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(57) **Abrégé/Abstract:**

The present invention provides liquid formulations of SYNAGIS® or an antigen-binding fragment thereof that immunospecifically bind to a respiratory syncytial virus (RSV) antigen, which formulations exhibit stability, low to undetectable levels of aggregation, and very little to no loss of the biological activities of SYNAGIS® or an antigen-binding fragment thereof, even during long periods of storage. In particular, the present invention provides liquid formulations of SYNAGIS® or an antigen-binding fragment thereof which immunospecifically binds to a RSV antigen, which formulations are substantially free of surfactant, inorganic salts, and/or other common excipients. Furthermore, the invention provides method of preventing, treating or ameliorating symptoms associated with RSV infection utilizing liquid formulations of the present invention.



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STABILIZED LIQUID ANTI-RSV ANTIBODY FORMULATIONS

1. INTRODUCTION

The present invention relates to liquid formulations of SYNAGIS® or an antigen-binding fragment thereof, which formulations exhibit stability, low to undetectable levels of antibody fragmentation, low to undetectable levels of aggregation, and very little to no loss of the biological activity (*e.g.*, therapeutic efficacy) SYNAGIS® or an antigen-binding fragment thereof, even during or after long periods of storage. In particular, the present invention relates to liquid formulations of SYNAGIS® or an antigen-binding fragment thereof, which formulations are substantially free of surfactant and/or inorganic salts. The present invention also relates to methods of preventing, treating or ameliorating symptoms associated with a respiratory syncytial virus (RSV) infection utilizing liquid formulations of SYNAGIS® or an antigen binding fragment thereof.

2. BACKGROUND OF THE INVENTION

Respiratory syncytial virus (RSV) is the leading cause of serious lower respiratory tract disease in infants and children (Feigen *et al.*, eds., 1987, *In: Textbook of Pediatric Infectious Diseases*, WB Saunders, Philadelphia at pages 1653-1675; New Vaccine Development, Establishing Priorities, Vol. 1, 1985, National Academy Press, Washington DC at pages 397-409; and Ruuskanen *et al.*, 1993, *Curr. Probl. Pediatr.* 23:50-79). The yearly epidemic nature of RSV infection is evident worldwide, but the incidence and severity of RSV disease in a given season vary by region (Hall, C.B., 1993, *Contemp. Pediatr.* 10:92-110). In temperate regions of the northern hemisphere, it usually begins in late fall and ends in late spring (Hall, C.B., 1995, *In: Mandell G.L., Bernnett J.E., Dolin R., eds., 1995, Principles and Practice of Infections Diseases. 4th ed., Churchill Livingstone, New York at pages 1501-1519*). It is estimated that RSV illness results in 90,000 hospitalizations and causes 4,500 deaths annually in the United States. Primary RSV infection occurs most often in children from 6 weeks to 2 years of age and uncommonly in

the first 4 weeks of life during nosocomial epidemics (Hall *et al.*, 1979, New Engl. J. Med. 300:393-396). RSV is estimated to cause as much as 75% of all childhood bronchiolitis and up to 40% of all pediatric pneumonias (Cunningham, C.K. *et al.*, 1991, *Pediatrics* 88:527-532). Children at increased risk from RSV infection include preterm infants (Hall *et al.*, 1979, New Engl. J. Med. 300:393-396) and children with bronchopulmonary dysplasia (Groothuis *et al.*, 1988, *Pediatrics* 82:199-203), congenital heart disease (MacDonald *et al.*, New Engl. J. Med. 307:397-400), congenital or acquired immunodeficiency (Ogra *et al.*, 1988, *Pediatr. Infect. Dis. J.* 7:246-249; and Pohl *et al.*, 1992, *J. Infect. Dis.* 165:166-169), and cystic fibrosis (Abman *et al.*, 1988, *J. Pediatr.* 113:826-830). The fatality rate in infants with heart or lung disease who are hospitalized with RSV infection is 3%-4% (Navas *et al.*, 1992, *J. Pediatr.* 121:348-354).

RSV infects adults as well as infants and children. In healthy adults, RSV causes predominantly upper respiratory tract disease. It has recently become evident that some adults, especially the elderly, have symptomatic RSV infections more frequently than had been previously reported (Evans, A.S., eds., 1989, *Viral Infections of Humans. Epidemiology and Control*, 3rd ed., Plenum Medical Book, New York at pages 525-544). Several epidemics also have been reported among nursing home patients and institutionalized young adults (Falsey, A.R., 1991, *Infect. Control Hosp. Epidemiol.* 12:602-608; and Garvie *et al.*, 1980, *Br. Med. J.* 281:1253-1254). Finally, RSV may cause serious disease in immunosuppressed persons, particularly bone marrow transplant patients (Hertz *et al.*, 1989, *Medicine* 68:269-281).

Treatment options for established RSV disease are limited. Severe RSV disease of the lower respiratory tract often requires considerable supportive care, including administration of humidified oxygen and respiratory assistance (Fields *et al.*, eds, 1990, *Fields Virology*, 2nd ed., Vol. 1, Raven Press, New York at pages 1045-1072). The only drug approved for treatment of infection is the antiviral agent ribavirin (American Academy of Pediatrics Committee on Infectious Diseases, 1993, *Pediatrics* 92:501-504). It has been shown to be effective in the treatment of RSV pneumonia and bronchiolitis, modifying the course of severe RSV disease in immunocompetent children (Smith *et al.*, 1991, New Engl. J. Med. 325:24-29). However, ribavirin has a number of limitations including high cost, need for prolonged aerosol administration and potential risk to pregnant women as well as to exposed health care personnel. The American Academy of Pediatrics Committee on Infectious Diseases revised their recommendation for use of ribavirin. The current recommendation is that the decision to use ribavirin should be based on the particular

clinical circumstances and physician's experience (American Academy of Pediatrics. Summaries of Infectious Diseases. *In*: Pickering L.K., ed., 2000 *Red Book: Report of the Committee on Infectious Diseases*. 25th ed., Elk Grove Village, IL, American Academy of Pediatrics, 2000, pp. 483-487).

5 While a vaccine might prevent RSV infection, no vaccine is yet licensed for this indication. A major obstacle to vaccine development is safety. A formalin-inactivated vaccine, though immunogenic, unexpectedly caused a higher and more severe incidence of lower respiratory tract disease due to RSV in immunized infants than in infants immunized with a similarly prepared trivalent parainfluenza vaccine (Kim *et al.*, 1969, *Am. J. Epidemiol.* 89:422-434; and Kapikian *et al.*, 1969, *Am. J. Epidemiol.* 89:405-421). Several
10 candidate RSV vaccines have been abandoned and others are under development (Murphy *et al.*, 1994, *Virus Res.* 32:13-36), but even if safety issues are resolved, vaccine efficacy must also be improved. A number of problems remain to be solved. Immunization would be required in the immediate neonatal period since the peak incidence of lower respiratory
15 tract disease occurs at 2-5 months of age. The immaturity of the neonatal immune response together with high titers of maternally acquired RSV antibody may be expected to reduce vaccine immunogenicity in the neonatal period (Murphy *et al.*, 1988, *J. Virol.* 62:3907-3910; and Murphy *et al.*, 1991, *Vaccine* 9:185-189). Finally, primary RSV infection and disease do not protect well against subsequent RSV disease (Henderson *et al.*, 1979, *New Engl. J. Med.* 300:530-534).
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 Currently, the only approved approach to prophylaxis of RSV disease is passive immunization. Initial evidence suggesting a protective role for IgG was obtained from observations involving maternal antibody in ferrets (Prince, G.A., Ph.D. diss., University of California, Los Angeles, 1975) and humans (Lambrecht *et al.*, 1976, *J. Infect. Dis.* 134:211-217; and Glezen *et al.*, 1981, *J. Pediatr.* 98:708-715). Hemming *et al.* (Morell *et al.*, eds., 1986, *Clinical Use of Intravenous Immunoglobulins*, Academic Press, London at pages 285-294) recognized the possible utility of RSV antibody in the treatment or prevention of RSV infection during studies involving the pharmacokinetics of an intravenous immune globulin (IVIG) in newborns suspected of having neonatal sepsis. They noted that one infant, whose
25 respiratory secretions yielded RSV, recovered rapidly after IVIG infusion. Subsequent analysis of the IVIG lot revealed an unusually high titer of RSV neutralizing antibody. This same group of investigators then examined the ability of hyperimmune serum or immune globulin, enriched for RSV neutralizing antibody, to protect cotton rats and primates against RSV infection (Prince *et al.*, 1985, *Virus Res.* 3:193-206; Prince *et al.*, 1990, *J. Virol.*
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64:3091-3092; Hemming *et al.*, 1985, J. Infect. Dis. 152:1083-1087; Prince *et al.*, 1983, Infect. Immun. 42:81-87; and Prince *et al.*, 1985, J. Virol. 55:517-520). Results of these studies suggested that RSV neutralizing antibody given prophylactically inhibited respiratory tract replication of RSV in cotton rats. When given therapeutically, RSV antibody reduced pulmonary viral replication both in cotton rats and in a nonhuman primate model. Furthermore, passive infusion of immune serum or immune globulin did not produce enhanced pulmonary pathology in cotton rats subsequently challenged with RSV.

A humanized antibody directed to an epitope in the A antigenic site of the F protein of RSV, SYNAGIS[®], comprising variable heavy (VH) complementarity determining regions (CDRs) having the amino acid sequences of SEQ ID NO:1-3 and variable light (VL) CDRs having the amino acid sequences of SEQ ID NO:4-6, is approved for intramuscular administration to pediatric patients for prevention of serious lower respiratory tract disease caused by RSV at recommended monthly doses of 15 mg/kg of body weight throughout the RSV season (November through April in the northern hemisphere). SYNAGIS[®] is a composite of human (95%) and murine (5%) antibody sequences. See, Johnson *et al.*, 1997, J. Infect. Diseases 176:1215-1224 and U.S. Patent No. 5,824,307.

The human heavy chain sequence was derived from the constant domains of human IgG₁ and the variable framework regions of the VH genes of Cor (Press *et al.*, 1970, Biochem. J. 117:641-660) and Cess (Takashi *et al.*, 1984, Proc. Natl. Acad. Sci. USA 81:194-198). The human light chain sequence was derived from the constant domain of C6 and the variable framework regions of the VL gene K104 with J6-4 (Bentley *et al.*, 1980, Nature 288:5194-5198). The murine sequences were derived from a murine monoclonal antibody, Mab 1129 (Beeler *et al.*, 1989, J. Virology 63:2941-2950), in a process which involved the grafting of the murine complementarity determining regions into the human antibody frameworks.

SYNAGIS[®] has high specific activity against RSV *in vitro* (approximately 50-100 times that of RespiGam[®]) and is known to neutralize a broad range of RSV isolates. Since it is not derived from human plasma, prophylactic treatment with SYNAGIS[®] does not carry potential risk of transmission of blood borne pathogens.

SYNAGIS[®] was initially formulated as a liquid for IV use, at a concentration of 10 mg/ml SYNAGIS[®] in phosphate buffered saline. A lyophilized formulation of SYNAGIS[®], which allows a higher concentration (100 mg/ml after reconstitution, in 50 mM histidine and 3.2 mM glycine buffer with 6% (w/v) mannitol at pH 6.0) of the antibody than this initial liquid formulation, was produced later to allow intramuscular use. The lyophilized

formulation of SYNAGIS[®] is prepared by lyophilizing SYNAGIS[®] at 54 mg/ml in an aqueous solution containing 25 mM histidine, 1.6 mM glycine, and 3% (w/v) mannitol at pH 6.0. The initial liquid formulation in PBS and the lyophilized formulation of SYNAGIS[®] have been tested in phase I clinical studies in healthy adults. The lyophilized
5 formulation was tested in phase I through phase IV studies in pediatric patients. SYNAGIS[®], at doses of 15 mg/kg to 30 mg/kg for adults is found to be well tolerated, and 15 mg/kg for children is found to be safe and efficacious for RSV prophylaxis. The lyophilized formulation was approved in 1998 by the FDA for use in the prevention of serious lower respiratory tract disease caused by RSV in children at high risk of RSV
10 disease.

However, the lyophilized formulation has a number of limitations, including a prolonged process for lyophilization and resulting high cost for manufacturing. In addition, the lyophilized formulation has to be reconstituted aseptically and accurately by healthcare practitioners prior to administering to patients. The reconstitution step itself requires certain
15 specific procedures: (1) a sterile diluent (*i.e.*, water or 5% dextrose in water for intravenous administration and water for intramuscular administration) is added to the vial containing lyophilized SYNAGIS[®], slowly and aseptically, and the vial must be swirled very gently for 30 seconds to avoid foaming; (2) the reconstituted SYNAGIS[®] needs to stand at room temperature for a minimum of 20 minutes until the solution clarifies; and (3) the
20 reconstituted preparation must be administered within six (6) hours after the reconstitution. Such reconstitution procedure is cumbersome and the time limitation after the reconstitution can cause a great inconvenience in administering the formulation to patients, leading to significant waste, if not reconstituted properly or if the reconstituted dose is not used within six (6) hours and must be discarded.

25 Thus, a need exists for a liquid formulation of SYNAGIS[®] at a concentration comparable to or higher than the reconstituted lyophilized formulation so that there is no need to reconstitute the formulation prior to administration. This allows health care practitioners much quicker and easier administration of SYNAGIS[®] to a patient.

Prior liquid antibody preparations have short shelf lives and may lose biological
30 activity of the antibodies resulting from chemical and physical instabilities during the storage. Chemical instability may be caused by deamidation, racemization, hydrolysis, oxidation, beta elimination or disulfide exchange, and physical instability may be caused by antibody denaturation, aggregation, precipitation or adsorption. Among those, aggregation, deamidation and oxidation are known to be the most common causes of the antibody

degradation (Wang *et al.*, 1988, *J. of Parenteral Science & Technology* 42(Suppl):S4-S26; Cleland *et al.*, 1993, *Critical Reviews in Therapeutic Drug Carrier Systems* 10(4):307-377). Thus, there is a need for a stable liquid formulation of SYNAGIS® or an antigen-binding fragment thereof effective to prevent RSV infection.

3. SUMMARY OF INVENTION

The present invention is based, in part, on the development of high concentration liquid formulations of SYNAGIS® or an antigen-binding fragment thereof, which formulations exhibit, in the absence of surfactant, inorganic salts, and/or other excipients, stability, low to undetectable levels of antibody fragmentation and/or aggregation, and very little to no loss of the biological activity(ies) of SYNAGIS® or an antigen-binding fragment thereof during manufacture, preparation, transportation, and storage. The liquid formulations of the present invention facilitate the administration of SYNAGIS® or an antigen-binding fragment thereof for the prevention, treatment, management and/or amelioration of a RSV infection or one or more symptoms thereof. In particular, the liquid formulations of the present invention enable a healthcare professional to quickly administer a sterile dosage of SYNAGIS® or an antigen-binding fragment thereof without having to accurately and aseptically reconstitute the antibody or antibody fragment prior to administration as required for the lyophilized dosage form. Such liquid formulations of SYNAGIS® can be also manufactured more easily and cost effectively than the lyophilized formulation since liquid formulations do not require a prolonged drying step, such as lyophilization, freeze-drying, etc. The liquid formulations are made by a process in which the antibody being formulated is in an aqueous phase throughout the purification and formulation process. Preferably, the liquid formulations are made by a process that does not include a drying step, for example, but not by way of limitation, a lyophilization, freeze-drying, spray-drying, or air-drying step.

The present invention provides liquid formulations of SYNAGIS® or an antigen-binding fragment thereof, substantially free of surfactant, inorganic salts, sugars, and/or other common excipients, said formulations comprising histidine and a concentration of about 15 mg/ml or higher of SYNAGIS® or an antigen-binding fragment thereof. Optionally, the formulation may further comprise glycine. Alternatively, the formulation of the present invention may further comprise other common excipients, such as saccharides, polyols and amino acids, including, but not limited to, arginine, lysine, and methionine. The present invention also provides liquid formulations substantially free of surfactant,

inorganic salts, sugars, and/or other commonly-known excipients, said formulation having a pH ranging from about 5.0 to about 7.0, preferably about 5.5 to 6.5, more preferably about 5.8 to about 6.2, and most preferably about 6.0, and comprising histidine and a concentration of about 15 mg/ml or higher of SYNAGIS[®] or an antigen-binding fragment thereof.

The present invention encompasses stable liquid formulations of SYNAGIS[®] which exhibit low to undetectable levels of antibody aggregation and/or fragmentation with very little to no loss of biological activity(ies) of SYNAGIS[®] or an antigen-binding fragment thereof during manufacture, preparation, transportation and long periods of storage. The present invention also encompasses stable liquid formulations of modified forms of SYNAGIS[®] or an antigen-binding fragment thereof that have increased *in vivo* half-lives relative to unmodified SYNAGIS[®] or an antigen-binding fragment thereof, said formulations exhibiting low to undetectable levels of antibody aggregation and/or fragmentation, and very little to no loss of biological activity(ies) of SYNAGIS[®] or an antigen-binding fragment thereof.

The present invention encompasses liquid formulations of SYNAGIS[®] or an antigen-binding fragment thereof, said formulations having stability at 38° - 42°C as assessed by high performance size exclusion chromatography (HPSEC). The liquid formulations of the present invention exhibits stability, as assessed by HSPEC, at the temperature ranges of 38°C-42°C for at least 60 days (in specific embodiments, not more than 120 days), of 20°C-24°C for at least 1 year, and of 2°C-8°C for at least 3 years. The present invention also encompasses liquid formulations of SYNAGIS[®] or an antigen-binding fragment thereof, said formulations having low to undetectable aggregation as measured by HPSEC. In a preferred embodiment, the liquid formulations of the present invention exhibit stability at 38°-42°C for at least 60 days and exhibit low to undetectable levels of antibody aggregation as measured by HPSEC, and further, exhibit very little to no loss of biological activity (ies) of SYNAGIS[®] or an antigen-binding fragment thereof compared to the reference antibodies as measured by antibody binding assays such as, *e.g.*, ELISAs.

The present invention provides methods for preparing liquid formulations of SYNAGIS[®] or an antigen-binding fragment thereof. The liquid formulations of the present invention are prepared by maintaining SYNAGIS[®] or an antigen-binding fragment thereof in an aqueous solution at any time during the preparation. In other words, the liquid formulations are prepared without involving any step of drying SYNAGIS[®] or an antigen-

binding fragment thereof or the formulations themselves by, for example, lyophilization, vacuum drying, etc.

The present invention provides methods for preparing liquid formulations of SYNAGIS[®] or an antigen-binding fragment thereof, said methods comprising concentrating
5 a fraction of purified SYNAGIS[®] or an antigen-binding fragment thereof to a final concentration of about 15 mg/ml, about 20 mg/ml, about 30 mg/ml, about 40 mg/ml, about 50 mg/ml, about 60 mg/ml, about 70 mg/ml, about 80 mg/ml, about 90 mg/ml, about 100 mg/ml, about 200 mg/ml, about 250 mg/ml, or about 300 mg/ml using a semi-permeable membrane with an appropriate molecular weight (mw) cutoff (*e.g.*, 30 kD cutoff for
10 SYNAGIS[®] and F(ab')₂ fragments thereof; and 10 kD cutoff for SYNAGIS[®] fragments such Fab fragments), and diafiltering the concentrated antibody fraction into the formulation buffer using the same membrane. The formulation buffer of the present invention comprises histidine at a concentration ranging from about 1 mM to about 100 mM, about 10 mM to about 50 mM, about 20 mM to about 30 mM, or about 23 mM to about 27 mM, and
15 is most preferably about 25 mM. To obtain an appropriate pH for SYNAGIS[®] or an antigen-binding fragment thereof, it is preferably that histidine (and glycine, if added) is first dissolved in water to obtain a buffer solution with higher pH than the desired pH and then the pH is brought down to the desired level by the addition of HCl. This way, the formation of inorganic salts (*e.g.*, the formation of NaCl when, *e.g.*, histidine hydrochloride
20 is used as the source of histidine and the pH is raised to the desired level by the addition of NaOH) can be avoided.

The liquid formulations of the present invention may be sterilized by sterile filtration using a 0.2 μ or a 0.22 μ filter. Sterilized liquid formulations of the present invention may be administered to a subject to prevent, treat or ameliorate one or more symptoms
25 associated with a RSV infection or a symptom thereof.

The present invention also provides kits comprising the liquid formulations of SYNAGIS[®] or an antigen-binding fragment thereof for use by, *e.g.* a healthcare professional. The present invention further provides methods of preventing, treating managing or ameliorating a RSV infection or one or more symptoms thereof by
30 administering the liquid formulations of the present invention.

3.1 Terminology

As used herein, all liquid formulations of SYNAGIS[®] and/or fragments thereof that immunospecifically bind to a RSV antigen described above are collectively referred to as

"liquid formulations of the invention," "SYNAGIS[®] liquid formulations of the invention," or "liquid formulations of SYNAGIS[®] or an antigen-binding fragment thereof."

As used herein, the term "cytokine receptor modulator" refers to an agent which modulates the phosphorylation of a cytokine receptor, the activation of a signal transduction pathway associated with a cytokine receptor, and/or the expression of a particular protein such as a cytokine. Such an agent may directly or indirectly modulate the phosphorylation of a cytokine receptor, the activation of a signal transduction pathway associated with a cytokine receptor, and/or the expression of a particular protein such as a cytokine. Thus, examples of cytokine receptor modulators include, but are not limited to, cytokines, fragments of cytokines, fusion proteins and antibodies that immunospecifically binds to a cytokine receptor or a fragment thereof. Further, examples of cytokine receptor modulators include, but are not limited to, peptides, polypeptides (*e.g.*, soluble cytokine receptors), fusion proteins and antibodies that immunospecifically binds to a cytokine or a fragment thereof.

As used herein, the terms "SYNAGIS[®] fragment", "antigen-binding fragment" and like terms used in the context of SYNAGIS[®] refer to a fragment of SYNAGIS[®] that immunospecifically binds to a RSV antigen. Fragments of SYNAGIS[®] may be generated by any technique known to those skilled in the art. For example, Fab and F(ab')₂ fragments may be produced by proteolytic cleavage of immunoglobulin molecules, using enzymes such as papain (to produce Fab fragments) or pepsin (to produce F(ab')₂ fragments). F(ab')₂ fragments contain the complete light chain, and the variable region, the CH1 region and the hinge region of the heavy chain. Preferably, the fragment also binds to a RSV antigen, more preferably to the same epitope as SYNAGIS[®].

As used herein, the term "effective amount" refers to the amount of a therapy (*e.g.*, a prophylactic or therapeutic agent), which is sufficient to reduce the severity, and/or duration of a RSV infection, ameliorate one or more symptoms thereof, prevent the advancement of a RSV infection, or cause regression of a RSV infection, or which is sufficient to result in the prevention of the development, recurrence, onset, or progression of a RSV infection or one or more symptoms thereof, or enhance or improve the prophylactic and/or therapeutic effect(s) of another therapy (*e.g.*, another therapeutic agent). In a specific embodiment, an effective amount of a therapeutic or a prophylactic agent reduces one or more of the following steps of a RSV life cycle: the docking of the virus particle to a cell, the introduction of viral genetic information into a cell, the expression of viral proteins, the production of new virus particles and the release of virus particles from a cell by at least

5%, preferably at least 10%, at least 15%, at least 20%, at least 25%, at least 30%, at least 35%, at least 40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, or at least 100%. In another specific embodiment, an effective amount of a therapeutic or a

5 prophylactic agent reduces the replication, multiplication or spread of a virus by at least 5%, preferably at least 10%, at least 15%, at least 20%, at least 25%, at least 30%, at least 35%, at least 40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, or at least 100%.

As used herein, the term "epitope" refers to a portion of a RSV polypeptide or
 10 protein having antigenic or immunogenic activity in an animal, preferably a mammal, and most preferably in a human. An epitope having immunogenic activity is a portion of a RSV polypeptide or protein that elicits an antibody response in an animal. An epitope having antigenic activity is a portion of a RSV polypeptide or protein to which an antibody immunospecifically binds as determined by any method well known in the art, for example,
 15 by the immunoassays described herein. Antigenic epitopes need not necessarily be immunogenic. Specifically, the epitope of SYNAGIS® is the A antigenic site of the F protein of RSV.

As used herein, the term "excipients" refers to inert substances which are commonly used as a diluent, vehicle, preservatives, binders, or stabilizing agent for drugs and includes,
 20 but not limited to, proteins (*e.g.*, serum albumin, etc.), amino acids (*e.g.*, aspartic acid, glutamic acid, lysine, arginine, glycine, histidine, etc.), fatty acids and phospholipids (*e.g.*, alkyl sulfonates, caprylate, etc.), surfactants (*e.g.*, SDS, polysorbate, nonionic surfactant, etc.), saccharides (*e.g.*, sucrose, maltose, trehalose, etc.) and polyols (*e.g.*, mannitol, sorbitol, etc.). *Also see* Remington's Pharmaceutical Sciences (by Joseph P. Remington,
 25 18th ed., Mack Publishing Co., Easton, PA).

Preferably, the excipients impart a beneficial physical property to the formulation, such as increased protein stability, increased protein solubility and decreased viscosity.

The term "fragment" as used herein refers to a peptide, polypeptide, or protein (including an antibody) comprising an amino acid sequence of at least 5 contiguous amino
 30 acid residues, at least 10 contiguous amino acid residues, at least 15 contiguous amino acid residues, at least 20 contiguous amino acid residues, at least 25 contiguous amino acid residues, at least 40 contiguous amino acid residues, at least 50 contiguous amino acid residues, at least 60 contiguous amino residues, at least 70 contiguous amino acid residues, at least contiguous 80 amino acid residues, at least contiguous 90 amino acid residues, at

least contiguous 100 amino acid residues, at least contiguous 125 amino acid residues, at least 150 contiguous amino acid residues, at least contiguous 175 amino acid residues, at least contiguous 200 amino acid residues, or at least contiguous 250 amino acid residues of the amino acid sequence of another polypeptide or protein. In a specific embodiment, a
5 fragment of a protein or polypeptide retains at least one function of the protein or polypeptide. In another embodiment, a fragment of a protein or polypeptide retains at least two, three or four functions of the protein or polypeptide. Preferably a fragment of an antibody that immunospecifically binds to a RSV antigen retains the ability to bind to a RSV antigen.

10 As used herein, the term "fusion protein" refers to a polypeptide or protein that comprises an amino acid sequence of a first protein, polypeptide or fragment, analog or derivative thereof, and an amino acid sequence of a heterologous protein or polypeptide (*i.e.*, a second protein, polypeptide or fragment, analog or derivative thereof different than the first protein or functional fragment, analog or derivative thereof). In one embodiment, a
15 fusion protein comprises a prophylactic or therapeutic agent fused to a heterologous protein, polypeptide or peptide. In accordance with this embodiment, the heterologous protein, polypeptide or peptide may or may not be a different type of prophylactic or therapeutic agent.

As used herein, the term "human infant" refers to a human less than 24 months, preferably less than 16 months, less than 12 months, less than 6 months, less than 3 months, less than 2 months, or less than 1 month of age.

As used herein, the term "human infant born prematurely" refers to a human born at less than 40 weeks gestational age, preferably less than 35 weeks gestational age, who is less than 6 months old, preferably less than 3 months old, more preferably less than 2
25 months old, and most preferably less than 1 month old.

As used herein, the term "high concentration" refers to a concentration of 50 mg/ml or higher, preferably 95 mg/ml or higher of an antibody or fragment thereof that immunospecifically binds to a RSV antigen, in an antibody formulation.

As used herein, the term "host cell" includes a subject cell transfected or
30 transformed with a nucleic acid molecule and the progeny or potential progeny of such a cell. Progeny of such a cell may not be identical to the parent cell transfected with the nucleic acid molecule due to mutations or environmental influences that may occur in succeeding generations or integration of the nucleic acid molecule into the host cell genome.

As used herein, the term "immunospecifically binds a RSV antigen" and analogous terms refer to antibodies or fragments thereof that specifically bind to a RSV antigen thereof and do not specifically bind to other polypeptides. Antibodies or fragments that immunospecifically bind a RSV antigen may be cross-reactive with related antigens.

5 Preferably, antibodies or fragments that immunospecifically bind a RSV antigen do not cross-react with other antigens. Antibodies or fragments that immunospecifically bind to and RSV antigen can be identified, for example, by immunoassays, BIAcore, isothermal titration calorimetry, or other techniques known to those of skilled in the art. An antibody or an antigen-binding fragment thereof binds specifically to a RSV antigen when it binds
10 with higher affinity than to any cross-reactive antigen as determined using experimental techniques, such as radioimmunoassays (RIA) and enzyme-linked immunosorbent assays (ELISAs). See, *e.g.*, Paul, ed., 1989, Fundamental Immunology Second Edition, Raven Press, New York at pages 332-336 for a discussion regarding antibody specificity.

The term "in combination" as used herein refers to the use of more than one
15 therapies (*e.g.*, prophylactic and/or therapeutic agents). The use of the term "in combination" does not restrict the order in which therapies (*e.g.*, prophylactic and/or therapeutic agents) are administered to a subject with a RSV infection. A first therapy (*e.g.*, a prophylactic or therapeutic agent) can be administered prior to (*e.g.*, 5 minutes, 15 minutes, 30 minutes, 45 minutes, 1 hour, 2 hours, 4 hours, 6 hours, 12 hours, 24 hours, 48
20 hours, 72 hours, 96 hours, 1 week, 2 weeks, 3 weeks, 4 weeks, 5 weeks, 6 weeks, 8 weeks, or 12 weeks before), concomitantly with, or subsequent to (*e.g.*, 5 minutes, 15 minutes, 30 minutes, 45 minutes, 1 hour, 2 hours, 4 hours, 6 hours, 12 hours, 24 hours, 48 hours, 72 hours, 96 hours, 1 week, 2 weeks, 3 weeks, 4 weeks, 5 weeks, 6 weeks, 8 weeks, or 12 weeks after) the administration of a second therapy (*e.g.*, a prophylactic or therapeutic
25 agent) to a subject with a RSV infection.

As used herein, the term "inorganic salt" refers to any compounds, containing no carbon, that result from replacement of part or all of the acid hydrogen or an acid by a metal or a group acting like a metal and are often used as a tonicity adjusting compound in pharmaceutical compositions and preparations of biological materials. The most common
30 inorganic salts are NaCl, KCl, NaH₂PO₄, etc.

An "isolated" or "purified" antibody is substantially free of cellular material or other contaminating proteins from the cell or tissue source from which the protein is derived, or substantially free of chemical precursors or other chemicals when chemically synthesized. The language "substantially free of cellular material" includes preparations of an antibody

in which the polypeptide/protein is separated from cellular components of the cells from which it is isolated or recombinantly produced. Thus, an antibody that is substantially free of cellular material includes preparations of the antibody having less than about 30%, 20%, 10%, 5%, 2.5%, or 1%, (by dry weight) of contaminating protein. When the antibody is
5 recombinantly produced, it is also preferably substantially free of culture medium, *i.e.*, culture medium represents less than about 20%, 10%, or 5% of the volume of the protein preparation. When antibody is produced by chemical synthesis, it is preferably substantially free of chemical precursors or other chemicals, *i.e.*, it is separated from chemical precursors or other chemicals which are involved in the synthesis of the protein. Accordingly, such
10 preparations of the antibody have less than about 30%, 20%, 10%, 5% (by dry weight) of chemical precursors or compounds other than antibody of interest. In a preferred embodiment of the present invention, antibodies are isolated or purified.

As used herein, the phrase "low to undetectable levels of aggregation" refers to samples containing no more than 5%, no more than 4%, no more than 3%, no more than
15 2%, no more than 1%, and most preferably no more than 0.5%, aggregation by weight protein as measured by high performance size exclusion chromatography (HPSEC).

As used herein, the term "low to undetectable levels of fragmentation" refers to samples containing equal to or more than 80%, 85%, 90%, 95%, 98%, or 99%, of the total protein, for example, in a single peak as determined by high performance size exclusion
20 chromatography (HPSEC), or in two (2) peaks (heavy- and light-chains) by reduced Capillary Gel Electrophoresis (rCGE), representing the non-degraded SYNAGIS[®] or a non-degraded fragment thereof which immunospecifically binds to a RSV antigen, and containing no other single peaks having more than 5%, more than 4%, more than 3%, more than 2%, more than 1%, or more than 0.5% of the total protein each. The term "Reduced-
25 Capillary Gel Electrophoresis (CGE)" as used herein refers to capillary gel electrophoresis under reducing conditions sufficient to reduce disulfide bonds in SYNAGIS[®] or an antigen-binding fragment thereof.

As used herein, the terms "manage", "managing" and "management" refer to the beneficial effects that a subject derives from a therapy (*e.g.*, SYNAGIS[®] or an antigen-
30 binding fragment thereof), which does not result in a cure of a RSV infection. In certain embodiments, a subject is administered one or more therapies (*e.g.*, prophylactic or therapeutic agents) to "manage" a RSV infection or a symptom thereof so as to prevent the progression or worsening of the infection.

As used herein, the term "modified" in the context of modified forms of SYNAGIS[®] or an antigen-binding fragment thereof refers to SYNAGIS[®] or an antigen-binding fragment thereof which has been altered by any method known in the art to increase its half life (see, e.g., Section 5.1.1., *infra*). SYNAGIS[®] and antigen-binding fragments thereof with improved *in vivo* half-lives and methods for preparing them are disclosed in International Publication No. WO 02/060919, filed December 12, 2001, and U.S. Patent No. 7,083,784, both entitled "Molecules with Extended Half-Lives, Compositions and Uses" and by L. Johnson *et al.*

The term "modified" in the context of SYNAGIS[®] or an antigen-binding fragment thereof also refers to SYNAGIS[®] or an antigen-binding fragment modified by covalent attachment of any type of molecule to SYNAGIS[®] or an antigen-binding fragment thereof. For example, but not by way of limitation, SYNAGIS[®] or an antigen-binding fragment thereof may be modified, e.g., by glycosylation, acetylation, pegylation, phosphorylation, amidation, derivatization by known protecting/blocking groups, proteolytic cleavage, linkage to a cellular ligand or other protein, etc.

As used herein, the term "pharmaceutically acceptable" means being approved by a regulatory agency of the Federal or a state government or listed in the U.S. Pharmacopia, European Pharmacopia or other generally recognized pharmacopia for use in animals, and more particularly in humans.

As used herein, the term "polyol" refers to a sugar that contains many -OH groups compared to a normal saccharide.

As used herein, the terms "prevent", "preventing", and "prevention" refer to the prevention or reduction of the recurrence, onset, development or progression of a RSV infection, or the prevention or reduction of the severity and/or duration of a RSV infection or one or more symptoms thereof.

As used herein, the terms "prophylactic agent" and "prophylactic agents" refer to any agent(s) which can be used in the prevention of a RSV infection or one or more symptoms thereof. In certain embodiments, the term "prophylactic agent" refers to SYNAGIS[®] or an antigen-binding fragment thereof. In accordance with these embodiments, the antibody or antibody fragment may be a component of a liquid formulation of the invention. In certain other embodiments, the term "prophylactic agent" does not refer to SYNAGIS[®] or an antigen-binding fragment thereof. In yet other embodiments, the term "prophylactic agent" does not refer to antibodies or fragments thereof other than SYNAGIS[®] that immunospecifically bind to a RSV antigen. Preferably,

a prophylactic agent is an agent which is known to be useful to, or has been or is currently being used to prevent or impede the onset, development, progression, and/or severity of a RSV infection or a symptom thereof.

As used herein, the term “prophylactically effective amount” refers to the amount of a liquid formulation of the invention which is sufficient to result in the prevention of the development, recurrence, onset or progression of a RSV infection. In a specific embodiment, a prophylactically effective amount of a prophylactic agent reduces one or more of the following steps of a RSV life cycle: the docking of the virus particle to a cell, the introduction of viral genetic information into a cell, the expression of viral proteins, the production of new virus particles and the release of virus particles from a cell by at least 5%, preferably at least 10%, at least 15%, at least 20%, at least 25%, at least 30%, at least 35%, at least 40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, or at least 100%. In another specific embodiment, a prophylactically effective amount of a prophylactic agent reduces the replication, multiplication or spread of a virus by at least 5%, preferably at least 10%, at least 15%, at least 20%, at least 25%, at least 30%, at least 35%, at least 40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, or at least 100%.

As used herein, the term “RSV antigen” refers to a RSV protein, polypeptide or peptide to which an antibody immunospecifically binds.

As used herein, the term “saccharide” refers to a class of molecules that are polyhydric alcohols. Saccharides are commonly referred to as carbohydrates and may contain different amounts of sugar (saccharide) units, e.g., monosaccharides, dissacharides and polysacchorides.

The term “small molecule” and analogous terms include, but are not limited to, peptides, peptidomimetics, amino acids, amino acid analogues, polynucleotides, polynucleotide analogues, nucleotides, nucleotide analogues, organic or inorganic compounds (*i.e.*, including heterorganic and/or ganometallic compounds) having a molecular weight less than about 10,000 grams per mole, organic or inorganic compounds having a molecular weight less than about 5,000 grams per mole, organic or inorganic compounds having a molecular weight less than about 1,000 grams per mole, organic or inorganic compounds having a molecular weight less than about 500 grams per mole, and salts, esters, and other pharmaceutically acceptable forms of such compounds.

As used herein, The terms "stability" and "stable" in the context of a liquid formulation comprising SYNAGIS[®] or an antigen-binding fragment thereof refer to the resistance of SYNAGIS[®] or an antigen-binding fragment thereof in the formulation to thermal and chemical unfolding, aggregation, degradation or fragmentation under given manufacture, preparation, transportation and storage conditions. The "stable" formulations of the invention retain biological activity equal to or more than 80%, 85%, 90%, 95%, 98%, 99%, or 99.5% under given manufacture, preparation, transportation and storage conditions. The stability of SYNAGIS[®] or an antigen-binding fragment thereof can be assessed by degrees of aggregation, degradation or fragmentation by methods known to those skilled in the art, including but not limited to reduced Capillary Gel Electrophoresis (rCGE), Sodium Dodecyl Sulfate Polyacrylamide Gel Electrophoresis (SDS-PAGE) and HPSEC, compared to a reference, that is, a commercially available lyophilized SYNAGIS[®] reconstituted to 100 mg/ml in 47 mM histidine/3 mM glycine buffer with 5.6% mannitol at pH 6.0. The reference regularly gives a single peak ($\geq 97\%$ area) by HPSEC. The overall stability of a liquid formulation comprising SYNAGIS[®] or an antigen-binding fragment thereof that immunospecifically binds to a RSV antigen can be assessed by various immunological assays including, for example, ELISA and radioimmunoassay using the specific epitope of RSV.

As used herein, the term "SYNAGIS[®] standard reference" or analogous terms refer to commercially available lyophilized SYNAGIS[®], as described in the Physicians' Desk Reference, 56th edition, 2002. Reconstituted SYNAGIS[®] may contain, *e.g.*, the following excipients: 47 mM histidine, 3.0 mM glycine and 5.6% manitol and the active ingredient, the antibody, at a concentration of 100 milligrams per ml solution.

The terms "subject" and "patient" are used interchangeably herein. As used herein, the terms "subject" and "subjects" refer to an animal, preferably a mammal including a non-primate (*e.g.*, a cow, pig, horse, cat, dog, rat, and mouse) and a primate (*e.g.*, a chimpanzee, a monkey such as a cynomolgous monkey, and a human), and more preferably a human.

As used herein, the term "substantially free of surfactant" refers to a formulation of SYNAGIS[®] or an antigen-binding fragment thereof, said formulation containing less than 0.0005%, less than 0.0003%, or less than 0.0001% of surfactants and/or less than 0.0005%, less than 0.0003%, or less than 0.0001% of surfactant.

As used herein, the term "substantially free of inorganic salts" refers to a formulation of SYNAGIS[®] or an antigen-binding fragment thereof, said formulation containing less than 0.0005%, less than 0.0003%, or less than 0.0001% of inorganic salts.

As used herein, the term "surfactant" refers to organic substances having amphipathic structures; namely, they are composed of groups of opposing solubility tendencies, typically an oil-soluble hydrocarbon chain and a water-soluble ionic group. Surfactants can be classified, depending on the charge of the surface-active moiety, into anionic, cationic, and nonionic surfactants. Surfactants are often used as wetting, emulsifying, solubilizing, and dispersing agents for various pharmaceutical compositions and preparations of biological materials.

As used herein, the terms "therapeutic agent" and "therapeutic agents" refer to any agent(s) which can be used in the treatment, management or amelioration a RSV infection or one or more symptoms thereof. In certain embodiments, the term "therapeutic agent" refers to SYNAGIS[®] or an antigen-binding fragment thereof. In accordance with these embodiments, the antibody or antibody fragment may be a component of a liquid formulation of the invention. In certain other embodiments, the term "therapeutic agent" does not refer to SYNAGIS[®] or an antigen-binding fragment thereof. In yet other embodiments, the term "therapeutic agent" does not refer to antibodies or fragments thereof, other than SYNAGIS[®] that immunospecifically bind to a RSV antigen. Preferably, a therapeutic agent is an agent which is known to be useful for, or has been or is currently being used for the treatment, management or amelioration of a RSV infection or one or more symptoms thereof.

As used herein, the term "therapeutically effective amount" refers to the amount of a liquid formulation of the invention that reduces or ameliorates the progression, severity, and/or duration of a RSV infection, and/or ameliorates one or more symptoms associated with a RSV infection. With respect to the treatment of a RSV infection, a therapeutically effective amount refers to the amount of a therapeutic agent sufficient to reduce or inhibit the replication of a virus, inhibit or reduce the infection of cell with the virus, inhibit or reduce the production of the viral particles, inhibit or reduce the release of viral particles, inhibit or reduce the spread of the virus to other tissues or subjects, or ameliorate one or more symptoms associated with the infection. In a specific embodiment, a therapeutically effective amount of a therapeutic agent reduces one or more of the following steps of a RSV life cycle: the docking of the virus particle to a cell, the introduction of viral genetic information into a cell, the expression of viral proteins, the production of new virus particles and the release of virus particles from a cell by at least 5%, preferably at least 10%, at least 15%, at least 20%, at least 25%, at least 30%, at least 35%, at least 40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%,

at least 85%, at least 90%, at least 95%, or at least 100%. In another specific embodiment, a therapeutically effective amount of a therapeutic agent reduces the replication, multiplication or spread of a virus by at least 5%, preferably at least 10%, at least 15%, at least 20%, at least 25%, at least 30%, at least 35%, at least 40%, at least 45%, at least 50%,
5 at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, or at least 100%.

As used herein, the terms “therapies” and “therapy” can refer to any protocol(s), method(s) and/or agent(s) that can be used in the prevention, treatment, management or amelioration of a RSV infection or one or more symptoms thereof. In certain embodiments,
10 the terms “therapy” and “therapies” refer to biological therapy, and/or other therapies useful for the treatment of a RSV infection known to medical personnel skilled.

As used herein, the terms “treat”, “treating” and “treatment” refer to the reduction or amelioration of the progression, severity, and/or duration of a RSV infection and/or reduces or ameliorates one or more symptoms of a RSV infection. In specific embodiments, such
15 terms refer to the reduction or inhibition of the replication of a respiratory syncytial virus (RSV), the inhibition or reduction in the spread of a respiratory syncytial virus (RSV) to other tissues or subjects, the inhibition or reduction of infection of a cell with a respiratory syncytial virus (RSV), or the amelioration of one or more symptoms associated with a respiratory syncytial virus (RSV) infection.

20 A used herein, a “protocol” includes dosing schedules and dosing regimens. The protocols herein are methods of use and include prophylactic and therapeutic protocols.

As used herein, the term “T cell receptor modulator” refers to an agent which modulates the phosphorylation of a T cell receptor, the activation of a signal transduction pathway associated with a T cell receptor and/or the expression of a particular protein such
25 as a cytokine. Such an agent may directly or indirectly modulate the phosphorylation of a T cell receptor, the activation of a signal transduction pathway associated with a T cell receptor, and/or the expression of a particular protein such as a cytokine. Examples of T cell receptor modulators include, but are not limited to, peptides, polypeptides, proteins, fusion proteins and antibodies which immunospecifically bind to a T cell receptor or a
30 fragment thereof. Further, examples of T cell receptor modulators include, but are not limited to, proteins, peptides, polypeptides (e.g., soluble T cell receptors), fusion proteins and antibodies that immunospecifically binds to a ligand for a T cell receptor or a fragment thereof.

As used herein, the term "very little to no loss of the biological activities" refers to antibody activities, including specific binding abilities of SYNAGIS[®] or an antigen-binding fragment as measured by various immunological assays, including, but not limited to ELISAs and radioimmunoassays. In one embodiment, SYNAGIS[®] or an antigen-binding
5 fragment of the liquid formulations of the invention retain approximately 50%, preferably 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95% or 98% of the ability to immunospecifically bind to a RSV antigen as compared to a reference antibody or antibody fragment as measured by an immunological assay known to one of skill in the art or described herein. For example, an ELISA based assay may be used to compare the ability
10 of a liquid formulation of SYNAGIS[®] or an antigen-binding fragment thereof to immunospecifically bind to a RSV antigen to a SYNAGIS[®] reference standard. In this assay, plates are coated with a RSV antigen and the binding signal of a set concentration of a SYNAGIS[®] reference standard is compared to the binding signal of the same concentration of a test antibody or antibody fragment.

4. BRIEF DESCRIPTION OF THE FIGURES

Fig. 1 is a schematic diagram showing the outline for preparing purified SYNAGIS[®].

Fig. 2 shows the clinical study flow chart for comparing the liquid formulation of SYNAGIS[®] with the lyophilized formulation of SYNAGIS[®].

5. DETAILED DESCRIPTION OF THE INVENTION

The liquid formulations of the present invention provide a ready-to-use preparation of SYNAGIS[®] or an antigen-binding fragment thereof for administering to a subject without having to reconstitute the preparation accurately and aseptically and waiting for a period of time until the solution clarifies before administering the formulation to the subject. It
25 simplifies the procedure of administering the formulation to a subject for a healthcare professional. Furthermore, due to its high stability during the storage, the formulations of the present invention can contain SYNAGIS[®] or an antigen-binding fragment thereof at concentrations in the range of about 15 mg/ml to about 300 mg/ml without causing an adverse effect on the biological activity(ies) of SYNAGIS[®] or an antigen-binding fragment
30 thereof due to protein aggregation and/or fragmentation during a prolonged storage. Such stability not only ensures the efficacy of SYNAGIS[®] or an antigen-binding fragment thereof but also reduces possible risks of causing adverse effects on a subject. In addition, the manufacturing process of the liquid formulations of the present invention is simplified and

more efficient than the manufacturing process for the lyophilized version because all stages of the manufacturing of the liquid formulations are carried out in an aqueous solution, involving no drying process, such as lyophilization and freeze-drying. Accordingly, it is more cost effective as well.

5 5.1 SYNAGIS[®] Liquid Formulations

The liquid formulations of the present invention provide antibody formulations which are substantially free of surfactant, inorganic salts, and/or other excipients and yet exhibit high stability during long periods of storage. In a specific embodiment, such antibody formulations are homogeneous. The formulations of the present invention
10 comprise histidine at concentrations between 1 and 100 mM and SYNAGIS[®] or an antigen-binding fragment thereof at concentrations of about 15 mg/ml to about 300 mg/ml. In one embodiment, the formulations of the invention do not comprise other ingredients except for water or suitable solvents. In another specific embodiment, a modified form of SYNAGIS[®] antibody or an antigen-binding fragment thereof having improved half-life and/or affinity is
15 used in the liquid formulations of the invention.

The concentration of SYNAGIS[®] or an antigen-binding fragment thereof which is included in the liquid formulations of the invention, is at least 15 mg/ml, at least 20 mg/ml, at least 25 mg/ml, at least 30 mg/ml, at least 35 mg/ml, at least 40 mg/ml, at least 45 mg/ml, at least 50 mg/ml, at least 55 mg/ml, at least 60 mg/ml, at least 65 mg/ml, at least 70 mg/ml,
20 at least 75 mg/ml, at least 80 mg/ml, at least 85 mg/ml, at least 90 mg/ml, at least 95 mg/ml, at least 100 mg/ml, at least 105 mg/ml, at least 110 mg/ml, at least 115 mg/ml, at least 120 mg/ml, at least 125 mg/ml, at least 130 mg/ml, at least 135 mg/ml, at least 140 mg/ml, at least 150 mg/ml, at least 200 mg/ml, at least 250 mg/ml, or at least 300 mg/ml.

The concentration of histidine which is included in the liquid formulations of the
25 invention ranges from about 1 mM to about 100 mM, about 10 mM to about 50 mM, about 20 mM to about 30 mM, or about 23 mM to about 27 mM, and is most preferably about 25 mM. Histidine can be in the form of L-histidine, D-histidine, or a mixture thereof, but L-histidine is the most preferable. Histidine can be also in the form of hydrates. Histidine may be used in a form of pharmaceutically acceptable salt, such as hydrochloride (e.g.,
30 monohydrochloride and dihydrochloride), hydrobromide, sulfate, acetate, etc. The purity of histidine should be at least 98%, preferably at least 99%, and most preferably at least 99.5%.

The pH of the formulation should not be equal to the isoelectric point of the particular antibody to be used in the formulation (*e.g.*, the isoelectric point of SYNAGIS® ranges from 8.65 to 9.43) and may range from about 5.0 to about 7, preferably about 5.5 to about 6.5, more preferably about 5.8 to about 6.2, and most preferably about 6.0.

5 In addition to histidine and SYNAGIS® or an antigen-binding fragment thereof, the formulations of the present invention may further comprise glycine at a concentration of less than 150 mM, less than 100 mM, less than 50 mM, less than 3.0 mM, less than 2.0 mM, or less than 1.8 mM, and most preferably 1.6 mM. The amount of glycine in the formulation should not cause a significant buffering effect so that antibody precipitation at
10 its isoelectric point can be avoided. Glycine may be also used in a form of a pharmaceutically acceptable salt, such as hydrochloride, hydrobromide, sulfate, acetate, etc. The purity of glycine should be at least 98%, preferably at least 99%, and most preferably 99.5%. In a specific embodiment, glycine is not included in the liquid formulations of the present invention.

15 Optionally, the formulations of the present invention may further comprise other excipients, such as saccharides (*e.g.*, sucrose, mannose, trehalose, etc.) and polyols (*e.g.*, mannitol, sorbitol, etc.). In one embodiment, the other excipient is a saccharide. In a specific embodiment, the saccharide is sucrose, which is at a concentration range between about 1% and about 20%, preferably about 5% and about 15%, more preferably about 8%
20 and 10%. In another embodiment, the other excipient is a polyol. Preferably, however, the liquid formulations of the present invention do not contain mannitol. In a specific embodiment, the polyol is polysorbate (*e.g.*, Tween™ 20), which is at a concentration range between about 0.001% and about 1%, preferably, about 0.01 to about 0.1.

 The liquid formulations of the present invention exhibit stability at the temperature
25 ranges of 38°C-42°C for at least 60 days and, in some embodiments, not more than 120 days, of 20°C-24°C for at least 1 year, of 2°C-8°C (in particular, at 4°C) for at least 3 years, at least 4 years, or at least 5 years and at -20°C for at least 3 years, at least 4 years, or at least 5 years, as assessed by high performance size exclusion chromatography (HPSEC). Namely, the liquid formulations of the present invention have low to undetectable levels of
30 aggregation and/or fragmentation, as defined herein, after the storage for the defined periods as set forth above. Preferably, no more than 5%, no more than 4%, no more than 3%, no more than 2%, no more than 1%, and most preferably no more than 0.5%, of SYNAGIS® or an antigen-binding fragment thereof forms an aggregate as measured by HPSEC, after the storage for the defined periods as set forth above. Furthermore, liquid formulations of the

present invention exhibit almost no loss in biological activity(ies) of SYNAGIS® or an antigen-binding fragment thereof during the prolonged storage under the condition described above, as assessed by various immunological assays including, for example, enzyme-linked immunosorbent assay (ELISA) and radioimmunoassay to measure the RSV antigen-binding ability of SYNAGIS® or an antigen-binding fragment thereof, or, for example, by a C3a/C4a assay to measure the complement activating ability of SYNAGIS® or an antigen-binding fragment thereof. The liquid formulations of the present invention retain after the storage for the above-defined periods more than 80%, more than 85%, more than 90%, more than 95%, more than 98%, more than 99%, or more than 99.5% of the initial biological activity(ies) prior to the storage.

The liquid formulations of the present invention can be prepared as unit dosage forms. For example, a unit dosage per vial may contain 1 ml, 2 ml, 3 ml, 4 ml, 5 ml, 6 ml, 7 ml, 8 ml, 9 ml, 10 ml, 15 ml, or 20 ml of different concentrations of SYNAGIS® or an antigen-binding fragment thereof ranging from about 15 mg/ml to about 300 mg/ml concentration of SYNAGIS® or an antigen-binding fragment thereof which immunospecifically binds to a RSV. If necessary, these preparations can be adjusted to a desired concentration by adding a sterile diluent to each vial.

5.1.1 SYNAGIS®

The invention relates to liquid formulations comprising SYNAGIS® or an antigen-binding fragment thereof. In a preferred embodiments, the invention provides liquid formulations of SYNAGIS®, a humanized monoclonal antibody which neutralizes a broad range of RSV isolates. The amino acid sequence of SYNAGIS® is disclosed, *e.g.*, in Johnson *et al.*, 1997, *J. Infectious Disease* 176:1215-1224, and U.S. Patent No. 5,824,307, and its V_HCDRs and V_LCDRs are shown in Table 1, *infra*. The properties and uses of SYNAGIS® are also disclosed in, *e.g.*, other applications, see, *e.g.*, U.S. Patent Application No. 09/724,396 filed November 28, 2000; U.S. Patent No. 6,855,493 and U.S. Patent No. 7,179,900.

Table 1. CDR Sequences of SYNAGIS®

CDR	Sequence	SEQ ID NO:
VH1	TSGMSVG	1
VH2	DIWWDD <u>KK</u> DYNPSLKS	2

VH3	<u>S</u> MITN <u>W</u> YFDV	3
VL1	<u>K</u> COLSVGYMH	4
VL2	DTS <u>K</u> LAS	5
VL3	FQGS <u>G</u> YP <u>F</u> T	6

In addition, the present invention also encompasses stable liquid formulations of modified forms of SYNAGIS[®] or an antigen-binding fragment thereof that have improved half-lives. In particular, the present invention encompasses a modified form of SYNAGIS[®] or an antigen-binding fragment thereof which has a half-life in a subject, preferably a human, of greater than 3 days, greater than 7 days, greater than 10 days, preferably greater than 15 days, greater than 25 days, greater than 30 days, greater than 35 days, greater than 40 days, greater than 45 days, greater than 2 months, greater than 3 months, greater than 4 months, or greater than 5 months. By prolonging the half-lives of SYNAGIS[®] and antigen-binding fragments thereof, it is possible to reduce the amount and/or frequency of dosing of the antibody or antigen-binding fragment.

To prolong the serum circulation of an antibody *in vivo*, various techniques can be used. For example, inert polymer molecules, such as high molecular weight polyethyleneglycol (PEG), can be attached to an antibody with or without a multifunctional linker either through site-specific conjugation of the PEG to the N- or C-terminus of the antibody or via epsilon-amino groups present on lysine residues. Linear or branched polymer derivatization that results in minimal loss of biological activity can be used. The degree of conjugation can be closely monitored by SDS-PAGE and mass spectrometry to ensure proper conjugation of PEG molecules to the antibodies. Unreacted PEG can be separated from antibody-PEG conjugates by size-exclusion or by ion-exchange chromatography. PEG-derivatized antibodies can be tested for binding activity as well as for *in vivo* efficacy using methods known to those of skilled in the art, for example, by immunoassays described herein.

An antibody having an increased half-life *in vivo* can also be generated by introducing one or more amino acid modifications (*i.e.*, substitutions, insertions or deletions) into an IgG constant domain, or FcRn binding fragment thereof (preferably a Fc or hinge Fc domain fragment). See, *e.g.*, International Publication No. WO 98/23289; International Publication No. WO 97/34631; and U.S. Patent No. 6,277,375.

SYNAGIS[®] and antigen-binding fragments thereof with improved *in vivo* half-lives and methods for preparing them are

disclosed in International Application WO 02/060919, filed December 12, 2001, and U.S. Patent No. 7,083,784, both entitled "Molecules with Extended Half-Lives, Compositions and Uses" and by L. Johnson *et al.*

5 Further, an antibody can be conjugated to albumin in order to make the antibody or an antigen-binding fragment thereof more stable *in vivo* or have a longer half life *in vivo*. The techniques are well known in the art, see *e.g.*, International Publication Nos. WO 93/15199, WO 93/15200, and WO 01/77137; and European Patent No. EP 413, 622.

10 The invention further comprises liquid formulations of SYNAGIS® or antigen-binding fragments thereof that have been modified, for example, by glycosylation, acetylation, pegylation, phosphorylation, amidation, derivatization by known protecting/blocking groups, proteolytic cleavage, linkage to a cellular ligand or other protein, etc., and retain RSV antigen-binding activity.

15 5.1.2 Antibody Conjugates

The present invention encompasses the use of liquid formulations of SYNAGIS® or an antigen-binding fragment thereof (including modified forms that have increased *in vivo* half-lives) that conjugated to one or more moieties, including but not limited to, peptides, polypeptides, proteins, fusion proteins, nucleic acid molecules, small molecules, mimetic
20 agents, synthetic drugs, inorganic molecules, and organic molecules.

The present invention encompasses the use of liquid formulations of SYNAGIS® recombinantly fused or chemically conjugated (including both covalent and non-covalent conjugations) to a heterologous protein or polypeptide (or antigen-binding fragment, preferably to a polypeptide of at least 10, at least 20, at least 30, at least 40, at least 50, at
25 least 60, at least 70, at least 80, at least 90 or at least 100 amino acids) to generate fusion proteins. The fusion does not necessarily need to be direct, but may occur through linker sequences. For example, an antibody may be used to target a heterologous polypeptide to a particular cell type, either *in vitro* or *in vivo*, by fusing or conjugating the antibody to another antibody specific for particular cell surface receptors. An antibody fused or
30 conjugated to a heterologous polypeptide may also be used in *in vitro* immunoassays and purification methods using methods known in the art. See *e.g.*, International publication No. WO 93/21232; European Patent No. EP 439,095; Naramura *et al.*, 1994, Immunol. Lett. 39:91-99; U.S. Patent No. 5,474,981; Gillies *et al.*, 1992, PNAS 89:1428-1432; and

Fell *et al.*, 1991, *J. Immunol.* 146:2446-2452.

The present invention further includes compositions comprising a heterologous protein, peptide or polypeptide fused or conjugated to an antigen-binding fragment of SYNAGIS[®]. For example, a heterologous polypeptides may be fused or conjugated to a Fab fragment, Fd fragment, Fv fragment, or F(ab)₂ fragment. Methods for fusing or conjugating a polypeptide to antibody portion are known in the art. See, *e.g.*, U.S. Patent Nos. 5,336,603, 5,622,929, 5,359,046, 5,349,053, 5,447,851, and 5,112,946; European Patent Nos. EP 307,434 and EP 367,166; International publication Nos. WO 96/04388 and WO 91/06570; Ashkenazi *et al.*, 1991, *Proc. Natl. Acad. Sci. USA* 88: 10535-10539; Zheng *et al.*, 1995, *J. Immunol.* 154:5590-5600; and Vil *et al.*, 1992, *Proc. Natl. Acad. Sci. USA* 89:11337- 11341.

Additional fusion proteins may be generated through the techniques of gene-shuffling, motif-shuffling, exon-shuffling, and/or codon-shuffling (collectively referred to as "DNA shuffling"). DNA shuffling may be employed to alter the activities of SYNAGIS[®] or fragments thereof (*e.g.*, an antibody or an antigen-binding fragment thereof with higher affinities and lower dissociation rates). See, generally, U.S. Patent Nos. 5,605,793; 5,811,238; 5,830,721; 5,834,252; and 5,837,458, and Patten *et al.*, 1997, *Curr. Opinion Biotechnol.* 8:724-33 ; Harayama, 1998, *Trends Biotechnol.* 16(2):76-82; Hansson *et al.*, 1999, *J. Mol. Biol.* 287:265-76; and Lorenzo and Blasco, 1998, *Biotechniques* 24(2):308- 313.

SYNAGIS[®] or an antigen-binding fragment thereof, or the nucleic acid encoding SYNAGIS[®] or an antigen-binding fragment thereof, may be altered by being subjected to random mutagenesis by error-prone PCR, random nucleotide insertion or other methods prior to recombination. SYNAGIS[®] or an antigen-binding fragment thereof may be recombined with one or more components, motifs, sections, parts, domains, fragments, etc. of one or more heterologous molecules.

Moreover, SYNAGIS[®] or an antigen-binding fragment thereof can be fused to a marker sequence, such as a peptide to facilitate purification. In preferred embodiments, the marker amino acid sequence is a hexa-histidine peptide, such as the tag provided in a pQE vector (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311), among others, many of which are commercially available. As described in Gentz *et al.*, 1989, *Proc. Natl. Acad. Sci. USA* 86:821-824, for instance, hexa-histidine provides for convenient purification of the fusion protein. Other peptide tags useful for purification include, but are not limited to,

the hemagglutinin "HA" tag, which corresponds to an epitope derived from the influenza hemagglutinin protein (Wilson *et al.*, 1984, Cell 37:767) and the "flag" tag.

The present invention also encompasses the liquid formulations of SYNAGIS® or an antigen-binding fragment thereof conjugated to a diagnostic or detectable agent or any other molecule for which serum half-life is desired to be increased. Such an antibody can be useful for monitoring or prognosing the development or progression of a RSV infection as part of a clinical testing procedure, such as determining the efficacy of a particular therapy. Such diagnosis and detection can be accomplished by coupling SYNAGIS® or an antigen-binding fragment thereof to a detectable substance including, but not limited to, various enzymes, such as but not limited to, horseradish peroxidase, alkaline phosphatase, beta-galactosidase, or acetylcholinesterase; prosthetic groups, such as but not limited to, streptavidin/biotin and avidin/biotin; fluorescent materials, such as but not limited to, umbelliferone, fluorescein, fluorescein isothiocyanate, rhodamine, dichlorotriazinylamine fluorescein, dansyl chloride or phycoerythrin; luminescent materials, such as but not limited to, luminol; bioluminescent materials, such as but not limited to, luciferase, luciferin, and aequorin; radioactive materials, such as but not limited to iodine (¹³¹I, ¹²⁵I, ¹²³I, ¹²¹I), carbon (¹⁴C), sulfur (³⁵S), tritium (³H), indium (¹¹⁵In, ¹¹³In, ¹¹²In, ¹¹¹In), and technetium (⁹⁹Tc), thallium (²⁰¹Tl), gallium (⁶⁸Ga, ⁶⁷Ga), palladium (¹⁰³Pd), molybdenum (⁹⁹Mo), xenon (¹³³Xe), fluorine (¹⁸F), ¹⁵³Sm, ¹⁷⁷Lu, ¹⁵⁹Gd, ¹⁴⁹Pm, ¹⁴⁰La, ¹⁷⁵Yb, ¹⁶⁶Ho, ⁹⁰Y, ⁴⁷Sc, ¹⁸⁶Re, ¹⁸⁸Re, ¹⁴²Pr, ¹⁰⁵Rh, ⁹⁷Ru, ⁶⁸Ge, ⁵⁷Co, ⁶⁵Zn, ⁸⁵Sr, ³²P, ¹⁵³Gd, ¹⁶⁹Yb, ⁵¹Cr, ⁵⁴Mn, ⁷⁵Se, ¹¹³Sn, and ¹¹⁷Tm; positron emitting metals using various positron emission tomographies, noradioactive paramagnetic metal ions, and molecules that are radiolabelled or conjugated to specific radioisotopes. The detectable substance may be coupled or conjugated either directly to SYNAGIS® or an antigen-binding fragment thereof or indirectly, through an intermediate (such as, for example, a linker known in the art) using techniques known in the art. See, *e.g.*, U.S. Patent No. 4,741,900 for metal ions which can be conjugated to antibodies for use as a diagnostics according to the present invention.

The present invention further encompasses uses of SYNAGIS® or an antigen-binding fragment thereof conjugated to a therapeutic moiety. An antibody or antigen-binding fragment may be conjugated to a therapeutic moiety such as a cytotoxin, *e.g.*, a cytostatic or cytotoxic agent, a therapeutic agent or a radioactive metal ion, *e.g.*, alpha-emitters. A cytotoxin or cytotoxic agent includes any agent that is detrimental to cells. Examples include paclitaxel, cytochalasin B, gramicidin D, ethidium bromide, emetine, mitomycin, etoposide, teniposide, vincristine, vinblastine, colchicin, doxorubicin,

daunorubicin, dihydroxy anthracin dione, mitoxantrone, mithramycin, actinomycin D, 1-dehydrotestosterone, glucocorticoids, procaine, tetracaine, lidocaine, propranolol, and puromycin and analogs or homologs thereof. Therapeutic moieties include, but are not limited to, antimetabolites (*e.g.*, methotrexate, 6-mercaptopurine, 6-thioguanine, cytarabine, 5-fluorouracil decarbazine), alkylating agents (*e.g.*, mechlorethamine, thioepa chlorambucil, melphalan, carmustine (BCNU) and lomustine (CCNU)), cyclophosphamide, busulfan, dibromomannitol, streptozotocin, mitomycin C, and cisdichlorodiamine platinum (II) (DDP) cisplatin)); anthracyclines (*e.g.*, daunorubicin (formerly daunomycin) and doxorubicin); antibiotics (*e.g.*, dactinomycin (formerly actinomycin), bleomycin, mithramycin, and anthramycin (AMC)); Auristatin molecules (*e.g.*, auristatin PHE, bryostatin 1, solastatin 10, see Woyke *et al.*, *Antimicrob. Agents Chemother.* 46:3802-8 (2002), Woyke *et al.*, *Antimicrob. Agents Chemother.* 45:3580-4 (2001), Mohammad *et al.*, *Anticancer Drugs* 12:735-40 (2001), Wall *et al.*, *Biochem. Biophys. Res. Commun.* 266:76-80 (1999), Mohammad *et al.*, *Int. J. Oncol.* 15:367-72 (1999));

anti-mitotic agents (*e.g.*, vincristine and vinblastine); hormones (*e.g.*, glucocorticoids, progestatins, androgens, and estrogens); DNA repair enzyme inhibitors (*e.g.*, etoposide or topotecan); kinase inhibitors (*e.g.*, compound ST1571, imatinib mesylate (Kantarjian *et al.*, *Clin Cancer Res.* 8(7):2167-76 (2002)), and those compounds disclosed in U.S. Pat. Nos. 6,245,759, 6,399,633, 6,383,790, 6,335,156, 6,271,242, 6,242,196, 6,218,410, 6,218,372, 6,057,300, 6,034,053, 5,985,877, 5,958,769, 5,925,376, 5,922,844, 5,911,995, 5,872,223, 5,863,904, 5,840,745, 5,728,868, 5,648,239, and 5,587,459); farnesyl transferase inhibitors (*e.g.*, R115777, BMS 214662, and those disclosed by, for example, U.S. Patent Nos: 6,458,935, 6,451,812, 6,440,974, 6,436,960, 6,432,959, 6,420,387, 6,414,145, 6,410,541, 6,410,539, 6,403,581, 6,399,615, 6,387,905, 6,372,747, 6,369,034, 6,362,188, 6,342,765, 6,342,487, 6,300,501, 6,268,363, 6,265,422, 6,248,756, 6,239,140, 6,232,338, 6,228,865, 6,228,856, 6,225,322, 6,218,406, 6,211,193, 6,187,786, 6,169,096, 6,159,984, 6,143,766, 6,133,303, 6,127,366, 6,124,465, 6,124,295, 6,103,723, 6,093,737, 6,090,948, 6,080,870, 6,077,853, 6,071,935, 6,066,738, 6,063,930, 6,054,466, 6,051,582, 6,051,574, and 6,040,305); topoisomerase inhibitors (*e.g.*, camptothecin, irinotecan, SN 38, topotecan, 9 aminocamptothecin, GG 211 (GI 147211), DX 8951f, IST 622, rubitecan, pyrazoloacridine, XR 5000, saintopin, UCE6, UCE1022, TAN 1518A, TAN 1518B, KT6006, KT6528, ED 110, NB 506, ED 110, NB 506, rebeccamycin, and bulgarein); DNA minor groove binders such as Hoescht dye 33342 and Hoechst dye 33258; nitidine; fagaronine; epiberberine; coralyne; beta lapachone; BC 4 1; and pharmaceutically

acceptable salts, solvates, clathrates, and prodrugs thereof (See, *e.g.*, Rothenberg, M.L., *Annals of Oncology* 8:837-855(1997); and Moreau *et al.*, *J. Med. Chem.* 41:1631-1640(1998)). Therapeutic moieties may also be antisense oligonucleotides (*e.g.*, those disclosed in the U.S. Pat. Nos. 6,277,832, 5,998,596, 5,885,834, 5,734,033, and 5,618,709); immunomodulators (*e.g.*, antibodies and cytokines); antibodies (*e.g.*, rituximab (Rituxan®), calicheamycin (Mylotarg®), ibritumomab tiuxetan (Zevalin®), and tositumomab (Bexxar®)); and adenosine deaminase inhibitors (*e.g.*, Fludarabine phosphate and 2-Chlorodeoxyadenosine).

Further, an antibody or an antigen-binding fragment thereof may be conjugated to a therapeutic moiety or drug moiety that modifies a given biological response. Therapeutic moiety or drug moieties are not to be construed as limited to classical chemical therapeutic agents. For example, the drug moiety may be a protein or polypeptide possessing a desired biological activity. Such proteins may include, for example, a toxin such as abrin, ricin A, pseudomonas exotoxin, cholera toxin, or diphtheria toxin; a protein such as tumor necrosis factor, α -interferon, β -interferon, nerve growth factor, platelet derived growth factor, tissue plasminogen activator, an apoptotic agent, *e.g.*, TNF- α , TNF- β , AIM I (see, International publication No. WO 97/33899), AIM II (see, International Publication No. WO 97/34911), Fas Ligand (Takahashi *et al.*, 1994, *J. Immunol.*, 6:1567-1574), and VEGF (see, International publication No. WO 99/23105); or, a biological response modifier such as, for example, a lymphokine (*e.g.*, interleukin-1 ("IL-1"), interleukin-2 ("IL-2"), interleukin-4 ("IL-4"), interleukin-6 ("IL-6"), interleukin-9 (IL-9), interleukin-10 (IL-10), interleukin-12 (IL-12), interferon- α , β , γ , granulocyte macrophage colony stimulating factor ("GM-CSF"), and granulocyte colony stimulating factor ("G-CSF")), or a growth factor (*e.g.*, growth hormone ("GH")).

Moreover, an antibody can be conjugated to therapeutic moieties such as a radioactive metal ion, *e.g.*, alpha-emitters such as ^{213}Bi or macrocyclic chelators useful for conjugating radiometal ions, including but not limited to, ^{131}In , ^{131}Lu , ^{131}Y , ^{131}Ho , ^{131}Sm , to polypeptides. In certain embodiments, the macrocyclic chelator is 1,4,7,10-tetraazacyclododecane-N,N',N'',N'''-tetraacetic acid (DOTA) which can be attached to the antibody via a linker molecule. Such linker molecules are commonly known in the art and described in Denardo *et al.*, 1998, *Clin Cancer Res.* 4(10):2483-90; Peterson *et al.*, 1999, *Bioconjug. Chem.* 10(4):553-7; and Zimmerman *et al.*, 1999, *Nucl. Med. Biol.* 26(8):943-50.

Techniques for conjugating therapeutic moieties to antibodies are well known, see, *e.g.*, Arnon *et al.*, "Monoclonal Antibodies For Immunotargeting Of Drugs In Cancer Therapy", in *Monoclonal Antibodies And Cancer Therapy*, Reisfeld *et al.* (eds.), pp. 243-56 (Alan R. Liss, Inc. 1985); Hellstrom *et al.*, "Antibodies For Drug Delivery", in *Controlled Drug Delivery* (2nd Ed.), Robinson *et al.* (eds.), pp. 623-53 (Marcel Dekker, Inc. 1987); Thorpe, "Antibody Carriers Of Cytotoxic Agents In Cancer Therapy: A Review", in *Monoclonal Antibodies 84: Biological And Clinical Applications*, Pinchera *et al.* (eds.), pp. 475-506 (1985); "Analysis, Results, And Future Prospective Of The Therapeutic Use Of Radiolabeled Antibody In Cancer Therapy", in *Monoclonal Antibodies For Cancer Detection And Therapy*, Baldwin *et al.* (eds.), pp. 303-16 (Academic Press 1985), and Thorpe *et al.*, 1982, *Immunol. Rev.* 62:119-58.

Alternatively, SYNAGIS® or an antigen-binding fragment thereof can be conjugated to a second antibody to form an antibody heteroconjugate as described by Segal in U.S. Patent No. 4,676,980.

SYNAGIS® or an antigen-binding fragment thereof may also be attached to solid supports, which are particularly useful for immunoassays or purification of the target antigen. Such solid supports include, but are not limited to, glass, cellulose, polyacrylamide, nylon, polystyrene, polyvinyl chloride or polypropylene.

The therapeutic moiety or drug conjugated to SYNAGIS® or an antigen-binding fragment thereof should be chosen to achieve the desired prophylactic or therapeutic effect(s) for a RSV infection in a subject. A clinician or other medical professional should consider the following when deciding on which therapeutic moiety or drug to conjugate to SYNAGIS® or an antigen-binding fragment thereof: the severity of the infection, and the condition of the subject.

SYNAGIS® or an antigen-binding fragment thereof, with or without a therapeutic moiety conjugated to it, can be used as a therapeutic.

5.2 Method of Preparing the Antibody Formulations

The present invention provides methods for preparing liquid formulations of SYNAGIS® or an antigen-binding fragment thereof. Figure 1 is a schematic diagram showing the outline for preparing purified SYNAGIS®. The methods for preparing liquid formulations of the present invention comprise: purifying the antibody from conditioned medium (either single lots or pooled lots of medium) and concentrating a fraction containing the purified SYNAGIS® to a final antibody concentration of from about 15

mg/ml, about 20 mg/ml, about 30 mg/ml, about 40 mg/ml, about 50 mg/ml, about 60 mg/ml, about 70 mg/ml, about 80 mg/ml, about 90 mg/ml, about 100 mg/ml, about 150 mg/ml, about 200 mg/ml, about 250 mg/ml, or about 300 mg/ml using a semipermeable membrane with an appropriate molecular weight (MW) cutoff (*e.g.*, 30 kD cutoff for whole antibody molecules and F(ab')₂ fragments; and 10 kD cutoff for antibody fragments, such as Fab fragments) and diafiltrating the concentrated antibody fraction into the formulation buffer using the same membrane. The formulation buffer of the present invention comprises histidine at a concentration from about 1 mM to about 100 mM, about 10 mM to about 50 mM, about 20 mM to about 30 mM, or about 23 mM to about 27 mM, and is most preferably about 25 mM. The formulations may further comprise glycine at a concentration of less than 100 mM, less than 50 mM, less than 3.0 mM, less than 2.0 mM, or less than 1.8 mM, and most preferably of 1.6 mM. The amount of glycine in the formulation should not cause a significant buffering in order to avoid antibody precipitation at its isoelectric point. The pH of the formulation may range from about 5.0 to about 7.0, preferably about 5.5 to about 6.5, more preferably about 5.8 to about 6.2, and most preferably about 6.0. To obtain an appropriate pH for a particular antibody, it is preferable that histidine (and glycine, if added) is first dissolved in water to obtain a buffer solution with higher pH than the desired pH and then the pH is brought down to the desired level by adding HCl. This way, the formation of inorganic salts (*e.g.*, formation of NaCl when, for example, histidine hydrochloride is used as histidine and pH is raised to a desired level by adding NaOH) can be avoided.

The liquid formulations of the present invention can be prepared as unit dosage forms by preparing a vial containing an aliquot of the liquid formulation for a one-time use. For example, a unit dosage per vial may contain 1 ml, 2 ml, 3 ml, 4 ml, 5 ml, 6 ml, 7 ml, 8 ml, 9 ml, 10 ml, 15 ml, or 20 ml of different concentrations of SYNAGIS[®] or an antigen-binding fragment thereof ranging from about 15 mg/ml to about 300 mg/ml concentration of SYNAGIS[®] or an antigen-binding fragment thereof which immunospecifically binds to a RSV. If necessary, these preparations can be adjusted to a desired concentration by adding a sterile diluent to each vial.

The liquid formulations of the present invention may be sterilized by various sterilization methods, including sterile filtration, radiation, etc. In a most preferred embodiment, the diafiltrated antibody formulation is filter-sterilized with a presterilized 0.2 or 0.22-micron filter. Sterilized liquid formulations of the present invention may be

administered to a subject to prevent, treat, manage or ameliorate a RSV infection or one or more symptoms thereof.

Although the invention is directed to liquid non-lyophilized formulations, it should be noted for the purpose of equivalents that the formulations of the invention may be lyophilized if desired. Thus, the invention encompasses lyophilized forms of the formulations of the invention although such lyophilized formulations are not necessary and thus not preferred.

5.3 Methods of Preparing SYNAGIS®

SYNAGIS® and an antigen-binding fragment thereof contained in the liquid formulations of the present invention can be prepared by any method known in the art for the synthesis of antibodies, in particular, by chemical synthesis or, preferably, by recombinant expression techniques.

The nucleotide sequence encoding the heavy and light chain variable domains of SYNAGIS® may be obtained from, for example, co-pending application 09/724,396, filed November 28, 2000 and U.S. Patent No. 6,855,493, both by Young *et al.*

See also, U.S. Patent No.

5,824,307 by Johnson *et al.* In certain embodiments, a nucleic acid encoding SYNAGIS® or an antigen-binding fragment thereof may be chemically synthesized or assembled from oligonucleotides as well known in the art, and then amplified by PCR, cloning or other method known in the art.

Recombinant expression of an antibody (such as SYNAGIS®) requires construction of an expression vector containing a nucleotide sequence that encodes the antibody. Once a nucleotide sequence encoding an antibody molecule or a heavy or light chain of an antibody, or an antigen-binding fragment thereof has been obtained, the vector for the production of the antibody molecule may be produced by recombinant DNA technology using techniques well known in the art as discussed in the previous sections. Methods which are well known to those skilled in the art can be used to construct expression vectors containing antibody coding sequences and appropriate transcriptional and translational control signals. These methods include, for example, *in vitro* recombinant DNA techniques, synthetic techniques, and *in vivo* genetic recombination. The nucleotide sequence encoding the heavy-chain variable region, light-chain variable region, both the heavy-chain and light-chain variable regions, an epitope-binding fragment of the heavy- and/or light-chain variable region, or one or more complementarity determining regions (CDRs) of an

antibody may be cloned into such a vector for expression. Thus-prepared expression vector can be then introduced into appropriate host cells for the expression of the antibody. Accordingly, the invention includes host cells containing a polynucleotide encoding SYNAGIS[®] or an antigen-binding fragment thereof.

5 The host cell may be co-transfected with two expression vectors of the invention, the first vector encoding a heavy chain derived polypeptide and the second vector encoding a light chain derived polypeptide. The two vectors may contain identical selectable markers which enable equal expression of heavy and light chain polypeptides or different selectable markers to ensure maintenance of both plasmids. Alternatively, a single vector may be used
10 which encodes, and is capable of expressing, both heavy and light chain polypeptides. In such situations, the light chain should be placed before the heavy chain to avoid an excess of toxic free heavy chain (Proudfoot, *Nature*, 322:52, 1986; and Kohler, *Proc. Natl. Acad. Sci. USA*, 77:2 197, 1980). The coding sequences for the heavy and light chains may comprise cDNA or genomic DNA.

15 For long-term, high-yield production of recombinant antibodies, stable expression is preferred. For example, cell lines which stably express the antibody molecule may be engineered. Rather than using expression vectors which contain viral origins of replication, host cells can be transformed with DNA controlled by appropriate expression control
20 elements (*e.g.*, promoter, enhancer, sequences, transcription terminators, polyadenylation sites, etc.), and a selectable marker. Following the introduction of the foreign DNA, engineered cells may be allowed to grow for 1-2 days in an enriched media, and then are switched to a selective media. The selectable marker in the recombinant plasmid confers resistance to the selection and allows cells to stably integrate the plasmid into their
25 chromosomes and grow to form foci which in turn can be cloned and expanded into cell lines. This method may advantageously be used to engineer cell lines which express the antibody molecule. Such engineered cell lines may be particularly useful in screening and evaluation of compositions that interact directly or indirectly with the antibody molecule.

 A number of selection systems may be used, including but not limited to, the herpes simplex virus thymidine kinase (Wigler *et al.*, *Cell*, 11:223, 1977), hypoxanthineguanine
30 phosphoribosyltransferase (Szybalska & Szybalski, *Proc. Natl. Acad. Sci. USA*, 48:202, 1992), and adenine phosphoribosyltransferase (Lowy *et al.*, *Cell*, 22:8-17, 1980) genes can be employed in tk⁻, hgp⁺ or apr⁺ cells, respectively. Also, antimetabolite resistance can be used as the basis of selection for the following genes: *dhfr*, which confers resistance to methotrexate (Wigler *et al.*, *Natl. Acad. Sci. USA*, 77:357, 1980 and O'Hare *et al.*, *Proc.*

Natl. Acad. Sci. USA, 78:1527, 1981); *gpt*, which confers resistance to mycophenolic acid (Mulligan & Berg, *Proc. Natl. Acad. Sci. USA*, 78:2072, 1981); *neo*, which confers resistance to the aminoglycoside G-418 (Wu and Wu, *Biotherapy*, 3:87-95, 1991; Tolstoshev, *Ann. Rev. Pharmacol. Toxicol.*, 32:573-596, 1993; Mulligan, *Science*, 260:926-932, 1993; and Morgan and Anderson, *Ann. Rev. Biochem.*, 62: 191-217, 1993; and May, *TIB TECH*, 11(5):155-215, 1993); and *hygro*, which confers resistance to hygromycin (Santerre *et al.*, *Gene*, 30:147, 1984). Methods commonly known in the art of recombinant DNA technology may be routinely applied to select the desired recombinant clone, and such methods are described, for example, in Ausubel *et al.* (eds.), 1993, *Current Protocols in Molecular Biology*, John Wiley & Sons, NY; Kriegler, 1990, *Gene Transfer and Expression, A Laboratory Manual*, Stockton Press, NY; in Chapters 12 and 13, Dracopoli *et al.* (eds), 1994, *Current Protocols in Human Genetics*, John Wiley & Sons, NY; and Colberre-Garapin *et al.*, *J. Mol. Biol.*, 150:1, 1981.

The expression levels of an antibody molecule can be increased by vector amplification (for a review, see Bebbington and Hentschel, 1987, *The use of vectors based on gene amplification for the expression of cloned genes in mammalian cells in DNA cloning*, Vol.3. Academic Press, New York). When a marker in the vector system expressing antibody is amplifiable, increasing in the level of inhibitor present in culture of host cell will increase the number of copies of the marker gene. Since the amplified region is associated with the antibody gene, production of the antibody will also increase (Crouse *et al.*, *Mol., Cell. Biol.*, 3:257, 1983).

Once an antibody molecule of the invention has been produced by recombinant expression, it may be purified by any method known in the art for purification of an immunoglobulin molecule, for example, by chromatography (*e.g.*, ion exchange, affinity, particularly by affinity for the specific antigen after Protein A purification, and sizing column chromatography), centrifugation, differential solubility, or by any other standard techniques for the purification of proteins. Further, SYNAGIS® or an antigen-binding fragment thereof may be fused to heterologous protein, polypeptide or peptide sequences described herein or otherwise known in the art to facilitate purification.

Antigen-binding fragments of SYNAGIS® that immunospecifically bind RSV may be generated by known techniques. For example, Fab and F(ab')₂ fragments may be produced by proteolytic cleavage of immunoglobulin molecules, using enzymes such as papain (to produce Fab fragments) or pepsin (to produce F(ab')₂ fragments). F(ab')₂

fragments contain the complete light chain, and the variable region, the CH1 region and the hinge region of the heavy chain.

5.4 Methods of Monitoring the Stability And Aggregation of Antibody Formulations

There are various methods available for assessing the stability of protein formulations, including antibody formulations, based on the physical and chemical structures of the proteins as well as on their biological activities. For example, to study denaturation of proteins, methods such as charge-transfer absorption, thermal analysis, fluorescence spectroscopy, circular dichroism, NMR, and HPSEC, are available. *See*, for example, Wang *et al.*, 1988, J. of Parenteral Science & Technology 42(supp):S4-S26. The rCGE, and HPSEC are the most common and simplest methods to assess the formation of protein aggregates, protein degradation and protein fragmentation. Accordingly, the stability of the liquid formulations of the present invention may be assessed by these methods.

For example, the stability of the liquid formulations of the present invention may be evaluated by HPSEC or rCGE, wherein the percentage area of the peaks represents the non-degraded SYNAGIS[®] or non-degraded antigen-binding fragments of SYNAGIS[®]. In particular, approximately 250 μ g of SYNAGIS[®] or an antigen-binding fragment thereof (approximately 25 μ l of a liquid formulation comprising 10 mg/ml of SYNAGIS[®] or an antigen-binding fragment thereof) is injected onto a TOSOH TSK G3000SW_{XL} column (7.8 mm x 30 cm) fitted with a TSK SW x1 guard column (6.0 mm x 4.0 cm), SYNAGIS[®] or an antigen-binding fragment thereof is eluted isocratically with 0.1 M disodium phosphate containing 0.1 M sodium sulfate and 0.05% sodium azide, at a flow rate of 0.8 to 1.0 ml/min. Eluted protein is detected using UV absorbance at 280 nm. SYNAGIS[®] reference standard is run in the assay as a control, and the results are reported as the area percent of the product monomer peak compared to all other peaks excluding the included volume peak observed approximately at 12 to 14 minutes. Peaks eluting earlier than the monomer peak are recorded as percent aggregate.

The liquid formulations of the present invention exhibit low to undetectable levels of aggregation as measured by HPSEC or rCGE, that is, no more than 5%, no more than 4%, no more than 3%, no more than 2%, no more than 1%, and most preferably no more than 0.5% aggregate by weight protein, and low to undetectable levels of fragmentation, that is, 80% or higher, 85% or higher, 90% or higher, 95% or higher, 98% or higher, or 99% or higher, or 99.5% or higher of the total peak area in the peak(s) representing intact antibodies

or fragments thereof. In the case of SDS-PAGE, the density or the radioactivity of each band stained or labeled with radioisotope can be measured and the % density or % radioactivity of the band representing non-degraded SYNAGIS® or antigen-binding fragments thereof can be obtained.

5 The stability of the liquid formulations of the present invention can be also assessed by any assays which measures the biological activity SYNAGIS® or an antigen-binding fragment thereof in the formulation. The biological activities of an antibody include, but not limited to, antigen-binding activity, complement-activation activity, Fc-receptor binding activity, and so forth. Antigen-binding activity of SYNAGIS® or an antigen-binding
10 fragment thereof can be measured by any method known to those skilled in the art, including but not limited to ELISA, radioimmunoassay, Western blot, and the like. Complement-activation activity can be measured by a C3a/C4a assay in the system where SYNAGIS® or an antigen-binding fragment thereof is reacted in the presence of the complement components with cells expressing a RSV antigen. *Also see Harlow et al.,*
15 *Antibodies: A Laboratory Manual*, (Cold Spring Harbor Laboratory Press, 2nd ed. 1988).

An ELISA based assay, *e.g.*, may be used
to compare the ability of a liquid formulation of SYNAGIS® or an antigen-binding fragment thereof to immunospecifically bind to a RSV antigen to a SYNAGIS® reference standard.
In this assay, plates are coated with RSV antigen (in particular, the A antigenic site of F
20 protein of RSV) and the binding signal of a set concentration of a SYNAGIS® reference standard is compared to the binding signal of the same concentration of the liquid formulation of SYNAGIS® or an antigen-binding fragment thereof.

The purity of the liquid antibody formulations of the invention may be measured by any method well-known to one of skilled in the art such as, *e.g.*, HPSEC. The sterility of
25 the liquid antibody formulations may be assessed as follows: sterile soybean-casein digest medium and fluid thioglycollate medium are inoculated with a test liquid antibody formulation by filtering the liquid antibody formulation through a sterile filter having a nominal porosity of 0.45 μm . When using the Sterisure™ or Steritest™ method, each filter device is aseptically filled with approximately 100 ml of sterile soybean-casein digest
30 medium or fluid thioglycollate medium. When using the conventional method, the challenged filter is aseptically transferred to 100 ml of sterile soybean-casein digest medium or fluid thioglycollate medium. The media are incubated at appropriate temperatures and observed three times over a 14 day period for evidence of bacterial or fungal growth.

5.5 Prophylactic And Therapeutic Utility Of The Antibody Formulations

The present invention is also directed to antibody-based therapies which involve administering to a subject, preferably a mammal, most preferably a human, the liquid antibody formulations of the present invention for preventing, treating, managing or
5 ameliorating a RSV infection or one or more symptoms thereof. Prophylactic and therapeutic formulations of the invention comprise SYNAGIS® or an antigen-binding fragment thereof at concentrations of from about 15 mg/ml to about 300 mg/ml in a solution containing histidine.

The liquid formulations of the invention may comprise modified SYNAGIS® or
10 antigen-binding fragments thereof that have improved *in vivo* half-lives compared to known antibodies that immunospecifically binds to a RSV antigen (*e.g.*, unmodified SYNAGIS®).

In one embodiment, the liquid formulations of the present invention are administered to a mammal, preferably a human, to prevent, treat, manage or ameliorate a RSV infection or one or more symptoms thereof. In another embodiment, the liquid
15 formulations of the invention are administered to a human with cystic fibrosis, bronchopulmonary dysplasia, congenital heart disease, congenital immunodeficiency or acquired immunodeficiency, or to a human who has had a bone marrow transplant to prevent, treat, manage or ameliorate a RSV infection or one or more symptoms thereof. In another embodiment, the liquid formulations of the invention are administered to a human
20 infant, preferably a human infant born prematurely or a human infant at risk of hospitalization for a RSV infection to prevent, treat, manage or ameliorate a RSV infection or one or more symptoms thereof. In another embodiment, the liquid formulations of the invention are administered to an elderly person to prevent, treat, manage or ameliorate a RSV infection or one or more symptoms thereof. In yet another embodiment, the liquid
25 formulations of the invention are administered to a subject in an institution or group home (*e.g.*, a nursing home or orphanage).

The liquid formulations of the present invention may be used locally or systemically in the body of a subject prophylactically or therapeutically. The formulations of the present invention may also be advantageously utilized in combination with other therapies useful in
30 the prevention, treatment, management or amelioration of a RSV infection (*e.g.*, a prophylactic or a therapeutic agent other than SYNAGIS®). Non-limiting examples of prophylactic or therapeutic agents that can be used in combination with the liquid formulations of the present invention, see Section 5.6, *infra*.

When one or more other therapies are used, they can be administered separately, in any appropriate form and by any suitable route. A liquid formulation of the invention may be administered to a mammal, preferably a human, concurrently with one or more other therapies (*e.g.*, one or more other prophylactic or therapeutic agents) useful for the prevention, treatment, management or amelioration of a RSV infection or one or more symptoms thereof. The term “concurrently” is not limited to the administration of therapies at exactly the same time, but rather it is meant that a liquid formulation of the invention and another therapy are administered to a mammal in a sequence and within a time interval such that SYNAGIS® or an antigen-binding fragment thereof contained in the liquid formulation can act together with the other therapy to provide an increased benefit than if they were administered otherwise. For example, a liquid formulation of the invention and one or more other prophylactic or therapeutic agents useful for prevention, treatment, management or amelioration of a RSV infection may be administered at the same time or sequentially in any order at different points in time; however, if not administered at the same time, they should be administered sufficiently close in time so as to provide the desired therapeutic or prophylactic effect.

In various embodiments, a liquid formulation of the invention and one or more other therapies (*e.g.*, one or more other prophylactic or therapeutic agents) useful for prevention, treatment, management or amelioration of a RSV infection or a symptom thereof are administered less than 1 hour apart, at about 1 hour apart, at about 1 hour to about 2 hours apart, at about 2 hours to about 3 hours apart, at about 3 hours to about 4 hours apart, at about 4 hours to about 5 hours apart, at about 5 hours to about 6 hours apart, at about 6 hours to about 7 hours apart, at about 7 hours to about 8 hours apart, at about 8 hours to about 9 hours apart, at about 9 hours to about 10 hours apart, at about 10 hours to about 11 hours apart, at about 11 hours to about 12 hours apart, no more than 24 hours apart or no more than 48 hours apart. In preferred embodiments, a liquid formulation of the invention and one or more other therapies (*e.g.*, one or more other prophylactic or therapeutic agents) useful for prevention, treatment, management or amelioration of a RSV infection or a symptom thereof are administered within the same patient visit. In other embodiments, a liquid formulation of the invention and one or more other therapies (*e.g.*, one or more other prophylactic or therapeutic agents) useful for prevention, treatment, management or amelioration of a RSV infection or a symptom thereof are administered at about 2 to 4 days apart, at about 4 to 6 days apart, at about 1 week apart, at about 1 to 2 weeks apart, or more than 2 weeks apart. In preferred embodiments, a liquid formulation of the invention and

one or more other prophylactic or therapeutic agents useful for prevention, treatment, management or amelioration of a RSV infection or a symptom thereof are administered in a time frame where both agents are still active. One skilled in the art would be able to determine such a time frame by determining the half-life of the administered agents.

5 In certain embodiments, a liquid formulation of the invention and one or more other therapies (e.g., one or more other prophylactic or therapeutic agents) useful for prevention, treatment, management or amelioration of a RSV infection or a symptom thereof are cyclically administered to a subject. Cycling therapy involves the administration of a first therapy for a period of time, followed by the administration of a second therapy and/or third
10 therapy for a period of time and repeating this sequential administration. Cycling therapy can reduce the development of resistance to one or more of the therapies, avoid or reduce the side effects of one of the therapies, and/or improves the efficacy of the treatment.

 In certain embodiments, a liquid formulation of the invention and one or more other therapies (e.g., one or more other prophylactic or therapeutic agents) useful for prevention,
15 treatment, management or amelioration of a RSV infection or a symptom thereof are administered in a cycle of less than about 3 weeks, about once every two weeks, about once every 10 days or about once every week. One cycle can comprise the administration of a therapy (e.g., a therapeutic or prophylactic agent) by infusion over about 90 minutes every cycle, about 1 hour every cycle, about 45 minutes every cycle. Each cycle can comprise at
20 least 1 week of rest, at least 2 weeks of rest, at least 3 weeks of rest. The number of cycles administered is from about 1 to about 12 cycles, more typically from about 2 to about 10 cycles, and more typically from about 2 to about 8 cycles.

 Generally, administration of products of a species origin or species reactivity (in the case of antibodies) that is the same species as that of the patient is preferred. Thus, in a
25 preferred embodiment, human or humanized antibodies, fragments derivatives, or analogs, are administered to a human patient for therapy or prophylaxis.

5.6 Agents Useful in Combination with SYNAGIS® Formulations

 The present invention provides methods for preventing, managing, treating, or ameliorating a RSV infection or one or more symptoms thereof comprising administering to
30 a subject in need thereof a liquid formulation of the invention alone or in combination with one or more therapies (e.g., one or more prophylactic or therapeutic agents) other than SYNAGIS®. The present invention provides methods for preventing, treating, managing or ameliorating a RSV infection or one or more symptoms thereof comprising administering to

a subject in need thereof a liquid formulation of the invention alone or in combination with one or more therapies (*e.g.*, one or more prophylactic or therapeutic agents) other than SYNAGIS®. The present invention also provides compositions comprising a liquid formulation of SYNAGIS® or an antigen-binding fragment thereof and one or more
 5 prophylactic or therapeutic agents other than SYNAGIS® and methods of preventing, treating, managing or ameliorating a RSV infection or one or more symptoms thereof utilizing said compositions. Therapeutic or prophylactic agents include, but are not limited to, small molecules, synthetic drugs, peptides, polypeptides, proteins, nucleic acids (*e.g.*, DNA and RNA nucleotides including, but not limited to, antisense nucleotide sequences,
 10 triple helices, RNA interference (RNAi), and nucleotide sequences encoding biologically active proteins, polypeptides or peptides) antibodies, synthetic or natural inorganic molecules, mimetic agents, and synthetic or natural organic molecules.

Any therapy which is known to be useful, or which has been used or is currently being used for the prevention, management, treatment, or amelioration of a RSV infection
 15 or one or more symptoms thereof can be used in combination with a liquid formulation in accordance with the invention described herein. See, *e.g.*, Gilman *et al.*, *Goodman and Gilman's: The Pharmacological Basis of Therapeutics*, 10th ed., McGraw-Hill, New York, 2001; *The Merck Manual of Diagnosis and Therapy*, Berkow, M.D. *et al.* (eds.), 17th Ed., Merck Sharp & Dohme Research Laboratories, Rahway, NJ, 1999; *Cecil Textbook of*
 20 *Medicine*, 20th Ed., Bennett and Plum (eds.), W.B. Saunders, Philadelphia, 1996, for information regarding therapies (*e.g.*, prophylactic or therapeutic agents) which have been or are currently being used for preventing, treating, managing, or ameliorating a RSV infection or one or more symptoms thereof. Examples of such agents include, but are not limited to, immunomodulatory agents, anti-inflammatory agents (*e.g.*, adrenocorticoids,
 25 corticosteroids (*e.g.*, beclomethasone, budesonide, flunisolide, fluticasone, triamcinolone, methylprednisolone, prednisolone, prednisone, hydrocortisone), glucocorticoids, steroids, non-steroidal anti-inflammatory drugs (*e.g.*, aspirin, ibuprofen, diclofenac, and COX-2 inhibitors), pain relievers, leukotriene antagonists (*e.g.*, montelukast, methyl xanthines, zafirlukast, and zileuton), beta2-agonists (*e.g.*, albuterol, biterol, fenoterol, isoetharie,
 30 metaproterenol, pirbuterol, salbutamol, terbutalin formoterol, salmeterol, and salbutamol terbutaline), anticholinergic agents (*e.g.*, ipratropium bromide and oxitropium bromide), sulphasalazine, penicillamine, dapsone, antihistamines, anti-malarial agents (*e.g.*, hydroxychloroquine)), and anti-viral agents.

In specific embodiments, a liquid formulation of the invention is used in combination with a monoclonal or chimeric antibody, or with a lymphokine or hematopoietic growth factor (such as, *e.g.*, IL-2, IL-3, IL-4, IL-7, IL-9, IL-10, IL-12, and interferon α , β , and γ), which, for example, serves to increase the number or activity of effector cells which interact with the antibody. A liquid formulation of the present invention may also be advantageously utilized in combination with other monoclonal or chimeric antibodies, or with lymphokines or hematopoietic growth factors (such as, *e.g.*, IL-2, IL-3, IL-4, IL-7, IL-9, IL-10, IL-12, and interferon α , β , and γ), which, for example, serve to increase the immune response. The liquid formulations of the present invention may also be advantageously utilized in combination with one or more drugs used to treat RSV infection such as, for example anti-viral agents. The liquid formulations of the present invention may be used in combination with one or more of the following drugs: NIH-351 (Gemini Technologies), recombinant RSV vaccine (MedImmune Vaccines, Inc. U.S. Application No. 60/358,934 filed February 21, 2002, U.S. Patent Application No. 2004-0005545 filed February 21, 2003, U.S. Patent No. 7,449,324, U.S. Patent No. 7,531,342, U.S. Patent Application Nos. 2003-0232061 filed February 21, 2003, 60/466,181 filed April 25, 2003 and 60/465,811 filed April 25, 2003), RSVf-2 (Intracel), F-50042 (Pierre Fabre), T-786 (Trimeris), VP-36676 (ViroPharma), RFI-641 (American Home Products), VP-14637 (ViroPharma), PFP-1 and PFP-2 (American Home Products), RSV vaccine (Avant Immunotherapeutics), and F-50077 (Pierre Fabre).

5.6.1 Immunomodulatory Agents

Any immunomodulatory agent well-known to one of skilled in the art may be used in accordance with the methods of the invention to prevent, treat, manage or ameliorate a RSV infection or one or more symptoms thereof. Immunomodulatory agents can affect one or more or all aspects of the immune response in a subject. Aspects of the immune response include, but are not limited to, the inflammatory response, the complement cascade, leukocyte and lymphocyte differentiation, proliferation, and/or effector function, monocyte and/or basophil counts, and the cellular communication among cells of the immune system. In certain embodiments of the invention, an immunomodulatory agent modulates one aspect of the immune response. In other embodiments, an immunomodulatory agent modulates more than one aspect of the immune response. In a preferred embodiment of the invention, the administration of an immunomodulatory agent to a subject inhibits or reduces one or more aspects of the subject's immune response capabilities. In an alternative embodiment

of the invention, the immunomodulatory agent enhances one or more aspects of a subject's immune response. In certain embodiments, an immunomodulatory agent is not an anti-inflammatory agent. In a specific embodiment, an immunomodulatory agent is an agent other than a chemotherapeutic agent.

5 Examples of immunomodulatory agents include, but are not limited to, proteinaceous agents such as cytokines, peptide mimetics, and antibodies (*e.g.*, human, humanized, chimeric, monoclonal, polyclonal, Fvs, ScFvs, Fab or F(ab)₂ fragments or epitope binding fragments), nucleic acid molecules (*e.g.*, antisense nucleic acid molecules and triple helices), small molecules, organic compounds, and inorganic compounds. In particular, immunomodulatory agents include, but are not limited to, methotrexate, 10 leflunomide, cyclophosphamide, cytoxan, Immuran, cyclosporine A, minocycline, azathioprine, antibiotics (*e.g.*, FK506 (tacrolimus)), methylprednisolone (MP), corticosteroids, steroids, mycophenolate mofetil, rapamycin (sirolimus), mizoribine, deoxyspergualin, brequinar, malononitriloamindes (*e.g.*, leflunamide), T cell receptor 15 modulators, and cytokine receptor modulators.

 Examples of T cell receptor modulators include, but are not limited to, anti-T cell receptor antibodies (*e.g.*, anti-CD4 antibodies (*e.g.*, cM-T412 (Boeringer), IDEC-CE9.1® (IDEC and SKB), mAB 4162W94, Orthoclone and OKTcdr4a (Janssen-Cilag)), anti-CD3 antibodies (*e.g.*, Nuvion (Product Design Labs), OKT3 (Johnson & Johnson), or Rituxan 20 (IDEC)), anti-CD5 antibodies (*e.g.*, an anti-CD5 ricin-linked immunoconjugate), anti-CD7 antibodies (*e.g.*, CHH-380 (Novartis)), anti-CD8 antibodies, anti-CD40 ligand monoclonal antibodies (*e.g.*, IDEC-131 (IDEC)), anti-CD52 antibodies (*e.g.*, CAMPATH 1H (Ilex)), anti-CD2 antibodies (*e.g.*, MEDI-507 (MedImmune, Inc., International Publication Nos. WO 02/098370 and WO 02/069904), anti-CD11a antibodies (*e.g.*, Xanelim (Genentech)), 25 and anti-B7 antibodies (*e.g.*, IDEC-114 (IDEC))), CTLA4-immunoglobulin, and LFA-3TIP (Biogen, International Publication No. WO 93/08656 and U.S. Patent No. 6,162,432).

 Examples of cytokine receptor modulators include, but are not limited to, soluble cytokine receptors (*e.g.*, the extracellular domain of a TNF- α receptor or an antigen-binding fragment thereof, the extracellular domain of an IL-1 β receptor or an antigen-binding 30 fragment thereof, and the extracellular domain of an IL-6 receptor or an antigen-binding fragment thereof), cytokines or fragments thereof (*e.g.*, interleukin IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, IL-8, IL-9, IL-10, IL-11, IL-12, IL-13, IL-15, IL-23, TNF- α , TNF- β , interferon (IFN)- α , IFN- β , IFN- γ , and GM-CSF), anti-cytokine receptor antibodies (*e.g.*, anti-IFN receptor antibodies, anti-IL-2 receptor antibodies (*e.g.*, Zenapax (Protein Design Labs)),

anti-IL-3 receptor antibodies, anti-IL-4 receptor antibodies, anti-IL-6 receptor antibodies, anti-IL-9 receptor antibodies, anti-IL-10 receptor antibodies, anti-IL-12 receptor antibodies, anti-IL-13 receptor antibodies, anti-IL-15 receptor antibodies, and anti-IL-23 receptor antibodies), anti-cytokine antibodies (*e.g.*, anti-IFN antibodies, anti-TNF- α antibodies, anti-IL-1 β antibodies, anti-IL-3 antibodies, anti-IL-6 antibodies, anti-IL-8 antibodies (*e.g.*, ABX-IL-8 (Abgenix)), anti-IL-9 antibodies, anti-IL-12 antibodies, anti-IL-13 antibodies, anti-IL-15 antibodies, and anti-IL-23 antibodies).

In a specific embodiment, a cytokine receptor modulator is IFN, IL-2, IL-3, IL-4, IL-10, IL-12 or an antigen-binding fragment thereof. In another embodiment, a cytokine receptor modulator is an anti-IL-1 β antibody, anti-IL-6 antibody, anti-IL-9 antibody, anti-IL-12 receptor antibody, or anti-TNF- α antibody. In another embodiment, a cytokine receptor modulator is the extracellular domain of a TNF- α receptor or an antigen-binding fragment thereof.

An immunomodulatory agent may be selected to interfere with the interactions between the T helper subsets (TH1 or TH2) and B cells to inhibit neutralizing antibody formation. Antibodies that interfere with or block the interactions necessary for the activation of B cells by TH (T helper) cells, and thus block the production of neutralizing antibodies, are useful as immunomodulatory agents in the methods of the invention. For example, B cell activation by T cells requires certain interactions to occur (Durie *et al.*, Immunol. Today, 15(9):406-410 (1994)), such as the binding of CD40 ligand on the T helper cell to the CD40 antigen on the B cell, and the binding of the CD28 and/or CTLA4 ligands on the T cell to the B7 antigen on the B cell. Without both interactions, the B cell cannot be activated to induce production of the neutralizing antibody.

The CD40 ligand (CD40L)-CD40 interaction is a desirable point to block the immune response because of its broad activity in both T helper cell activation and function as well as the absence of redundancy in its signaling pathway. Thus, in a specific embodiment of the invention, the interaction of CD40L with CD40 is transiently blocked at the time of administration of one or more of the immunomodulatory agents. This can be accomplished by treating with an agent which blocks the CD40 ligand on the TH cell and interferes with the normal binding of CD40 ligand on the T helper cell with the CD40 antigen on the B cell. An antibody to CD40 ligand (anti-CD40L) (available from Bristol-Myers Squibb Co; see, *e.g.*, European patent application 555,880, published Aug. 18, 1993) or a soluble CD40 molecule can be selected and used as an immunomodulatory agent in accordance with the methods of the invention.

An immunomodulatory agent may be selected to inhibit the interaction between TH1 cells and cytotoxic T lymphocytes ("CTLs") to reduce the occurrence of CTL-mediated killing. An immunomodulatory agent may be selected to alter (*e.g.*, inhibit or suppress) the proliferation, differentiation, activity and/or function of the CD4⁺ and/or CD8⁺ T cells. For
5 example, antibodies specific for T cells can be used as immunomodulatory agents to deplete, or alter the proliferation, differentiation, activity and/or function of CD4⁺ and/or CD8⁺ T cells.

In one embodiment, an immunomodulatory agent which reduces or inhibits one or more biological activities (*e.g.*, the differentiation, proliferation, and/or effector functions)
10 of TH0, TH1, and/or TH2 subsets of CD4⁺ T helper cells is administered to a subject with a RSV infection in accordance with the methods of the invention. One example of such an immunomodulatory agent is IL-4. IL-4 enhances antigen-specific activity of TH2 cells at the expense of the TH1 cell function (see, *e.g.*, Yokota *et al*, 1986 Proc. Natl. Acad. Sci., USA, 83:5894-5898; and U.S. Pat. No. 5,017,691). Other examples of immunomodulatory
15 agents that affect the biological activity (*e.g.*, proliferation, differentiation, and/or effector functions) of T-helper cells (in particular, TH1 and/or TH2 cells) include, but are not limited to, IL-2, IL-4, IL-5, IL-6, IL-10, IL-12, IL-13, IL-15, IL-23, and interferon (IFN)- γ .

In another embodiment, an immunomodulatory agent administered to a subject with a RSV infection in accordance with the methods of the invention is a cytokine that prevents
20 antigen presentation. In a specific embodiment, an immunomodulatory agent used in the methods of the invention is IL-10. IL-10 also reduces or inhibits macrophage action which involves bacterial elimination.

In accordance with the invention, one or more immunomodulatory agents are administered to a subject with a RSV infection prior to, subsequent to, or concomitantly
25 with a liquid formulation of SYNAGIS[®] or an antigen-binding fragment thereof. Preferably, one or more immunomodulatory agents are administered in combination with a liquid formulation of SYNAGIS[®] or an antigen-binding fragment thereof to a subject with a RSV infection to reduce or inhibit one or more aspects of the immune response as deemed necessary by one of skilled in the art. Any technique well-known to one skilled in the art
30 can be used to measure one or more aspects of the immune response in a particular subject, and thereby determine when it is necessary to administer an immunomodulatory agent to said subject. In a preferred embodiment, a mean absolute lymphocyte count of approximately 500 cells/mm³, preferably 600 cells/mm³, 650 cells/mm³, 700 cells/mm³, 750 cells/mm³, 800 cells/mm³, 900 cells/mm³, 1000 cells/mm³, 1100 cells/mm³, or 1200

cells/mm³ is maintained in a subject. In another preferred embodiment, a subject with a RSV infection is not administered an immunomodulatory agent if their absolute lymphocyte count is 500 cells/mm³ or less, 550 cells/mm³ or less, 600 cells/mm³ or less, 650 cells/mm³ or less, 700 cells/mm³ or less, 750 cells/mm³ or less, or 800 cells/mm³ or less.

5 In a specific embodiment, one or more immunomodulatory agents are administered in combination with a liquid formulation of SYNAGIS[®] or an antigen-binding fragment thereof to a subject with a RSV infection so as to transiently reduce or inhibit one or more aspects of the immune response. Such a transient inhibition or reduction of one or more aspects of the immune system can last for hours, days, weeks, or months. Preferably, the
10 transient inhibition or reduction in one or more aspects of the immune response lasts for a few hours (*e.g.*, 2 hours, 4 hours, 6 hours, 8 hours, 12 hours, 14 hours, 16 hours, 18 hours, 24 hours, 36 hours, or 48 hours), a few days (*e.g.*, 3 days, 4 days, 5 days, 6 days, 7 days, or 14 days), or a few weeks (*e.g.*, 3 weeks, 4 weeks, 5 weeks or 6 weeks). The transient reduction or inhibition of one or more aspects of the immune response enhances the
15 prophylactic and/or therapeutic effect(s) of a liquid formulation of SYNAGIS[®] or an antigen-binding fragment thereof.

In a preferred embodiment, proteins, polypeptides or peptides (including antibodies) that are utilized as immunomodulatory agents are derived from the same species as the recipient of the proteins, polypeptides or peptides so as to reduce the likelihood of an
20 immune response to those proteins, polypeptides or peptides. In another preferred embodiment, when the subject is a human, the proteins, polypeptides, or peptides that are utilized as immunomodulatory agents are human or humanized.

Nucleic acid molecules encoding proteins, polypeptides, or peptides with immunomodulatory activity or proteins, polypeptides, or peptides with immunomodulatory
25 activity can be administered to a subject with a RSV infection in accordance with the methods of the invention. Further, nucleic acid molecules encoding derivatives, analogs, or fragments of proteins, polypeptides, or peptides with immunomodulatory activity, or derivatives, analogs, or fragments of proteins, polypeptides, or peptides with immunomodulatory activity can be administered to a subject with a RSV infection in
30 accordance with the methods of the invention. Preferably, such derivatives, analogs, and fragments retain the immunomodulatory activity of the full-length, wild-type protein, polypeptide, or peptide.

Preferably, agents that are commercially available and known to function as immunomodulatory agents are used in the methods of the invention. The

immunomodulatory activity of an agent can be determined *in vitro* and/or *in vivo* by any technique well-known to one skilled in the art, including, *e.g.*, by CTL assays, proliferation assays, and immunoassays (*e.g.* ELISAs) for the expression of particular proteins such as co-stimulatory molecules and cytokines.

5.6.2 Anti-inflammatory Agents

Any anti-inflammatory agent, including agents useful in therapies for inflammatory disorders, well-known to one of skilled in the art can be used in accordance with methods of the invention to prevent, treat, manage, or ameliorate a RSV infection or one or more symptoms thereof. Non-limiting examples of anti-inflammatory agents include non-steroidal anti-inflammatory drugs (NSAIDs), steroidal anti-inflammatory drugs, anticholinergics (*e.g.*, atropine sulfate, atropine methylnitrate, and ipratropium bromide (ATROVENT™)), beta2-agonists (*e.g.*, albuterol (VENTOLIN™ and PROVENTIL™), bitolterol (TORNALATE™), levalbuterol (XOPONEX™), metaproterenol (ALUPENT™), pirbuterol (MAXAIR™), terbutaline (BRETHAIRE™ and BRETHINE™), albuterol (PROVENTIL™, REPETABS™, and VOLMAX™), formoterol (FORADIL AEROLIZER™), and salmeterol (SEREVENT™ and SEREVENT DISKUS™)), and methylxanthines (*e.g.*, theophylline (UNIPHYL™, THEO-DUR™, SLO-BID™, and TEHO-42™)). Examples of NSAIDs include, but are not limited to, aspirin, ibuprofen, celecoxib (CELEBREX™), diclofenac (VOLTAREN™), etodolac (LODINE™), fenoprofen (NALFON™), indomethacin (INDOCIN™), ketoralac (TORADOL™), oxaprozin (DAYPRO™), nabumentone (RELAFEN™), sulindac (CLINORIL™), tolmetin (TOLECTIN™), rofecoxib (VIOXX™), naproxen (ALEVE™, NAPROSYN™), ketoprofen (ACTRON™) and nabumetone (RELAFEN™). Such NSAIDs function by inhibiting a cyclooxygenase enzyme (*e.g.*, COX-1 and/or COX-2). Examples of steroidal anti-inflammatory drugs include, but are not limited to, glucocorticoids, dexamethasone (DECADRON™), corticosteroids (*e.g.*, methylprednisolone (MEDROL™)), cortisone, hydrocortisone, prednisone (PREDNISON™ and DELTASONE™), prednisolone (PRELONE™ and PEDIAPRED™), triamcinolone, azulfidine, and inhibitors of eicosanoids (*e.g.*, prostaglandins, thromboxanes, and leukotrienes (see Table 2, *infra*, for non-limiting examples of leukotriene and typical dosages of such agents)).

5.6.3 Anti-Viral Agents

Any anti-viral agent well-known to one of skilled in the art (in particular, one useful for the treatment, prevention, management, or amelioration of a RSV infection) can be used

in accordance with the methods of the invention to prevent, treat, manage or ameliorate a RSV infection or one or more symptoms thereof. Non-limiting examples of anti-viral agents include proteins, polypeptides, peptides, fusion proteins antibodies, nucleic acid molecules, organic molecules, inorganic molecules, and small molecules that inhibit and/or
5 reduce the attachment of a virus to its receptor, the internalization of a virus into a cell, the replication of a virus, or release of virus from a cell. In particular, anti-viral agents include, but are not limited to, nucleoside analogs (*e.g.*, zidovudine, acyclovir, gangcyclovir, vidarabine, idoxuridine, trifluridine, and ribavirin), foscarnet, amantadine, rimantadine, saquinavir, indinavir, ritonavir, alpha-interferons and other interferons, and AZT.

10 In specific embodiments, the anti-viral agent is an antibody agent other than SYNAGIS[®] that is immunospecific for a viral antigen. As used herein, the term "viral antigen" includes, but is not limited to, any RSV peptide, polypeptide and protein (*e.g.*, RSV F glycoprotein and RSV G glycoprotein) that is capable of eliciting an immune response.

15 In preferred embodiments, the viral infection is RSV and the anti-viral antigen is an antibody other than SYNAGIS[®] that immunospecifically binds to an antigen of RSV. In certain embodiments, the anti-RSV-antigen antibody immunospecifically binds to a RSV antigen of the Group A of RSV. In other embodiments, the anti-RSV-antigen antibody immunospecifically binds to a RSV antigen of the Group B of RSV. In other embodiments,
20 the anti-RSV antigen antibody immunospecifically binds to an antigen of RSV of one Group and cross reacts with the analogous antigen of the other Group. In particular embodiments, the anti-RSV-antigen antibody immunospecifically binds to a RSV nucleoprotein, RSV phosphoprotein, RSV matrix protein, RSV small hydrophobic protein, RSV RNA-dependent RNA polymerase, RSV F protein, and/or RSV G protein. In
25 additional specific embodiments, the anti-RSV-antigen antibody binds to allelic variants of a RSV nucleoprotein, a RSV nucleocapsid protein, a RSV phosphoprotein, a RSV matrix protein, a RSV attachment glycoprotein, a RSV fusion glycoprotein, a RSV nucleocapsid protein, a RSV matrix protein, a RSV small hydrophobic protein, a RSV RNA-dependent RNA polymerase, a RSV F protein, a RSV L protein, a RSV P protein, and/or a RSV G
30 protein.

Anti-viral therapies and their dosages, routes of administration and recommended usage are known in the art and have been described in such literature as the *Physician's Desk Reference* (57th ed., 2003). Additional information on respiratory viral infections is available in *Cecil Textbook of Medicine* (18th ed., 1988).

5.7 Methods of Administering the SYNAGIS® Formulations

The invention provides methods of treatment, prophylaxis, and amelioration of a RSV infection or one or more symptoms thereof by administering to a subject of an effective amount of liquid formulations of the invention. The subject is preferably a mammal such as non-primate (*e.g.*, cows, pigs, horses, cats, dogs, rats etc.) and a primate (*e.g.*, monkey such as a cynomolgous monkey and a human). In a preferred embodiment, the subject is a human. In another preferred embodiment, the subject is a human infant or a human infant born prematurely.

Various delivery systems are known and can be used to administer a liquid formulation of the present invention. Methods of administering SYNAGIS® liquid formulations of the present invention include, but are not limited to, parenteral administration (*e.g.*, intradermal, intramuscular, intraperitoneal, intravenous and subcutaneous), epidural administration, topical administration, pulmonary administration, and mucosal administration (*e.g.*, intranasal and oral routes). In a specific embodiment, liquid formulations of the present invention are administered intramuscularly, intravenously, or subcutaneously and, preferably, intramuscularly. The formulations may be administered by any convenient route, for example by infusion or bolus injection, by absorption through epithelial or mucocutaneous linings (*e.g.*, oral mucosa, rectal and intestinal mucosa, etc.) and may be administered together with other biologically active agents. Administration can be systemic or local. In addition, pulmonary administration can be employed, *e.g.*, by use of an inhaler or nebulizer.

The invention also provides that a liquid formulation of the present invention is packaged in a hermetically sealed container such as an ampoule or sachette indicating the quantity of the SYNAGIS® or antigen-binding fragments thereof. Preferably, the liquid formulations of the present invention are in a hermetically sealed container indicating the quantity and concentration of the antibody or antibody fragment. Preferably, the liquid formulation of the present invention is supplied in a hermetically sealed container at least 15 mg/ml, 20 mg/ml, 30 mg/ml, 40 mg/ml, 50 mg/ml, 60 mg/ml, 70 mg/ml, 80 mg/ml, 90 mg/ml, 100 mg/ml, 150 mg/ml, 200 mg/ml, 250 mg/ml, or 300 mg/ml and, most preferably, 105 mg/ml, in a quantity of 1 ml, 2 ml, 3 ml, 4 ml, 5 ml, 6 ml, 7 ml, 8 ml, 9 ml, 10 ml, 15 ml, or 20 ml and, most preferably, 1.2 ml.

The amount of the liquid formulations of the present invention which will be effective in the prevention, treatment, management or amelioration of a RSV infection or one or more symptoms thereof can be determined by standard clinical techniques. For

example, the dosage of the composition which will be effective in the treatment, prevention or amelioration of symptoms associated with a RSV infection can be determined by administering the formulation to a cotton rat, measuring the RSV titer after challenging the cotton rat with 10^5 pfu of RSV and comparing the RSV titer to that obtain for a cotton rat not administered the formulation. Accordingly, a dosage that results in a 2 log decrease or a 99% reduction in RSV titer in the cotton rat challenged with 10^5 pfu of RSV relative to the cotton rat challenged with 10^5 pfu of RSV but not administered the formulation is the dosage of the formulation that can be administered to a human for prevention, treatment, management or amelioration of a RSV infection or one or more symptoms thereof. The dosage of the formulation which will be effective in prevention, treatment, management or amelioration of a RSV infection or one or more symptoms thereof can be determined by administering the formulation to an animal model (*e.g.*, a cotton rat or monkey) and measuring the serum titer of SYNAGIS[®] or antigen-binding fragments thereof. Accordingly, a dosage of the formulation that results in a serum titer of at least 1 $\mu\text{g/ml}$, preferably 2 $\mu\text{g/ml}$, 5 $\mu\text{g/ml}$, 10 $\mu\text{g/ml}$, 20 $\mu\text{g/ml}$, 25 $\mu\text{g/ml}$, at least 35 $\mu\text{g/ml}$, at least 40 $\mu\text{g/ml}$, at least 50 $\mu\text{g/ml}$, at least 75 $\mu\text{g/ml}$, at least 100 $\mu\text{g/ml}$, at least 125 $\mu\text{g/ml}$, at least 150 $\mu\text{g/ml}$, at least 200 $\mu\text{g/ml}$, at least 250 $\mu\text{g/ml}$, at least 300 $\mu\text{g/ml}$, at least 350 $\mu\text{g/ml}$, at least 400 $\mu\text{g/ml}$, or at least 450 $\mu\text{g/ml}$ can be administered to a human for prevention, treatment, management or amelioration of a RSV infection or one or more symptoms thereof. In addition, *in vitro* assays may optionally be employed to help identify optimal dosage ranges.

The precise dose to be employed in the formulation will also depend on the route of administration, and the seriousness of the RSV infection, and should be decided according to the judgment of the practitioner and each patient's circumstances. Effective doses may be extrapolated from dose-response curves derived from *in vitro* or animal model (*e.g.*, the cotton rat or Cynomolgous monkey) test systems.

For antibodies (*e.g.*, SYNAGIS[®]), proteins, polypeptides, peptides and fusion proteins, the dosage administered to a patient is typically about 1 mg/kg to 30 mg/kg of the patient's body weight. Preferably, the dosage administered to a patient is between 10 mg/kg and 20 mg/kg of the patient's body weight, more preferably 15 mg/kg of the patient's body weight. Generally, human antibodies have a longer half-life within the human body than antibodies from other species due to the immune response to the foreign polypeptides. Thus, lower dosages of human antibodies and less frequent administration is often possible. Further, the dosage, volume and frequency of administration of liquid formulations of the

present invention may be reduced by increasing the concentration of SYNAGIS[®] or an antigen-binding fragment thereof in the formulations, increasing affinity and/or avidity of SYNAGIS[®] or an antigen-binding fragment thereof, and/or increasing the half-life of SYNAGIS[®] or an antigen-binding fragment thereof.

5 Exemplary doses of a small molecule include milligram or microgram amounts of the small molecule per kilogram of subject or sample weight (*e.g.*, about 1 microgram per kilogram to about 500 milligrams per kilogram, about 100 micrograms per kilogram to about 5 milligrams per kilogram, or about 1 microgram per kilogram to about 50 micrograms per kilogram).

10 In a specific embodiment, a mammal, preferably a human, is administered a stable liquid formulation of the present invention for the prevention, treatment, management or amelioration of a RSV infection or one or more symptoms thereof in an amount effective for decreasing RSV titers. In accordance with this embodiment, an effective amount of the liquid formulations of the present invention reduces the RSV titers in the lung as measured,
15 for example, by the concentration of RSV in sputum samples or a lavage from the lungs from a mammal. In another embodiment, a mammal, preferably a human, is administered a liquid formulation of the present invention for the prevention, treatment, management or amelioration of a RSV infection or one or more symptoms thereof in an amount effective for inducing an immune response in the mammal.

20 In another embodiment, a mammal, preferably a human, is administered a first dose of a liquid formulation of the present invention comprising 30 mg/kg or less, 15 mg/kg or less, 10 mg/kg or less, 5 mg/kg or less, 3 mg/kg or less, 1 mg/kg or less, or 0.5 mg/kg or less of SYNAGIS[®] or an antigen-binding fragment thereof for the prevention, treatment, management or amelioration of a RSV infection or one or more symptoms thereof in an
25 amount effective to induce a serum titer of at least 1 µg/ml, preferably at least 2 µg/ml, at least 5 µg/ml, at least 10 µg/ml, at least 15 µg/ml, at least 20 µg/ml, at least 25 µg/ml, at least 30 µg/ml, at least 35 µg/ml, at least 40 µg/ml 20 days (preferably 25, 30, 35, 40 days) after the administration of the first dose and prior to the administration of a subsequent dose. In a specific embodiment, a liquid formulation of the present invention comprises
30 SYNAGIS[®] or an antigen-binding fragment thereof and is administered to a subject a first dose of about 1 mg/kg to about 30 mg/kg to induce a serum titer of about 40 µg/ml or higher 30 days after the administration of the first dose and prior to the administration of a subsequent dose. Preferably, the serum titer of said SYNAGIS[®] or an antigen-binding

fragment thereof is less than 50 µg/ml 30 days after the administration of the first dose and prior to the administration of a subsequent dose.

In another embodiment, a mammal, preferably a human, is administered a first dose of a liquid formulations of the present invention comprising 30 mg/kg or less, 15 mg/kg or less, 10 mg/kg or less, 5 mg/kg or less, 3 mg/kg or less, 1 mg/kg or less or 0.5 mg/kg or less of SYNAGIS[®] or an antigen-binding fragment thereof for the prevention, treatment, management or amelioration of a RSV infection or one or more symptoms thereof in an amount effective to induce a serum titer of at least 1 µg/ml, preferably at least 2 µg/ml, at least 5 µg/ml, at least 10 µg/ml, at least 15 µg/ml, at least 20 µg/ml, or at least 25 µg/ml 20 days (preferably 25, 30, 35, 40 days) after the administration of the first dose and prior to the administration of subsequent dose. Preferably, the serum titer of said SYNAGIS[®] or an antigen-binding fragment thereof is less than 30 µg/ml 30 days after the administration of the first dose and prior to the administration of a subsequent dose.

In another embodiment, a mammal, preferably a human, is administered a first dose of a liquid formulation of the present invention comprising 30 mg/kg or less, 15 mg/kg or less, 5 mg/kg or less, 3 mg/kg or less, 1 mg/kg or less or 0.5 mg/kg or less of a modified form of SYNAGIS[®] or an antigen-binding fragment thereof for the prevention, treatment, management or amelioration of a RSV infection or one or more symptoms thereof which has an increased *in vivo* half-life in an amount effective to induce a serum titer of at least 1 µg/ml, preferably at least 2 µg/ml, at least 5 µg/ml, at least 10 µg/ml, at least 15 µg/ml, at least 20 µg/ml, or at least 25 µg/ml 25 days (preferably 30, 35, or 40 days) after the administration of the first dose and prior to the administration of a subsequent dose. Preferably, the serum titer of said SYNAGIS[®] or an antigen-binding fragment thereof is less than 30 µg/ml 30 days after the administration of the first dose and prior to the administration of a subsequent dose.

In another embodiment, a mammal, preferably a human, is administered a first dose of a liquid formulation of the present invention comprising 30 mg/kg or less, 15 mg/kg or less, 5 mg/kg or less, 3 mg/kg or less, 1 mg/kg or less, or 0.5 mg/kg or less of a modified SYNAGIS[®] or an antigen-binding fragment thereof for the prevention, treatment, management or amelioration of a RSV infection or one or more symptoms thereof which has an increased *in vivo* half-life in an amount effective to induce a serum titer of at least 1 µg/ml, preferably at least 2 µg/ml, at least 5 µg/ml, at least 10 µg/ml, at least 15 µg/ml, at least 20 µg/ml, or at least 25 µg/ml 25 days (preferably 30, 35, or 40 days) after the administration of the first dose and prior to the administration of a subsequent dose.

Preferably, the serum titer of said SYNAGIS[®] or an antigen-binding fragment thereof is less than 30 µg/ml 30 days after the administration of the first dose and prior to the administration of a subsequent dose.

5 In another embodiment, a mammal, preferably a human, is administered a first dose of the liquid formulation comprising approximately 30 mg/kg or less, 15 mg/kg or less (preferably 10 mg/kg or less, 5 mg/kg or less, 3 mg/kg or less, 1 mg/kg or less, or 0.5 mg/kg or less) of a modified form of SYNAGIS[®] or an antigen-binding fragment thereof which has an increased *in vivo* half-life for the prevention, treatment, management or amelioration of a RSV infection or one or more symptoms thereof in an amount effective to induce a serum
10 titer of at least 30 µg/ml, preferably at least 35 µg/ml, at least 40 µg/ml, or at least 50 µg/ml 25 days (preferably 30, 35, or 40 days) after the administration of the first dose and prior to the administration of a subsequent dose.

In one embodiment, a mammal, preferably a human, is administered a first dose of a liquid formulation of the present invention for pulmonary delivery comprising 30 mg/kg or
15 less, 15 mg/kg or less, 5 mg/kg or less, 3 mg/kg or less, 1 mg/kg or less, 0.5 mg/kg or less, or 0.01 mg/kg or less of SYNAGIS[®] or an antigen-binding fragment thereof for the prevention, treatment, management or amelioration of a RSV infection or one or more symptoms thereof in an amount effective to induce a titer of at least 20 ng per mg of lung protein (preferably at least 40 ng/mg, at least 60 ng/mg, at least 80 ng/mg, at least 50 ng/mg,
20 at least 75 ng/mg, at least 100 ng/mg, or at least 150 ng/mg) in an intubation sample or lavage from the lungs of said mammal 20 days (preferably 25, 30, 35, or 40 days) after the administration of the first dose and prior to the administration of a subsequent dose. Preferably, the serum titer of said SYNAGIS[®] or an antigen-binding fragment thereof is less than 100 ng/ml of protein 30 days after the administration of the first dose and prior to the
25 administration of a subsequent dose.

In another embodiment, a mammal, preferably a human, is administered a first dose of a liquid formulation of the present invention 10 mg/kg or less, 5 mg/kg or less, 3 mg/kg or less, 1 mg/kg or less, or 0.5 mg/kg or less of SYNAGIS[®] or an antigen-binding fragment thereof for the prevention, treatment, management or amelioration of a RSV infection or
30 one or more symptoms thereof in an amount effective to induce a serum titer of at least 35 µg/ml, at least 40 µg/ml, at least 50 µg/ml, at least 80 µg/ml, at least 100 µg/ml, at least 120 µg/ml, at least 150 µg/ml, at least 200 µg/ml, at least 250 µg/ml, or at least 300 µg/ml 20 days (preferably 25, 30, 35 or 40 days) after the administration of the first dose. In another embodiment, a mammal, preferably a human, is administered a first dose of a liquid

formulation of the present invention comprising approximately 15 mg/kg of SYNAGIS[®] or an antigen-binding fragment thereof for the prevention, treatment, management or amelioration of a RSV infection or one or more symptoms thereof in an amount effective to induce a serum titer of at least 100 µg/ml, at least 125 µg/ml, at least 150 µg/ml, at least 200 µg/ml, at least 250 µg/ml, at least 300 µg/ml, at least 350 µg/ml, at least 400 µg/ml, or at least 450 µg/ml 20 days (preferably 25, 30, 35 or 40 days) after the administration of the first dose. The term "approximately 15 mg/kg" as used herein refers to a range of between 14 mg/kg and 16 mg/kg.

In another embodiment, a mammal, preferably a human, is administered a dose of a liquid formulation of the present invention comprising SYNAGIS[®] or an antigen-binding fragment thereof for the prevention of a RSV infection or a symptom thereof in an amount effective to induce a prophylactically effective serum titer of less than 10 µg/ml, less than 8 µg/ml, less than 5 µg/ml, less than 3 µg/ml, less than 1 µg/ml, or less than 0.5 µg/ml 30 days after the administration of the dose, wherein said prophylactically effective serum titer is the serum titer that reduces the incidence of RSV infection in the human or the serum titer in a cotton rat that results in a RSV titer 5 days after challenge with 10⁵ pfu RSV that is 99% lower than the RSV titer in the cotton rat 5 days after challenge with 10⁵ pfu of RSV in a cotton rat not administered the dose prior to challenge. Preferably, the dose of the therapeutic or pharmaceutical composition comprises 10 mg/kg or less, 5 mg/kg or less, 3 mg/kg or less, 1 mg/kg or less, or 0.5 mg/kg or less of SYNAGIS[®] or an antigen-binding fragment thereof.

In yet another embodiment, a mammal, preferably a human, is administered a dose of a liquid formulation of the present invention comprising SYNAGIS[®] or an antigen-binding fragment thereof for the treatment, management or amelioration of a RSV infection or one or more symptoms thereof in an amount effective to induce a therapeutically effective serum titer of less than 10 µg/ml, less than 8 µg/ml, less than 5 µg/ml, less than 3 µg/ml, less than 1 µg/ml, or less than 0.5 µg/ml 30 days after the administration of the dose, wherein said therapeutically effective serum titer is the serum titer that reduces the severity or length of RSV infection or is the serum titer in a cotton rat that results in a RSV titer in the rat 5 days after challenge with 10⁵ pfu RSV that is 99% lower than the RSV titer 5 days after challenge with 10⁵ pfu of RSV in a cotton rat not administered the dose prior to challenge. Preferably, the dose of the liquid formulation of the present invention comprises 12 mg/kg or less, 10 mg/kg or less, 5 mg/kg or less, 3 mg/kg or less, 1 mg/kg or less, or 0.5 mg/kg or less of an antibody or an antigen-binding fragment thereof.

In a specific embodiment, formulations of the present invention are administered once a month just prior to or during the RSV season. In another embodiment, the formulations are administered every two months just prior to or during the RSV season. In yet another embodiment, the stable formulations of the present invention are administered
5 once just prior to or during the RSV season. The term "RSV season" refers to the season when RSV infection is most likely to occur. Typically, the RSV season in the northern hemisphere commences in November and lasts through April.

In one embodiment, a liquid formulation comprising approximately 5 mg/kg or less (preferably 1.5 mg/kg or less) of SYNAGIS® or an antigen-binding fragment thereof is
10 administered five times, 3 times, or 1 to 2 times during a RSV season to a mammal, preferably a human. In another embodiment, approximately 1.5 mg/kg of SYNAGIS® or an antigen-binding fragment thereof, in the liquid formulations of the present invention is administered monthly five times during a RSV season to a mammal, preferably a human, intramuscularly. In another embodiment, 3 mg/kg of SYNAGIS® or an antigen-binding
15 fragment thereof in the liquid formulation of the invention is administered monthly three times during a RSV season to a mammal, preferably a human, intramuscularly. In yet another embodiment, 5 mg/kg of an SYNAGIS® or an antigen-binding fragment thereof in a liquid formulation of the invention is administered monthly one to two times during a RSV season to a mammal, preferably a human, intramuscularly.

20 In a specific embodiment, 15 mg/kg of SYNAGIS® or an antigen-binding fragment thereof in the liquid formulation of the present invention is administered to a mammal, preferably a human, intramuscularly five times during a RSV season, wherein said SYNAGIS® or antibody fragment has an increased *in vivo* half-life. In another embodiment, approximately 5 mg/kg or less (preferably 1.5 mg/kg or less) of SYNAGIS®
25 or an antigen-binding fragment thereof in the liquid formulation of the present invention is administered five times, 3 times, or 1 to 2 times during a RSV season to a mammal, preferably a human. In another embodiment, 3 mg/kg of SYNAGIS® or an antigen-binding fragment thereof, which has an increased *in vivo* half-life, in the liquid formulation of the present invention is administered monthly three times during a RSV season to a mammal,
30 preferably a human, intramuscularly. In another embodiment, 5 mg/kg of SYNAGIS® or an antigen-binding fragment thereof, which has an increased *in vivo* half life, in the liquid formulation of the present invention is administered to a mammal, preferably a human, intramuscularly twice times during a RSV season.

5.8 Biological Assays

5.8.1 Immunospecificity of the Antibodies of the Invention

Antibodies of the present invention or fragments thereof may be characterized in a variety of ways well-known to one of skill in the art. In particular, antibodies of the invention or antigen-binding fragments thereof in a liquid formulation of the present invention may be assayed for the ability to immunospecifically bind to an epitope of a respiratory syncytial virus. Such an assay may be performed in solution (*e.g.*, Houghten, 1992, *Bio/Techniques* 13:412-421), on beads (Lam, 1991, *Nature* 354:82-84), on chips (Fodor, 1993, *Nature* 364:555-556), on bacteria (U.S. Patent No. 5,223,409), on spores (U.S. Patent Nos. 5,571,698; 5,403,484; and 5,223,409), on plasmids (Cull *et al.*, 1992, *Proc. Natl. Acad. Sci. USA* 89:1865-1869) or on phage (Scott and Smith, 1990, *Science* 249:386-390; Cwirla *et al.*, 1990, *Proc. Natl. Acad. Sci. USA* 87:6378-6382; and Felici, 1991, *J. Mol. Biol.* 222:301-310).

SYNAGIS[®] or an antigen-binding fragment thereof in a liquid formulation of the present invention can be assayed for its specificity and affinity.

SYNAGIS[®] or an antigen-binding fragment thereof of the present invention may be assayed for immunospecific binding to a RSV antigen and cross-reactivity with other antigens by any method known in the art. Immunoassays which can be used to analyze immunospecific binding and cross-reactivity include, but are not limited to, competitive and non-competitive assay systems using techniques such as western blots, radioimmunoassays, ELISA (enzyme linked immunosorbent assay), "sandwich" immunoassays, immunoprecipitation assays, precipitin reactions, gel diffusion precipitin reactions, immunodiffusion assays, agglutination assays, complement-fixation assays, immunoradiometric assays, fluorescent immunoassays, protein A immunoassays, to name but a few. Such assays are routine and well-known in the art (see, *e.g.*, Ausubel *et al.*, eds., 1994, *Current Protocols in Molecular Biology*, Vol. 1, John Wiley & Sons, Inc., New York).

5.8.2 *In vitro* and *in vivo* Assays

A liquid formulation or a combination therapy of the present invention can be tested *in vitro* and/or *in vivo* in various assays or suitable animal model systems for its activity.

A liquid formulation of the present invention for treating, managing, preventing, or ameliorating a RSV infection or one or more symptoms thereof can be tested for its ability to inhibit viral replication or reduce viral load in *in vitro* assays. For example, viral

replication can be assayed by a plaque assay such as described, *e.g.*, by Johnson *et al.*, 1997, Journal of Infectious Diseases 176:1215-1224 176:1215-1224. A liquid formulation of the invention administered according to the methods of the invention can also be assayed for their ability to inhibit or down-regulate the expression of viral polypeptides. Techniques
5 known to those of skill in the art, including, but not limited to, western blot analysis, northern blot analysis, and RT-PCR can be used to measure the expression of viral polypeptides and/or viral titers.

A liquid formulation of the invention can be tested in suitable animal model systems prior to use in humans. Such animal model systems include, but are not limited to, rats,
10 mice, chicken, cows, monkeys, pigs, dogs, rabbits, etc. Any animal system well-known in the art may be used. Several aspects of the procedure may vary; said aspects include, but are not limited to, the temporal regime of administering the therapies (*e.g.*, prophylactic and/or therapeutic agents) whether such therapies are administered separately or as an admixture, and the frequency of administration of the therapies.

15 Animal models can be used to assess the efficacy of the methods of the invention for treating, managing, preventing, or ameliorating a RSV infection or one or more symptom thereof. Animal models for RSV infection include, but are not limited to, those as described by, *e.g.*, Piedimonte *et al.*, Am J Physiol 1999, 277:L831-L840; McArthur-Vaughan *et al.*, J. Med. Primatol. 2002, 31(2):61-73; and Byrd *et al.*, Clin. Infect. Dis. 1997, 25(6):1363-8.

20 In a specific embodiment, cotton rats are administered a liquid formulation comprising SYNAGIS[®] according to the methods of the invention, challenged with 10⁵ pfu of RSV, and four or more days later, the rats are sacrificed and RSV titer and SYNAGIS[®] serum titer is determined. Accordingly, a dosage that results in a 2 log decrease or a 99% reduction in RSV titer in the cotton rat challenged with 10⁵ pfu of RSV relative to the cotton rat
25 challenged with 10⁵ pfu of RSV but not administered the formulation is the dosage of the formulation that can be administered to a human for the treatment, prevention or amelioration of a RSV infection or one or more symptoms thereof. Further, this embodiment, the tissues (*e.g.*, the lung tissues) from the sacrificed rats can be examined for histological changes.

30 The administration of a liquid formulation of the invention according to the methods of the present invention can be tested for its ability to decrease the time course of a RSV infection by at least 25%, preferably at least 50%, at least 60%, at least 75%, at least 85%, at least 95%, or at least 99%. A liquid formulation of the invention can also be tested for its ability to increase the survival period of humans suffering from a RSV infection by at least

25%, preferably at least 50%, at least 60%, at least 75%, at least 85%, at least 95%, or at least 99%. Further, a liquid formulation of the invention can be tested for its ability reduce the hospitalization period of a human suffering from RSV infection by at least 60%, preferably at least 75%, at least 85%, at least 95%, or at least 99%. Techniques known to those of skill in the art can be used to analyze the function of a liquid formulation of the invention *in vivo*.

Further, any *in vitro* or *in vivo* assays known to those skilled in the art can be used to evaluate the prophylactic and/or therapeutic utility of a liquid formulation of the invention disclosed herein for a RSV infection or one or more symptoms thereof.

5.8.3 Toxicity Assays

The toxicity and/or efficacy of the prophylactic and/or therapeutic protocols of the instant invention can be determined by standard pharmaceutical procedures in cell cultures or experimental animals, *e.g.*, for determining the LD50 (the dose lethal to 50% of the population) and the ED50 (the dose therapeutically effective in 50% of the population). The dose ratio between toxic and therapeutic effects is the therapeutic index and it can be expressed as the ratio LD50/ED50. Therapies that exhibit large therapeutic indices are preferred. While therapies that exhibit toxic side effects may be used, care should be taken to design a delivery system that targets such agents to the site of affected tissue in order to minimize potential damage to uninfected cells and, thereby, reduce side effects.

The data obtained from the cell culture assays and animal studies can be used in formulating a range of dosage of the prophylactic and/or therapeutic agents for use in humans. The dosage of such agents lies preferably within a range of circulating concentrations that include the ED50 with little or no toxicity. The dosage may vary within this range depending upon the dosage form employed and the route of administration utilized. For any therapy used in the method of the invention, the therapeutically effective dose can be estimated initially from cell culture assays. A dose may be formulated in animal models to achieve a circulating plasma concentration range that includes the IC50 (*i.e.*, the concentration of the test compound that achieves a half-maximal inhibition of symptoms) as determined in cell culture. Such information can be used to more accurately determine useful doses in humans. Levels in plasma may be measured, for example, by high performance liquid chromatography.

5.9 Kits

The invention provides a pharmaceutical pack or kit comprising one or more containers filled with a liquid formulation of the invention for the prevention, treatment, management or amelioration of a RSV infection or one or more symptoms thereof. In a specific embodiment, the liquid formulations of the invention comprise SYNAGIS[®] or an antigen-binding fragment thereof recombinantly fused or chemically conjugated to another moiety, including but not limited to, a heterologous protein, a heterologous polypeptide, a heterologous peptide, a large molecule, a small molecule, a marker sequence, a diagnostic or detectable agent, a therapeutic moiety, a drug moiety, a radioactive metal ion, a second antibody, and a solid support.

The present invention provides kits that can be used in the above methods. In one embodiment, a kit comprises a liquid formulation of the invention, in one or more containers. In another embodiment, a kit comprises a liquid formulation of the invention, in one or more containers, and one or more other prophylactic or therapeutic agents useful for the prevention, management or treatment of a RSV infection or one or more symptoms thereof, in one or more other containers. Preferably, the kit further comprises instructions for preventing, treating, managing or ameliorating a RSV infection (*e.g.*, using the liquid formulations of the invention alone or in combination with another prophylactic or therapeutic agent), as well as side effects and dosage information for method of administration. Optionally associated with such container(s) can be a notice in the form prescribed by a governmental agency regulating the manufacture, use or sale of pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use or sale for human administration.

6. EXAMPLES

6.1 Example 1 (Stability Study)

An antibody formulation of the present invention comprising, in an aqueous carrier, 25 mM of histidine, 1.6 mM of glycine, and SYNAGIS[®] at pH 6 was prepared according to the following protocol:

For a 1 kg solution of buffer: In 800 g water, 3.875 g histidine (free base) and 0.12 g glycine were dissolved. The pH was adjusted with 6 N HCl to 6.0 ± 0.2 . Water was added to bring the total mass up to 1.0 kg (qs).

For the diafiltration: After the chromatography steps in the purification process, SYNAGIS[®] was concentrated to a target of 150 g/L. The concentrated product is diafiltered

into formulation buffer. The formulated product was diluted to a target manufacturing concentration of 103 ± 3 g/L.

For a stability study, two formulations were prepared: one contained 105 mg/ml of SYNAGIS[®] and the other contained 160 mg/ml of SYNAGIS[®]. The stability of each formulation was measured using HPSEC in terms of degrees of aggregate formation and fragmentation during the storage at 2-8°C for up to 15 months and at 38-42°C for up to 1 year. For the HPSEC analysis, typically, TosohHaas G3000WXL column with a mobile phase containing 0.1 M sodium phosphate and 0.1 M sodium sulfate, pH 6.8, is used at a flow rate of 0.8 ml/min. A sample containing 250 mg of protein in an appropriate volume is injected into the column and protein peaks are detected by 280 nm UV and/or fluorescence (280 nm excitation and 340 nm emission).

The data showed that there was no detectable increase in aggregation when each formulation of SYNAGIS[®] was stored at 2-8°C for 15 months as shown in Table 3.

Table 3. Percent Aggregates during Storage at 2-8°C.

Month	% Aggregates	
	105 mg/ml	160 mg/ml
0	0.3	0.4
5	0.3	0.3
8	0.4	---
12	0.4	---
15	0.4	0.5

$\pm 0.1\%$ error.

When the formulations were stored at 38-42°C for 60 days, about 1.5% increase in aggregate was observed with the formulation containing 105 mg/ml of SYNAGIS[®] and 2.0% increase was observed with the formulation containing 160 mg/ml of SYNAGIS[®].

6.2 Example 2 (Clinical Study)

The liquid formulation of the present invention comprising 100 mg/ml of SYNAGIS[®] in an aqueous solution containing 25 mM of histidine and 1.6 mM of glycine at pH 6 is tested for safety and tolerability study in a Phase I, parallel group, double-blind, randomized study at two sites. The study drugs are a liquid (Liq) formulation of

SYNAGIS[®] and the currently licensed lyophilized (Lyo) formulation of SYNAGIS[®]. A total of 48 volunteers will be randomized to one of four treatment groups:

	GROUP 1:	N=12	3 mg/kg SYNAGIS [®] (Liq) at Study Days 0 and 30 (IM)
	GROUP 2:	N=12	3 mg/kg SYNAGIS [®] (Lyo) at Study Days 0 and 30 (IM)
5	GROUP 3:	N=12	15 mg/kg SYNAGIS [®] (Liq) at Study Day 0 (IV)
	GROUP 4:	N=12	15 mg/kg SYNAGIS [®] (Lyo) at Study Day 0 (IV)

Vital signs will be obtained before and 30 minutes after each dose of study drug. Adverse events will be monitored through 30 days after the last dose of study drug and serious adverse events will be monitored through Study Day 60.

10 On Study Day 0, all volunteers will have blood collected for SYNAGIS[®] serum concentration before dosing, at the end of infusion (IV dose groups only), and at 0.25, 0.5, 1, 4, 8, and 12 hours after IM injection or end of infusion. Subsequently, blood samples for determination of SYNAGIS[®] levels will be collected daily through Study Day 5 and on Study Days 7, 14, 21, 30, 37 (IM groups only), and 60. Serum samples for anti-SYNAGIS[®]
15 antibodies will be collected on Study Days 0, 7, 14, 21, 30, 37 (IM groups only), and 60. Samples for serum chemistry and CBC with differential and platelets, and urine samples for urinalysis will be collected on Study Day 0 as well as 7 days after each dose of SYNAGIS[®] (Study Day 7 for the IV groups and Study Days 7 and 37 for the IM groups). Urine β HCG tests will be performed on the day of dosing before each dose of SYNAGIS[®] (Study Day 0
20 for the IV groups and Study Days 0 and 30 for the IM groups). A study flow diagram is shown in Fig. 2.

6.2.1 Study procedures

A. Volunteer Selection

25 The volunteers in this study will be healthy male or female adults. The volunteer will be counseled by an investigator (physician) who will address the questions and concerns of the volunteer and secure written informed consent for participation in the study. Written informed consent will be obtained prior to conducting study procedures or administration of study drug.

a. Inclusion Criteria

30 Volunteers must meet *all* of the following criteria:

1. Male or female.
2. Age 18 through 49 years at the time of the first dose of study drug.

3. Weight ≤ 150 kg.

4. Written informed consent obtained from the volunteer.

5. Sexually active females, unless surgically sterile, must have used an effective method of avoiding pregnancy (including oral or implanted contraceptives, IUD, female condom, diaphragm with spermicide, cervical cap, abstinence, use of a condom by the sexual partner or sterile sexual partner) for 14 days prior to the first dose of study drug, must agree to continue using such precautions for 30 days after the final dose of study drug, and must have a negative serum pregnancy test within 2 days prior to the first dose of study drug.

10 6. Healthy by medical history and physical examination.

7. Ability to complete follow-up period of 60 days as required by the protocol.

b. Exclusion Criteria

Volunteers must have *none* of the following:

1. Acute illness at the start of the study

15 2. Fever $\geq 99.5^{\circ}\text{F}$ at the start of the study

3. Any drug therapy within 7 days prior to Study Day 0 (except for contraceptives)

4. Blood donation in excess of 400 mL within 6 months of study start

5. Receipt of immunoglobulin or blood products within 60 days before entry into the study

20 6. Receipt of any investigational drug therapy or standard vaccine within 120 days before the first dose of study drug in this protocol through 60 days after the final dose of study drug

7. History of immunodeficiency

25 8. History of allergic disease or reactions likely to be exacerbated by any component of the study drug

9. Previous medical history or evidence of an intercurrent illness that may compromise the safety of the volunteer in the study

10. Evidence of infection with hepatitis A, B, or C virus or HIV-1

30 11. At screening (must be within 21 days before entry into the study) any of the following: CBC: Hgb < 12.0 gm/dl; WBC $< 4,000/\text{mm}^3$; platelet count $< 120,000/\text{mm}^3$ (or laboratory normal values); AST, ALT, BUN, creatinine $>$ upper limit of normal; other abnormal laboratory values in the screening panel which in the opinion of the principal investigator are judged to be clinically significant; other abnormal laboratory values in the

screening panel which in the opinion of the principal investigator are judged to potentially confound analysis of study results

12. Nursing mother

13. History of alcohol or drug abuse within the past 2 years

5 14. Evidence of any systemic disease on physical examination

B. Randomization

a. Volunteer Randomization Procedures and Treatment Allocation

At the screening visit, volunteers will be evaluated by the principal investigator to assess eligibility for entry into the study. A master log will be maintained for all screened
 10 volunteers. Volunteers who sign an informed consent and who meet eligibility criteria will be entered into the study. When a volunteer arrives in the study site for randomization (Study Day 0), the investigator will confirm that the volunteer meets all inclusion and exclusion criteria. The investigator will then assign a patient identification number (PID). Patient identification numbers will be assigned sequentially within each of the two study
 15 sites beginning with #101 in site 1 and with #201 in site 2. Volunteers will be considered to have entered the study when the PID is assigned. A randomization list provided to the study pharmacist at each study site will contain assignments to each of the four treatment groups for volunteers at that site. The investigator will notify MedImmune by facsimile transmission (fax) at 301-527-4217 that a volunteer has been randomized. Volunteers who
 20 have been assigned a PID and do not receive any study drug, who receive an incomplete infusion of study drug, who do not receive both IM injections of study drug, or who do not complete at least 50% of the study visits may be replaced at the discretion of the sponsor. Volunteers who withdraw due to an adverse event or whose status cannot be ascertained will not be replaced. Volunteers who withdraw consent for reasons other than an adverse
 25 event may be replaced. Replacement volunteers will be assigned a new PID. Volunteers who are replaced will continue to be followed for safety according to the protocol.

b. Blinding

This is a double-blind study. Blinding will be maintained for assignment of volunteers to lyophilized or liquid formulation within IM or IV groups. In order to maintain
 30 blinding during administration of the two formulations of SYNAGIS[®], the study pharmacist at each site will prepare the study drug at a site physically removed from the treatment station and shielded from the observation of the principal investigator or any study personnel directly involved in the conduct of the study. For IM injection, the pharmacist

will prepare identical appearing 5 mL syringes containing the calculated volume of either liquid or reconstituted lyophilized SYNAGIS[®]. For IV infusion, the pharmacist will prepare identical appearing 200 mL infusion bags containing the calculated volume of either liquid or reconstituted lyophilized SYNAGIS[®]. Labels will not identify whether the
 5 syringe/bag contains liquid or reconstituted lyophilized SYNAGIS[®]. An independent monitor who will review the pharmacy record only, the statistician and clinical supplies manager at MedImmune, and the study pharmacist at the study site are the only individuals who will have access to the randomization list which identifies a volunteer's study treatment allocation. These individuals must not reveal randomization information to anyone. In the
 10 event that the study treatment for a volunteer becomes known to the investigator, MedImmune must be notified immediately by the investigator. All instances of unblinding will be documented in the study report.

C. Study Drug

a. Study Drug Supplies and Accountability

15 The sponsor will provide the investigator with adequate quantities of liquid SYNAGIS[®], lyophilized SYNAGIS[®], and diluent (sterile water for injection). Study drug should be stored at 2°C to 8°C (36°F to 46°F) and must not be frozen.

Liquid SYNAGIS[®] will be provided in 3 mL vials containing 100 mg of sterile liquid product in a volume of 1 mL (25 mM histidine, 1.6 mM glycine, at pH 6.0).

20 Lyophilized SYNAGIS[®] will be provided in 5 mL vials containing 100 mg of sterile lyophilized product which when formulated (before lyophilization) contains 25 mM histidine, 1.6 mM glycine, and 3% (w/v) mannitol at pH 6.0.

The study pharmacist is required to maintain accurate drug accountability records. Upon completion of the study, all study drug accountability records will be returned to the
 25 sponsor. All unused study drug will be returned to the sponsor.

b. Treatment Regimens

The following regimens are employed in the study:

GROUP 1:	N=12	3 mg/kg SYNAGIS [®] (Liq) at Study Days 0 and 30 (IM)
GROUP 2:	N=12	3 mg/kg SYNAGIS [®] (Lyo) at Study Days 0 and 30 (IM)
GROUP 3:	N=12	15 mg/kg SYNAGIS [®] (Liq) at Study Day 0 (IV)
GROUP 4:	N=12	15 mg/kg SYNAGIS [®] (Lyo) at Study Day 0 (IV)

c. Ordering and Preparation of Study Drug

The dose of study drug for administration must be prepared by the study pharmacist. The study drug prescription form indicating the PID and the volunteer's body weight will be sent to the pharmacist by the investigator (or designee). The study pharmacist will then use this information to prepare the study drug.

To prepare liquid or lyophilized SYNAGIS[®] for administration, the pharmacist should remove the tab portion of the vial cap and clean the rubber stopper with 70% ethanol or equivalent. The vial should be vented. Doses for each volunteer will be calculated as described below based on the volunteer's weight (to the nearest 0.1 kilogram) on the day SYNAGIS[®] is administered. The dose should be rounded to the nearest 0.1 mL. All preparations of study drug must be administered within 6 hours after entering the vial of SYNAGIS[®]. If it is not administered within 6 hours a new vial or vials must be used.

Preparation of Liquid SYNAGIS[®]

(1) **For IM injections:** The required volume of liquid SYNAGIS[®] (100 mg/mL) will be obtained by pooling the contents of as many vials as necessary with a 5 mL syringe.

$$\text{Dose (mL)} = [\text{Volunteer Weight (kg)} \times \text{Dose Level (3 mg/kg)}] \div \text{Drug Concentration (100 mg/mL)}$$

Example: A volunteer who weighs 75.6 kg receives 2.3 mL of SYNAGIS[®]

$$(75.6 \text{ kg} \times 3 \text{ mg/kg}) \div 100 \text{ mg/mL} = 2.268 \text{ mL (rounded to 2.3 mL)}$$

(2) **For IV infusions:** The required volume of liquid SYNAGIS[®] (100 mg/mL) will be obtained by pooling the contents of as many vials as necessary with a 20 mL (or larger) syringe.

$$\text{Dose (mL)} = [\text{Volunteer Weight (kg)} \times \text{Dose Level (15 mg/kg)}] \div \text{Drug Concentration (100 mg/mL)}$$

This volume of liquid SYNAGIS[®] will then be injected into an empty 200 mL infusion bag and diluted with diluent 1:4 by adding four volumes of diluent to the bag for a final concentration of 20 mg/mL SYNAGIS[®].

Example: A volunteer who weighs 71.4 kg receives 10.7 mL of SYNAGIS[®]

$$[(71.4 \text{ kg} \times 15 \text{ mg/kg}) \div 100 \text{ mg/mL} = 10.71 \text{ mL (rounded to 10.7 mL)}]$$

and 42.8 mL of diluent (4 x 10.7) for a total infusion volume of 53.5 mL.

Preparation of Lyophilized SYNAGIS[®]

(1) **For IM injections:** One (1) ml of diluent should be added slowly to the vial of lyophilized SYNAGIS[®] for a final concentration of 100 mg/mL SYNAGIS[®]. The vial should then be gently swirled for 30 seconds to avoid foaming. Reconstituted SYNAGIS[®] should stand at room temperature for a minimum of 20 minutes until SYNAGIS[®] clarifies.

5 The required volume of reconstituted lyophilized SYNAGIS[®] (100 mg/mL) will be obtained by pooling the contents of as many vials as necessary with a 5 mL syringe.

Dose (mL) = [Volunteer Weight (kg) x Dose Level (3 mg/kg)] ÷ Drug Concentration (100 mg/mL)

Example: A volunteer who weighs 75.6 kg receives 2.3 mL of SYNAGIS[®]

10 *(75.6 kg x 3 mg/kg) ÷ 100 mg/mL = 2.268 mL (rounded to 2.3 mL)*

(2) **For IV infusions:** Five (5) ml of diluent should be added slowly to the vial of lyophilized SYNAGIS[®] for a final concentration of 20 mg/mL SYNAGIS[®]. The vial should then be gently swirled for 30 seconds to avoid foaming. Reconstituted SYNAGIS[®] should stand at room temperature for a minimum of 20 minutes until SYNAGIS[®] clarifies.

15 The required volume of reconstituted lyophilized SYNAGIS[®] (20 mg/mL) will be obtained by pooling the contents of as many vials as necessary with a 20 mL (or larger) syringe and injecting this volume into an empty 200 mL infusion bag.

Dose (mL) = [Volunteer Weight (kg) x Dose Level (15 mg/kg)] ÷ Drug Concentration (20 mg/mL)

20 *Example: A volunteer who weighs 71.4 kg receives 53.6 mL of SYNAGIS[®]*

(71.4 kg x 15 mg/kg) ÷ 20 mg/mL = 53.55 mL (rounded to 53.6 mL)

d. Administration of Study Drug

Study drug for the IM and IV treatment groups will be dispensed from the pharmacy in identical appearing 5 mL syringes and identical appearing 200 mL infusion bags, respectively.

IM Injection

The study drug will be administered by IM injection into the deltoid muscle (after confirming the needle is not in a blood vessel) using standard aseptic technique. Volunteers will remain under observation in the study site for at least 30 minutes after the injection.

IV Infusion

Prior to drug administration, an IV catheter will be placed in an accessible vein using standard insertion techniques. Patency of the IV catheter will be maintained by a

continuous IV infusion of 5% Dextrose for Injection USP at a rate of 10 to 25 mL/h. The dextrose infusion will be interrupted, and SYNAGIS[®] will be infused through a low protein binding 0.22 µm filter at a rate of approximately 1-2 mL/minute. After SYNAGIS[®] has been administered, the IV tubing should be flushed and kept open with 5% Dextrose for Injection USP at 10 to 25 mL/h until the IV catheter is removed.

e. Concomitant Medications

All concomitant medications used by the volunteer from Study Day 0 through Study Day 60 will be recorded on the case report form. Volunteers may not receive the following:

1. Immunosuppressive medication (inhaled and topical corticosteroids are permitted).
2. Investigational agents from 120 days before study entry through Study Day 60.

The sponsor must be notified if any volunteer receives prohibited concomitant medications. Volunteers may receive medications to treat adverse events as deemed necessary by the investigator or the volunteer's physician.

D. Schedule of Volunteer Evaluations

All volunteers who are assigned a PID and receive any study drug will be followed according to the protocol regardless of the number of doses of study drug received, unless consent for follow-up is withdrawn. The sponsor must be notified of all deviations from protocol visits or evaluations and these evaluations, if applicable, must be rescheduled or performed at the nearest possible time to the original schedule.

Volunteers will be instructed to call study personnel to report any abnormalities during the intervals between study visits and to come to the study site if medical evaluation is needed and the urgency of the situation permits. For emergency and other unscheduled visits to a medical facility other than the study site, medical records will be obtained by the investigator. A schedule of screening and on-study visit procedures is presented in Table 4, followed by a detailed description of each visit.

Table 4 Schedule of Volunteer Evaluations

	Scn	Study Day											
		0	1	2	3	4	5	7	14	21	30	37	60
Written Informed Consent	x												
Verify Eligibility Criteria	x	x											
Medical History	x												
Physical Examination	x												
Height and body weight ^g	x	x									x ^d		
Urinalysis	x	x						x				x ^d	
Hepatitis A, B, C	x												
Serum β HCG ^b	x												
Urine β HCG ^b		x									x ^d		
Serum Chemistry ^c	x	x						x				x ^d	
CBC, Differential, Platelets	x	x						x				x ^d	
SYNAGIS [®] Serum Level		x ^a	x	x	x	x	x	x	x	x	x	x ^d	x
Anti-SYNAGIS [®] Antibody		x						x	x	x	x	x ^d	x
Vital Signs ^e		x									x ^d		
Randomization/Assignment of PID		x											
Study Drug Injection (IM Groups)		x									x ^d		
Study Drug Infusion (IV Groups)		x											
Assessment of Adverse Events ^f		x	x	x	x	x	x	x	x	x	x	x ^d	x

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- a. Blood will be sampled for SYNAGIS[®] serum concentration before dosing, at the end of infusion (IV dose groups only), and at 0.25, 0.5, 1, 4, 8, and 12 hours after injection/end of infusion.
- b. Females only.
- 5 c. ALT, AST, BUN, Creatinine.
- d. IM dose groups only.
- e. Vital signs obtained before and 30 minutes after each dose.
- f. Adverse events through 30 days after each dose. Serious adverse events through Study Day 60.
- 10 g. Body weight only at Study Day 30.

a. Screening

Note: All screening laboratory assessments must be performed within 21 days before study entry (Study Day 0). The screening evaluations may be carried out over more than one visit.

- 15 1. Written informed consent
2. Verify eligibility criteria
3. Screening medical history
4. Screening physical examination
5. Height and body weight
- 20 6. Urinalysis
7. Blood collection for screening
 - Serum for hepatitis A antibody, hepatitis B surface antigen, hepatitis C antibody
 - Serum β HCG (female volunteers only)
 - 25 • Chemistry panel (AST, ALT, BUN, creatinine)
 - CBC with differential and platelet count

Study Day 0: Dose 1

Visit 1

- 30 1. Verify eligibility criteria
2. Height and body weight
3. Urinalysis
4. Urine β HCG (female volunteers only)

5. Baseline blood collection

- Chemistry panel (AST, ALT, BUN, creatinine)
- CBC with differential and platelet count
- Serum for SYNAGIS[®] level
- Serum for anti-SYNAGIS[®] antibody

6. Randomization and assignment of PID

7. Vital signs before administration of study drug (temperature, blood pressure, pulse rate, respiratory rate)

Note: All of the above must be completed before administration of study drug.

1. Administration of study drug

2. Monitor vital signs 30 minutes after injection or end of infusion

3. Monitor for adverse events and serious adverse events

4. Post-dose blood collection

- Serum for SYNAGIS[®] levels immediately after completion of infusion (IV dose groups only) and at 0.25, 0.5, 1, 4, 8, and 12 hours after injection/end of infusion

Study Day 1: Dose 1 Pharmacokinetic Sampling**Visit 2**1. Post-dose blood collection for 24 hour SYNAGIS[®] serum level

2. Monitor for adverse events and serious adverse events

Study Day 2: Dose 1 Pharmacokinetic Sampling**Visit 3**1. Post-dose blood collection for 48 hour SYNAGIS[®] serum level

2. Monitor for adverse events and serious adverse events

Study Day 3: Dose 1 Pharmacokinetic Sampling**Visit 4**1. Post-dose blood collection for 72 hour SYNAGIS[®] serum level

2. Monitor for adverse events and serious adverse events

Study Day 4: Dose 1 Pharmacokinetic Sampling**Visit 5**

1. Post-dose blood collection for 96 hour SYNAGIS[®] serum level
2. Monitor for adverse events and serious adverse events

Study Day 5: Dose 1 Pharmacokinetic Sampling**Visit 6**

- 5 1. Post-dose blood collection for 120 hour SYNAGIS[®] serum level
2. Monitor for adverse events and serious adverse events

Study Day 7:**Visit 7**

1. Urinalysis
- 10 2. Blood collection
 - Chemistry panel (AST, ALT, BUN, creatinine)
 - CBC with differential and platelet count
 - Serum for SYNAGIS[®] level
 - Serum for anti-SYNAGIS[®] antibody
- 15 3. Monitor for adverse events and serious adverse events

Study Day 14±1:**Visit 8**

1. Blood collection
 - Serum for SYNAGIS[®] level
 - 20 • Serum for anti-SYNAGIS[®] antibody
2. Monitor for adverse events and serious adverse events

Study Day 21±1:**Visit 9**

1. Blood collection
 - 25 • Serum for SYNAGIS[®] level
 - Serum for anti-SYNAGIS[®] antibody
2. Monitor for adverse events and serious adverse events

Study Day 30±1:**Visit 10**

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All volunteers

1. Blood collection

- Serum for SYNAGIS[®] level
- Serum for anti-SYNAGIS[®] antibody

5 2. Monitor for adverse events and serious adverse events

Volunteers in IM groups only

3. Body weight

4. Urine β HCG

5. Vital signs before administration of study drug

10 6. Administration of study drug

7. Monitor vital signs 30 minutes after injection

Study Day 37 $\pm$ 1:**Visit 11**

Volunteers in IM groups only

15 1. Urinalysis

2. Blood collection

- Chemistry panel (AST, ALT, BUN, creatinine)
- CBC with differential and platelet count
- Serum for SYNAGIS[®] level
- Serum for anti-SYNAGIS[®] antibody

20 3. Monitor for adverse events and serious adverse events

Study Day 60 $\pm$ 2:**Visit 12**

25 1. Blood collection

- Serum for SYNAGIS[®] level
- Serum for anti-SYNAGIS[®] antibody

2. Monitor for serious adverse events

E. Volunteer Evaluation Methods30 **a. Routine Laboratory Evaluations**

Routine laboratory tests during screening and during the study will be performed at each study site's local clinical laboratory. Urine pregnancy tests during the study will be

performed in the study site using a licensed test. Abnormal laboratory results should be repeated as soon as possible (preferably within 24-48 hours).

b. Pharmacokinetic and Immunologic Evaluations

SYNAGIS[®] serum concentration and anti-SYNAGIS[®] antibodies will be measured
5 by MedImmune, Inc. by ELISA.

F. Completion of Study and Loss to Follow-up

Volunteers will be considered to have completed the study if they were followed up through Study Day 60. It should be specified on the case report form whether or not the
10 volunteer completed the study follow-up procedures through Study Day 60. Volunteers will be considered lost-to-follow-up only if no contact has been established by the time the study is completed such that there is insufficient information to determine the volunteer's status at Study Day 60. The investigator should document attempts to re-establish contact with missing volunteers throughout the study period. If contact with a missing volunteer is re-
15 established, follow-up should resume according to the protocol.

Pharmacokinetic and immunologic evaluations will be made based on SYNAGIS[®] serum concentration and anti-SYNAGIS[®] antibodies measured by ELISA.

Equivalents

20 Those skilled in the art will recognize, or be able to ascertain using no more than routine experimentation, many equivalents to the specific embodiments of the invention described herein.

25

Citation or discussion of a reference herein shall not be construed as an admission that such is prior art to the present invention.

30

SEQUENCE LISTING

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<120> STABILIZED LIQUID ANTI-RSV ANTIBODY FORMULATIONS

<130> PCA17161

<140> PCT/US03/18913

<141> 2003-06-16

<150> 60/388,921

<151> 2002-06-14

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<170> PatentIn version 3.1

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The embodiments of the present invention for which an exclusive property or privilege is claimed are defined as follows:

1. An aqueous palivizumab formulation comprising, in an aqueous carrier: (a) at least 75 mg/ml of palivizumab, or an antigen-binding fragment thereof; (b) about 10 mM to about 50 mM of histidine, and (c) less than 3mM glycine, wherein said palivizumab or palivizumab antigen-binding fragment in the formulation is stable at 38°C to 42°C for at least 60 days as determined by high performance size exclusion chromatography (HPSEC), and wherein the formulation has a pH of about 5.0 to about 7.0.
2. The formulation of claim 1, wherein the formulation does not comprise mannitol.
3. The formulation of claim 1 or 2, wherein said palivizumab or palivizumab antigen-binding fragment is at a concentration of about 80 mg/ml.
4. The formulation of claim 1 or 2, wherein said palivizumab or palivizumab antigen-binding fragment is at a concentration of at least 80 mg/ml.
5. The formulation of claim 1 or 2, wherein said palivizumab or palivizumab antigen-binding fragment is at a concentration of at least 85 mg/ml.
6. The formulation of claim 1 or 2, wherein said palivizumab or palivizumab antigen-binding fragment is at a concentration of at least 90 mg/ml.
7. The formulation of claim 1 or 2, wherein said palivizumab or palivizumab antigen-binding fragment is at a concentration of at least 95 mg/ml.
8. The formulation of claim 1 or 2, wherein said palivizumab or palivizumab antigen-binding fragment is at a concentration of at least 100 mg/ml.
9. The formulation of claim 1 or 2, wherein said palivizumab or palivizumab antigen-binding fragment is at a concentration of about 100 mg/ml.

10. The formulation of claim 8, wherein said palivizumab or palivizumab antigen-binding fragment is at a concentration of 103 ± 3 mg/ml.
11. The formulation of any one of claims 1 to 10, wherein the histidine is at a concentration of about 20 mM to about 30 mM.
12. The formulation of any one of claims 1 to 10, wherein the histidine is at a concentration of about 23 mM to about 27 mM.
13. The formulation of any one of claims 1 to 10, wherein the histidine is at a concentration of about 25 mM.
14. The formulation of claim 1, wherein the glycine is at a concentration of less than 2 mM.
15. The formulation of claim 1, wherein the glycine is at a concentration of less than 1.8 mM.
16. The formulation of claim 1, wherein the glycine is at a concentration of 1.6 mM.
17. The formulation of claim 1, further comprising a saccharide.
18. The formulation of claim 17, wherein the saccharide is sucrose, mannose, or trehalose.
19. The formulation of claim 1, further comprising a polyol.
20. The formulation of claim 19, wherein the polyol is sorbitol.
21. The formulation of claim 19, wherein the polyol is polysorbate.
22. The formulation of claim 21, wherein the polysorbate is TweenTM 20.

23. An aqueous palivizumab formulation comprising, in an aqueous carrier: (a) 103 ± 3 mg/ml of palivizumab or a palivizumab antigen-binding fragment, (b) about 25 mM histidine, and (c) about 1.6 mM glycine, wherein said palivizumab or palivizumab antigen-binding fragment in the formulation is stable at 2°C to 8°C at 15 months as determined by high performance size exclusion chromatography (HPSEC).
24. The formulation of claim 23, wherein the formulation does not comprise mannitol.
25. The formulation of claim 23 or 24, wherein said palivizumab or palivizumab antigen-binding fragment is at a concentration of 100 mg/ml.
26. The formulation of any one of claims 1 to 25, wherein said palivizumab or palivizumab antigen-binding fragment in the formulation is stable at 2°C to 8°C at 15 months as determined by HPSEC.
27. The formulation of any one of claims 23 to 25, wherein said palivizumab or palivizumab antigen-binding fragment in the formulation is stable at 38°C to 42°C for at least 60 days as determined by HPSEC.
28. The formulation of any one of claims 1 to 27, wherein the formulation is substantially free of surfactants and inorganic salts.
29. The formulation of any one of claims 1 to 27, wherein said formulation is substantially free of surfactants, inorganic salts and other excipients.
30. The formulation of any one of claims 1 to 29, wherein the formulation has a pH of about 5.5 to about 6.5.
31. The formulation of any one of claims 1 to 29, wherein the formulation has a pH of about 5.8 to about 6.2.
32. The formulation of any one of claims 1 to 29, wherein the formulation has a pH of about 6.0.

33. The formulation of any one of claims 1 to 32, wherein the formulation contains no more than 3% aggregation by weight protein as measured by HPSEC.
34. The formulation of any of claims 1 to 33, wherein the formulation contains equal to or more than 98% of the total protein in a single peak as determined by HPSEC and contains no other single peaks having more than 2% of total protein each.
35. The formulation of any one of claims 1 to 34, wherein the formulation is sterile.
36. The formulation of any one of claims 1 to 35, wherein the formulation is homogenous.
37. The formulation of any one of claims 1 to 36, which has been prepared by a process in which, for each step of the process, said palivizumab or palivizumab antigen-binding fragment is in an aqueous phase.
38. The formulation of any one of claims 1 to 37, wherein the aqueous carrier is water.
39. The formulation of any one of claims 1 to 37, wherein the aqueous carrier is distilled water.
40. The formulation of any one of claims 1-39, wherein the histidine is L-histidine.
41. The formulation of any one of claims 1-40, wherein the formulation further comprises chloride.
42. The formulation of any one of claims 1 to 41, wherein said palivizumab or palivizumab antigen-binding fragment is conjugated to a therapeutic moiety.
43. The formulation of any one of claims 1 to 41, wherein said palivizumab or palivizumab antigen-binding fragment is conjugated to a detectable substance.
44. A pharmaceutical unit dosage comprising the formulation of any one of claims 1 to 43, which dosage is in a form for administration to a human and is in a container.

45. The pharmaceutical unit dosage of claim 44, wherein said palivizumab or palivizumab antigen-binding fragment is in 1 ml of the aqueous carrier.
46. The pharmaceutical unit dosage of claim 44 or 45, wherein the formulation is in a form for parenteral administration to a human.
47. The pharmaceutical unit dosage of claim 44 or 45, wherein the formulation is in a form for intramuscular administration to a human.
48. The pharmaceutical unit dosage of claim 44 or 45, wherein the formulation is in a form for aerosol, subcutaneous, or intranasal administration to a human.
49. The pharmaceutical unit dosage of claim 44 or 45, wherein the formulation is in a form for intravenous administration to a human.
50. A sealed container comprising the formulation of any one of claims 1 to 43.
51. A sealed container comprising aqueous palivizumab formulation comprising 103 ± 3 mg/ml of palivizumab or a palivizumab antigen-binding fragment, about 25 mM histidine, and about 1.6 mM glycine in 1 ml of distilled water, wherein said palivizumab or palivizumab antigen-binding fragment in the formulation is stable at 2°C to 8°C at 15 months as determined by HPSEC.
52. The sealed container of claim 51, wherein said palivizumab or palivizumab antigen-binding fragment is at a concentration of 100 mg/ml.
53. The sealed container of claim 51 or 52, wherein the histidine is L-histidine.
54. The sealed container of any one of claims 51-53, wherein the formulation further comprises chloride.
55. The formulation of any one of claims 1 to 42 for use in the treatment of a respiratory syncytial virus (RSV) infection or one or more symptoms caused by or associated with the RSV infection in a subject.

56. The formulation of any one of claims 1 to 42 for use in preventing a respiratory syncytial virus (RSV) infection or one or more symptoms caused by or associated with the RSV infection in a subject.
57. The formulation of any one of claims 1 to 42 for use in managing or ameliorating a respiratory syncytial virus (RSV) infection or one or more symptoms caused by or associated with the RSV infection in a subject.
58. The formulation of any one of claims 1 to 41 and 43 for use in the diagnosis of a RSV infection.
59. The formulation of claim 58 for use in the diagnosis of the RSV infection in a subject.
60. The formulation of any one of claims 55 to 57 and 59, wherein the subject is a human.
61. The formulation of any one of claims 55 to 57 and 59, wherein the subject is a human with cystic fibrosis.
62. The formulation of any one of claims 55 to 57 and 59, wherein the subject is a human with bronchopulmonary dysplasia.
63. The formulation of any one of claims 55 to 57 and 59, wherein the subject is a human with congenital heart disease.
64. The formulation of any one of claims 55 to 57 and 59, wherein the subject is a human with congenital immunodeficiency or acquired immunodeficiency.
65. The formulation of any one of claims 55 to 57 and 59, wherein the subject is a human who has had a bone marrow transplant.
66. The formulation of any one of claims 55 to 57 and 59, wherein the subject is a human infant.
67. The formulation of any one of claims 55 to 57 and 59, wherein the subject is a human infant born prematurely.

68. The formulation of any one of claims 55 to 57 and 59, wherein the subject is a human infant born at less than 35 weeks gestational age who is less than 6 months old.

69. The formulation of any one of claims 55 to 57 and 59, wherein the subject is a human infant at risk of hospitalization for a RSV infection.

70. The formulation of any one of claims 55 to 57 and 59, wherein the subject is a human in a nursing home or orphanage.

71. The formulation of any one of claims 55 to 57 and 60 to 70, wherein the formulation is in a form for intramuscular administration.

72. The formulation of any one of claims 55 to 57 and 60 to 70, wherein the formulation is in a form for subcutaneous administration.

73. The formulation of any one of claims 55 to 57 and 60 to 70, wherein the formulation is in a form for intranasal administration to a human.

74. The formulation of any one of claims 55 to 57 and 60 to 70, wherein the formulation is in a form for intravenous administration.

75. Use of the formulation of any one of claims 1 to 43 in the manufacture of a medicament in the treatment of a respiratory syncytial virus (RSV) infection or one or more symptoms caused by or associated with the RSV infection in a subject.

76. Use of the formulation of any one of claims 1 to 43 in the manufacture of a medicament in the prevention of a respiratory syncytial virus (RSV) infection or one or more symptoms caused by or associated with the RSV infection in a subject.

77. Use of the formulation of any one of claims 1 to 43 in the manufacture of a medicament in the management or amelioration of a respiratory syncytial virus (RSV) infection or one or more symptoms caused by or associated with the RSV infection in a subject.

78. Use of the formulation of claim 43 in the manufacture of a medicament in the diagnosis of a respiratory syncytial virus (RSV) infection.
79. The use of claim 78 in the manufacture of a medicament in the diagnosis of the RSV infection in a subject.
80. The use of any one of claims 75 to 77 and 79, wherein the subject is a human.
81. The use of any one of claims 75 to 77, wherein the subject is a human with cystic fibrosis.
82. The use of any one of claims 75 to 77, wherein the subject is a human with bronchopulmonary dysplasia.
83. The use of any one of claims 75 to 77, wherein the subject is a human with congenital heart disease.
84. The use of any one of claims 75 to 77, wherein the subject is a human with congenital immunodeficiency or acquired immunodeficiency.
85. The use of any one of claims 75 to 77, wherein the subject is a human who has had a bone marrow transplant.
86. The use of any one of claims 75 to 77, wherein the subject is a human infant.
87. The use of any one of claims 75 to 77, wherein the subject is a human infant born prematurely.
88. The use of any one of claims 75 to 77 and 79, wherein the subject is a human infant born at less than 35 weeks gestational age who is less than 6 months old.
89. The use of any one of claims 75 to 77 and 79, wherein the subject is a human infant at risk of hospitalization for and RSV infection.

90. The use of any one of claims 75 to 77 and 79, wherein the subject is a human in a nursing home or orphanage.
91. The use of any one of claims 75 to 77 and 80 to 90, wherein the formulation is in a form for intramuscular administration.
92. The use of any one of claims 75 to 77 and 80 to 90, wherein the formulation is in a form for subcutaneous administration.
93. The use of any one of claims 75 to 77 and 80 to 90, wherein the formulation is in a form for intranasal administration to a human.
94. The use of any one of claims 75 to 77 and 80 to 90, wherein the formulation is in a form for intravenous administration.
95. A process for the preparation of a high concentration stable aqueous palivizumab formulation, comprising diafiltering about 70 mg/ml, about 80 mg/ml, about 90 mg/ml, about 100 mg/ml or 150 mg/ml of palivizumab or an antigen-binding fragment thereof into a formulation buffer comprising about 10 mM to about 50 mM of histidine and less than 3 mM of glycine, wherein the palivizumab or palivizumab antigen-binding fragment is in an aqueous phase during the preparation of the stable aqueous palivizumab formulation.
96. The process of claim 95, wherein the formulation buffer comprises about 20 mM to about 30 mM of histidine.
97. The process of claim 95, wherein the formulation buffer comprises about 23 mM to about 27 mM of histidine.
98. The process of claim 95, wherein the formulation buffer comprises about 25 mM of histidine.
99. The process of any one of claims 95 to 98, wherein the formulation buffer comprises less than 2.0 mM of glycine.

100. The process of any one of claims 95 to 98, wherein the formulation buffer comprises less than 1.8 mM of glycine.

101. The process of any one of claims 95 to 98, wherein the formulation buffer comprises 1.6 mM of glycine.

102. The process of any one of claims 95 to 101, wherein the stable aqueous formulation has a pH of about 5.0 to about 7.0.

103. The process of any one of claims 95 to 101, wherein the stable aqueous formulation has a pH of about 5.5 to about 6.5.

104. The process of any one of claims 95 to 101, wherein the stable aqueous formulation has a pH of about 5.8 to about 6.2.

105. The process of any one of claims 95 to 101, wherein the stable aqueous formulation has a pH of about 6.0.

106. The process of any one of claims 95 to 105, wherein the process further comprises sterilization of the high concentration stable aqueous palivizumab formulation.

107. The process of any one of claims 95 to 106, wherein the process comprises purification of palivizumab or palivizumab antigen-binding fragment from conditioned medium and concentration of the palivizumab or palivizumab antigen-binding fragment before diafiltering the palivizumab or palivizumab antigen-binding fragment into the formulation buffer.

108. The process of any one of claims 95 to 107, wherein the palivizumab or palivizumab antigen-binding fragment is diafiltrated into 1 ml of the formulation buffer.

109. The process of any one of claims 95 to 107, wherein the palivizumab or palivizumab antigen-binding fragment is diafiltrated into 1.2 ml of the formulation buffer.

110. The process of any one of claims 95 to 109, wherein the high concentration stable aqueous palivizumab formulation comprises at least 65 mg/ml, at least 70 mg/ml, at least 75 mg/ml, at least 80 mg/ml, at least 85 mg/ml, or at least 90 mg/ml of the palivizumab or palivizumab antigen-binding fragment.

111. The process of any one of claims 95 to 109, wherein the high concentration stable aqueous palivizumab formulation comprises at least 95 mg/ml or at least 100 mg/ml of the palivizumab or palivizumab antigen-binding fragment.

112. The process of claim 111, wherein the high concentration stable aqueous palivizumab formulation comprises 103 ± 3 mg/ml of the palivizumab or palivizumab antigen-binding fragment.

113. The process of any one of claims 95 to 112, wherein the high concentration stable aqueous palivizumab formulation is substantially free of surfactants and inorganic salts.

114. The process of any one of claims 95 to 113, wherein the high concentration stable aqueous palivizumab formulation is substantially free of other excipients.

115. The process of any one of claims 95 to 113, wherein the high concentration stable aqueous palivizumab formulation does not comprise mannitol.

116. The process of any one of claims 95 to 115, wherein the high concentration stable aqueous palivizumab formulation is stable at 2°C to 8°C at 15 months as assessed by high performance size exclusion chromatography (HPSEC).

117. The process of any one of claims 95 to 116, wherein the high concentration stable aqueous palivizumab formulation is stable at 38°C to 42°C for at least 60 days as assessed by HPSEC.

118. The process of any one of claims 95 to 117, wherein the high concentration stable aqueous palivizumab formulation contains no more than 3% aggregation by weight protein as measured by HPSEC.

119. The process of any one of claims 95 to 118, wherein the high concentration stable aqueous palivizumab formulation contains equal to or more than 95% of the total protein in a single peak as determined by HPSEC and contains no other single peaks having more than 2% of total protein each.
120. The process of any one of claims 95 to 119 which does not involve a drying step.
121. The process of any one of claims 95-120, wherein the histidine is a L-histidine.
122. The process of any one of claims 95-121, wherein the formulation further comprises chloride.
123. A high concentration stable aqueous palivizumab formulation produced by the process of any one of claims 95 to 122.
124. A unit dosage form comprising the high concentration stable aqueous palivizumab formulation of claim 123.
125. The unit dosage form of claim 124 for use in the prevention of a respiratory syncytial virus (RSV) infection or one or more symptoms caused by or associated with the RSV infection in a human subject.
126. The unit dosage form of claim 124 for use in the treatment of a respiratory syncytial virus (RSV) infection or one or more symptoms caused by or associated with the RSV infection in a human subject.
127. A sealed container comprising the high concentration stable aqueous palivizumab formulation of claim 123.
128. The high concentration stable aqueous palivizumab formulation of claim 123 in the manufacture of a medicament for use in the prevention of a respiratory syncytial virus (RSV) infection or one or more symptoms caused by or associated with the RSV infection in a human subject.
129. The high concentration stable aqueous palivizumab formulation of claim 123 in the manufacture of a medicament for use in the treatment of a respiratory syncytial virus (RSV) infection or one or more symptoms caused by or associated with the RSV infection in a human subject.

130. The high concentration stable aqueous palivizumab formulation of claim 123 in the manufacture of a medicament for use in the management of a respiratory syncytial virus (RSV) infection or one or more symptoms caused by or associated with the RSV infection in a human subject.

131. The high concentration stable aqueous palivizumab formulation of claim 123 in the manufacture of a medicament for use in the amelioration of a respiratory syncytial virus (RSV) infection or one or more symptoms caused by or associated with the RSV infection in a human subject.

132. Use of the formulation of any one of claims 1 to 43 in the treatment of a respiratory syncytial virus (RSV) infection or one or more symptoms caused by or associated with the RSV infection in a subject.

133. Use of the formulation of any one of claims 1 to 43 in the prevention of a respiratory syncytial virus (RSV) infection or one or more symptoms caused by or associated with the RSV infection in a subject.

134. Use of the formulation of any one of claims 1 to 43 in the management or amelioration of a respiratory syncytial virus (RSV) infection or one or more symptoms caused by or associated with the RSV infection in a subject.

135. Use of the formulation of claim 43 in the diagnosis of a respiratory syncytial virus (RSV) infection.

136. The use of claim 135 in the diagnosis of the RSV infection in a subject.

137. The use of any one of claims 132 to 134 and 136 wherein the subject is a human.

138. The use of any one of claims 132 to 134, wherein the subject is a human with cystic fibrosis.

139. The use of any one of claims 132 to 134, wherein the subject is a human with bronchopulmonary dysplasia.

140. The use of any one of claims 132 to 134, wherein the subject is a human with congenital heart disease.

141. The use of any one of claims 132 to 134, wherein the subject is a human with congenital immunodeficiency or acquired immunodeficiency.
142. The use of any one of claims 132 to 134, wherein the subject is a human who has had a bone marrow transplant.
143. The use of any one of claims 132 to 134 and 136, wherein the subject is a human infant.
144. The use of any one of claims 132 to 134 and 136, wherein the subject is a human infant born prematurely.
145. The use of any one of claims 132 to 134 and 136, wherein the subject is a human infant born at less than 35 weeks gestational age.
146. The use of any one of claims 132 to 134 and 136, wherein the subject is a human infant at risk of hospitalization for the RSV infection.
147. The use of any one of claims 132 to 134 and 136, wherein the subject is a human in a nursing home or orphanage.
148. The use of any one of claims 132 to 134 and 136 to 147, wherein the formulation is in a form for intramuscular administration.
149. The use of any one of claims 132 to 134 and 136 to 147, wherein the formulation is in a form for subcutaneous administration.
150. The use of any one of claims 132 to 134 and 136 to 147, wherein the formulation is in a form for intranasal administration to a human.
151. The use of any one of claims 132 to 134 and 136 to 147, wherein the formulation is in a form for intravenous administration.
152. An aqueous palivizumab formulation comprising, in an aqueous carrier: (a) about 100 mg/ml of palivizumab, (b) about 10 mM to about 50mM histidine, and (c) less than 3 mM glycine, wherein said palivizumab in the formulation is stable at 2°C to 8°C at 15 months as determined by high performance size exclusion chromatography (HPSEC).

153. The aqueous palivizumab formulation of claim 152 comprising, in an aqueous carrier: (a) about 100 mg/ml of palivizumab, (b) about 23 mM to about 27 mM histidine, and (c) less than 1.8 mM glycine.

154. The aqueous palivizumab formulation of claim 152 comprising, in an aqueous carrier: (a) about 100 mg/ml of palivizumab, (b) about 25 mM histidine, and (c) about 1.6 mM glycine.

155. The aqueous palivizumab formulation of any one of claims 152-154, wherein the histidine is L-histidine.

156. The aqueous palivizumab formulation of any one of claims 152-155, wherein the formulation further comprises chloride.

157. A dosage form comprising a solution for injection, said solution comprising in an aqueous carrier: (a) about 100 mg/ml of palivizumab, (b) about 10 mM to about 50mM histidine, and (c) less than 3 mM glycine, wherein said palivizumab in the formulation is stable at 2°C to 8°C at 15 months as determined by high performance size exclusion chromatography (HPSEC).

158. The dosage form of claim 157 comprising a solution for injection, said solution comprising in an aqueous carrier: (a) about 100 mg/ml of palivizumab, (b) about 23 mM to about 27 mM histidine, and (c) less than 1.8 mM glycine.

159. The dosage form of claim 157 comprising a solution for injection, said solution comprising in an aqueous carrier: (a) about 100 mg/ml of palivizumab, (b) about 25 mM histidine, and (c) about 1.6 mM glycine.

160. The dosage form of any one of claims 157-159, wherein the histidine is L-histidine.

161. The dosage form of any one of claims 157-160, wherein the formulation further comprises chloride.

162. A solution for injection, said solution comprising in an aqueous carrier: (a) about 100 mg/ml of palivizumab, (b) about 10 mM to about 50mM histidine, and (c) less than 3 mM glycine, wherein said palivizumab in the formulation is stable at 2°C to 8°C at 15 months as determined by high performance size exclusion chromatography (HPSEC).

163. The solution for injection of claim 162, said solution comprising in an aqueous carrier: (a) about 100 mg/ml of palivizumab, (b) about 23 mM to about 27 mM histidine, and (c) less than 1.8 mM glycine.

164. The solution for injection of claim 162, said solution comprising in an aqueous carrier: (a) about 100 mg/ml of palivizumab, (b) about 25 mM histidine, and (c) about 1.6 mM glycine.

165. The solution for injection of any one of claims 162-164, wherein the histidine is L-histidine.

166. The solution for injection of any one of claims 162-165, wherein the formulation further comprises chloride.

167. Use of the formulation of any one of claims 1 to 43 in the prevention of lower respiratory tract disease caused by respiratory syncytial virus (RSV) in pediatric patients at high risk of RSV disease.

168. The use of claim 167, wherein the pediatric patient is an infant with bronchopulmonary dysplasia (BPD).

169. The use of claim 167, wherein the pediatric patient is an infant with a history of prematurity (≤ 35 weeks gestational age).

170. The use of claim 167, wherein the pediatric patient is a child with congenital heart disease.

171. Use of about 100 mg/ml of palivizumab in an aqueous solution for the treatment of lower respiratory tract disease caused by respiratory syncytial virus (RSV) in pediatric patients at high risk of RSV disease, wherein the aqueous solution comprises about 25 mM histidine and about 1.6 mM glycine, and wherein said palivizumab in the aqueous solution is stable at 2°C to 8°C at 15 months as determined by high performance size exclusion chromatography (HPSEC).

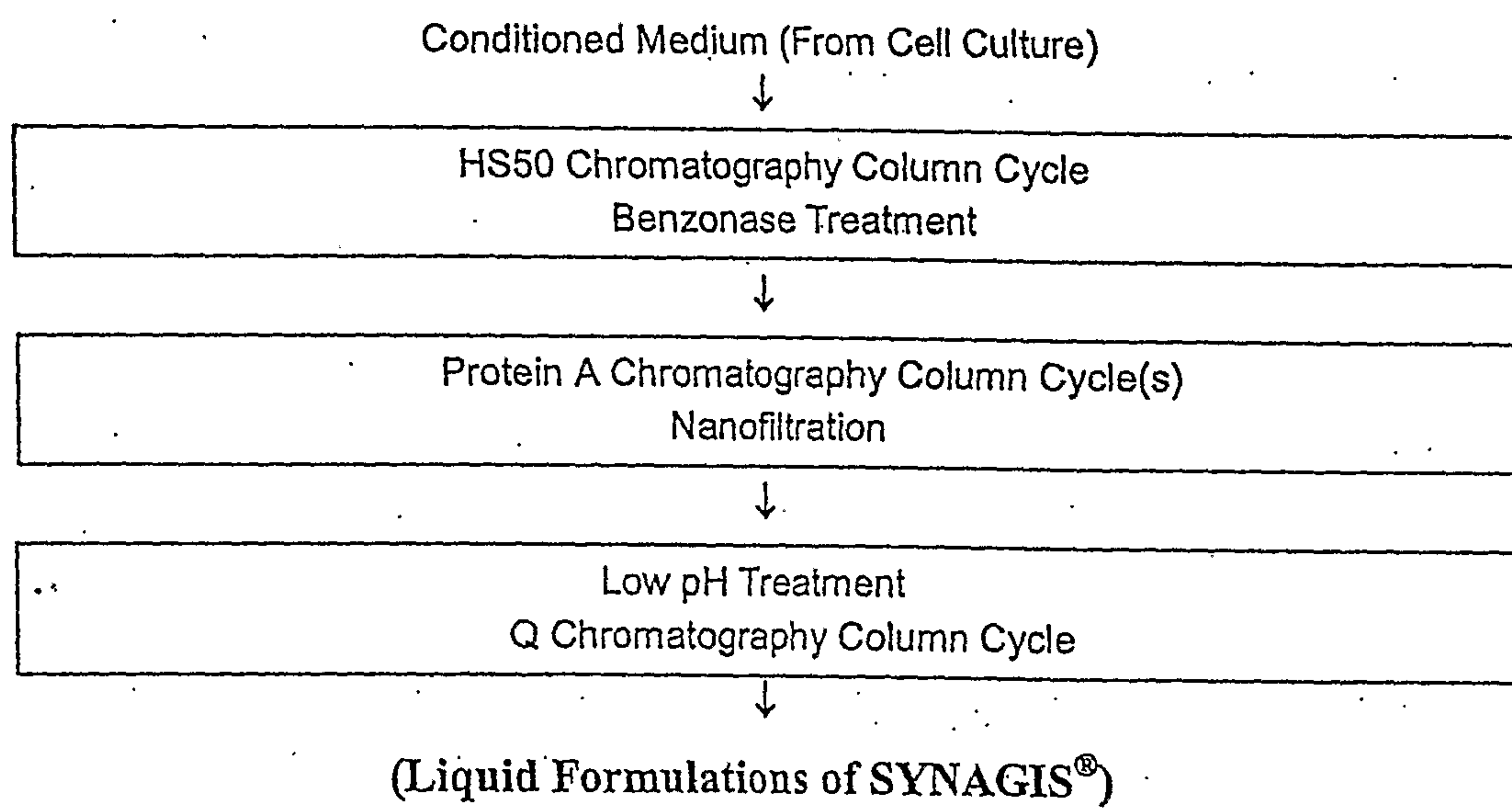
172. The use of claim 171, wherein the pediatric patient is an infant with bronchopulmonary dysplasia (BPD).

173. The use of claim 171, wherein the pediatric patient is an infant with a history of prematurity (≤ 35 weeks gestational age).
174. The use of claim 171, wherein the pediatric patient is a child with congenital heart disease.
175. The use of any one of claims 171-174, wherein the histidine is L-histidine.
176. The use of any one of claims 171-175, wherein the formulation further comprises chloride.
177. Use of the formulation of any one of claims 1 to 43 in the treatment of lower respiratory tract disease caused by respiratory syncytial virus (RSV) in pediatric patients at risk of hospitalization for RSV disease.
178. The use of claim 177, wherein the pediatric patient is an infant with bronchopulmonary dysplasia (BPD).
179. The use of claim 177, wherein the pediatric patient is an infant with a history of prematurity (≤ 35 weeks gestational age).
180. The use of claim 177, wherein the pediatric patient is a child with congenital heart disease.
181. Use of about 100 mg/ml of palivizumab in an aqueous solution for the treatment of lower respiratory tract disease caused by respiratory syncytial virus (RSV) in pediatric patients at risk for hospitalization of RSV disease, wherein the aqueous solution comprises about 25 mM histidine and about 1.6 mM glycine, and wherein said palivizumab in the aqueous solution is stable at 2°C to 8°C at 15 months as determined by high performance size exclusion chromatography (HPSEC).
182. The use of claim 181, wherein the pediatric patient is an infant with bronchopulmonary dysplasia (BPD).
183. The use of claim 181, wherein the pediatric patient is an infant with a history of prematurity (≤ 35 weeks gestational age).

184. The use of claim 181, wherein the pediatric patient is a child with congenital heart disease.

185. The use of any one of claims 181-184, wherein the histidine is L-histidine.

186. The use of any one of claims 181-185, wherein the formulation further comprises chloride.

**Fig. 1**

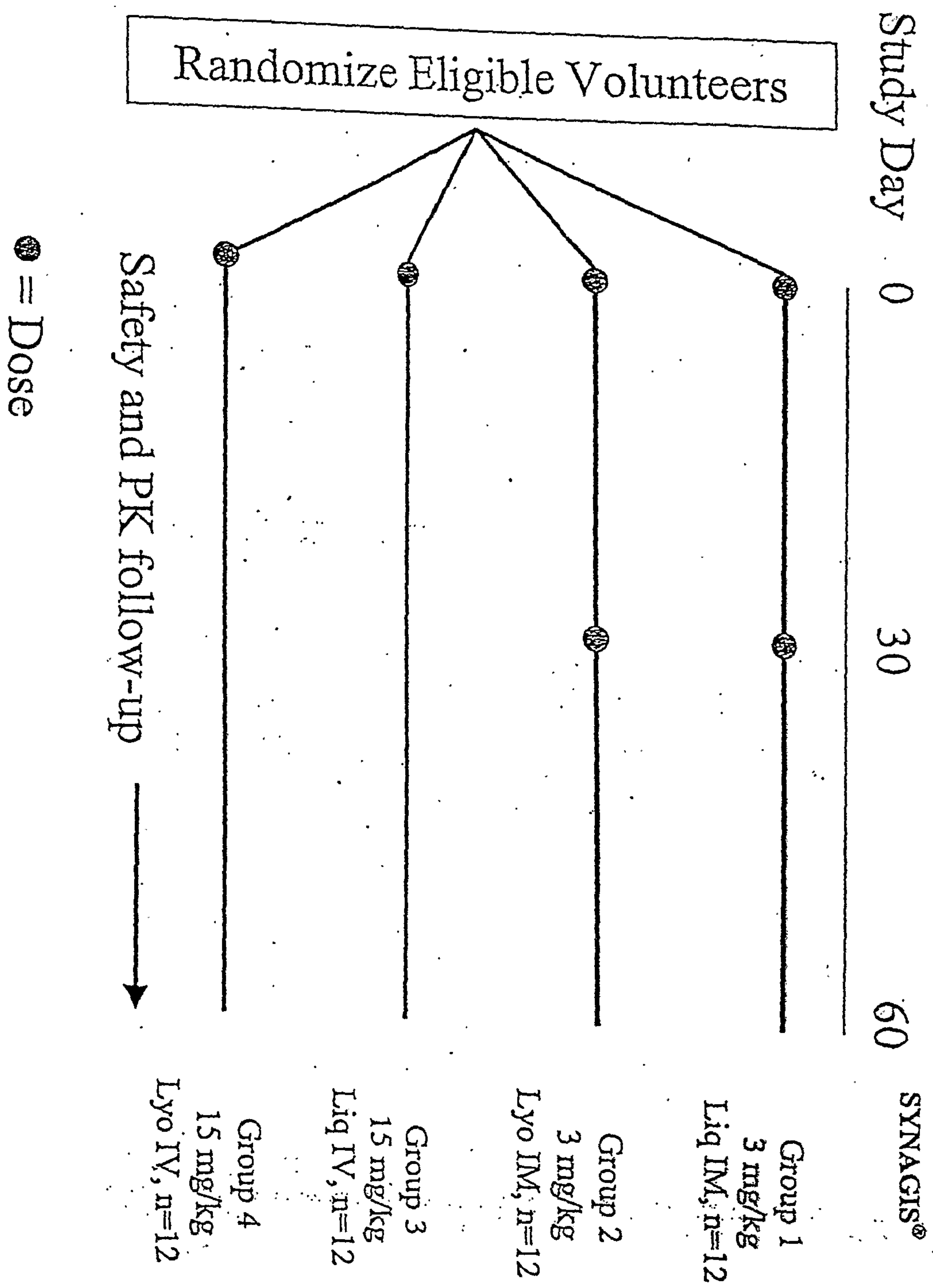


Fig. 2