate 80 distributions for Subfamily A and B proteins are shown in Figure 15 and Figure 16, respectively. The Relative Potency (%) for Subfamily B was compared to the Bound Molar Ratio as shown in Figure 17. 40

Results

45 [0097] As shown in Figure 14, the total % Polysorbate 80 and the % Polysorbate 80 in the supernatant were the same (0.017%), which indicates that Polysorbate 80 did not bind to the aluminum or get trapped in the pellet. In addition, Polysorbate 80 was stable after 5 months at both 2-8°C and 25°C.

50 [0098] The distribution of Polysorbate 80 in the bound (pellet), unbound (supernatant) and total of rLP2086 Subfamily A and Subfamily B samples are shown in Figure 15 and Figure 16, respectively. While the % Polysorbate 80 in the supernatant and pellet for Subfamily A at 2-8°C and 25°C at 5 months time point did not change. More Polysorbate 80 was observed, however, in the pellet for Subfamily B sample at 25°C at 5 months time point. Despite the different concentrations of Polysorbate 80 in the supernatant and pellet at 2-8 °C and 25°C, an accurate mass balance was achieved for both Subfamilies. As rLP2086 proteins bind 100% to aluminum phosphate at this matrix, Polysorbate 80 associated in the pellet was most likely bound to the protein molecules.

55 [0099] While both protein A and B bound to Polysorbate 80, protein A binding was the same for samples stored at 2-8°C and 25°C, and protein B binding was almost double for samples stored at 25°C compared to samples stored at 2-8°C. The Relative Potency for Subfamily B was determined at both 2-8°C and 25°C at T₀ and 5 months time points and was found to behave inversely to the Bound Molar Ratio as described in Figure 17. The % Potency dropped from 120 at T₀ to 16% at 5M/25°C, while the Bound Molar Ratio increased from 5.3 to 13.9 in the same time period.

Example 3: Critical Molar Ratio Study

[0100] To determine the critical concentration of Polysorbate 80 required for rLP2086 stability, forty (40) rLP2086 formulations were prepared containing Subfamily A only, Subfamily B only, and both Subfamily A and B at 200 µg/mL and 400 µg/mL with different Polysorbate 80 concentrations as described in Table 1. The total and bound proteins were determined for each sample as well as the % Polysorbate 80 in the total, supernatant and pellet at time zero (T_0), 14 days, and 1 Month at both 2-8°C and 25°C. The results from this study are shown in Figure 18 to Figure 24 .

Table 1

Total Protein Antigen	Protein Subfamily	% Polysorbate 80						
		0.003	X	0.005	X	X	X	0.02
0	Placebo	0.003	X	0.005	X	X	X	0.02
200	A	X	0.004	0.005	0.006	0.008	0.01	0.02
200	B	0.003	0.004	0.005	0.006	0.008	0.01	0.02
400	A+B	X	X	X	0.007	X	0.01	0.02
400	A	X	X	X	0.007	X	0.01	0.02
400	B	X	X	X	0.007	X	0.01	0.02

Results

[0101] The Polysorbate 80 concentrations in the supernatant, pellet, and total were determined for all 40 rLP2086 formulation samples with aluminum phosphate. The Total and Bound Molar Ratios were determined for both Subfamily A and B and appear to be similar for both Subfamilies @ 200 µg/mL containing 0.005% Polysorbate 80 (5.4 Molar ratio) or less as shown in Figure 18 and Figure 19, respectively. The Total Molar Ratio for Subfamily B, however, was much higher than the Bound Molar Ratio for samples containing 0.0065% Polysorbate 80 (7.0 Molar ratio) or more. The data for Total and Bound Molar Ratios for Subfamily A, Subfamily B, and Subfamily A+B at 400 µg/mL each were also close for the formulations containing 0.008% Polysorbate 80 (8.6 Molar ratio) or less, however, the Total Molar Ratio was much higher than the Bound Molar Ratio for formulations containing 0.017% Polysorbate 80 (18.4 Molar ratio) as described in Figure 20.

Example 4: Polysorbate 80 Binding Over Time

[0102] The percentage (%) Polysorbate 80 in the Supernatant and Pellet for Subfamily A and B formulation samples with $AlPO_4$ was determined at T_0 , 14 Days/25 °C, 1 Month/4°C, and 1 Month/25°C. The % Polysorbate 80 in Supernatant for both Subfamily A and B formulation samples was relatively the same for samples stored at 2-8°C. The % Polysorbate 80 in the supernatant, however, decreased dramatically for samples stored at 25°C even after only 14 days. The % Polysorbate 80 in Pellet for both Subfamily A and B were relatively similar at T_0 /5°C and 1 Month/5°C C. The % Polysorbate 80 in the supernatant, however, increased significantly for the samples stored at 25°C, especially for Subfamily B containing 0.008% Polysorbate 80 (8.6 Molar ratio) or higher. The % Polysorbate 80 was also determined in the Supernatant and Pellet for rLP2086 Subfamily A and B formulations with $AlPO_4$ at T_0 , 14 Days/25°C, 1 Month/4°C, and 1 Month/25°C. As shown in Figure 21, the Polysorbate 80 concentrations for samples containing 0.008% were approximately the same for all 4 time points. The Polysorbate 80 concentrations, however, increased for the sample containing 0.017% Polysorbate 80 stored at 25°C. No Polysorbate 80 was found in the supernatant of samples containing 0.008% Polysorbate 80 or less. As shown in Figure 22, the Bound Molar Ratio was stable for samples containing 0.008% Polysorbate 80 or less at all 4 time points. The Bound Molar Ratio, however, increased for the sample containing 0.017% Polysorbate 80 stored at 25°C.

[0103] The Potency for Subfamily A and B formulation samples with $AlPO_4$ was determined at T_0 and 14 Days/25°C (Figure 23 and Figure 25, respectively). As described in Figure 23, the Potency for Subfamily A at different total molar ratios ranged from 91 to 102 at both 5°C and 25°C. While the Bound Molar Ratio results were also relatively the same at either temperature, a slight increase in the Potency was seen as the total/Bound Molar Ratio increased.

[0104] The Potency for Subfamily B for the 5°C samples was about 95% for total molar ratios up to 9.0. The Subfamily B potency, however, decreased to 79% as the total molar ratio increased to 18.1. Further, the sample with Total molar ratio at 18.1 had higher bound molar ratio compared to the other sample. At 25°C, the Subfamily B Potency exhibited a significant drop from 83% to 5% as the total molar ratio increased from 5.3 to 18.1. The bound molar ratio values for the 25°C samples increased from 5.3 to 13.8 as the total molar ratio increased. Thus, the Potency for Subfamily B is

inversely proportional to the Bound Molar Ratio.

[0105] Both Subfamily A and Subfamily B proteins bound to Polysorbate 80. Subfamily A binding was the same for samples stored at 2-8°C and 25°C, but Subfamily B binding was almost double for samples stored at 25°C. Further, the Critical Molar Ratio Study indicated that the 200 µg/mL formulation samples were stable when containing 0.008% Polysorbate 80 or less, which is equivalent to a Total Molar Ratio of 4.2 or less.

Example 5: Detergent concentration and rLP2086 Subfamily B Antigen Potency

[0106] Additional stability studies with varying concentrations of Polysorbate 80 corroborated the criticality of the molar ratio of Polysorbate 80 to protein for maintaining potency. In one experiment, the immunogenic composition was formulated at the 200 µg dosage (total protein concentration 400 µg/mL) at pH 6.3 in 10 mM histidine buffered saline (HBS) with 0.5 mg/mL aluminum (as aluminum phosphate) and spiked with 0.01%, 0.02%, 0.05% or 0.1% Polysorbate 80 (corresponding molar ratio of Polysorbate 80 to rLP2086 protein at 5.3, 10.7, 26.7 and 53.4). The formulated samples were incubated at 25°C and control samples were stored at 2-8°C. There was no significant change in potency at time "0" at Polysorbate 80 concentrations up to 0.1%. For longer periods at 2-8°C and 25°C, however, a reduction in potency was observed as a function of temperature and Polysorbate 80 concentration. As the concentration of Polysorbate 80 was increased from 0.01% to 0.1% in the immunogenic composition, the 3-month stability point demonstrated a reduction in potency of the Subfamily B protein to less than 10% and 25% at 25°C and 2-8°C, respectively (Figure 1).

[0107] An additional stability study (Figure 2) was performed evaluating the Subfamily B protein at a concentration of 4 mg/mL in HBS and spiked with Polysorbate 80 to a final concentration of 0.06, 0.5 and 1% (corresponding molar ratios of 3.3, 26.7 and 53.4). The control contained 0.09% Polysorbate 80. The Subfamily B protein in 0.06% Polysorbate 80 (molar ratio of 3.3) was stable. The same samples containing increased concentration of Polysorbate 80 to 0.5% and 1% (molar ratios 26.7 and 53.4 respectively) were unstable. For 400 µg/mL immunogenic composition formulations, instability of Subfamily B protein was noted in all formulations containing 0.01% Polysorbate 80 concentration (5.3 molar ratio) or higher. At 4 mg/mL protein and 0.06% Polysorbate 80 concentrations, however, there was no reduction in potency because the ratio of Polysorbate 80 to protein (3.3) is lower than that at 400 µg/mL protein plus 0.01% Polysorbate 80 concentrations (molar ratio 5.3). The reduction in potency of Subfamily B protein by Polysorbate 80 is thus correlated to the molar ratio of the Polysorbate 80 detergent to protein and not to the absolute concentration of Polysorbate 80 in the matrix.

[0108] Accordingly, Polysorbate 80 concentration must be reduced in the immunogenic composition in order to maintain the stability of the Subfamily B protein in the vaccine and during subsequent storage at 2-8°C. An accelerated 28-day stability study was designed for the immunogenic composition with varying molar ratios of Polysorbate 80 (0, 1.1, 2.7 and 5.3) at 20 and 200 µg dosages (Figure 3 and Figure 4). A bivalent (Subfamily A and Subfamily B) formulation was prepared in 10 mM histidine buffered saline pH 6.0, 0.5 mg/mL aluminum as aluminum phosphate with various Polysorbate 80 concentrations. Samples were incubated at 25°C along with a 2-8°C control group. Samples were analyzed for potency at 0, 7, 14 and 28 days. Both Subfamily A (data not shown) and B proteins were stable for all groups containing less than a 5.3 molar ratio of Polysorbate 80 to protein. A potency value of greater than 80% is considered to be within the assay variability. At the 5.3 molar ratio, a decreasing trend for the Subfamily B protein potency was observed for 25°C samples.

[0109] A comprehensive study evaluated all the potential clinical dosages (20, 60, 120 and 200 µg dosage) formulated with varying Polysorbate 80-to-protein molar ratios under accelerated storage stability conditions to investigate the effects of Polysorbate 80-to-protein molar ratios on the stability of MnB rLP2086 proteins. Bivalent MnB rLP2086 immunogenic compositions formulated at Polysorbate 80-to-protein molar ratios ranging from approximately 1.4 to 10.7 were used. To generate immunogenic compositions formulated at increasing Polysorbate 80-to-protein molar ratios (1.4, 2.4, 3.4, 3.9, 4.3, 4.7 and 10.7), antigens were adjusted to variable molar ratios by adding Polysorbate 80 such that during immunogenic composition formulation, additional Polysorbate 80 was not needed. There were two sets of antigen lots used in this study. One set of Subfamily A and B lots were generated with a Polysorbate 80-to-protein molar ratio of 1.4 and the other set at 2.4. The set of proteins with a molar ratio of 2.4 was used to adjust the molar ratios of 3.4, 3.9, 4.3, and 10.7 by spiking with additional Polysorbate 80. The final matrix of immunogenic composition was 10 mM histidine, 150 mM NaCl, pH 6.0, 0.5 mg/mL aluminum phosphate with the Polysorbate 80-to-protein molar ratios mentioned above. After storage at 2-8°C or 25°C for specific intervals, gentle mixing was applied with a rocker 24 hours prior to testing. Total protein by IEX-HPLC, potency, appearance, optical density at 320nm of the supernatant fraction and pH were tested.

[0110] Potency results of 200 and 20 µg doses are shown in Figure 5 and Figure 6, respectively. The potency assay was more sensitive than other tests used in the study. Overall, there was no significant reduction in potency observed either for Subfamily A or B antigens as compared to the initial time point for all dosages with molar ratios of 4.3 and lower. Formulations with a molar ratio of 4.7 were considered at marginal due to a slight reduction in potency for Subfamily B proteins stored at 25°C. The potency results for Subfamily B antigen for formulations at a molar ratio of 10.7 were significantly lower for samples stored at 25°C than those stored at 2-8°C.

Example 6: Aluminum concentration and rLP2086 Subfamily A and B Antigen Potency

[0111] A number of experiments were conducted to determine the optimal level of aluminum phosphate to ensure greater than 95% binding of both Subfamily A and B proteins. Initial studies focused on optimization of the formulation at pH 6.5. Formulations were prepared with a target dosage of 200 µg/mL of each protein from Subfamily A and B proteins in 10 mM histidine buffer at pH 6.5 with 0.02% Polysorbate 80 and either 0.25 or 0.5 mg/ mL aluminum (as aluminum phosphate). Subfamily B protein bound to aluminum phosphate to a lesser extent than did Subfamily A protein (Figure 7). Increasing the aluminum content from 0.25 mg/mL to 0.5 mg/mL increased binding of Subfamily B protein to >80%. Since the binding mechanism between protein and aluminum suspension is mostly an ionic interaction, the pH of the suspension is a factor that influences binding.

[0112] The formulation pH was optimized to ensure greater than 90-95% binding of the Subfamily B protein. Multiple formulations at 200 µg/mL of each A and B proteins with pH ranging from 5.6 through 6.5 with different lots of immunogenic compositions were examined (Figure 8). Greater than 90-95% binding of both proteins occurred at formulations with pH ranging from 5.6 to 6.4. As the pH of the formulations increased to 6.5 and above, the binding of Subfamily B protein was significantly reduced. The recommended target pH is 6.0 to ensure greater than 90% binding of both Subfamily A and B proteins.

[0113] The robustness of the formulation under formulation variables and/or limits by varying the pH, buffer, protein, and Polysorbate 80 concentrations was also evaluated (Figure 9). While the binding of Subfamily A protein was consistently high (≥95%) with total protein concentration up to 500 µg/mL (250 µg/mL each protein), Subfamily B protein binding was more sensitive to protein concentration and pH. As commercial formulations at a 200 µg dosage are used, the results from this experiment further supported the recommended formulation at a pH of 6.0 with 0.5 mg/mL aluminum phosphate.

[0114] Formulations with and without aluminum phosphate were evaluated to investigate the feasibility of providing a stable formulation without aluminum phosphate at concentrations of Polysorbate 80 low enough for Subfamily B protein stability. Immunogenic compositions were formulated at 20 and 200 µg dosages in histidine buffered saline buffer with Polysorbate 80 concentration ranging from 0 to 5.3 molar ratios. Half of the samples were subjected to agitation with a digital multi-tube vortexer set at 500 rpm under pulse mode (2 seconds on and one second off) for 24 hours prior to testing. This condition was adopted to simulate the ISTA tests (International Safe Transit Association) typically performed at the final immunogenic composition shipping package stage to mimic extreme vibrations during shipping conditions.

[0115] With agitation, formulations without aluminum phosphate precipitated which eventually led to potency loss of both Subfamily A and B antigens. An appearance test (Figure 10) and absorbance measurements at $\lambda = 320$ nm (Figure 11) demonstrated the formation of aggregates and/or precipitates when formulations without aluminum phosphate were agitated. Potency testing of these samples (Figure 12 and Figure 13) demonstrated significant loss of potency for both Subfamily A and B proteins at all time points tested. The loss of potency was most pronounced in formulations containing low amounts of Polysorbate 80. Since low amounts of Polysorbate 80 are necessary to maintain Subfamily B protein stabilization, the inclusion of aluminum phosphate in the formulation is required to preserve stability. rLP2086 immunogenic compositions may be formulated with aluminum phosphate, which will function to enhance potency stability as measured by the in vitro potency assay.

Example 7: Succinate and Histidine as Buffers

[0116] A series of formulations were prepared to compare binding of rLP2086 Subfamily A and B proteins in succinate and histidine, as well as effects of pH, Polysorbate 80, and MgCl₂ on binding (Table 2). The robustness of the formulation under formulation variables and or limits by varying the pH, buffer, protein, and polysorbate concentrations was evaluated (Figure 25 and 26). The binding of aluminum to Subfamily A and Subfamily B protein was similar regardless of the buffer (histidine or succinate) used.

Table 2: Formulations to Evaluate Histidine and Succinate buffers, MgCl₂, Polysorbate 80, and pH 5.6 - 6.0 on Binding of rLP2086 with AlPO₄¹

rLP2086 A (µg/mL)	rLP2086 B (µg/mL)	Histidine (mM)	Succinate (mM)	PS 80 (%)	Saline (%)	MgCl ₂ (mM)	Target pH
200	200	0	5	0.020	0.9	0	6.0
200	200	0	5	0.020	0.9	0	5.8
200	200	0	5	0.020	0.9	0	5.6
200	200	0	5	0.010	0.9	0	6.0
200	200	0	5	0.005	0.9	0	6.0

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(continued)

	rLP2086 A ($\mu\text{g/mL}$)	rLP2086 B ($\mu\text{g/mL}$)	Histidine (mM)	Succinate (mM)	PS 80 (%)	Saline (%)	MgCl ₂ (mM)	Target pH
5	250	250	0	5	0.020	0.9	0	6.0
	250	250	0	5	0.020	0.9	0	5.8
	250	250	0	5	0.020	0.9	0	5.6
	200	200	0	10	0.020	0.9	0	6.0
10	200	200	0	20	0.020	0.9	0	6.0
	200	200	0	5	0.020	0.9	10	6.0
	200	200	10	0	0.020	0.9	0	6.0
	200	200	10	0	0.020	0.9	0	5.8
	200	200	10	0	0.020	0.9	0	5.6
15	200	200	10	0	0.010	0.9	0	6.0
	200	200	10	0	0.005	0.9	0	6.0
	250	250	10	0	0.020	0.9	0	6.0
	250	250	10	0	0.020	0.9	0	5.8
20	250	250	10	0	0.020	0.9	0	5.6
	200	200	5	0	0.020	0.9	0	6.0
	200	200	20	0	0.020	0.9	0	6.0
	200	200	10	0	0.020	0.9	10	6.0

¹All formulations described in Table 2 contain 0.5 mg Al/mL.

[0117] The effect of buffer salt and mixing time on aluminum binding were evaluated with three commonly used buffer salts, chosen because their pKa's are in the physiologic range and because these salts are generally regarded as safe. rLP2086 Subfamily A and B proteins were formulated with one of the three buffer salts: 5 mM succinate, 10 mM histidine, or 10 mM phosphate at a pH suitable for the pKa of each salt to determine the extent of binding at each condition. The time required for binding to reach completion was evaluated by allowing the samples to mix for either 5 or 120 min before measuring the amount of protein bound.

[0118] As shown in Figure 27, Subfamily B protein exhibited reduced binding at pH 6.8 in phosphate buffer, while Subfamily A protein was not significantly affected at the same conditions. The amount of protein bound to aluminum was similar in samples formulated with histidine or succinate. Thus, these two buffer salts were chosen for further evaluation. While not wishing to be bound by theory, it is possible that the reduced binding in phosphate buffer results from competition for binding sites on AlPO₄ with the added phosphate ions.

[0119] At these conditions and concentrations of protein and AlPO₄, binding was complete after 5 min of mixing at room temperature as similar results were obtained after mixing for 2 hours.

[0120] To further examine whether the reduced binding of Subfamily B protein in phosphate buffer at pH 6.8 was due to pH or differences between buffer salts, binding was measured over a pH range of 5.3 to 7.0 in either histidine- or succinate-buffered formulations. Bivalent formulations were prepared containing 0.2 mg/mL of each subfamily protein (0.4 mg/mL total protein), 0.02% PS80, 0.5 mg Al/mL, and 150 mM NaCl. Samples were formulated in either 10 mM histidine or 5 mM succinate to compare the effect of buffer salt. After formulation, the pH of each sample was individually verified.

[0121] The binding profile from pH 5.3 to 7.0 is shown for Subfamily A protein in Figure 28 and for Subfamily B protein in Figure 28. Subfamily A protein exhibited little change in the amount of protein bound, with binding remaining above 95% across the pH range tested. A formulation containing histidine with a target pH 7.0 resulted in a pH of 6.8. The pH was not adjusted to 7.0 (e.g. by addition of base) to avoid possible effects on the protein or AlPO₄ and results for this datapoint are therefore not available.

[0122] The binding profile of Subfamily B protein (shown in Figure 29) exhibited a pH-dependent trend. Whether binding was performed in histidine or succinate buffered formulations, however, the amount of protein bound to aluminum was similar. Binding was dependent on the pH of the formulation rather than the buffering salt. Binding remained at 95% up to pH 6.5 (94% in histidine, 95% in succinate), but decreased when pH was greater than 6.5. At pH 7.0, binding decreased to about 82%, with minor differences between buffering salts.

[0123] To obtain robust binding of Subfamily B protein with AlPO₄ at these concentrations, a pH of 6.5 or less is preferred.

Example 8: Safety, Tolerability And Immunogenicity Study

[0124] A study is conducted to assess the safety, tolerability, and immunogenicity of rLP2086 vaccine administered in a healthy adolescent population, according to regimens of either 0 and 2 months; 0, 2, and 6 months; 0 and 2 months followed by a 12-month booster dose.

[0125] The immunogenic composition is a rLP2086 vaccine (recombinant lipidated). The immunogenic composition includes a *N. meningitidis* serogroup B recombinant ORF2086 protein that was expressed in *Escherichia coli* and formulated in a bivalent vaccine composed of one subfamily A strain and one subfamily B strain of rLP2086. In particular, the immunogenic composition is a 0.5 mL dose formulated to contain 60 µg, 120 µg, or 200 µg each of a purified subfamily A and a purified subfamily B rLP2086 protein, a 2.8 molar ratio of polysorbate 80, and 0.25 mg of Al³⁺ as AlPO₄, 10 mM histidine-buffered saline at pH 6.0. A control composition includes a normal saline solution (0.9% sodium chloride) in a 0.5 mL dose.

[0126] Subjects are randomly assigned to 5 groups. See Table 3. The subjects are stratified into two age groups, ≥11 to <14 and ≥14 to <19 years of age.

Table 3: Study Design

	Vaccinati on 1	Vaccinati on 2	Post-vaccinati on 2 Blood draw	Vaccinati on 3	Post-vaccinati on 3 Blood draw	Vaccinati on 4	Post-vaccinati on 4 Blood Draw
Visit number	1	2	3	4	5	6	7
Approximate month	0	2	3	6	7	12	13
Group 1	rLP2086	rLP2086		Saline		Saline	
Group 2	rLP2086	rLP2086		rLP2086		Saline	
Group 3	rLP2086	rLP2086		Saline		rLP2086	
Group 4	rLP2086	Saline		rLP2086		Saline	
Group 5	Saline	Saline		rLP2086		rLP2086	
Blood Draw	20 mL		20 mL		20 mL	20 mL	20 mL

[0127] Saline is used as a placebo because there is no proven safe, immunogenic, and effective vaccine against MnB that could serve as an active control.

[0128] Subjects receive one dose of rLP2086 vaccine or saline at each of the vaccination visits (e.g., visits 1, 2, 4, and 6) according to Table 3. Standard vaccination practices are observed and the vaccine is not injected into blood vessels. The rLP2086 vaccine is administered intramuscularly by injecting 0.5 mL into the upper deltoid muscle. Saline is administered intramuscularly into the upper deltoid muscle.

A. Visit 1

[0129] On Visit 1, day 1, vaccination 1, the subject first has blood drawn and then receives a vaccination. The visit 1 blood draw and vaccination 1 occurs on the same day. Before vaccination, a blood sample (approximately 20 mL) from the subject is collected. For subjects randomized to group 1, 2, 3, and 4, a single 0.5-mL intramuscular injection of rLP2086 vaccine is administered into the upper deltoid muscle. For subjects in group 5, a single 0.5-mL intramuscular injection of saline is administered into the upper deltoid muscle.

B. Visit 2 (42 to 70 days after Visit 1), Vaccination 2

[0130] For groups 1, 2, and 3, a single 0.5-mL intramuscular injection of rLP2086 vaccine is administered into the upper deltoid muscle. For groups 4 and 5, a single 0.5-mL intramuscular injection of saline is administered into the upper deltoid muscle.

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C. Visit 3 (28 to 42 days after Visit 2), Post vaccination 2 Blood Draw

[0131] A blood sample (approximately 20 mL) from the subject is collected.

5 D. Visit 4 (105 to 126 days after Visit 2), Vaccination 3

[0132] For groups 2, 4, and 5, a single 0.5-mL intramuscular injection of rLP2086 vaccine is administered into the upper deltoid muscle. For groups 1 and 3, a single 0.5-mL intramuscular injection of saline is administered into the upper deltoid muscle.

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E. Visit 5 (28 to 42 days after Visit 4), Post vaccination 3 Blood Draw

[0133] A blood sample (approximately 20 mL) from the subject is collected.

15 F. Visit 6 (161 to 175 days after Visit 4), Vaccination 4

[0134] At visit 6, the subject first has blood drawn and then receives a vaccination. The visit 6 blood draw and vaccination 4 occurs on the same day. Before vaccination, a blood sample (approximately 20 mL) from the subject is collected. For groups 3 and 5, a single 0.5-mL intramuscular injection of rLP2086 vaccine is administered into the upper deltoid muscle. For subjects in groups 1, 2, and 4, a single 0.5-mL intramuscular injection of saline is administered into the upper deltoid muscle.

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G. Visit 7 (28 to 42 days after Visit 6), Post vaccination 4 Blood Draw

25 **[0135]** A blood sample (approximately 20 mL) from the subject is collected.

Immunogenicity Results

[0136] The primary objective of this study was to assess the immunogenicity of 60 µg, 120 µg, and 200 µg rLP2086 vaccine as measured by SBA performed with MnB strains expressing LP2086 subfamily A and B proteins.

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[0137] The secondary objective of this study was to assess the immunogenicity of 60 µg, 120 µg, and 200 µg rLP2086 vaccine as determined by quantitation of Ig binding to rLP2086 vaccine subfamily A and B proteins.

[0138] SBA activity was assessed using 3 subfamily A and 3 subfamily B strains as shown in Table 4.

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Table 4: Analysis of Subjects Achieving SBA Titer of Fold Rise ≥ 4 From Predose 1 - mITT Population (Study 6108A1-2001-WW/B1971005)

Strain	Randomized Vaccine Group	N ^a	n ^b (%)	(95%CI) ^c	p-Value ^d
1-month postdose 2					
Subfamily A Strain 1	Control	80	1 (1.3)	(0.0, 6.8)	>0.9999
	60 µg rLP2086 Vaccine	18	16 (88.9)	(65.3, 98.6)	0.0007
	120 µg rLP2086 Vaccine	115	96 (83.5)	(75.4, 89.7)	<0.0001
Subfamily B Strain 1	200 µg rLP2086 Vaccine	106	93 (87.7)	(79.9, 93.3)	<0.0001
	Control	84	0 (0.0)	(0.0, 4.3)	>0.9999
	60 µg rLP2086 Vaccine	21	15 (71.4)	(47.8, 88.7)	0.0392
	120 µg rLP2086 Vaccine	121	72 (59.5)	(50.2, 68.3)	0.0225
	200 µg rLP2086 Vaccine	114	68 (59.6)	(50.1, 68.7)	0.0244
	1-month postdose 3				
Subfamily A Strain 1	Control	73	4 (5.5)	(1.5, 13.4)	>0.9999
	60 µg rLP2086 Vaccine	19	17 (89.5)	(66.9, 98.7)	0.0004

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Strain	Randomized Vaccine Group	N ^a	n ^b (%)	(95%CI) ^c	p-Value ^d
Subfamily B Strain 1	120 µg rLP2086 Vaccine	111	103 (92.8)	(86.3, 96.8)	<0.0001
	200 µg rLP2086 Vaccine	100	94 (94.0)	(87.4, 97.8)	<0.0001
	Control	79	1 (1.3)	(0.0, 6.9)	>0.9999
200	60 µg rLP2086 Vaccine	21	17 (81.0)	(58.1, 94.6)	0.0036
	120 µg rLP2086 Vaccine	112	97 (86.6)	(78.9, 92.3)	<0.0001
	µg rLP2086 Vaccine	105	89 (84.8)	(76.4, 91.0)	<0.0001

Abbreviation: CI = confidence interval; SBA = serum bactericidal assay.

Note: The assay validation supports a lower limit of quantitation (LLOQ) of Subfamily A strain 1 = 9 and Subfamily B Strain 1 = 10. SBA titers above the LLOQ are considered accurate and their quantitated values will be reported. Values below the LLOQ or denoted as below LLOQ will be set to 0.5*LLOQ for analysis.

[0139] The proportions of subjects with titers achieving a defined level are presented in Table 5. For both subfamilies, the proportions of subjects achieving defined SBA titer levels were greater at postdose 3 than at postdose 2.

Table 5: Subjects Achieving Defined SBA Titer Levels in Stage 1 -mITT Population (Study 6108A1-2001-WW/B1971005)

Strain	Sampling Time Point	SBA Titer	Vaccine Group (as Randomized)															
			Control				60 µg				120 µg				200 µg			
			N ^a	n ^b	%	(95% CI) ^c	N ^a	n ^b	%	(95% CI) ^c	N ^a	n ^b	%	(95% CI) ^c	N ^a	n ^b	%	(95% CI) ^c
Subfamily A Strain 2	Predose 1	32	64	7	10.9	(5.3, 21.2)	15	0	0.0	(0.2, 35.8)	90	9	10.0	(5.3, 18.1)	92	11	12.0	(6.7, 20.3)
		64	64	2	3.1	(0.8, 11.7)	15	0	0.0	(0.2, 35.8)	90	7	7.8	(3.8, 15.4)	92	5	5.4	(2.3, 12.4)
		128	64	0	0.0	(0.0, 11.2)	15	0	0.0	(0.2, 35.8)	90	1	1.1	(0.2, 7.5)	92	0	0.0	(0.0, 8.0)
	1-month postdose 2	32	69	3	4.3	(1.4, 12.6)	21	20	95.2	(72.9, 99.3)	115	113	98.3	(93.3, 99.6)	115	105	91.3	(84.6, 95.3)
		64	69	1	1.4	(0.2, 9.6)	21	18	85.7	(63.9, 95.3)	115	95	82.6	(74.6, 88.5)	115	83	72.2	(63.3, 79.6)
		128	69	1	1.4	(0.2, 9.6)	21	11	52.4	(31.8, 72.1)	115	51	44.3	(35.5, 53.5)	115	49	42.6	(33.9, 51.8)
	1-month postdose 3	32	57	5	8.8	(3.7, 19.4)	19	18	94.7	(70.6, 99.3)	108	108	100.0	(93.1, 100.0)	99	98	99.0	(93.2, 99.9)
		64	57	3	5.3	(1.7, 15.1)	19	18	94.7	(70.6, 99.3)	108	103	95.4	(89.4, 98.1)	99	90	90.9	(83.4, 95.2)
		128	57	2	3.5	(0.9, 13.0)	19	13	68.4	(45.2, 85.1)	108	73	67.6	(58.2, 75.7)	99	67	67.7	(57.9, 76.1)
Subfamily A Strain 1	Predose 1	16	81	10	12.3	(6.8, 21.4)	21	1	4.8	(0.7, 27.1)	122	11	9.0	(5.1, 15.6)	114	7	6.1	(3.0, 12.3)
		32	81	6	7.4	(3.4, 15.5)	21	1	4.8	(0.7, 27.1)	122	7	5.7	(2.8, 11.5)	114	5	4.4	(1.8, 10.1)
		64	81	4	4.9	(1.9, 12.4)	21	0	0.0	(0.1, 28.2)	122	3	2.5	(0.8, 7.3)	114	2	1.8	(0.4, 6.7)
	1-month postdose 2	128	81	1	1.2	(0.2, 8.2)	21	0	0.0	(0.1, 28.2)	122	1	0.8	(0.1, 5.6)	114	0	0.0	(0.0, 6.6)
		16	83	6	7.2	(3.3, 15.2)	18	16	88.9	(64.8, 97.2)	118	105	89.0	(81.9, 93.5)	110	100	90.9	(83.9, 95.0)
		32	83	3	3.6	(1.2, 10.6)	18	16	88.9	(64.8, 97.2)	118	101	85.6	(78.0, 90.9)	110	90	81.8	(73.5, 88.0)
	1-month postdose 3	64	83	1	1.2	(0.2, 8.1)	18	15	83.3	(59.1, 94.5)	118	70	59.3	(50.2, 67.8)	110	71	64.5	(55.2, 72.9)
		128	83	0	0.0	(0.0, 8.9)	18	6	33.3	(15.8, 57.1)	118	33	28.0	(20.6, 36.7)	110	44	40.0	(31.3, 49.4)
		16	76	9	11.8	(6.3, 21.2)	20	18	90.0	(67.6, 97.5)	114	110	96.5	(91.0, 98.7)	104	100	96.2	(90.2, 98.5)

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Vaccine Group (as Randomized)														
rLP2086 Vaccine														
200 µg														
Strain	Sampling Time Point	SBA Titer	Control			60 µg			120 µg			200 µg		
			N ^a	n ^b	% (95% CI ^c)	N ^a	n ^b	% (95% CI ^c)	N ^a	n ^b	% (95% CI ^c)	N ^a	n ^b	% (95% CI ^c)
		32	76	6	7.9 (3.6, 16.5)	20	18	90.0 (67.6, 97.5)	114	108	94.7 (88.8, 97.6)	104	99	95.2 (89.0, 98.0)
		64	76	3	3.9 (1.3, 11.5)	20	17	85.0 (62.4, 95.1)	114	102	89.5 (82.4, 93.9)	104	91	87.5 (79.7, 92.6)
Control														
Strain	Sampling Time Point	SBA Titer	Control			60 µg			120 µg			200 µg		
			N ^a	n ^b	% (95% CI ^c)	N ^a	n ^b	% (95% CI ^c)	N ^a	n ^b	% (95% CI ^c)	N ^a	n ^b	% (95% CI ^c)
Subfamily A Strain 3	Predose 1	128	76	1	1.3 (0.2, 8.8)	20	8	40.0 (21.4, 62.0)	114	78	68.4 (59.3, 76.3)	104	63	60.6 (50.9, 69.5)
		16	80	8	10.0 (5.1, 18.7)	21	1	4.8 (0.7, 27.1)	119	9	7.6 (4.0, 13.9)	115	7	6.1 (2.9, 12.2)
		32	80	6	7.5 (3.4, 15.7)	21	1	4.8 (0.7, 27.1)	119	6	5.0 (2.3, 10.8)	115	5	4.3 (1.8, 10.0)
		64	80	2	2.5 (0.6, 9.4)	21	0	0.0 (0.1, 28.2)	119	1	0.8 (0.1, 5.7)	115	4	3.5 (1.3, 8.9)
		128	80	1	1.3 (0.2, 8.3)	21	0	0.0 (0.1, 28.2)	119	0	0.0 (0.0, 6.3)	115	0	0.0 (0.0, 6.5)
	1-month postdose 2	16	81	10	12.3 (6.8, 21.4)	20	20	85.0 (62.4, 95.1)	111	111	(89.1, 97.7)	107	107	96.3 (90.5, 98.6)
						17	17		94.9			103	103	
		32	81	9	11.1 (5.9, 20.0)	20	17	85.0 (62.4, 95.1)	117	106	90.6 (83.8, 94.7)	107	95	88.8 (81.3, 93.5)
		64	81	8	9.9 (5.0, 18.5)	20	16	80.0 (57.2, 92.3)	117	94	80.3 (72.1, 86.6)	107	76	71.0 (61.8, 78.8)
		128	81	2	2.5 (0.6, 9.3)	20	9	45.0 (25.3, 66.4)	117	46	39.3 (30.9, 48.4)	107	49	45.8 (36.6, 55.3)
	1-month postdose 3	16	78	9	11.5 (6.1, 20.7)	21	20	95.2 (72.9, 99.3)	114	111	97.4 (92.2, 99.1)	112	107	95.5 (89.7, 98.1)
		32	78	7	9.0 (4.3, 17.6)	21	18	85.7 (63.9, 95.3)	114	107	93.9 (87.7, 97.0)	112	105	93.8 (87.5, 97.0)
Vaccine Group (as Randomized)														
rLP2086 Vaccine														
200 µg														
Strain	Sampling Time Point	SBA Titer	Control			60 µg			120 µg			200 µg		
			N ^a	n ^b	% (95% CI ^c)	N ^a	n ^b	% (95% CI ^c)	N ^a	n ^b	% (95% CI ^c)	N ^a	n ^b	% (95% CI ^c)
		64	78	3	3.8 (1.2, 11.3)	21	15	71.4 (49.2, 86.6)	114	104	91.2 (84.5, 95.2)	112	95	84.8 (76.9, 90.4)
		128	78	2	2.6 (0.6, 9.7)	21	13	61.9 (40.2, 79.7)	114	85	74.6 (65.8, 81.7)	112	75	67.0 (57.8, 75.0)

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Vaccine Group (as Randomized)														
rLP2086 Vaccine														
Strain	Sampling Time Point	SBA Titer	Control			60 µg			120 µg			200 µg		
			N ^a	n ^b	% (95% CI ^c)	N ^a	n ^b	% (95% CI ^c)	N ^a	n ^b	% (95% CI ^c)	N ^a	n ^b	% (95% CI ^c)
Subfamily B Strain 1	Predose 1	16	84	3	3.6 (1.2, 10.5)	22	0	0.0 (0.1, 27.3)	124	2	1.6 (0.4, 6.2)	118	3	2.5 (0.8, 7.6)
		32	84	1	1.2 (0.2, 8.0)	22	0	0.0 (0.1, 27.3)	124	1	0.8 (0.1, 5.5)	118	2	1.7 (0.4, 6.5)
		64	84	0	0.0 (0.0, 8.8)	22	0	0.0 (0.1, 27.3)	124	1	0.8 (0.1, 5.5)	118	0	0.0 (0.0, 6.4)
		128	84	0	0.0 (0.0, 8.8)	22	0	0.0 (0.1, 27.3)	124	1	0.8 (0.1, 5.5)	118	0	0.0 (0.0, 6.4)
	1-month postdose 2	16	84	1	1.2 (0.2, 8.0)	21	16	76.2 (54.0, 89.7)	122	84	68.9 (60.1, 76.4)	114	75	65.8 (56.6, 73.9)
		32	84	0	0.0 (0.0, 8.8)	21	12	57.1 (36.0, 76.0)	122	56	45.9 (37.3, 54.8)	114	55	48.2 (39.2, 57.4)
		64	84	0	0.0 (0.0, 8.8)	21	9	42.9 (24.0, 64.0)	122	31	25.4 (18.5, 33.9)	114	25	21.9 (15.3, 30.4)
		128	84	0	0.0 (0.0, 8.8)	21	3	14.3 (4.7, 36.1)	122	9	7.4 (3.9, 13.6)	114	10	8.8 (4.8, 15.5)
	1-month postdose 3	16	79	3	3.8 (1.2, 11.1)	21	18	85.7 (63.9, 95.3)	113	102	90.3 (83.3, 94.5)	105	90	85.7 (77.6, 91.2)
		32	79	1	1.3 (0.2, 8.4)	21	17	81.0 (58.8, 92.7)	113	89	78.8 (70.3, 85.3)	105	83	79.0 (70.2, 85.8)

Vaccine Group (as Randomized)														
rLP2086 Vaccine														
Strain	Sampling Time Point	SBA Titer	Control			60 µg			120 µg			200 µg		
			N ^a	n ^b	% (95% CI ^c)	N ^a	n ^b	% (95% CI ^c)	N ^a	n ^b	% (95% CI ^c)	N ^a	n ^b	% (95% CI ^c)
Subfamily B Strain 2	Predose 1	64	79	0	0.0 (0.0, 9.3)	21	12	57.1 (36.0, 76.0)	113	59	52.2 (43.0, 61.2)	105	63	60.0 (50.4, 68.9)
		128	79	0	0.0 (0.0, 9.3)	21	9	42.9 (24.0, 64.0)	113	22	19.5 (13.2, 27.8)	105	20	19.0 (12.6, 27.7)
		16	83	2	2.4 (0.6, 9.1)	22	0	0.0 (0.1, 27.3)	118	3	2.5 (0.8, 7.6)	117	3	2.6 (0.8, 7.6)
		32	83	2	2.4 (0.6, 9.1)	22	0	0.0 (0.1, 27.3)	118	3	2.5 (0.8, 7.6)	117	3	2.6 (0.8, 7.6)
	1-month postdose 2	64	83	1	1.2 (0.2, 8.1)	22	0	0.0 (0.1, 27.3)	118	0	0.0 (0.0, 6.4)	117	1	0.9 (0.1, 5.8)
		128	83	0	0.0 (0.0, 8.9)	22	0	0.0 (0.1, 27.3)	118	0	0.0 (0.0, 6.4)	117	0	0.0 (0.0, 6.4)
		16	84	2	2.4 (0.6, 9.0)	19	4	21.1 (8.1, 44.6)	102	33	32.4 (24.0, 42.0)	96	29	30.2 (21.9, 40.1)
		32	84	2	2.4 (0.6, 9.0)	19	4	21.1 (8.1, 44.6)	102	31	30.4 (22.3, 40.0)	96	27	28.1 (20.0, 37.9)
	1-month postdose 2	64	84	1	1.2 (0.2, 8.0)	19	1	5.3 (0.7, 29.4)	102	23	22.5 (15.5, 31.7)	96	19	19.8 (13.0, 29.0)

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Strain	Sampling Time Point	SBA Titer	Control			60 µg			120 µg			200 µg					
			N ^a	n ^b	%	N ^a	n ^b	%	N ^a	n ^b	%	N ^a	n ^b	%			
	16	83	1	1.2	(0.2, 8.1)	21	16	76.2	(54.0, 91.3)	115	99	86.1	(78.5, 91.3)	111	95	85.6	(77.8, 91.3)
	32	83	0	0.0	(0.0, 8.9)	21	16	76.2	(54.0, 91.3)	115	93	80.9	(72.6, 87.1)	111	89	80.2	(71.7, 87.1)
	64	83	0	0.0	(0.0, 8.9)	21	10	47.6	(27.9, 62.8)	115	62	53.9	(44.8, 62.8)	111	68	61.3	(51.9, 62.8)
	128	83	0	0.0	(0.0, 8.9)	21	2	9.5	(2.4, 31.1)	115	26	22.6	(15.9, 31.1)	111	22	19.8	(13.4, 31.1)

Abbreviations: CI = confidence interval; SBA = serum bactericidal assay; LLOQ = lower limit of quantitation.
 Note: LLOQ for A/1 = 9, A/2 = 18, A/3 = 12, B/1 = 10, B/2 = 9, B/3 = 7.

[0140] The immunogenicity data show that the vaccine can generate antibodies with significant SBA activity against subfamily A and subfamily B strains of MnB. For Subfamily A Strain 2, after dose 2, SBA response rates ranged from 88.9% to 90.9% and after dose 3 the SBA response rates ranged from 90.0% to 97.4%. For Subfamily A Strain 1 variant, after both dose 2 and dose 3, 100.0% of the subjects had SBA responses to this variant at both the 60- μ g and 120- μ g dose levels. At the 200- μ g dose level, 96.5% and 99.0% of the subjects had SBA responses after dose 2 and dose 3, respectively. For Subfamily A Strain 1 variant, the SBA response rates ranged from 85.0% to 96.3% after dose 2 and from 95.2% to 97.4% after dose 3.

[0141] For Subfamily B Strain 1 variant, after dose 2 the SBA response rates ranged from 76.2% to 81.0%, and after dose 3 the SBA response rates ranged from 89.5% to 92.0%. For Subfamily B Strain 2 variant after dose 2, the percentage of subjects with SBA response rates ranged from 21.1% to 33.3%. However, after the third dose, 53.3%, 75.6%, and 67.9% of the subjects had SBA responses at the 60- μ g, 120- μ g, and 200- μ g dose levels, respectively. For Subfamily B Strain 3 variant, the SBA response rates ranged from 61.9% to 70.8% after dose 2 and from 76.2% to 88.7% after dose 3.

[0142] Overall, a high proportion of subjects responded with an SBA titer \geq LLOQ irrespective of the subfamily A or subfamily B strain tested. The hSBA data showed robust immune responses at doses of 60 μ g to 200 μ g without a clear dose-response relationship. The frequency of response, irrespective of the analysis examined, was highest numerically in the 120- μ g group. The 200- μ g group did not have improved immune responses over the 120- μ g dose level.

Claims

1. A method for stabilizing the potency of a LP2086 (fHBP) Subfamily A polypeptide and a LP2086 (fHBP) Subfamily B polypeptide in an immunogenic composition by storing the LP2086 (fHBP) Subfamily A polypeptide and the LP2086 (fHBP) Subfamily B polypeptide in a buffer with between about 0.1 mg/ml and about 1 mg/ml aluminum and a molar ratio of detergent to protein of between about 0.5:1 and about 10:1, wherein the immunogenic composition does not include further active immunogens.
2. The method according to claim 1, wherein the molar ratio of detergent to protein is between about 1:1 and about 5:1.
3. The method according to claim 2, wherein the molar ratio of detergent to protein is between about 1.4:1 and about 4.2:1.
4. The method according to claim 3, wherein the molar ratio of detergent to protein is about 2.8:1.
5. The method according to any one of claims 1-4, wherein the amount of detergent is sufficient to reduce polypeptide binding to silicon in a container.
6. The method according to claim 5, wherein the container is a syringe or a vial.
7. The method according to any one of claims 1-6, wherein the detergent is a polysorbate detergent.
8. The method according to claim 7, wherein the detergent is Polysorbate 80.
9. The method according to any one of claims 1 to 8 wherein the concentration of aluminum is about 0.5 mg/ml.
10. The method according to any one of claims 1-9, wherein aluminum is present as AlPO_4 , $\text{Al}(\text{OH})_3$, $\text{Al}_2(\text{SO}_4)_3$ or alum.
11. The method according to any one of claims 1-10, wherein the buffer further comprises histidine.
12. The method according to claim 11, wherein the concentration of histidine is between about 2mM and about 20mM or is between about 5mM and about 15mM.
13. The method according to any one of claims 1-10, wherein the buffer further comprises succinate.
14. The method according to claim 13, wherein the concentration of succinate is between about 2mM and about 10mM or is between about 3mM and about 7mM.
15. The method according to any one of claims 1-14, wherein the pH is between about 5.0 and about 8.0.

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16. The method according to claim 15, wherein the pH is between about 5.8 and 6.0.
17. The method according to claim 12, wherein the concentration of histidine is about 10mM, pH 6.0.
- 5 18. The method according to claim 14, wherein the concentration of succinate is about 5mM, pH 6.0.
19. The method according to any one of claims 1-18, further comprising the step of (b) lyophilizing the immunogenic composition.
- 10 20. The method according to claim 1, wherein the buffer consists essentially of:
- about a 2.8:1 molar ratio of Polysorbate 80 to protein, 0.5 mg/mL aluminum as AlPO_4 , 10mM histidine pH 6.0, and 150 mM NaCl or;
 - about a 2.8:1 molar ratio of Polysorbate 80 to protein, 0.5 mg/mL aluminum as AlPO_4 , 5mM succinate pH 6.0, and 150 mM NaCl.
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Patentansprüche

- 20 1. Verfahren zur Stabilisierung der Wirksamkeit eines LP2086 (fHBP)-Polypeptids der Unterfamilie A und eines LP2086 (fHBP)-Polypeptids der Unterfamilie B in einer immunogenen Zusammensetzung durch Lagerung des LP2086 (fHBP)-Polypeptids der Unterfamilie A und des LP2086 (fHBP)-Polypeptids der Unterfamilie B in einem Puffer mit zwischen etwa 0,1 mg/ml und etwa 1 mg/ml Aluminium und einem molaren Verhältnis von Detergens zu Protein von etwa 0,5:1 bis etwa 10:1,
- 25 wobei die immunogene Zusammensetzung keine weiteren aktiven Immunogene enthält.
2. Verfahren nach Anspruch 1, wobei das molare Verhältnis von Detergens zu Protein zwischen etwa 1:1 und etwa 5:1 liegt.
- 30 3. Verfahren nach Anspruch 2, wobei das molare Verhältnis von Detergens zu Protein zwischen etwa 1,4:1 und etwa 4,2:1 liegt.
4. Verfahren nach Anspruch 3, wobei das molare Verhältnis von Detergens zu Protein etwa 2,8:1 beträgt.
- 35 5. Verfahren nach einem der Ansprüche 1 bis 4, wobei die Menge des Detergens ausreicht, um die Bindung des Polypeptids an Silicium in einem Behälter zu verringern.
6. Verfahren nach Anspruch 5, wobei der Behälter eine Spritze oder ein Fläschchen ist.
- 40 7. Verfahren nach einem der Ansprüche 1 bis 6, wobei das Detergens ein Polysorbat-Detergens ist.
8. Verfahren nach Anspruch 7, wobei das Detergens Polysorbat 80 ist.
9. Verfahren nach einem der Ansprüche 1 bis 8, wobei die Konzentration von Aluminium etwa 0,5 mg/ml beträgt.
- 45 10. Verfahren nach einem der Ansprüche 1 bis 9, wobei das Aluminium als AlPO_4 , $\text{Al}(\text{OH})_3$, $\text{Al}_2(\text{SO}_4)_3$ oder Alaun vorliegt.
11. Verfahren nach einem der Ansprüche 1-10, wobei der Puffer außerdem Histidin umfasst.
- 50 12. Verfahren nach Anspruch 11, wobei die Konzentration von Histidin zwischen etwa 2 mM und etwa 20 mM oder zwischen etwa 5 mM und etwa 15 mM liegt.
13. Verfahren nach einem der Ansprüche 1-10, wobei der Puffer weiterhin Succinat umfasst.
- 55 14. Verfahren nach Anspruch 13, wobei die Konzentration von Succinat zwischen etwa 2 mM und etwa 10 mM oder zwischen etwa 3 mM und etwa 7 mM liegt.
15. Verfahren nach einem der Ansprüche 1 bis 14, wobei der pH-Wert zwischen etwa 5,0 und etwa 8,0 liegt.

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16. Verfahren nach Anspruch 15, wobei der pH-Wert zwischen etwa 5,8 und 6,0 liegt.
17. Verfahren nach Anspruch 12, wobei die Konzentration von Histidin etwa 10 mM, pH 6,0, beträgt.
- 5 18. Verfahren nach Anspruch 14, wobei die Konzentration von Succinat etwa 5 mM, pH 6,0, beträgt.
19. Verfahren nach einem der Ansprüche 1 bis 18, das ferner den Schritt (b) des Lyophilisierens der immunogenen Zusammensetzung umfasst.
- 10 20. Verfahren nach Anspruch 1, wobei der Puffer im Wesentlichen besteht aus:
- etwa einem molaren Verhältnis von 2,8:1 von Polysorbat 80 zu Protein, 0,5 mg/ml Aluminium als AlPO_4 , 10 mM Histidin pH 6,0 und 150 mM NaCl oder;
 - etwa einem molaren Verhältnis von 2,8:1 von Polysorbat 80 zu Protein, 0,5 mg/ml Aluminium als AlPO_4 , 5
- 15 mM Succinat pH 6,0 und 150 mM NaCl.

Revendications

- 20 1. Procédé pour stabiliser la puissance d'un polypeptide de la sous-famille A de LP2086 (fHBP) et d'un polypeptide de la sous-famille B de LP2086 (fHBP) dans une composition immunogène en stockant le polypeptide de la sous-famille A de LP2086 (fHBP) et le polypeptide de la sous-famille B de LP2086 (fHBP) dans un tampon avec entre environ 0,1 mg/ml et environ 1 mg/ml d'aluminium et un rapport molaire détergent sur protéine entre environ 0,5:1 et environ 10:1,
- 25 dans lequel la composition immunogène ne comporte pas d'autres immunogènes actifs.
2. Procédé selon la revendication 1, dans lequel le rapport molaire détergent sur protéine est entre environ 1:1 et environ 5:1.
- 30 3. Procédé selon la revendication 2, dans lequel le rapport molaire détergent sur protéine est entre environ 1,4:1 et environ 4,2:1.
4. Procédé selon la revendication 3, dans lequel le rapport molaire détergent sur protéine est d'environ 2,8:1.
- 35 5. Procédé selon l'une quelconque des revendications 1 à 4, dans lequel la quantité de détergent est suffisante pour réduire une liaison du polypeptide à du silicium dans un récipient.
6. Procédé selon la revendication 5, dans lequel le récipient est une seringue ou un flacon.
- 40 7. Procédé selon l'une quelconque des revendications 1 à 6, dans lequel le détergent est un détergent de polysorbate.
8. Procédé selon la revendication 7, dans lequel le détergent est le Polysorbate 80.
9. Procédé selon l'une quelconque des revendications 1 à 8, dans lequel la concentration en aluminium est d'environ
- 45 0,5 mg/ml.
10. Procédé selon l'une quelconque des revendications 1 à 9, dans lequel de l'aluminium est présent sous forme d' AlPO_4 , $\text{Al}(\text{OH})_3$, $\text{Al}_2(\text{SO}_4)_3$ ou d'alun.
- 50 11. Procédé selon l'une quelconque des revendications 1 à 10, dans lequel le tampon comprend en outre de l'histidine.
12. Procédé selon la revendication 11, dans lequel la concentration en histidine est entre environ 2 mM et environ 20 mM ou est entre environ 5 mM et environ 15 mM.
- 55 13. Procédé selon l'une quelconque des revendications 1 à 10, dans lequel le tampon comprend en outre du succinate.
14. Procédé selon la revendication 13, dans lequel la concentration en succinate est entre environ 2 mM et environ 10 mM ou est entre environ 3 mM et environ 7 mM.

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15. Procédé selon l'une quelconque des revendications 1 à 14, dans lequel le pH est entre environ 5,0 et environ 8,0.

16. Procédé selon la revendication 15, dans lequel le pH est entre environ 5,8 et 6,0.

5 17. Procédé selon la revendication 12, dans lequel la concentration en histidine est d'environ 10 mM, pH 6,0.

18. Procédé selon la revendication 14, dans lequel la concentration en succinate est d'environ 5 mM, pH 6,0.

10 19. Procédé selon l'une quelconque des revendications 1 à 18, comprenant en outre l'étape consistant à (b) lyophiliser la composition immunogène.

20. Procédé selon la revendication 1, dans lequel le tampon est constitué essentiellement de :

15 - un rapport molaire Polysorbate 80 sur protéine d'environ 2,8:1, 0,5 mg/mL d'aluminium sous forme d' AlPO_4 , 10 mM d'histidine pH 6,0, et 150 mM de NaCl ou ;

- un rapport molaire Polysorbate 80 sur protéine d'environ 2,8:1, 0,5 mg/mL d'aluminium sous forme d' AlPO_4 , 5 mM de succinate pH 6,0, et 150 mM de NaCl.

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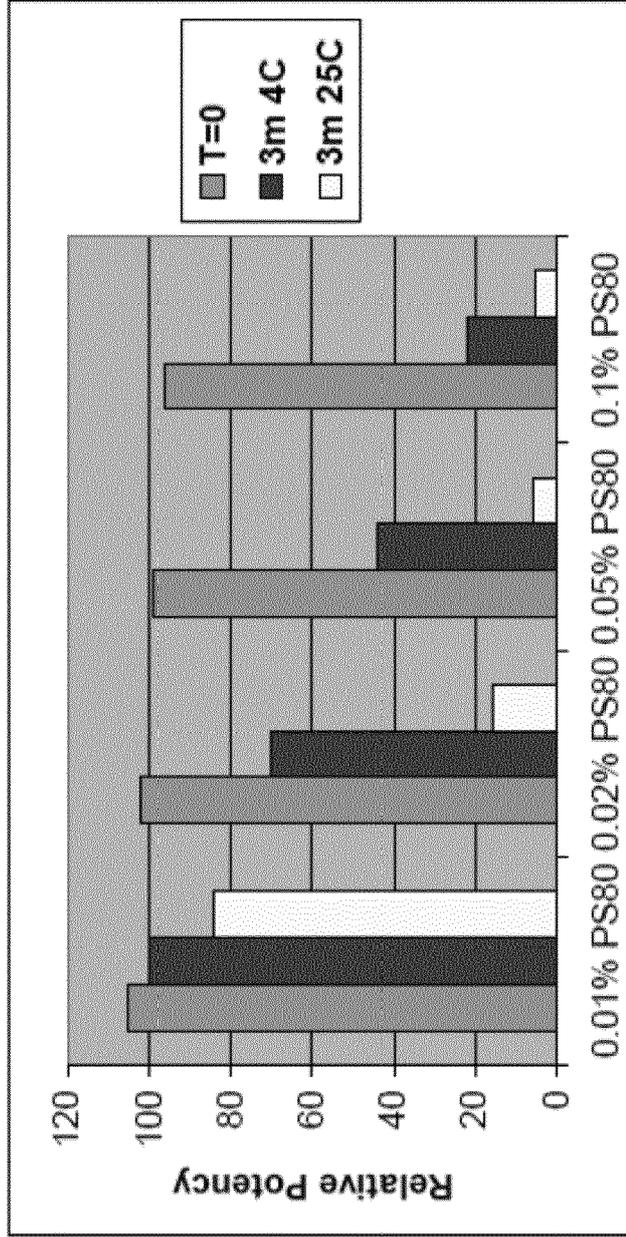
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FIGURE 1

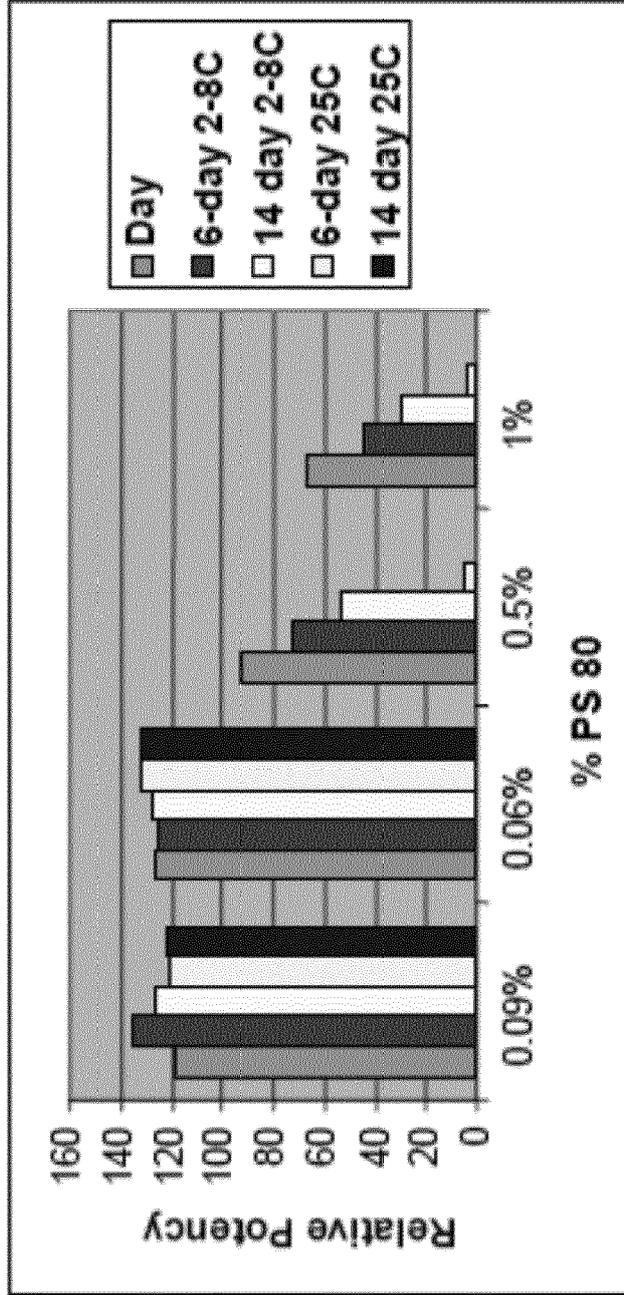
Stability of Subfamily B in Formulations with Various Polysorbate 80 Concentrations



200 µg/ mL each of subfamily A and B were formulated in 10 mM Histidine buffer at pH 6.3 with 0.5 mg/ mL Aluminum and variable Polysorbate 80 concentrations. Formulations were filled in BD syringes and subjected to storage at 2-8°C or 25°C. Potency values for subfamily B at initial and 3 months time point are shown.

FIGURE 2

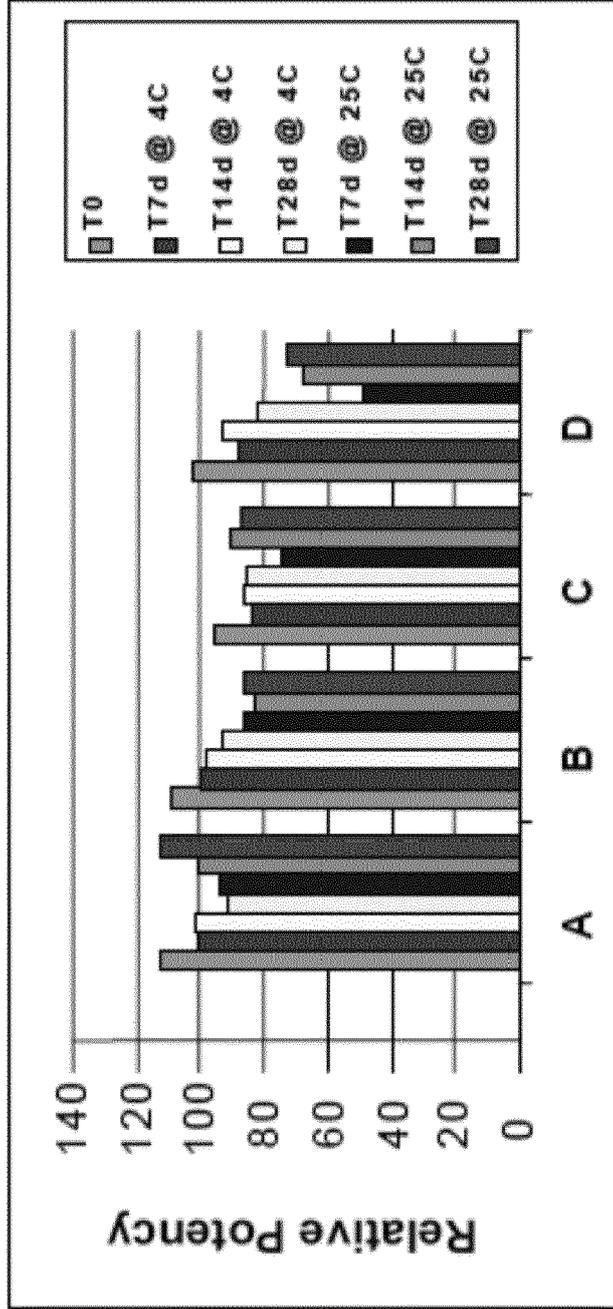
Accelerated Stability of Subfamily B with Various Polysorbate 80 Concentrations



Drug Substance subfamily B formulated in 0.09% Polysorbate 80, 10 mM Histidine, pH 6.5 was spiked with variable Polysorbate 80 concentrations to monitor the effect of Polysorbate 80 on stability. Samples were subjected to storage at 2-8°C or 25°C, and potency was measured at initial, 6-day and 14-day time points.

FIGURE 3

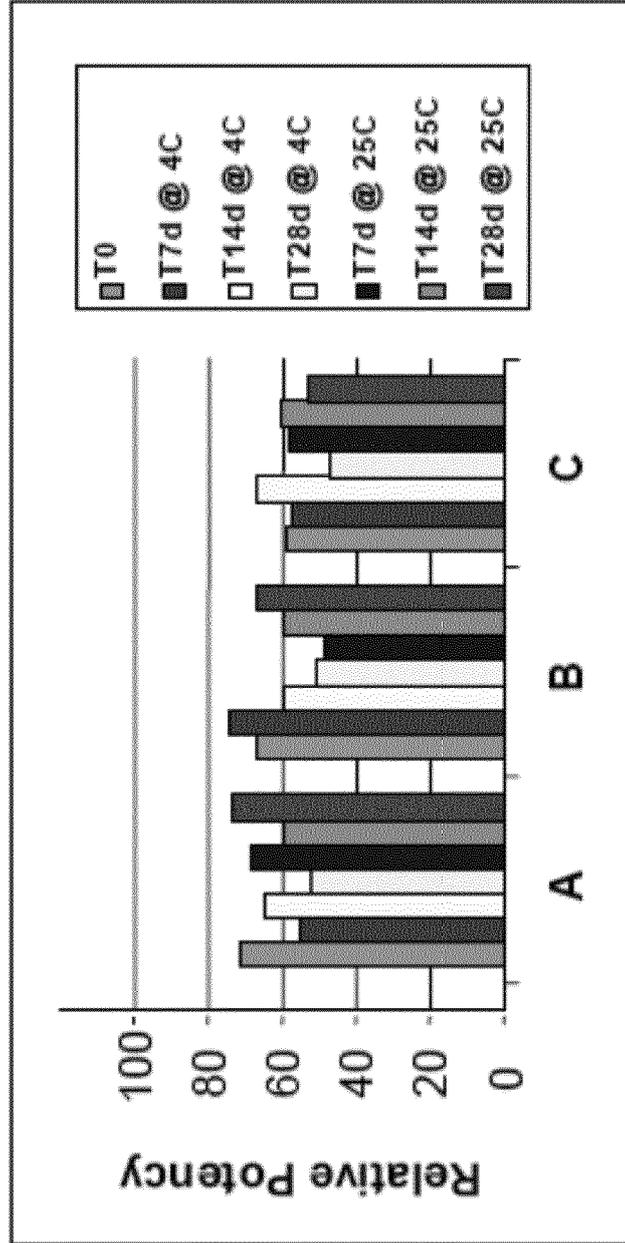
Potency of Subfamily B at 200 µg/mL for 28 days



MmB rLP2086 subfamily A + B formulated at 200 µg/mL dosage each with 0.5 mg/mL aluminum as aluminum phosphate in the presence of varying concentrations of Polysorbate 80 (A = 0%, B = 0.002%, C = 0.005% and D = 0.01%) and stored at either 2-8°C or 25°C. The Polysorbate 80: protein molar ratios of A, B, C and D are 0, 1.1, 2.7 and 5.3, respectively. The potency of subfamily B protein was then tested at 0, 7, 14 and 28 days. At each time point, the samples were agitated as described for 24 hours prior to the testing.

FIGURE 4

Potency of Subfamily B at 20 µg/mL for 28 days



MnB rLP2086 subfamily A + B formulated at 20 µg/mL dosage each with 0.5 mg/ mL aluminum as aluminum phosphate in the presence of varying concentrations of Polysorbate-80 (A = 0%, B = 0.0005% and C = 0.001%) and stored at either 2-8°C or 25°C. The Polysorbate 80: protein molar ratios of A, B and C are 0, 2.7 and 5.3, respectively. The potency of subfamily B protein was then tested at 0, 7, 14 and 28 days. At each time point, the samples were agitated as described for 24 hours prior to the testing.

Potency Results for 200 µg/mL with Different Molar Ratios

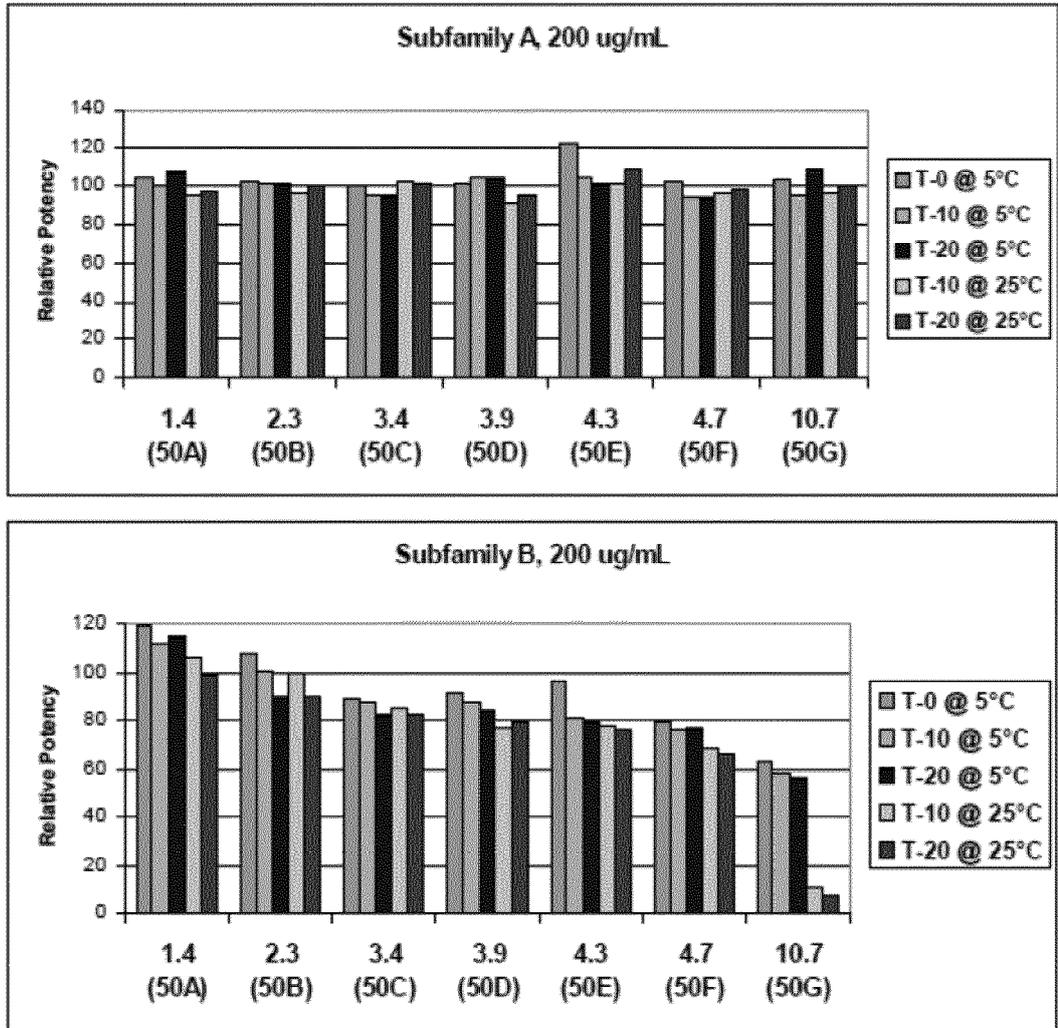
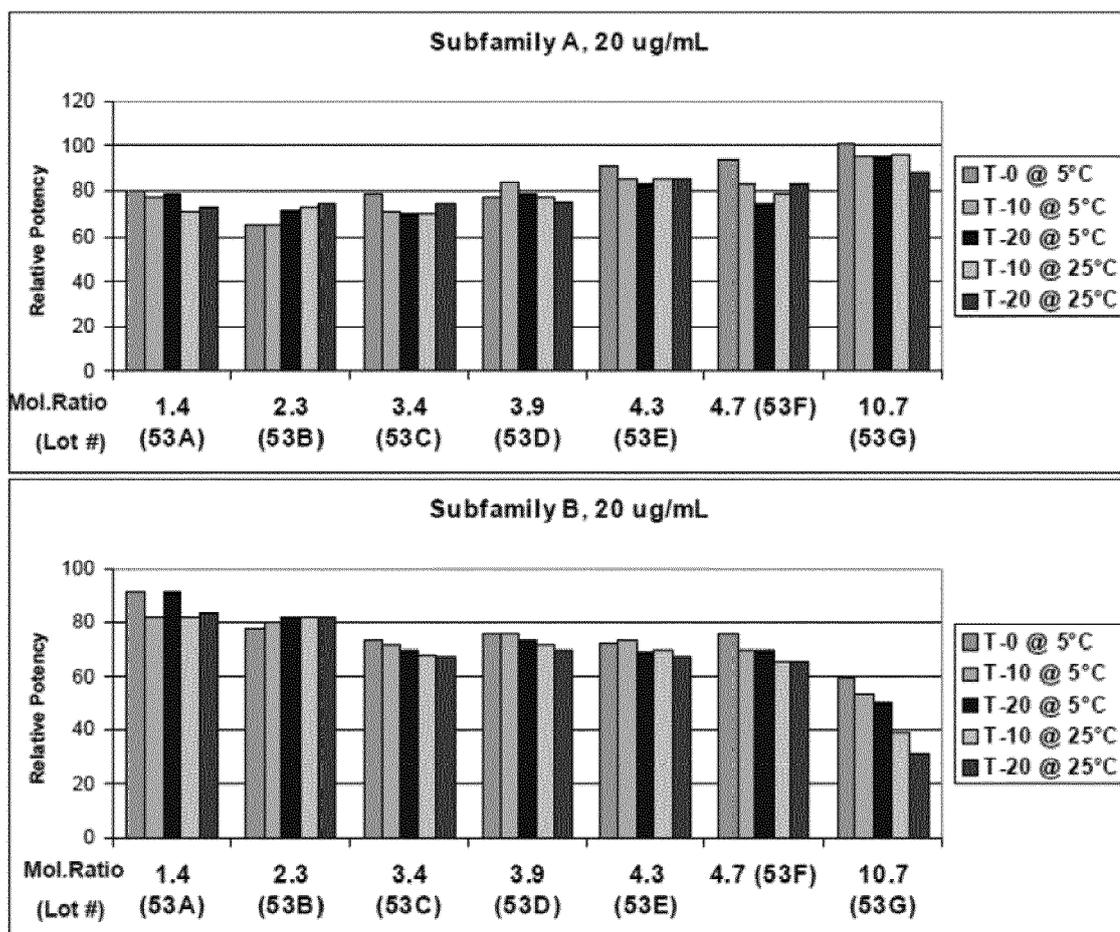


FIGURE 5

MnB rLP2086 subfamily A + B formulated at 200 µg/mL with Polysorbate-80:Protein molar ratios 1.4, 2.3, 3.4, 3.9, 4.3, 4.7, 10.7 for lots 50A, 50B, 50C, 50D, 50E, 50F, 50G, respectively. T-0, T-10, and T-20 represent time zero, 10, and 20 days, respectively. Top graph is potency results for subfamily A and bottom graph is potency results for subfamily B protein. At each time point, the samples were agitated as described for 24 hours prior to the testing.

FIGURE 6

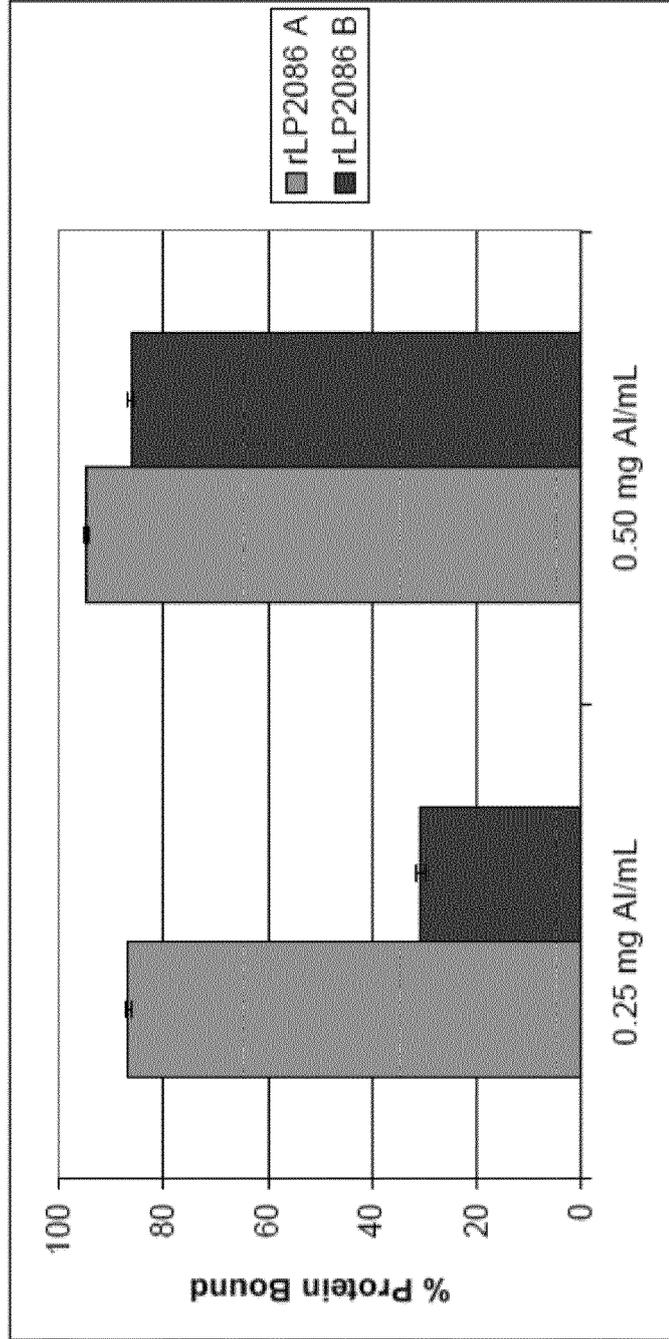
Potency Results for 20 µg/mL with Different Molar Ratios



MnB rLP2086 subfamily A + B formulated at 20 µg/mL with Polysorbate 80:protein molar ratios 1.4, 2.3, 3.4, 3.9, 4.3, 4.7, 10.7 for lots 53A, 53B, 53C, 53D, 53E, 53F, 53G, respectively. T-0, T-10, and T-20 represents time zero, 10, and 20 days, respectively. Top graph is potency results for subfamily A and bottom graph is potency results for subfamily B protein. At each time point, the samples were agitated as described for 24 hours prior to the testing.

FIGURE 7

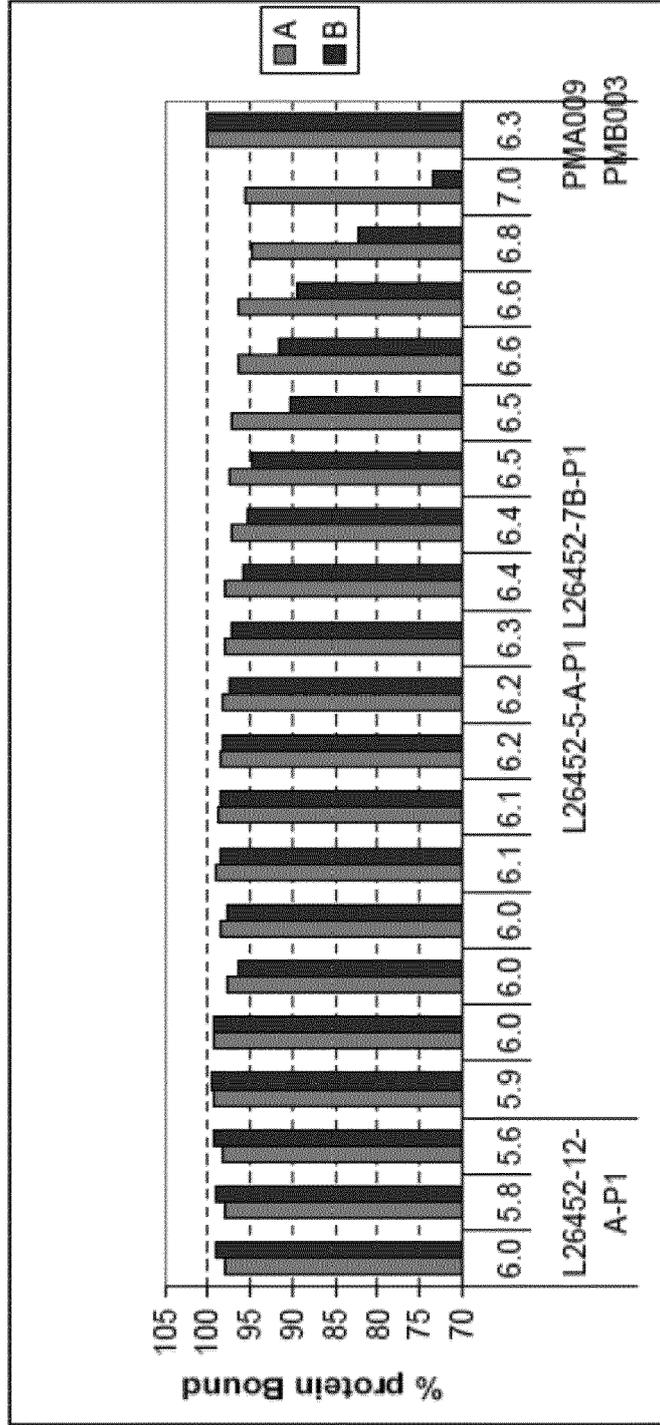
Protein Binding to Aluminum Phosphate at pH 6.5



MnB rLP2086 subfamily A + B was formulated at 400 µg/mL in 10mM Histidine buffer with 150 mM NaCl and POLYSORBATE 80 concentration at 0.02% at pH 6.5. The X-axis represents aluminum content in the formulation and Y-axis is the percent protein bound measured by IEX HPLC assay.

FIGURE 8

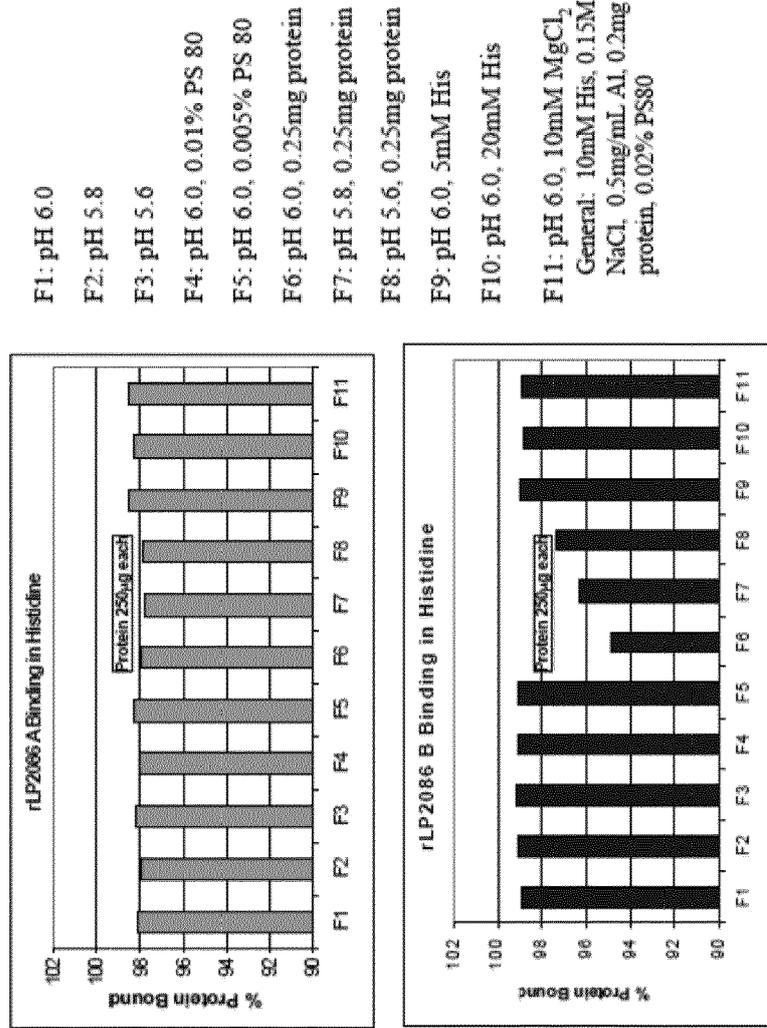
Binding of MnB rLP2086 Subfamily A and B as a Function of pH



MnB subfamily A + B formulations of subfamily A and B proteins at 200 µg/mL each were formulated with 0.5 mg/mL aluminum as aluminum phosphate in 10mM Histidine buffer with 150 mM NaCl varying pH. The multiple lots represent formulations prepared with different lots of DS as indicated in the figure.

FIGURE 9

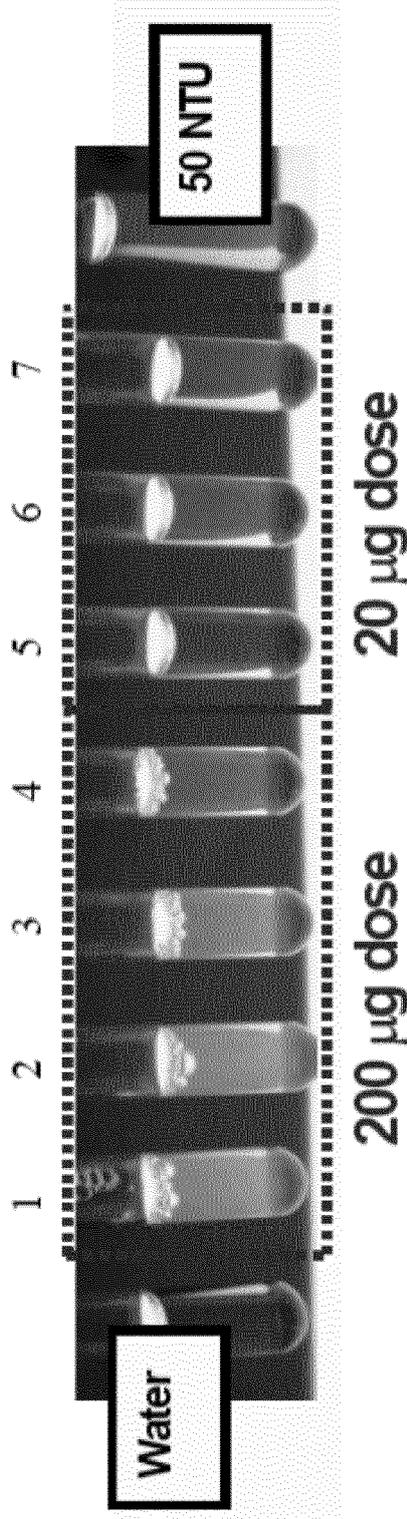
The Effect of pH, Buffer and Protein Concentration on Binding of rLP2086 Subfamily A and B



Eleven formulations of MnB subfamily A + B formulations with subfamily A and B proteins were formulated at 200µg/mL each with 0.5 mg/ mL aluminum as aluminum phosphate in 10mM Histidine buffer with 150 mM NaCl and 0.02% PS 80 varying pH (F1 through F3 with 200µg/mL protein each and 0.02% Polysorbate 80); varying Polysorbate concentrations of Polysorbate 80 from 0.01% to 0.005% (F4-F5); varying protein concentration to 250µg/mL for each subfamily protein (F6-F8), varying Histidine buffer concentration 5mM and 20mM (F9 and F10); additional 10mM MgCl₂ (F11).

FIGURE 10

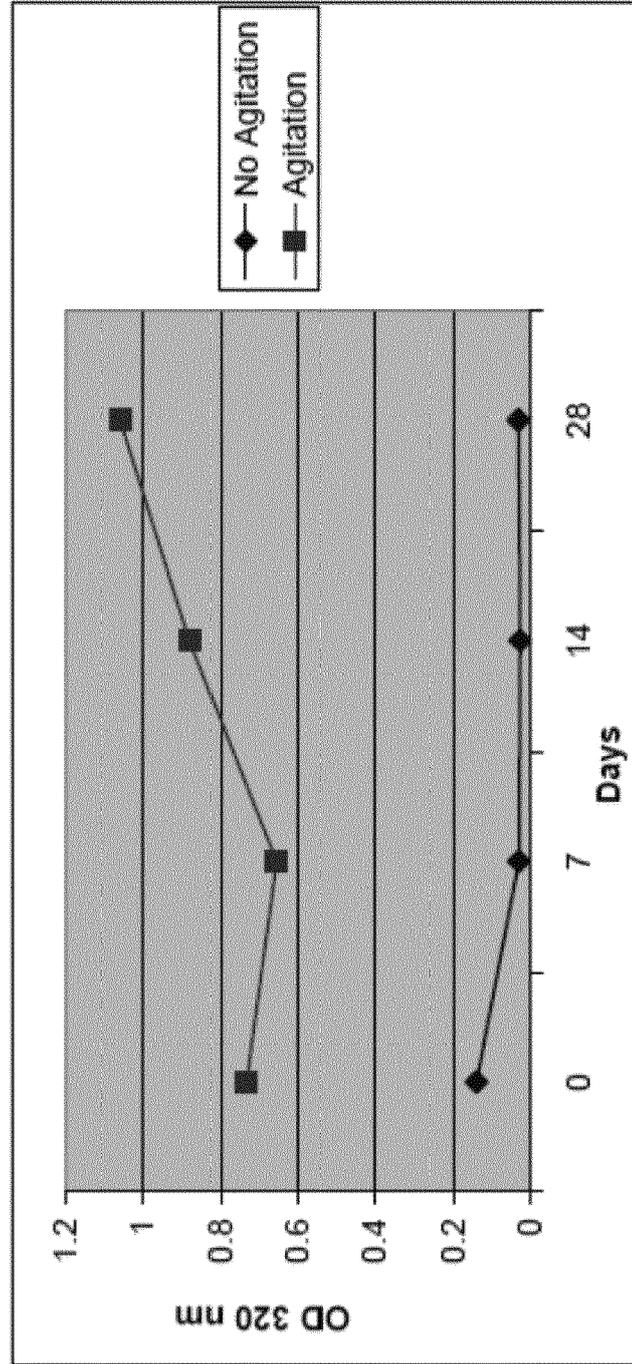
Visual Appearance of rLP2086 Formulations without Aluminum Phosphate



Photograph of two formulations of rLP2086 A + B proteins at 200 (left) and 20 (right) µg/mL without Aluminum Phosphate with various concentrations of Polysorbate-80, expressed as a percentage, indicated across the top (1 = 0%, 2 = 0.002%, 3 = 0.005%, 4 = 0.01%, 5 = 0%, 6 = 0.0005%, 7 = 0.001%). All tubes were incubated at 5°C for 14 days and agitated for 24 hours prior to the testing. Water and the 50 NTU (nephelometric turbidity units) turbidity standard were included for visual comparison on each end; water showing clarity and 50 NTU showing the appearance of a turbid solution. Note: similar results were also obtained after incubation for 1, 7 and 28 days.

FIGURE 11

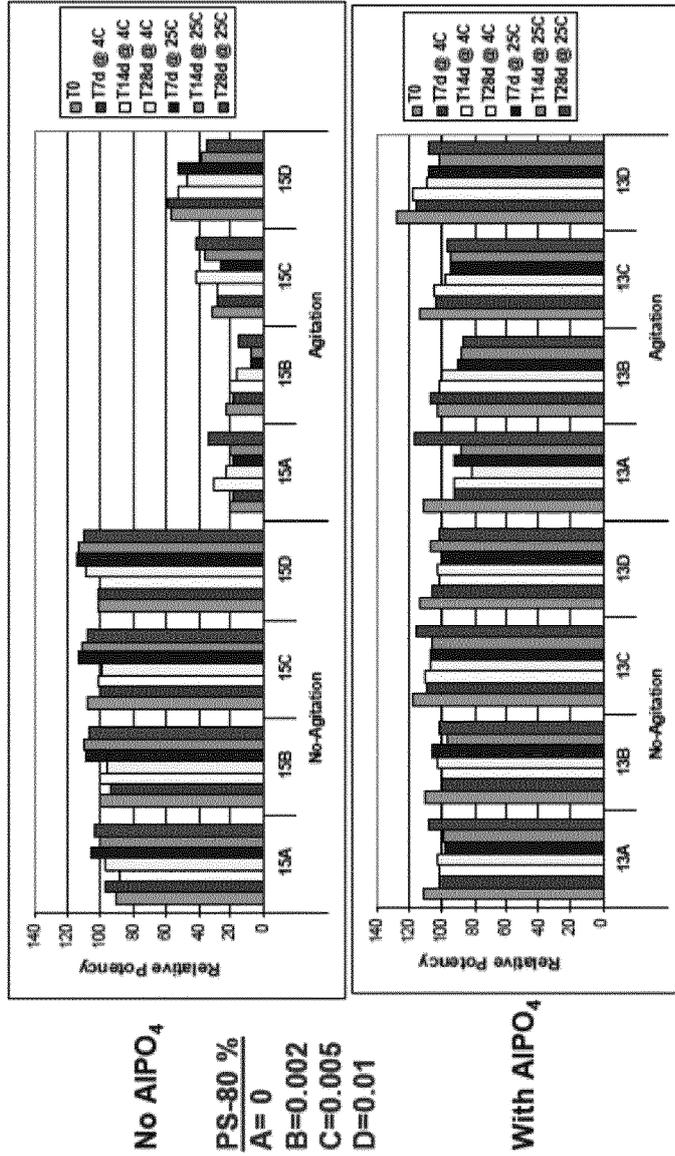
OD Measurements of Appearance Samples, 2-8°C



Optical Density measurements of samples from formulations without aluminum phosphate, at $\lambda = 320$ nm for rLP2086 200 $\mu\text{g}/\text{mL}$ formulation, designation 15C (0.005% Polysorbate 80), over one month at 5°C. Agitated samples, red squares, and no-agitation samples are indicated in blue diamonds.

FIGURE 12

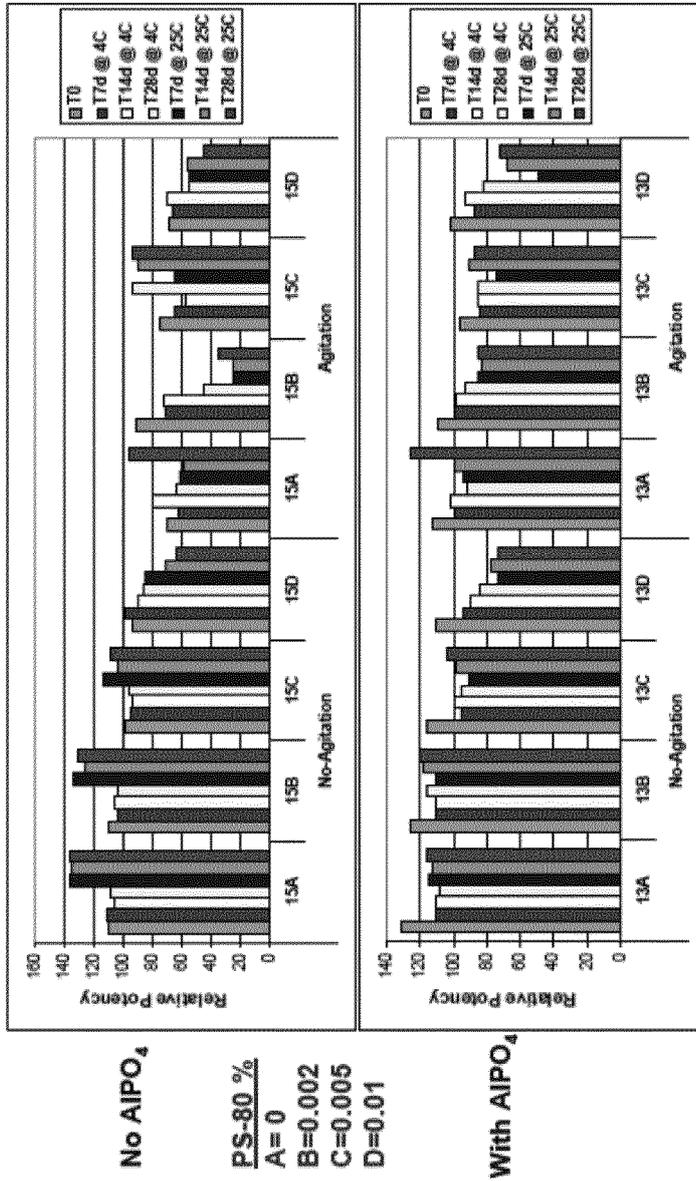
Potency Results for Subfamily A for Formulation with and without AlPO₄



MnB rLP2086 subfamily A + B formulated at 200 µg/mL of each subfamily. Formulations were made in histidine buffered saline with and without AlPO₄ and with varying levels of Polysorbate 80. Top graph represents formulation without AlPO₄ and bottom graph represents formulations with 0.5 mg/mL aluminum as aluminum phosphate. Four groups on the left half of the graph represent no-agitation and the other four groups on the right represent the agitation as labeled on the X-axis at the bottom of each graph. The Y-axis represents the percent relative potency. The Polysorbate 80 concentrations in the final formulations are 0%, 0.0005%, 0.001%, and 0.01% for lot 15A or 13A, 15B or 13B, 15C or 13C, and 15D or 13D, respectively. The corresponding Polysorbate 80:Protein molar ratios for these Polysorbate 80 concentrations are 0, 1.1, 2.7, and 5.3, respectively. The samples were stored at 2-8°C and 25°C. T0, T7d, T14d and T28d represent time zero, 7, 14 and 28 days. Each bar represents each time point data.

FIGURE 13

Potency Results for Subfamily B for Formulation with and without AlPO₄



MnB rLP2086 subfamily A + B formulated at 200 µg/mL of each subfamily. Formulations were made in histidine buffered saline with and without AlPO₄ and with varying levels of Polysorbate 80. Top graph represents formulation without AlPO₄ and bottom graph represents formulations with 0.5 mg/mL aluminum as aluminum phosphate. Four groups on the left half of the graph represent no-agitation and the other four groups on the right represent the agitation as labeled on the X-axis at the bottom of each graph. The Y-axis represents the percent relative potency. The Polysorbate 80 concentrations in the final formulations are 0%, 0.0005%, 0.001%, and 0.01% for lot 15A or 13A, 15B or 13B, 15C or 13C, and 15D or 13D, respectively. The corresponding Polysorbate 80:Protein molar ratios for these Polysorbate 80 concentrations are 0, 1.1, 2.7, and 5.3, respectively. The samples were stored at 2-8°C and 25°C. T0, T7d, T14d and T28d represent time zero, 7, 14 and 28 days. Each bar represents each time point data.

FIGURE 14

Polysorbate 80 Results in rLP2086 Placebo with 0.5 mg/mL Aluminum

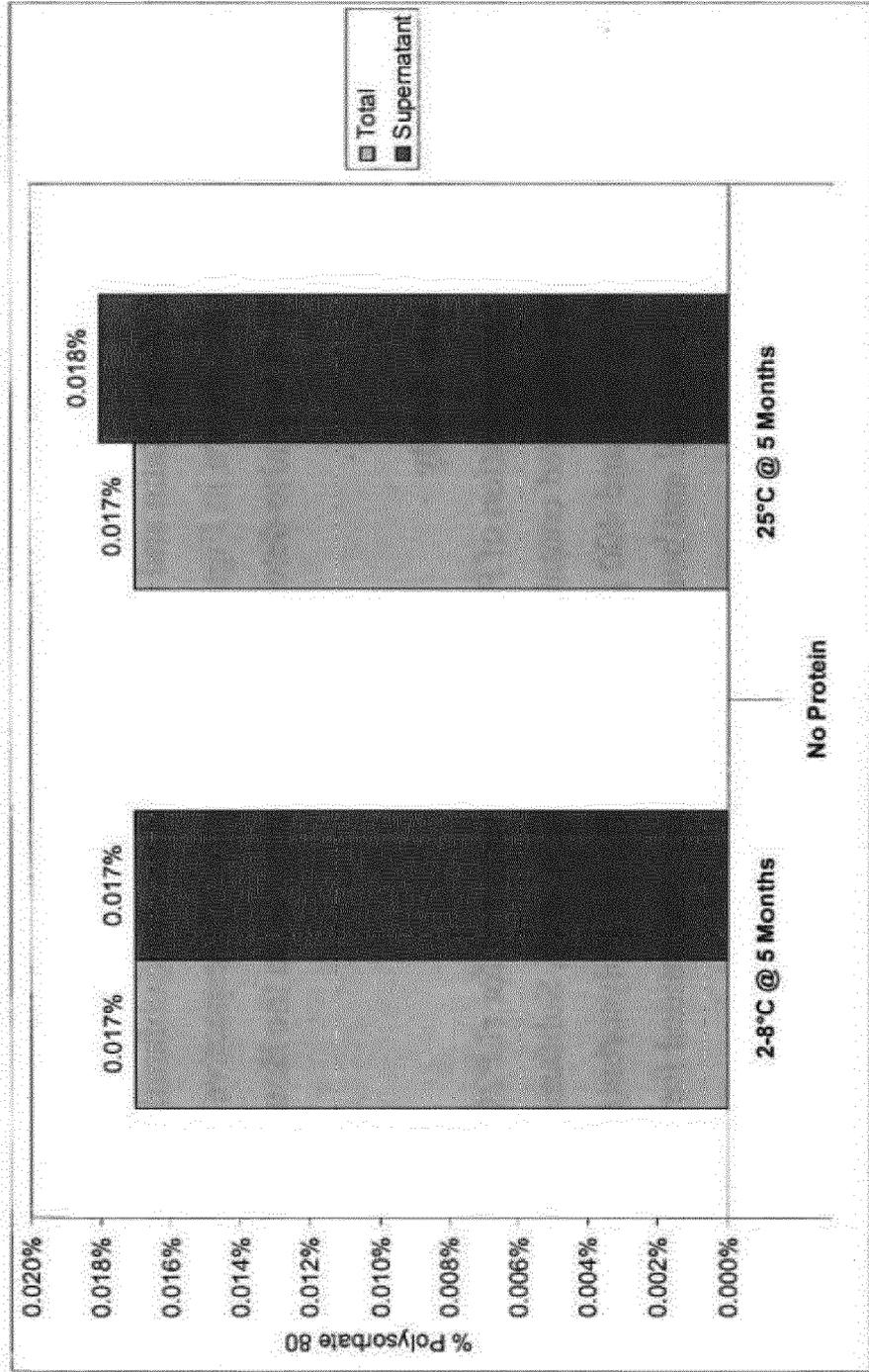


FIGURE 15

Polysorbate 80 Results for Subfamily A

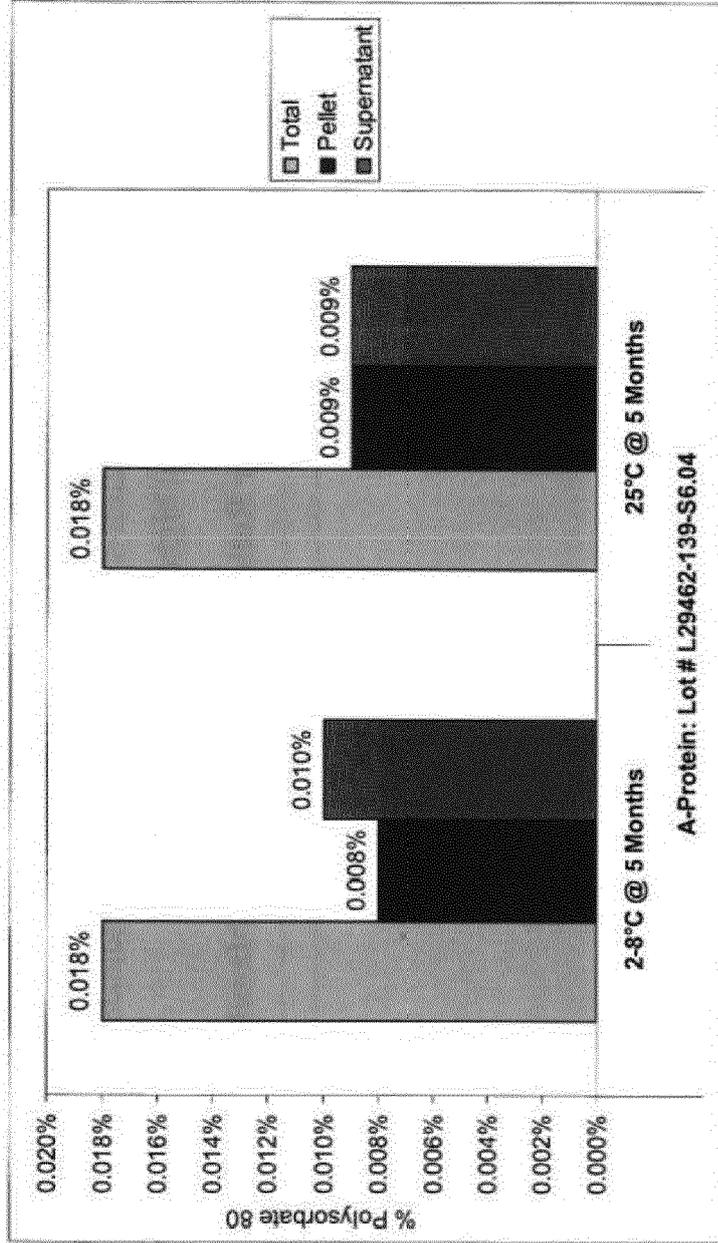


FIGURE 16

Polysorbate 80 Results for Subfamily B

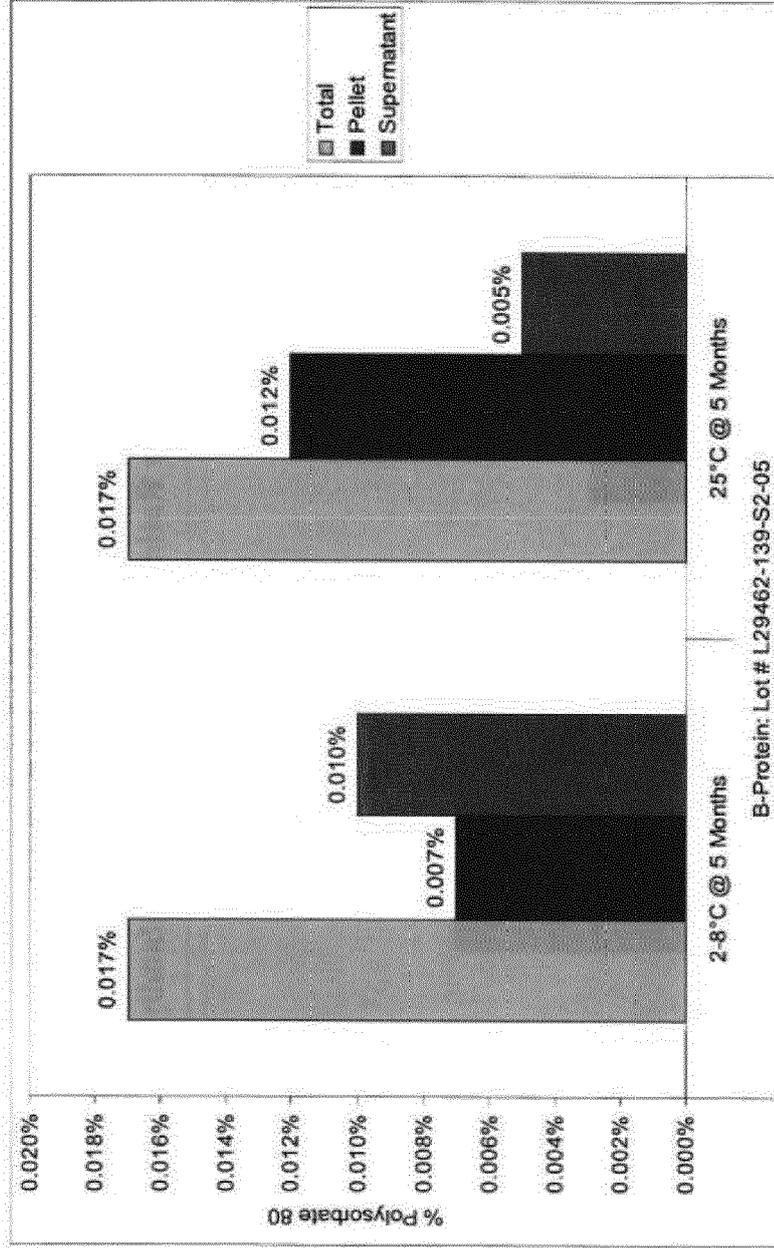


FIGURE 17

Correlation of Potency and Bound Molar Ratio for Subfamily B

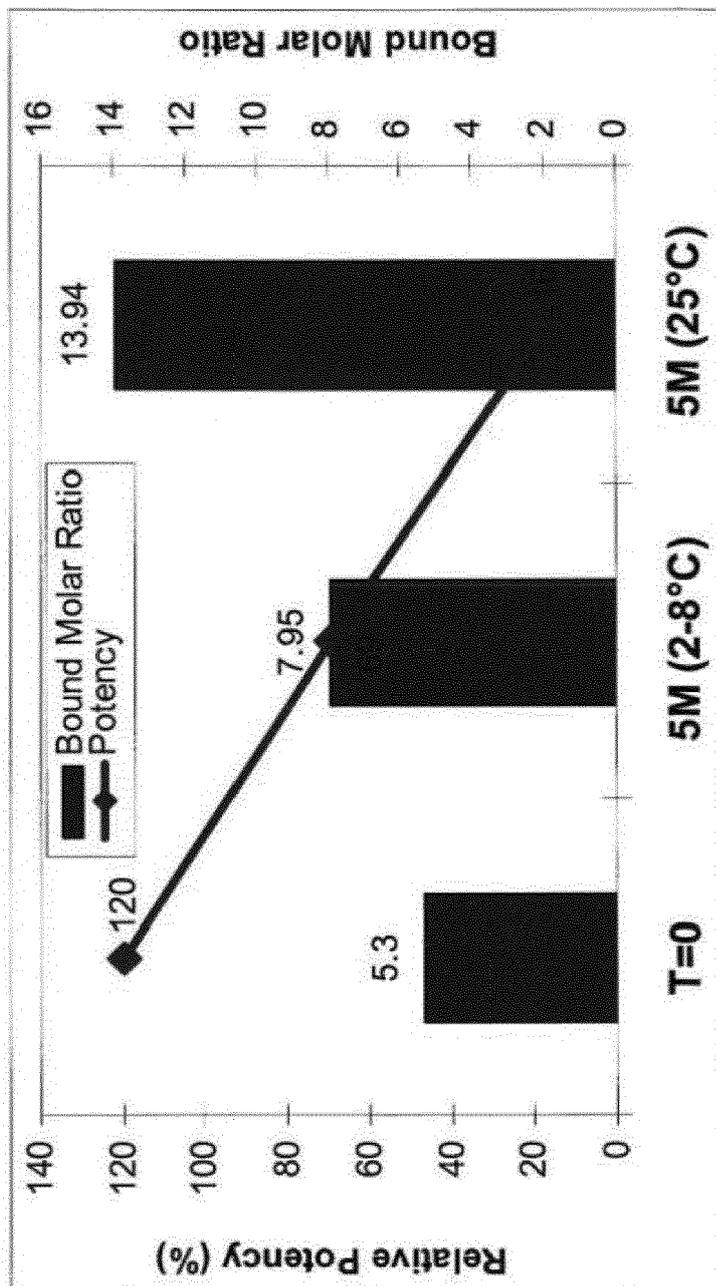


FIGURE 18

Molar Ratio Results for Subfamily A

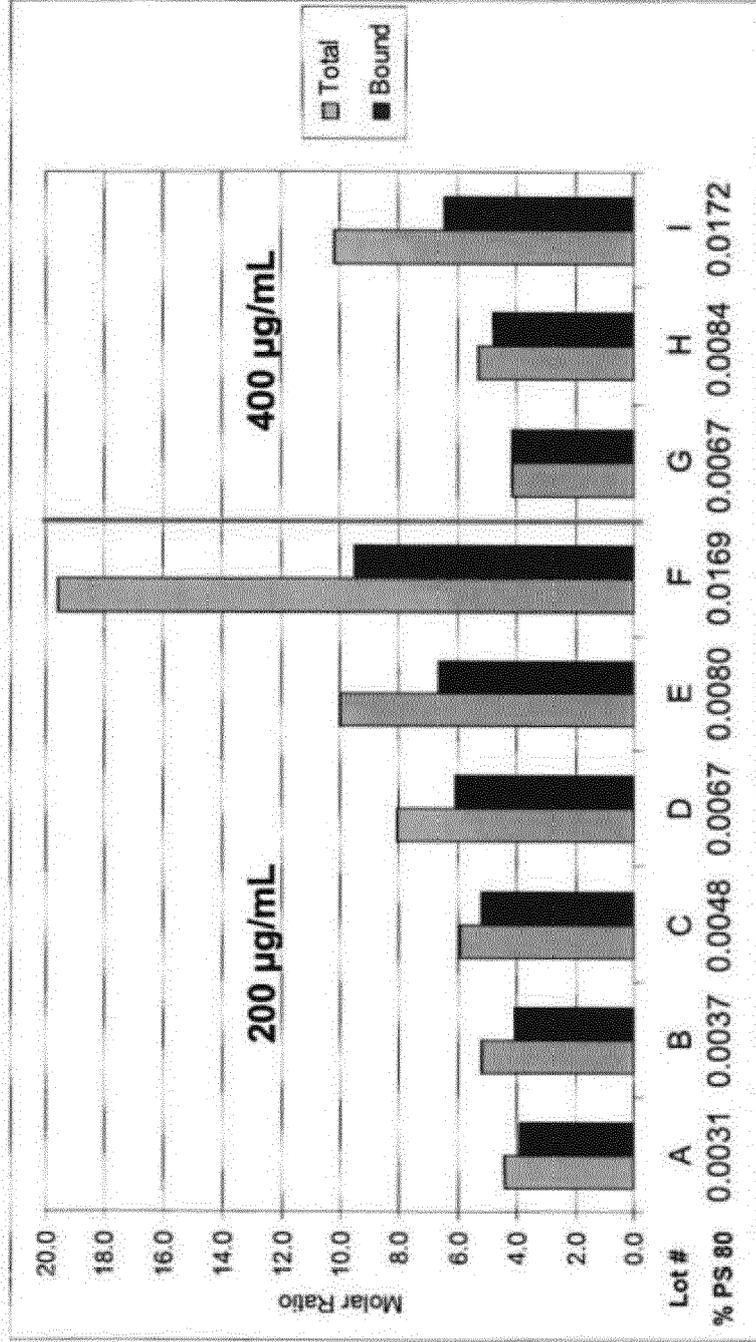


FIGURE 19

Molar Ratio Results for Subfamily B

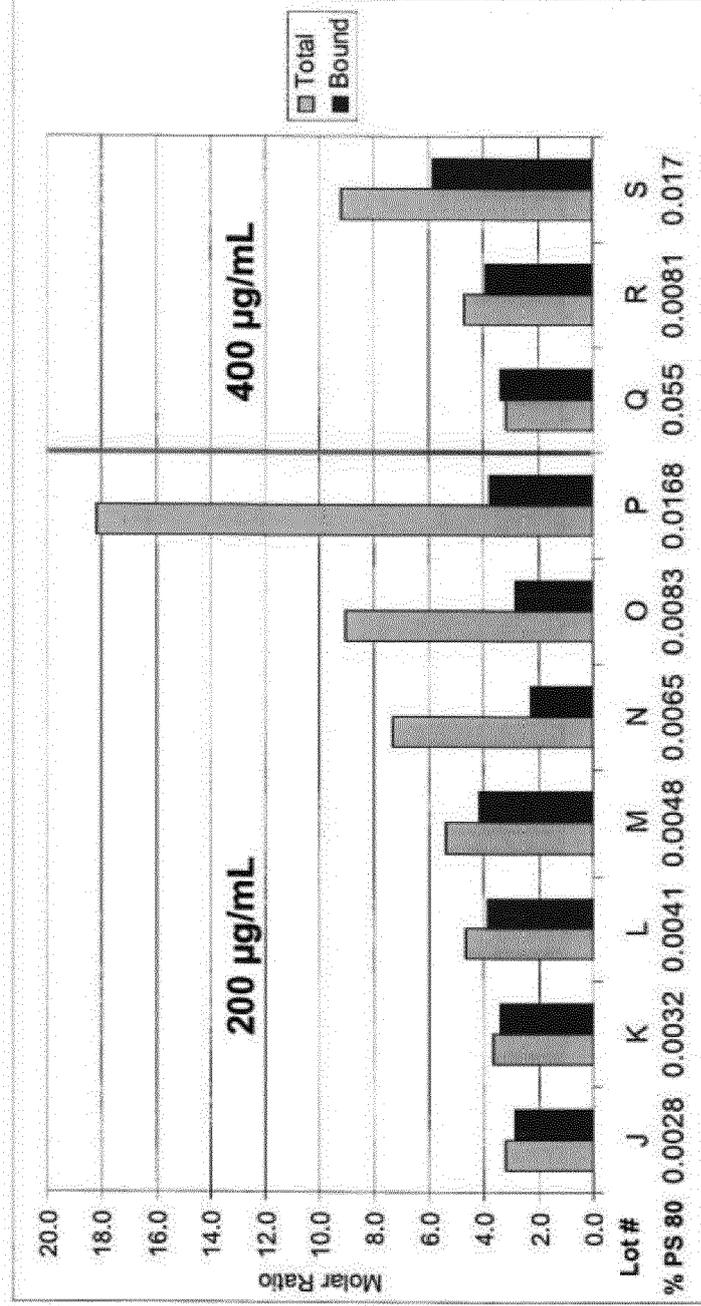


FIGURE 20

Molar Ratio Results for rLP2086 Formulations @ 400 µg/mL

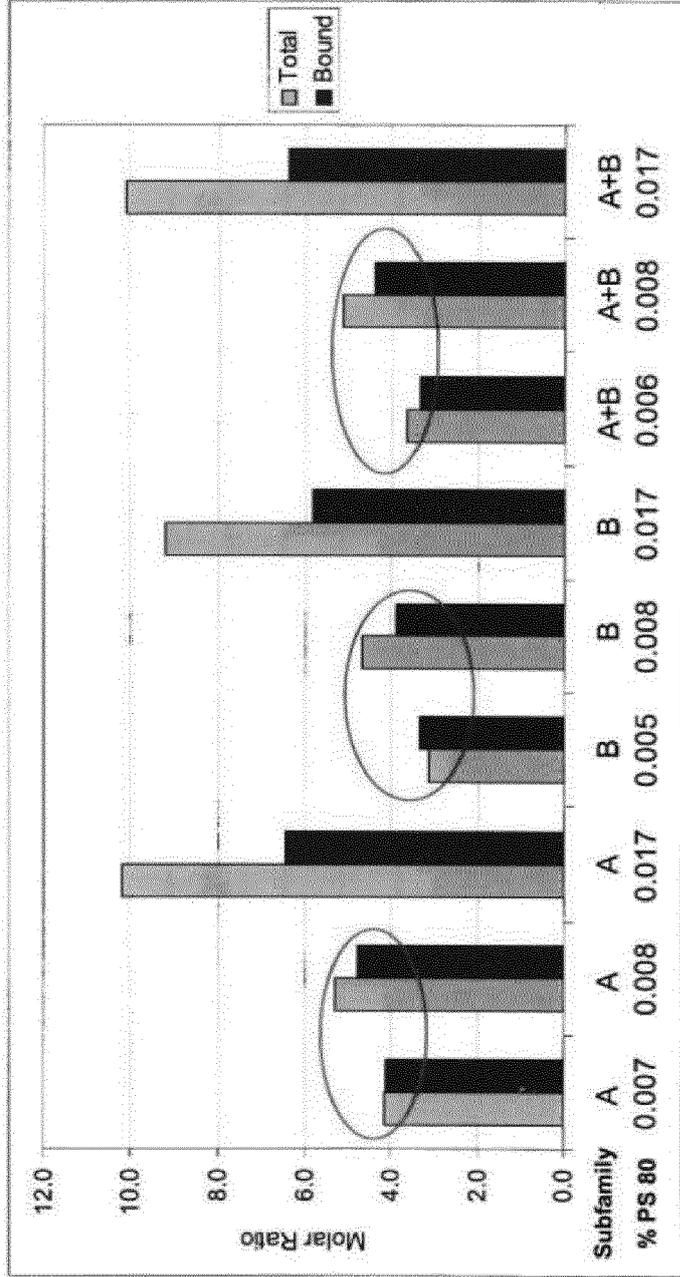


FIGURE 21

Polysorbate 80 Results for rLP2086 Drug Product at Different Time Points

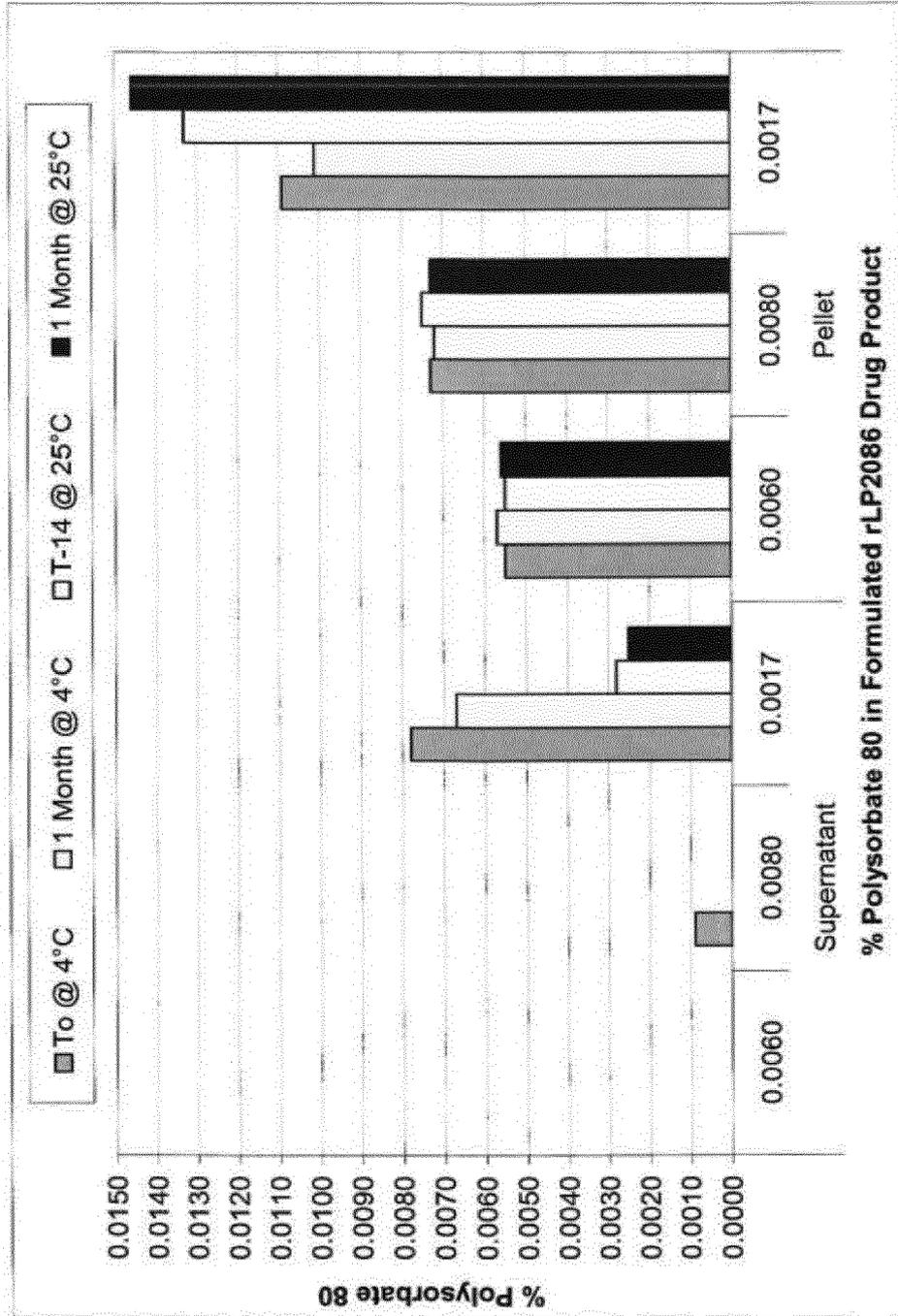


FIGURE 22

Bound Molar Ratio Results for rLP2086 Drug Product at Different Time Point

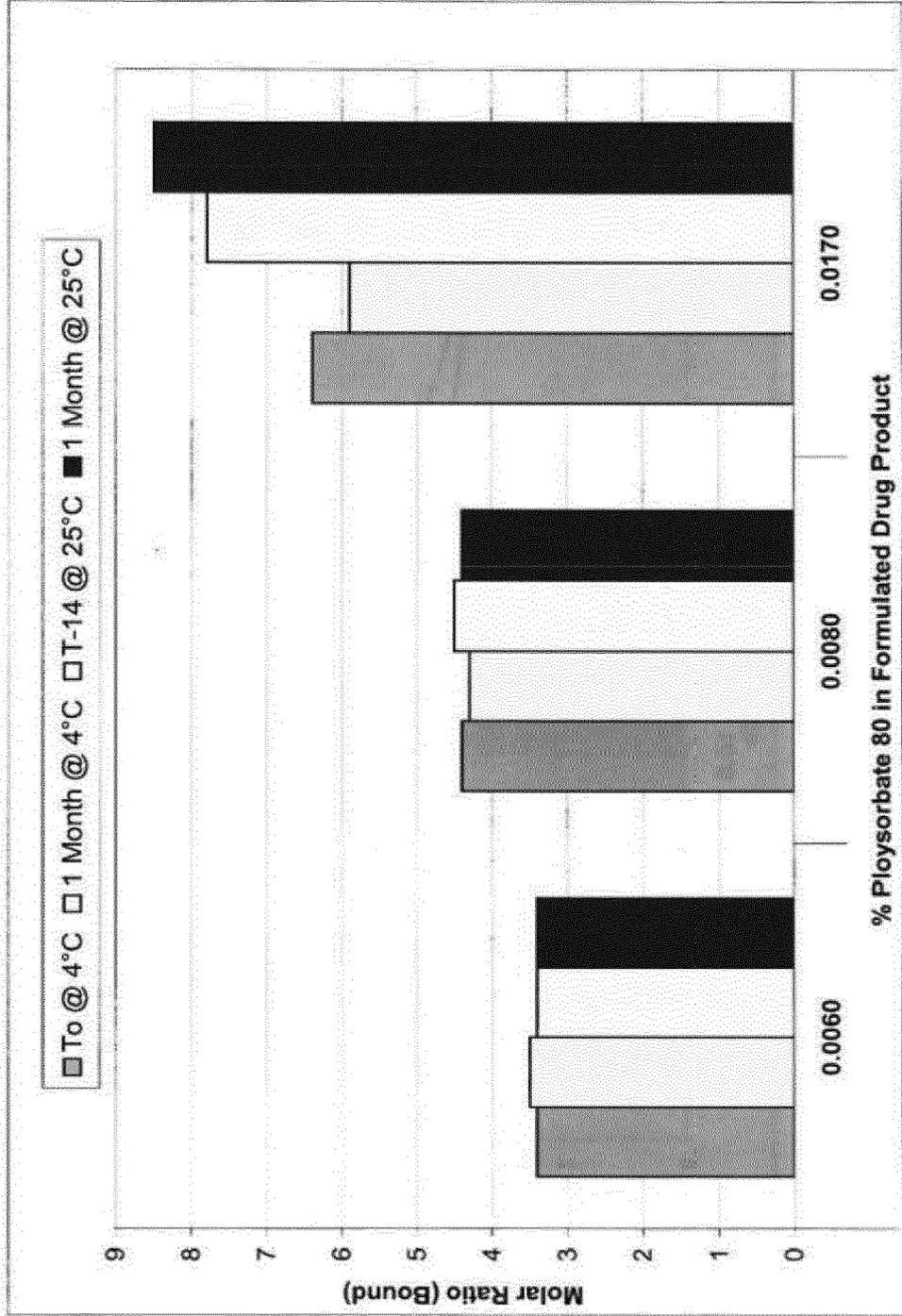


FIGURE 23

Potency and Bound Molar Ratio Results for Subfamily A

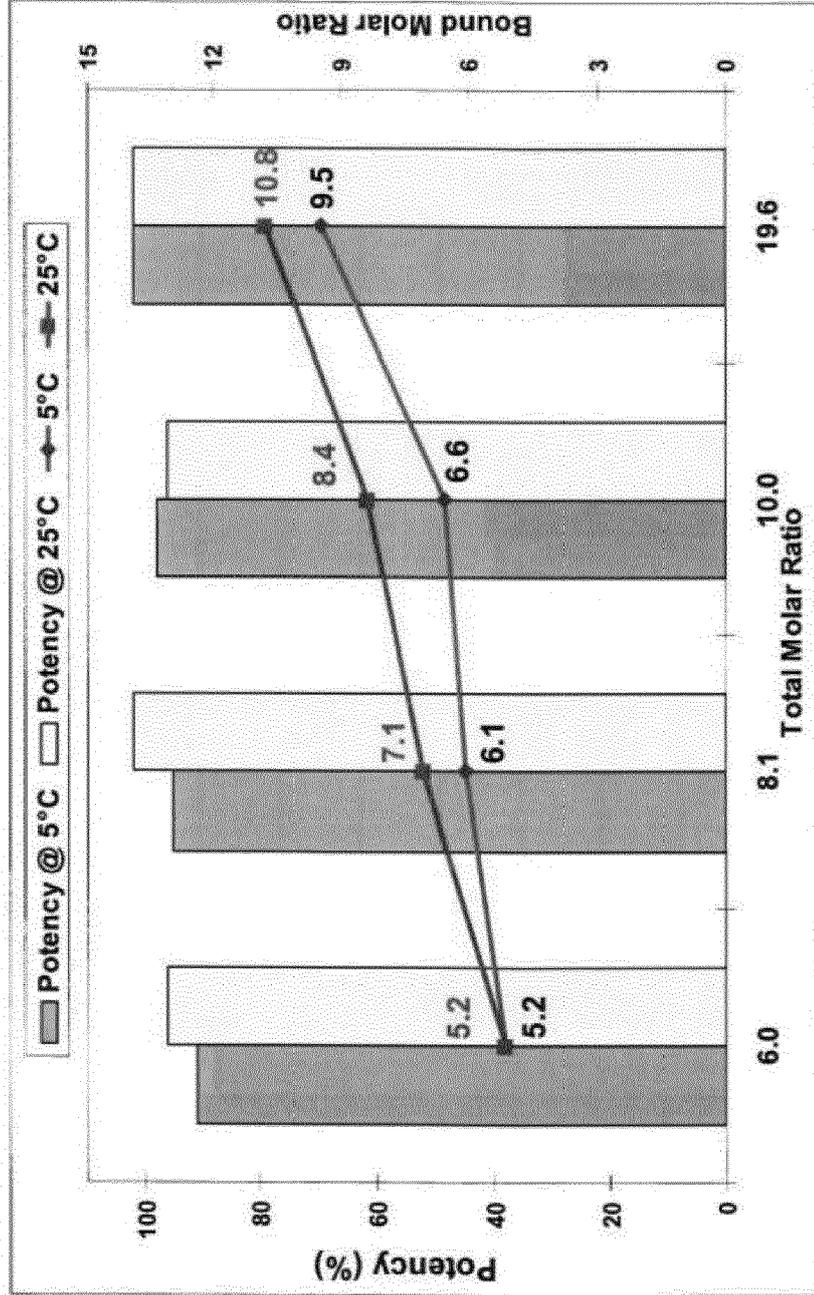


FIGURE 24

Potency and Bound Molar Ratio Results for Subfamily B

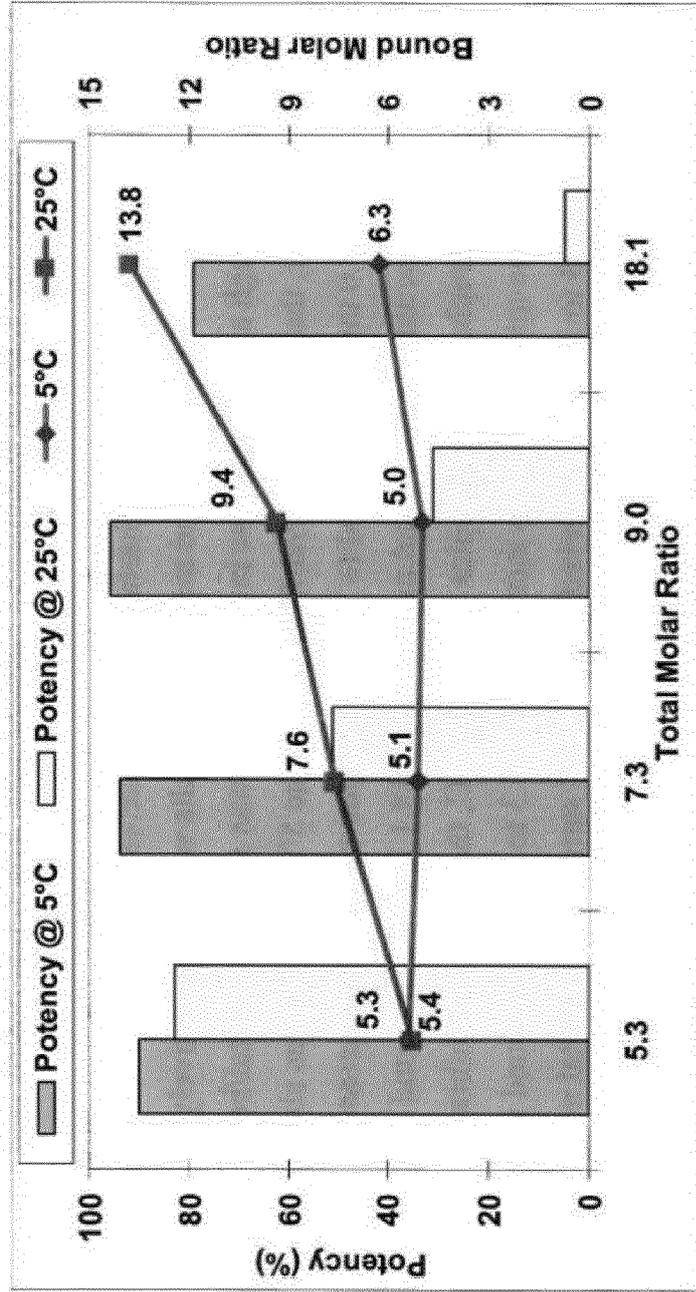
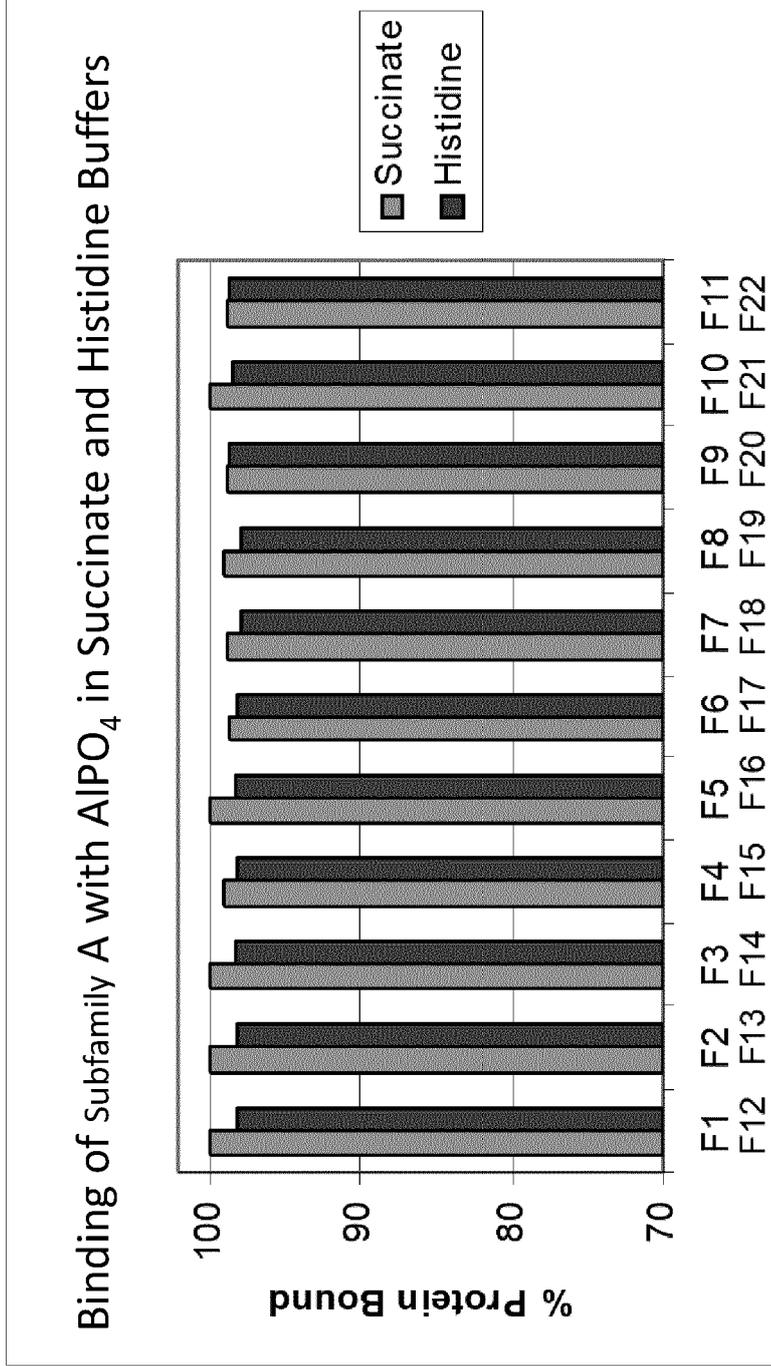


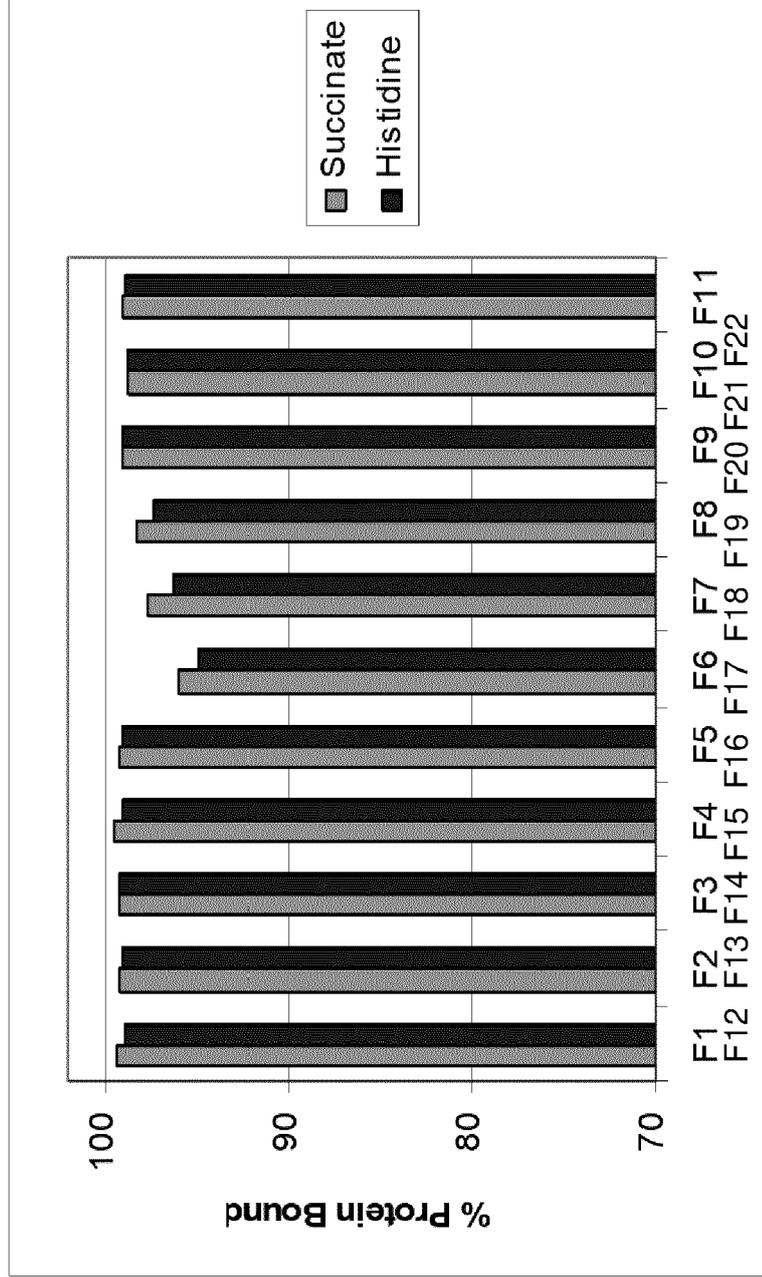
FIGURE 25



Binding of Subfamily A protein from formulations of bivalent MnB formulations formulated at 200mg/mL of each protein with 0.5 mg/ mL aluminum as aluminum phosphate in 10mM Histidine buffer with 150 mM NaCl and 0.02% PS 80 varying pH (F1 through F3 with 200mg/mL protein each and 0.02% PS 80); varying polysorbate concentrations of PS-80 from 0.01, 0.05 (F4-F5); varying protein concentration to 250mg/mL (F6-F8), varying Histidine buffer concentration 5mM and 20mM (F9 and F10); additional 10mM MgCl2 (F11).

FIGURE 26

Binding of Subfamily B with $AlPO_4$ in Succinate and Histidine Buffers



Binding of Subfamily B protein from formulations of bivalent MnB formulations formulated at 200mg/mL of each protein with 0.5 mg/ mL aluminum as aluminum phosphate in 10mM Histidine buffer with 150 mM NaCl and 0.02% PS 80 varying pH (F1 through F3 with 200mg/mL protein each and 0.02% PS 80); varying polysorbate concentrations of PS-80 from 0.01, 0.05 (F4-F5); varying protein concentration to 250mg/mL (F6-F8), varying Histidine buffer concentration 5mM and 20mM (F9 and F10); additional 10mM MgCl2 (F11).

FIGURE 27

Comparison of Binding in Succinate, Histidine, and Phosphate buffer

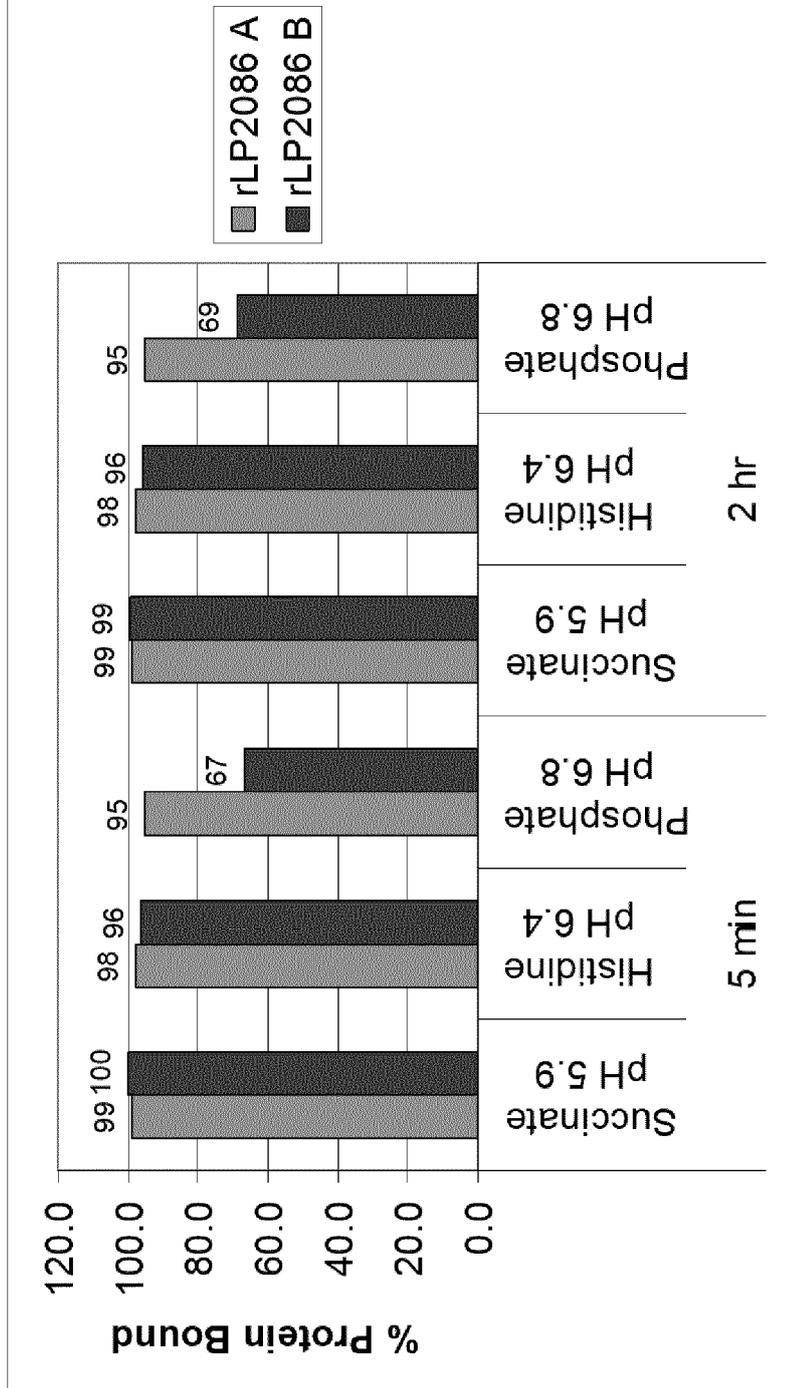


FIGURE 28

pH-Dependent Binding of Subfamily A with $AlPO_4$

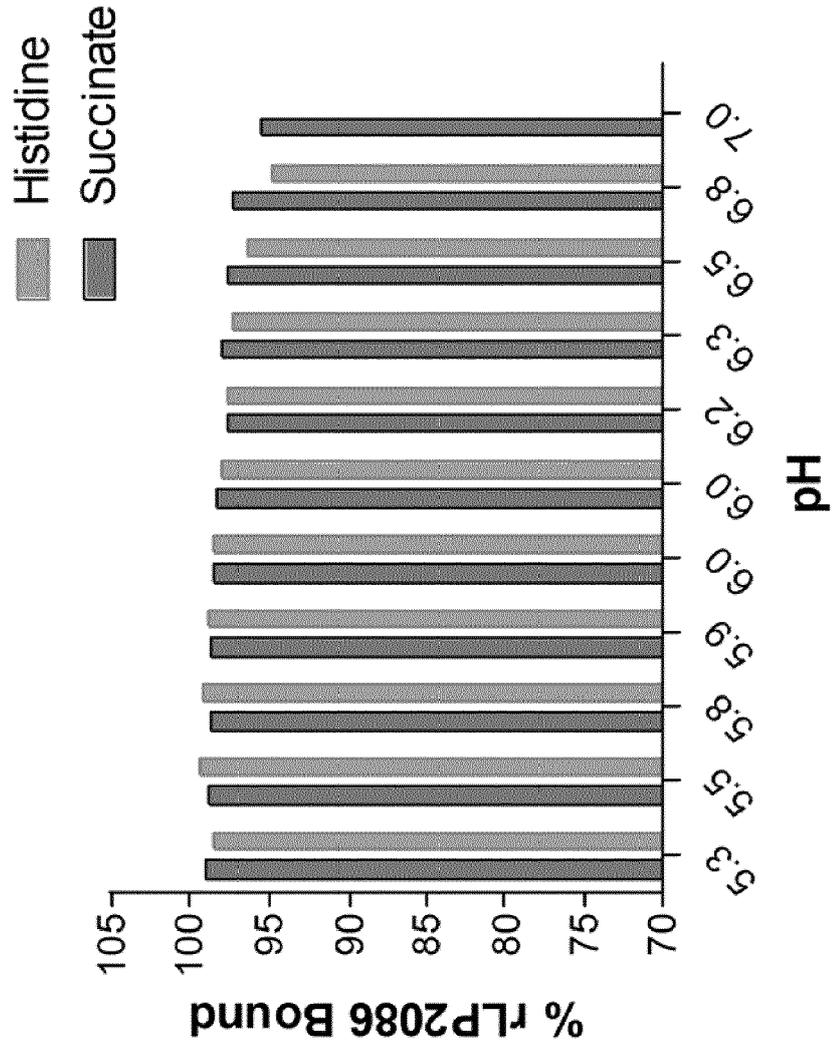
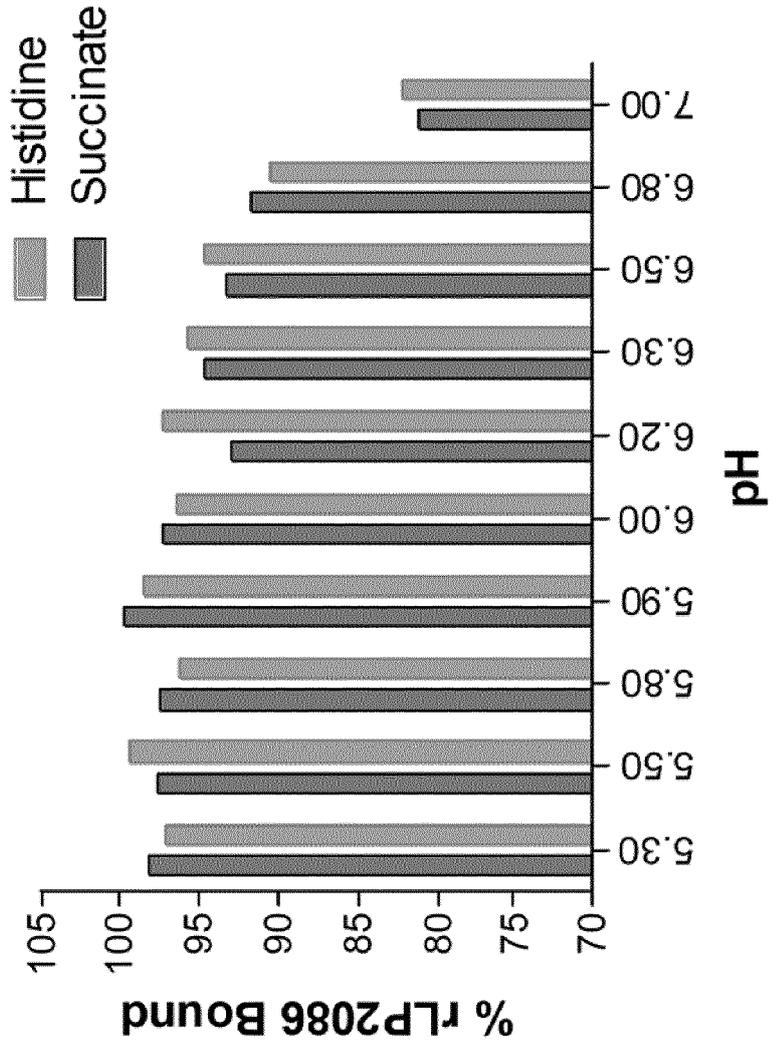


FIGURE 29

pH-Dependent Binding of Subfamily B with $AlPO_4$



REFERENCES CITED IN THE DESCRIPTION

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Patent documents cited in the description

- US 20060257413 A [0028] [0041]
- US 20090202593 A [0028] [0041]
- WO 0018434 A [0057]

Non-patent literature cited in the description

- **MCNEIL et al.** *Vaccine*, 2009, vol. 27, 3417-3421 [0035]
- **WOLFF et al.** *Biotechniques*, 1991, vol. 11 (4), 474-85 [0053]
- **SEDEGAH et al.** *PNAS*, 1994, vol. 91, 9866-9870 [0053]
- **EGAN et al.** *Vaccine*, 2009, vol. 27 (24), 3175-3180 [0080]