

US Patent & Trademark Office

Patent Public Search | Text View

United States Patent Application Publication

20250257109

Kind Code

A1

Publication Date

August 14, 2025

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FORMULATION OF MODIFIED INTERLEUKIN-7 FUSION PROTEIN

Abstract

Provided is a pharmaceutical formulation comprising a modified IL-7 protein. More particularly, it comprises (a) a modified IL-7 fusion protein; (b) a basal buffer with a concentration of 10 to 50 mM; (c) a sugar with a concentration of 2.5 to 5w/v %; and (d) a surfactant with a concentration of 0.05 to 6w/v %.

Such pharmaceutical formulation of a modified IL-7 fusion protein does not show aggregates formation, but shows protective effects on proteins under stress conditions such as oxidation or agitation, and thus can effectively be used for the treatment of a patient.

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Appl. No.: 19/028674

Filed: January 17, 2025

Foreign Application Priority Data

KR 10-2015-0156124

Nov. 06, 2015

Related U.S. Application Data

parent US continuation 18053671 20221108 PENDING child US 19028674

parent US continuation 15773273 20180503 parent-grant-document US 11548927 WO

continuation PCT/KR2016/012495 20161102 child US 18053671

Publication Classification

Int. Cl.: **C07K14/54** (20060101); **A61K9/08** (20060101); **A61K38/20** (20060101); **A61K47/10** (20170101); **A61K47/12** (20060101); **A61K47/18** (20170101); **A61K47/26** (20060101); **C07K5/062** (20060101); **C07K5/083** (20060101); **C07K5/103** (20060101); **C07K7/06** (20060101)

U.S. Cl.:

CPC **C07K14/5418** (20130101); **A61K9/08** (20130101); **A61K38/2046** (20130101); **A61K47/10** (20130101); **A61K47/12** (20130101); **A61K47/183** (20130101); **A61K47/26** (20130101); **C07K7/06** (20130101); **C07K14/54** (20130101); C07K5/06026 (20130101); C07K5/0606 (20130101); C07K5/0806 (20130101); C07K5/081 (20130101); C07K5/1008 (20130101); C07K5/1013 (20130101); C07K2319/30 (20130101)

Background/Summary

CROSS REFERENCE TO RELATED APPLICATIONS [0001] This application is a continuation application of U.S. patent application Ser. No. 18/053,671, filed Nov. 8, 2022, which is a continuation application of U.S. patent application Ser. No. 15/773,273 (issued as U.S. Pat. No. 11,548,927), which is a U.S. National Stage of International Application No. PCT/KR2016/012495, filed on Nov. 2, 2016, which claims the priority benefit of Korean Patent Application No. 10-2015-0156124, filed on Nov. 6, 2015, each of which is hereby incorporated by reference in its entirety.

REFERENCE TO SEQUENCE LISTING SUBMITTED ELECTRONICALLY VIA EFS-WEB [0002] The content of the electronically submitted sequence listing in ASCII text file (Name: 4241_0330002_Seglisting_ST26.xml; Size: 62,076 bytes; and Date of Creation: Oct. 31, 2022) filed with the application is incorporated herein by reference in its entirety.

TECHNICAL FIELD

[0003] The present invention relates to a formulation of a pharmaceutical composition comprising a modified interleukin-7 fusion protein.

BACKGROUND ART

[0004] Interleukin-7 (IL-7) is a cytokine stimulating immune responses mediated by T cell and B cell, and specifically, it plays an important role in an adaptive immune system. Although IL-7 is mainly secreted by stromal cells in the bone marrow and the thymus, it is also produced in keratinocytes, dendritic cells, hepatocytes, neural and epithelial cells (Heufler C et al., 1993, *J. Exp. Med.* 178 (3): 1109-14; Kroncke R et al., 1996, *Eur. J. Immunol.* 26 (10): 2541-4; Sawa Y et al., 2009, *Immunity* 30 (3): 447-57; Watanabe M et al., 1995, *J. Clin. Invest.* 95 (6): 2945-53).

[0005] IL-7 activates the immune system by influencing the survival and differentiation of T cell and B cell, and stimulating the activity of NK (natural killer) cell, and so on, and specifically, it plays an important role in the development of B cell. IL-7 enhances the immune response in the human body by stimulating the secretion of IL-2, a type of cytokine, and interferon- γ .

[0006] In other words, IL-7 is a cytokine which promotes the survival and proliferation of T-cell, B-cell and other immune cells, and is an excellent candidate for an immune therapeutic agent applicable to various disorders such as viral infection, cancer and immune system damage. Recently a clinical study has been conducted to elucidate the effect by IL-7 on various malignancies and human immune deficiency virus infection, and as a result, an immune enhancing effect in human of IL-7 was reported (Fry T J et al., 2002, *Blood* 99 (11): 3892-904; Muegge K et al., 1993, *Science* 261 (5117): 93-5; Rosenberg S A et al., *J. Immunother.* 29 (3): 313-9). In

addition, IL-7 has been reported to be helpful in the immune recovery after allogeneic stem cell transplantation (Snyder K M, 2006, *Leuk Lymphoma* 47 (7). 1222-8), and is used for the treatment of lymphopenia as well. Accordingly, IL-7 reorganizes immune cells, enhances T cell function, and competes with the substances which induce immunosuppression, and thus it can be used for treating cancer or chronic infection.

[0007] However, in order to utilize proteins such as IL-7 for a pharmaceutical use, it is necessary to prepare a suitable formulation in consideration of the structure and the physiochemical stability in surrounding environment of the protein. In addition, the physical and chemical stability of the protein is affected by external factors such as manufacturing of a pharmaceutical formulation, purification of a protein, and storage. In general, a protein is structurally and thermodynamically unstable, and thus is easily subject to aggregates formation or physiochemical degradation. In addition, when Fc region of an immunoglobulin is used as a fusion partner, the physiochemical property of the entire fusion protein such as thermodynamic property, Zeta potential and stress stability, etc., can vary greatly since the Fc region has a large molecular weight and a complex structure. Specifically, little is known with respect to a stabilized formulation for the modified IL-7 fusion protein. Thus, it is necessary to optimize the formulation of a protein in order to increase the stability of the protein for a pharmaceutical use.

DISCLOSURE OF INVENTION

Technical Problem

[0008] Therefore, the present inventors have established an appropriate formulation to increase the stability of a modified IL-7 fusion protein for use in treating a variety of disorders.

[0009] An object of the present invention is to provide a pharmaceutical formulation comprising a modified IL-7 fusion protein with increased stability.

Solution to Problem

[0010] In accordance with the object of the present invention, there is provided a pharmaceutical formulation comprising: (a) a modified IL-7 fusion protein; (b) a basal buffer with a concentration of 10 to 50 mM; (c) a sugar with a concentration of 2.5 to 5 w/v %; and (d) a surfactant with a concentration of 0.05 to 6 w/v %.

Advantageous Effects of Invention

[0011] A pharmaceutical formulation of a modified IL-7 fusion protein according to the present invention does not show aggregates formation, and maintains stability of the modified IL-7 fusion protein under stress conditions such as oxidation or agitation, and thus can be used for the treatment of a patient.

Description

BRIEF DESCRIPTION OF DRAWINGS

[0012] FIG. 1 is a schematic diagram showing the structure of a modified IL-7 fusion protein according to the present invention.

[0013] FIGS. 2A-2C present graphs showing the results of differential scanning calorimetry (DSC) of a modified IL-7 fusion protein under various basal buffer conditions.

[0014] FIGS. 3A-3C present graphs showing the results of dynamic light scattering (DLS) analysis of a modified IL-7 fusion protein under various basal buffer conditions.

[0015] FIGS. 4A-4C present graphs showing the results of DLS analysis of a modified IL-7 fusion protein under the conditions in which various excipients were added to a histidine-acetate buffer.

[0016] FIGS. 5A-5C present graphs showing the results of DLS analysis of a modified IL-7 fusion protein under the conditions in which various excipients were added to a sodium citrate buffer.

MODE FOR THE PRESENT INVENTION

[0017] Hereinafter, the present invention will be described in detail.

[0018] The present invention provides a pharmaceutical formulation comprising (a) a modified IL-7 fusion protein; (b) a basal buffer with a concentration of 10 to 50 mM; (c) a sugar with a concentration of 2.5 to 5 w/v %; and (d) a surfactant with a concentration of 0.05 to 6 w/v %.

[0019] As used herein, the term “interleukin-7 (IL-7)” refers to a hematopoietic growth factor, which is expressed in or secreted by various stromal cells, keratin-producing cells, dendritic cells, neurons, epidermal cells, etc. IL-7 may promote an immune response by binding to a receptor, and activate a variety of immune cells such as T cell, B cell, mononuclear cell involved in the immune system.

[0020] Such IL-7 is either interleukin-7 (IL-7) or a polypeptide having a similar activity to IL-7.

[0021] IL-7 the present invention includes the polypeptides consisting of the amino acid sequences represented by SEQ ID NOs:1 to 6. In addition, the amino acid sequence of the IL-7 fusion protein may have a sequence homology of about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more to the sequences of SEQ ID NOs: 1 to 6. IL-7 may be an IL-7 fusion protein or a fusion protein comprising a fragment thereof. In addition, IL-7 may be derived from human, rat, mouse, monkey, cow or sheep.

[0022] Specifically, human-derived IL-7 may have the amino acid sequence represented by SEQ ID NO:1 (Genbank Accession No. P13232); rat-derived IL-7 may have the amino acid sequence represented by SEQ ID NO:2 (Genbank Accession No. P56478); mouse-derived IL-7 may have the amino acid sequence represented by SEQ ID NO:3 (Genbank Accession No. P10168); monkey-derived IL-7 may have the amino acid sequence represented by SEQ ID NO:4 (Genbank Accession No. NP_001279008); bovine IL-7 may have the amino acid sequence represented by SEQ ID NO:5 (Genbank Accession No. P26895); and sheep-derived IL-7 may have the amino acid sequence represented by SEQ ID NO:6 (Genbank Accession No. Q28540).

[0023] As used herein, the term “modified IL-7” refers to a molecule wherein an oligopeptide composed of 1 to 10 amino acids is present at the terminal of IL-7, thereby to have a different sequence from the wild type IL-7.

[0024] Preferably, a modified IL-7 according to the present invention has the following structure:

[0025] A-IL-7; [0026] wherein said A may be bound to the N-terminal of the IL-7. The nucleic acid encoding said A may be present with IL-7 genes sequentially on a DNA construct, which may be expressed as a protein. Further, said A may be expressed or synthesized independently, and then bound to IL-7 by a chemical bond.

[0027] Said A may comprise 1 to 10, specifically 2 to 8, more specifically 1 to 5 amino acids, wherein the comprised amino acid may be selected from the group consisting of methionine (M), glycine (G), and a combination thereof. Examples of said A include M, G, MM, MG, GM, GG, MMM, GMM, MGM, MMG, GGM, GMG, MGG, GGG, MMMM (SEQ ID NO:9), GMMM (SEQ ID NO:10), MGMM (SEQ ID NO:11), MMGM (SEQ ID NO:12), MMMG (SEQ ID NO:13), GGMM (SEQ ID NO:14), MGGM (SEQ ID NO:15), MMGG (SEQ ID NO:16), GMGM (SEQ ID NO:17), MGMG (SEQ ID NO:18), GMMG (SEQ ID NO:19), GGGM (SEQ ID NO:20), MGGM (SEQ ID NO:21), GMGM (SEQ ID NO:22), GGMG (SEQ ID NO:23), GGGG (SEQ ID NO:24), MMMMM (SEQ ID NO:25), GMMMM (SEQ ID NO:26), GGMMM (SEQ ID NO:27), GGGMM (SEQ ID NO:28), GGGGM (SEQ ID NO:29), MGMMM (SEQ ID NO:30), MGGMM (SEQ ID NO:31), MGGGM (SEQ ID NO:32), MGGGG (SEQ ID NO:33), MMGMM (SEQ ID NO:34), MMGGM (SEQ ID NO:35), MMGGG (SEQ ID NO:36), MMMGM (SEQ ID NO:37), MMMGG (SEQ ID NO:38), MMMMG (SEQ ID NO:39), MGGGM (SEQ ID NO:40), MGMGM (SEQ ID NO:41), GMGMG (SEQ ID NO:42), GMMMG (SEQ ID NO:43), GGMGM (SEQ ID NO:44), GGMMG (SEQ ID NO:45), MGGMG (SEQ ID NO:46), MGMGG (SEQ ID NO:47), GMMGM (SEQ ID NO:48), MGMMG (SEQ ID NO:49), GMGGM (SEQ ID NO:50), MMGMG (SEQ ID NO:51), GMMGG (SEQ ID NO:52), GMGGG (SEQ ID NO:53), GGMGG (SEQ ID NO:54), GGGMG (SEQ ID NO:55) and GGGGG (SEQ ID NO:56), etc. According to an example of the present invention, said A may be M, MM, GM, MGM or MMM.

[0028] A modified IL-7 fusion protein according to the present invention may further comprise the Fc region of a modified immunoglobulin at the C-terminal thereof.

[0029] As used herein, the term “modified IL-7 fusion protein” refers to a molecule wherein a modified immunoglobulin Fc region is bound to the C-terminal of a modified IL-7.

[0030] Said modified immunoglobulin Fc region may be one whose binding activity to a Fc receptor or to a complement is modified, resulting in weakened antibody-dependent cellular cytotoxicity (ADCC) or complement dependent cytotoxicity (CDC). Such modification may be achieved by genetic mutation (i.e., substitution, deletion of the sequence) of the binding site for Fc gamma receptor (FcγR), C1q, and Fc binding receptor (FcRn) in the Fc region.

[0031] In addition, such modification may be achieved by mixing different types of immunoglobulin sequences, that is, by preparing a hybrid Fc.

[0032] As used herein, the term “Fc region”, “Fc fragment” or “Fc” may include a hinge region, a heavy chain constant region 2 (CH2) domain, and a heavy chain constant region 3 (CH3) domain in the direction from the N-terminal to the C-terminal. Such region may further comprise a hinge region, in which case the hinge region may serve as a linker, and may appropriately be modified to improve the property of the molecule.

[0033] The Fc region or a fragment thereof referred to as “hFc” or “hyFc” is a type of hybrid Fc, which is prepared by combining different types of Fc regions. Such hyFc may include a portion of a human IgD hinge region, an amino acid residue of a human IgD CH2 domain, an amino acid residue of a human IgG4 CH2 domain, and an amino acid residue of a human IgG4 CH3 domain.

[0034] The Fc region variant may be modified so as to prevent truncation at the hinge region. Specifically, the amino acid at position 144 and/or the amino acid at position 145 of SEQ ID NO: 7 can be modified. Preferably, K, the amino acid at position 144 of SEQ ID NO:7, may be substituted with G or S, and E, the amino acid at position 145 of SEQ ID NO:7, may be substituted with G or S, to form a variant.

[0035] In addition, two fusion proteins may form one dimer. For example, when the third domain is Fc region, the Fc regions may bind to each other to form a dimer.

[0036] According to an example of the present invention, the modified immunoglobulin Fc region may be a polypeptide consisting of the amino acid sequence represented by SEQ ID NO:7. The Fc region of a modified immunoglobulin according to the present invention may be one described in U.S. Pat. No. 7,867,491, and can be produced with reference to the description in U.S. Pat. No. 7,867,491.

[0037] According to an example of the present invention, the modified IL-7 may have the amino acid sequence of SEQ ID NO:8. In addition, the modified IL-7 may have a sequence homology of 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% to the amino acid sequence of SEQ ID NO:8,

[0038] As used herein, the term “basal buffer” refers to a solution which endures a pH change by the action of an acid-base conjugate component contained therein.

[0039] A basal buffer according to the present invention may be histidine-acetate or sodium citrate solution. The basal buffer may be used in a concentration of 10 to 50 mM, 15 to 40 mM, or 20 to 30 mM. According to an example of the present invention, it may be used in a concentration of 20 mM.

[0040] The histidine-acetate solution is a buffer containing histidine ions. The histidine-acetate solution is prepared by titrating L-histidine with liquid acetic acid. Meanwhile, the sodium citrate solution is prepared by titrating the sodium citrate solution with hydrous citric acid regarding the final concentration and pH of the solution.

[0041] A sugar stabilizes the fusion protein in the present invention. In addition, a sugar is widely used as an excipient for pharmaceutical formulations as it is easily dissolved in water and has a sweet taste.

[0042] In a formulation according to the present invention, the sugar may be comprised in a

concentration of 2.5 to 5 w/v %, 3.5 to 5 w/v %, or 4.5 to 5 w/v %, and particularly 5 w/v % in an example of the present invention.

[0043] In a formulation according to the present invention, the sugar may be sucrose, trehalose, dextrose, or a mixture thereof. In one example of the present invention, the sugar may be sucrose. Sucrose is a disaccharide formed of glucose and fructose by 1, 2-binding, which is relatively stable in an alkaline environment but may be easily hydrolyzed by an acid.

[0044] In a pharmaceutical formulation according to the present invention, a sugar has the effect of protecting a fusion protein from the oxidation stress, and a sugar alcohol may be used as a sugar substitute or an auxiliary substance. The sugar alcohol may be sorbitol, xylitol, maltitol, mannitol or a mixture thereof. In one example of the present invention, the sugar alcohol may be sorbitol.

[0045] As used herein, the term “surfactant” is a surface activating agent, and specifically, it refers to a non-ionic surfactant. In the pharmaceutical formulation according to the present invention, the surfactant inhibits aggregates formation of a fusion protein, thereby increasing the stability of the fusion protein.

[0046] In the formulation according to the present invention, a surfactant may be added in a concentration of 0.05 to 6 w/v %, 0.1 to 5.5 w/v %, or 0.1 to 5.0 w/v %.

[0047] The surfactant may be polysorbate, polyoxyethylene alkyl ether, polyoxyethylene stearate, alkyl sulfate, polyvinyl pyridone, poloxamer or a mixture thereof. Specifically, the surfactant may be polysorbate or poloxamer.

[0048] The polysorbate according to the present invention is a polyoxyethylene higher aliphatic alcohol formed by combining sorbitan fatty acid ester with ethylene oxide, which may be polysorbate 20 (monolaurate), 40 (monopalmitate), 60 (monostearate), 65 (tristearate), or 80 (mono-oleate) based on the number of the polyoxyethylene group and the kind of the fatty acid. In addition, poloxamer is a polyoxypropylene-polyoxyethylene block copolymer, which may vary depending on the length of the polymer.

[0049] In one example of the present invention, the surfactant may be polysorbate 20, polysorbate 80 or poloxamer 188.

[0050] A pharmaceutical formulation according to the present invention may further comprise an amino acid such as arginine, glutamate, glycine, histidine or a mixture thereof, etc. The amino acid is comprised as an excipient in order to stabilize a protein, and various amino acids may be comprised depending on the type of the protein. The amino acid may be arginine, glutamate, glycine or histidine, and according to an example of the present invention, it may be glutamate. The amino acid may be added in a concentration of 40 to 60 mM, according to an example of the present invention, 50 mM.

[0051] When using a pharmaceutical formulation of the present invention as an injection, it may further comprise a sugar alcohol to adjust the osmotic pressure. The sugar alcohol refers to a long-chain alcohol in which a carbonyl group of a sugar is reduced to a hydroxyl group (—OH). The sugar alcohol may be added such that the concentration thereof reaches 1 to 2 w/v %, and according to an example of the present invention, 1.5 w/v %.

[0052] Examples of sugar alcohol for use in the present invention include sorbitol, xylitol, maltitol, mannitol or a mixture thereof. According to an example of the present invention, the sugar alcohol may be sorbitol. Sorbitol is a sugar alcohol having 6 hydroxyl groups formed by high pressure addition and reduction of glucose, which is also referred to as D-sorbitol or D-glucitol.

[0053] The pharmaceutical formulation of the present invention may have a pH of 5.0 to 6.0, specifically 5.0. The conformational stability of a fusion protein decreases if the pH of the basal buffer is lower than 5.0, while the fusion protein may show aggregates formation depending on a specific composition if the pH exceeds 6.0.

[0054] The formulation may be a liquid formulation.

[0055] The inventors have prepared a modified IL-7 fusion protein in which an oligopeptide is bound to the N-terminal of IL-7 and the Fc region of an immunoglobulin is bound to the C-terminal

of IL-7, and selected a basal buffer and an excipient to prepare a pharmaceutical formulation of the fusion protein

[0056] First, stability of the modified IL-7 fusion protein depending on a basal buffer was examined at various pH conditions. When sodium citrate was used as a basal buffer, stability of the fusion protein increased as the pH increased (Table 2 and FIG. 2), but the zeta potential decreased and aggregates of the fusion protein were formed as the pH increased. When histidine-acetate was used as a basal buffer, similar results were obtained, and thus, the experiment was conducted at the condition of pH 5.0 first (Table 3 and FIG. 3).

[0057] In addition, to examine the protective effect on the fusion protein against the stress environments such as agitation and oxidation, a sugar or a sugar alcohol, a surfactant, and an amino acid were added as excipients. As a result, sucrose and sorbitol showed protective effect on the fusion protein against oxidation stress, while surfactants such as tween 20, Tween 80 and poloxamer 188 showed protective effect on the fusion protein against agitation stress, irrespective of the type of the surfactant. In the cases of further using an amino acid, the protective effect on the fusion protein according to the present invention was observed when arginine, glutamine, glycine, or histidine was added, among which glutamate showed an excellent protective effect (Table 4 and Table 5).

[0058] By the above method, Tween 80 and sucrose were selected, as showing a protective effect on the fusion protein, and the protective effect of the combination of the selected excipients was examined. As a result, a protective effect on the fusion protein against agitation or oxidation stress was shown when Tween 80 and sucrose were mixed in the histidine-acetate basal buffer. Also, the average change of about 1% was observed in a variety of stress conditions when sucrose, sorbitol, and Tween 80 were mixed in the histidine-acetate basal buffer, verifying that such formulation is excellent (Table 7, Table 8 and Table 10).

Best Mode for Carrying Out the Present Invention

[0059] Hereinafter, the present invention is explained in detail by Examples. The following Examples are intended to further illustrate the present invention without limiting its scope.

Example 1: Preparation of Modified Interleukin-7 Fusion Proteins

[0060] Prepared was an IL-7 fusion protein in which an oligopeptide and an immunoglobulin Fc region were bound to the N-terminal and the C-terminal, respectively. The sequence of a human IL-7 (SEQ ID NO:1) was used for IL-7, and the combination sequence of methionine (M) and glycine (G) was used for the oligopeptide. The Fc region is a hybrid of the Fc region of human IgD and the Fc region of human IgG4, which can increase in vivo half-life greatly when binding to a bioactive fusion protein, compared to known modified Fc regions of immunoglobulins.

[0061] First, gene sequence which encodes the amino acid sequence of M-IL-7-hyFc, G-IL-7-hyFc, MM-IL-7-hyFc, MG-IL-7-hyFc, GM-IL-7-hyFc, GG-IL-7-hyFc, MMM-IL-7-hyFc, GMM-IL-7-hyFc, MGM-IL-7-hyFc, MMG-IL-7-hyFc, GGM-IL-7-hyFc, GMG-IL-7-hyFc, MGG-IL-7-hyFc, GGG-IL-7-hyFc, MMMM-IL-7-hyFc, GMMM-IL-7-hyFc, MGMM-IL-7-hyFc, MMGM-IL-7-hyFc, MMMG-IL-7-hyFc, GGMM-IL-7-hyFc, MGGM-IL-7-hyFc, MMGG-IL-7-hyFc, GMGM-IL-7-hyFc, MGMG-IL-7-hyFc, GMMG-IL-7-hyFc, GGGM-IL-7-hyFc, MGGG-IL-7-hyFc, GMGG-IL-7-hyFc, GGMG-IL-7-hyFc, GGGG-IL-7-hyFc, MMMMM-IL-7-hyFc, GMMMM-IL-7-hyFc, GGMMM-IL-7-hyFc, GGGMM-IL-7-hyFc, GGGGM-IL-7-hyFc, MGMMM-IL-7-hyFc, MGGMM-IL-7-hyFc, MGGGM-IL-7-hyFc, MGGGG-IL-7-hyFc, MMGMM-IL-7-hyFc, MMGGM-IL-7-hyFc, MMGGG-IL-7-hyFc, MMMGM-IL-7-hyFc, MMMGG-IL-7-hyFc, MMMMG-IL-7-hyFc, MGMGM-IL-7-hyFc, GMGMG-IL-7-hyFc, GMMMG-IL-7-hyFc, GGMMG-IL-7-hyFc, MGMMG-IL-7-hyFc, MGMGG-IL-7-hyFc, GMMGG-IL-7-hyFc, GMGGG-IL-7-hyFc, GGMGG-IL-7-hyFc, GGGMG-IL-7-hyFc or GGGGG-IL-7-hyFc was inserted into pAD15 expression vector. In order to insert a fusion protein gene into an expression vector pAD15, EcoRI site was generated at the 5'-end of the fusion protein gene

sequence, and XbaI site was generated at the 3'-end of the hyFc gene sequence. Expression vector pAD15 was obtained from RcCMV backbone (available at Invitrogen, Carlsbad). The pAD15 contains a cytomegalovirus (CMV)-derived promoter, fetal bovine growth hormone-derived poly (A) sequence, rabbit beta globin-derived gIVS (globin intervening sequence) (Mol Cell Biol. 1988, 8: 4395), and other factors. The pAD15 vector was prepared by modifying various parts of RcCMV vector (Invitrogen). First, neomycin resistance site was removed by XhoI enzyme, and gIVS was added to the 3'-end of the CMV promoter region. In addition, mouse dihydrofolate reductase (DHFR) gene (GeneBank Accession No. NM 010049) was added to the 5'-end of the CMV promoter. In order to prepare the connection site between the 3'-end of IL-7 and the 5'-end of hyFc in the frame, NheI site was generated at the 3'-end of IL-7 coding sequence and the 5'-end of hyFc coding sequence. The final expression vector was prepared through a subcloning process using each restriction enzyme site. For more information regarding the preparation of the modified Fc region, U.S. Pat. No. 7,867,491 may be referred to (indicated as "hybrid (Fc) protein"). The hyFc protein of the present Example contains 9 amino acids of the C-terminal of IgD CH1 domain (90-98), 30 amino acids of IgD hinge region (133-162), 8 amino acids of the N-terminal of IgD CH2 domain (SHTQPLGV 163-170), 100 amino acids of IgG4 CH2 domain (121-220), and 107 amino acids of IgG4 CH3 domain (221-327).

[0062] Regarding the hyFc protein composed of a modified Fc region, the present invention takes reference to every preparation method for hFc-2, hFc-3, hFc-4, hFc-5 and hFc-6 described in the Example of the Korean Patent Publication No. 10-0897938. The hyFc protein used in the Examples of the present invention is a hybrid protein identical to hFc-5 described in the Korean Patent Publication No. 10-0897938.

[0063] PCR was carried out under the condition shown in Table 1. Denaturation, annealing and extension processes were carried out for 15 seconds at 98° C., for 30 seconds at 55° C., and for 60 seconds at 72° C., respectively.

TABLE-US-00001

TABLE 1	pAD15 template	comprising the gene	2 µl of a
modified IL-7 fusion protein	Forward primer	1.5 µl (5'-accgaattcatgttcacgtgagcttcag-3')	Reverse primer
	1.5 µl (5'-gggtctagattagtgtccttggtgcccatc-3')	5 × HF buffer	20 µl
2.5 mM dNTP	8 µl DMSO	3 µl Phusion (Thermo Scientific)	1 µl Tertiary distilled water
64 µl			

[0064] The prepared expression vector, which is M-IL-7-hyFc, G-IL-7-hyFc, MM-IL-7-hyFc, MG-IL-7-hyFc, GM-IL-7-hyFc, GG-IL-7-hyFc, MMM-IL-7-hyFc, GMM-IL-7-hyFc, MGM-IL-7-hyFc, MMG-IL-7-hyFc, GGM-IL-7-hyFc, GMG-IL-7-hyFc, MGG-IL-7-hyFc, GGG-IL-7-hyFc, MMMM-IL-7-hyFc, GMMM-IL-7-hyFc, MGMM-IL-7-hyFc, MMGM-IL-7-hyFc, MMMG-IL-7-hyFc, GGMM-IL-7-hyFc, MGGM-IL-7-hyFc, MMGG-IL-7-hyFc, GMGM-IL-7-hyFc, MGMG-IL-7-hyFc, GMMG-IL-7-hyFc, GGGM-IL-7-hyFc, MGGG-IL-7-hyFc, GMGG-IL-7-hyFc, GGGG-IL-7-hyFc, MMMMM-IL-7-hyFc, GMMMM-IL-7-hyFc, GGMMM-IL-7-hyFc, GGGMM-IL-7-hyFc, GGGGM-IL-7-hyFc, MGMMM-IL-7-hyFc, MGGMM-IL-7-hyFc, MGGGM-IL-7-hyFc, MGGGG-IL-7-hyFc, MMGMM-IL-7-hyFc, MMGGM-IL-7-hyFc, MMGGG-IL-7-hyFc, MMMGM-IL-7-hyFc, MMMMG-IL-7-hyFc, MGMGM-IL-7-hyFc, GMGMG-IL-7-hyFc, GMMMG-IL-7-hyFc, GGMMG-IL-7-hyFc, MGMMG-IL-7-hyFc, GMGMG-IL-7-hyFc, MGMMG-IL-7-hyFc, GMMMG-IL-7-hyFc, GMGGG-IL-7-hyFc, GGMGG-IL-7-hyFc, GGGMG-IL-7-hyFc or GGGGG-IL-7-hyFc, was transfected into CHO DG44 cells (from Dr. Chasm, Columbia University, U.S.A.) by an electroporation method. After 48 hours, HT selection was carried out using 10% dFBS+MEM a medium without HT, and MTX amplification was carried out using HT selected clones to amplify the productivity. The obtained single cells were used for the preparation of a modified IL-7 fusion protein after testing the long-term stability thereof.

Example 2: Purification of Modified IL-7 Fusion Protein

[0065] After obtaining the culture solution containing the modified IL-7 fusion protein prepared by the method of Example 1, the productivity was measured by performing size-exclusion (SE) HPLC according to the conditions shown in Table 2 below. SE-HPLC was carried out for 40 minutes using TSK-GEL G3000WxL column.

[0066] First, each of the sample solutions of the modified IL-7 fusion protein was diluted with a buffer to a concentration of 1 mg/ml. Each diluent was filtered with 1 ml syringe and 0.2 μ m filter and then put into an insert. The insert was inserted into a vial and the cap was kept closed. Then, each of the test solutions was injected into SE-HPLC system in an amount of 20 μ l. The purity and relative productivity were measured based on the peak value.

[0067] As a result, the modified IL-7 fusion protein with the structure shown in FIG. 1, which has high productivity and purity, was purified (preparation and purification of the modified IL-7 fusion protein were carried out according to Korea Patent Application No. 10-2015-0082793).

Experimental Example 1: Selection of Basal Buffer

1.1. Selection Based on Thermodynamic Property

[0068] To establish a formulation of the modified IL-7 fusion protein purified in Example 2 above, the thermodynamic property of the fusion proteins in three types of buffer was examined by differential scanning calorimetry (DSC) method.

[0069] First, the modified IL-7 fusion protein purified in Example 2 was subjected to dialysis at 4° C. using 20 mM histidine-acetate buffer or sodium citrate buffer with a pH of 5.0, 5.5, or 6.0, or tris buffer with a pH of 8.0.

[0070] Meanwhile, DSC was carried out using Microcal VP-DSC (Northampton, U.S.A.) having twin cells with the volume of 0.5147 cm³ regarding the samples and a control solution. The scanning was conducted at the speed of 1.25° C./min at the temperature from 15° C. to 110° C. Before measuring the values, every solution was agitated in a vacuum state to remove air. Then, the standard value for a solution without the fusion protein was first determined, and the temperature record of a modified IL-7 fusion protein was shown by the values determined by subtracting the standard value from the measured values.

[0071] The resulting graph was shown by using Microcal LLC DSC (Northampton, U.S.A.) which was connected with Origin 7.0 software package provided with the device. In addition, from the results, the transition temperature (T_m) and enthalpy (Δ H) values of the fusion protein were obtained by using Origin 7.0 software.

[0072] As shown in FIG. 2, in the case of MGM-IL-7-hyFc fusion protein, T_m 1 and T_m 2, the transition temperatures of IL-7 and hyFc, respectively, and two peaks representing the fusion protein structure unfolded by heat were identified in the typical DSC thermogram regarding the three types of buffers. These results indicate that each of IL-7 and hyFc exists in an unfolded state due to heat energy absorption, and they are structurally stable.

[0073] In general, hyFc is a design platform for the preparation of a long-acting protein in which IgD (hinge-C.sub.H2) and IgG4 (C.sub.H2-C.sub.H3) are hybridized, and has single or double transitions from IgG4 according to various environmental factors. However, the result of the present experiment showed that apparently hyFc had a single transitional peak.

[0074] In addition, as shown in Table 2 below, T_m 1 and T_m 2 respectively increased from 58.89° C. to 62.92° C. and from 66.93° C. to 69.91° C., as the pH of the histidine-acetate buffer increased from 5.0 to 6.0. Moreover, Δ H also increased along with T_m. These results indicate that as pH increases, higher temperature and more heat energy are necessary for unfolding the structure of IL-7 and hyFc. That is, conformational stability of IL-7 and hyFc protein might increase as pH increases.

[0075] Even when sodium citrate with the pH of 5.0, 5.5, or 6.0 was used as a basal buffer, T_m 1 and T_m 2 increased as the pH increased, which results were identical to the results obtained by using a histidine-acetate buffer. It should be noted that T_m 1 and T_m 2, respectively, increased from 57.23° C. to 65.12° C. and from 66.74° C. to 71.34° C. as the pH increased from 5.0 to 6.0.

[0076] Regarding ΔH value when using sodium citrate as a basal buffer, ΔH values of Tm 1 and Tm 2 were the lowest when pH was 5.0. This result indicates that the suitable pH of a basal buffer for stably sustaining the structure of the fusion protein is 5.5 or 6.0.

[0077] When tris buffer of pH 8.0 was used, Tm 1, Tm 2 and ΔH value of Tm 1 were higher than those obtained with other buffers. This result was consistent with the aforementioned results which showed that the conformational stability of a modified IL-7 fusion protein increased as pH increased. In the case of tris buffer with a high pH, Tm 2 value increased compared to other buffers with a pH of 6.0, but the ΔH value of Tm 2 decreased, indicating that it might have bigger influence on the conformational stability of IL-7 than on the stability of hyFc.

TABLE-US-00002 TABLE 2 Buffer/pH Transition Enthalpy (MGM-IL-7-hyFc temperature ($^{\circ}$ C.) (kJ/mol) concentration, mg/ml) Tm 1 Tm 2 $\Delta H1$ $\Delta H2$ Histidine-acetate pH 5.0 (8.6) 58.89 66.93 263.47 102.91 pH 5.5 (8.8) 61.11 68.72 290.68 104.17 pH 6.0 (8.6) 62.92 69.91 332.06 115.41 Sodium citrate pH 5.0 (9.3) 57.23 66.74 242.15 110.48 pH 5.5 (9.6) 62.17 70.28 305.27 124.77 pH 6.0 (9.8) 65.12 71.34 292.10 126.19 Tris pH 8.0 (7.8) 71.53 66.06 384.27 104.58

1.2. Selection by Dynamic Light Scattering (DLS) Analysis

[0078] DLS analysis was carried out in order to evaluate the electrostatic interaction between proteins, and Z-average size, polydispersity index (PDI) and zeta potential, and the effect of pH and a buffer on the aggregates formation. The particle size of the protein and zeta potential were measured using Zetasizer Nano ZS90 (Malvern Instruments, UK).

[0079] Z-average size refers to the average of elements such as a fragment, a monomer and an aggregate, which is very sensitive to even small changes such as the presence of a small aggregate portion. If the measured PDI value is lower than 0.05, it is shown as a monodisperse standard, and if the value is higher than 0.7, it indicates that the sample has a wide size distribution. Zeta potential is an electrical repulsion of proteins surrounding a hydrodynamic surface. If the absolute value of the zeta potential is high, the protein is less subject to aggregation and instability in a solution.

[0080] First, for the DLS analysis, a modified IL-7 fusion protein sample was equilibrated at the measurement temperature of 10° C. Using 1 ml of the sample, particle size was measured 5 times in a disposable sizing cuvette, and zeta potential was measured 3 times in a disposable capillary cell with the angle fixed to 90° . Intensity of the peak, average particle size, PDI and the zeta potential were calculated using Zetasizer software version 7.11 (Malvern Instruments, UK), which was provided with the device.

[0081] The protein size distribution represented by the intensity of protein peaks when using histidine-acetate or sodium citrate of pH 5.0, or tris of pH 8.0 as a basal buffer was shown in FIG. 3. The Z-average size, PDI and zeta potential of the three buffers in different pH conditions were shown in Table 3.

[0082] As shown in FIG. 3, when histidine-acetate or sodium citrate of pH 5.0 was used as a basal buffer, the resulting graph showed a single narrow peak throughout 5 consecutive measurements. Meanwhile, when tris buffer of pH 8.0 was used, a new peak was identified, which suggested aggregation at 100 to 1,000 nm.

[0083] As shown in Table 3, when histidine-acetate was used as a basal buffer, new aggregation appeared as pH increased from 5.0 to 6.0, and the Z-average size and PDI, respectively, increased from 10.43 nm to 10.64 nm, and 0.02 to 0.07. In addition, as pH increased, the absolute value of zeta potential decreased from 4.42 to 1.28.

[0084] The aggregation phenomenon according to the pH change as described above was also observed when sodium citrate was used as a basal buffer: that is, Z-average size and PDI increased and zeta potential decreased in a sodium citrate basal buffer.

[0085] However, Z-average size and PDI in a tris basal buffer highly increased to 12.38 nm and 0.34, respectively, as compared to those in other buffers. In addition, protein aggregation took place in the tris basal buffer. Also, even though aggregation took place, the absolute value of zeta

potential in the tris buffer was 20.27, which was higher than those in other buffers.

[0086] Therefore, in order to reduce the aggregation of the modified IL-7 fusion protein of the present invention, histidine-acetate or sodium citrate may be suitable as a basal buffer, preferably in the pH condition with low zeta potential. In a high pH condition, the buffer can be used in combination with protein aggregation inhibitor.

TABLE-US-00003 TABLE 3 Buffer/pH Z-average Absolute (MGM-IL-7-hyFc size value of zeta concentration, mg/ml) (d .Math. nm*) PDI potential (mV) Histidine-acetate pH 5.0 (8.6) 10.43 ± 0.06 0.02 ± 0.01 4.42 ± 0.32 pH 5.5 (8.8) 10.58 ± 0.09 0.03 ± 0.02 1.11 ± 0.43 pH 6.0 (8.6) 10.64 ± 0.24 0.07 ± 0.05 1.28 ± 0.47 Sodium citrate pH 5.0 (9.3) 10.73 ± 0.09 0.02 ± 0.01 5.18 ± 0.74 pH 5.5 (9.6) 10.82 ± 0.10 0.03 ± 0.02 3.38 ± 0.57 pH 6.0 (9.8) 11.03 ± 0.17 0.08 ± 0.03 2.46 ± 0.42 Tris pH 8.0 (7.8) 12.38 ± 0.84 0.34 ± 0.05 20.27 ± 0.42 *d .Math. nm: diameter in nanometer

Experimental Example 2: Selection of Excipients

2.1. Selection according to the protective effect on stress conditions

[0087] In order to examine the protective effect of excipients against stress environments such as agitation and oxidation, various excipients including a surfactant with a concentration of 5 w/v %, a sugar with a concentration of 5 w/v % and an amino acid with a concentration of 50 mM, were subjected to a size-exclusion chromatography. In the control group, only modified IL-7 fusion protein was added to a basal buffer. For the basal buffer, histidine-acetate or sodium citrate basal buffer of pH 5.0 was used.

[0088] Specifically, the modified IL-7 fusion proteins were dialyzed at 4° C. with the solutions prepared by adding various types of excipients to histidine-acetate or sodium citrate basal buffers of pH 5.0. For the agitation stress environment, 0.5 ml of the modified IL-7 fusion protein as dialyzed was put into a microtube, which was then put into a box and agitated for 2 hours at room temperature using Vortex-Genie 2 (Scientific Industries, Inc., U.S.A.) at strength 7.

[0089] Meanwhile, for the oxidation stress environment, 0.5 ml of the dialyzed MGM-IL-7-hyFc fusion protein was added with hydrogen peroxide to the final concentration of 1.0% (v/v) and stored for 20 hours at room temperature. Size-exclusion chromatography was carried out using high performance liquid chromatography (HPLC) system (Waters e2695, U.S.A.) at the UV wavelength of 214 nm of absorption spectrum.

<HPLC Condition>

[0090] Column: TSKgel G3000SWXL SEC column (300×7.8 mm) (TOSOH Bioscience, U.S.A.)

[0091] Mobile phase: solution containing 100 mM NaCl and 10% acetonitrile in 50 mM sodium phosphate (pH 6.8) [0092] Flow rate: 0.5 ml/min

[0093] As a result, several elements including water-soluble aggregation, monomers and fragments were observed, and the content of each element was calculated by the mathematical equation 1.

[00001] $\% \text{content of specific element} = C_t / C_s \times 100$ [Equation 1]

[0094] Herein, C_t refers to the area of each element (water-soluble aggregation, monomer and fragment), and C_s refers the sum of the peaks of the elements when a sample is obtained.

[0095] As a result, as shown in Tables 4 and 5, the monomer content in a sodium citrate buffer in the control group was 96.34%, which decreased to 78.18% by agitation and 48.16% by oxidation.

[0096] It was found that MGM-IL-7-hyFc fusion protein was protected from the agitation stress by adding widely-known non-ionic surfactant such as Tween or poloxamer to a histidine-acetate or sodium citrate buffer. Such non-ionic surfactants have an excellent effect of protecting the protein from uncoiling in a hydrophobic interface. When Twin 20, Twin 80 or poloxamer 188 were added to a buffer, the monomer content did not significantly decrease. However, the protective effects of these surfactants were substantially similar, indicating that the surfactants protected the fusion protein of the present invention in a hydrophobic environment which constantly changes due to agitation.

[0097] Meanwhile, the surfactants did not show any protective effect on MGM-IL-7-hyFc fusion

protein against oxidation stress, but rather caused more reduction of the monomer content than the control group. Moreover, Twin 20 and Twin 80 are known to oxidate a protein. Poloxamer 188 also made the fusion protein of the present invention unstable in the oxidation condition, as Twin 20 and Twin 80 did.

[0098] A sugar or a sugar alcohol such as sucrose and sorbitol is often used as a pharmaceutical excipient for stabilizing MGM-IL-7-hyFc fusion protein. However, sucrose and sorbitol caused lower monomer content than the control group, which indicated that they do not have a protective effect on the fusion protein of the present invention against agitation stress.

[0099] After agitation in the sodium citrate buffer, based on the reference value not exposed to a stress, the monomer content decreased by 18.16% in the control group, while the monomer contents decreased by 39.29% and 53.51% when sucrose and sorbitol, respectively, were added. Meanwhile, after agitation in the histidine-acetate buffer, based on the reference value not exposed to a stress, the monomer content decreased by 20.23% in the control group, while the monomer contents decreased by 11.53% and 130.77% when sucrose and sorbitol, respectively, were added.

[0100] It is known that the addition of a sugar and a sugar alcohol to a formulation stimulates aggregates formation of IgG during agitation. When sucrose or sorbitol were added to a formulation, the aggregates formation increased during agitation, which led to the decrease in the stability of MGM-IL-7-hyFc fusion protein, irrespective of the kind of a basal buffer.

[0101] Meanwhile, sucrose and sorbitol increased the stability of a fusion protein under the oxidation stress. In a sodium citrate buffer, the monomer content decreased by 48.18% in the control group based on the reference value not exposed to a stress, while the monomer contents decreased by 2.55% and 2.79% when sucrose and sorbitol, respectively, were added. Similarly, in a histidine-acetate buffer, the monomer content decreased by 35.36% in the control group based on the reference value not exposed to a stress, while the monomer contents decreased by 1.68% and 1.47% when sucrose and sorbitol, respectively, were added. This result indicates that sucrose and sorbitol have a protective effect on protein from oxidation. The two substances are reported to be able to stabilize a protein by preferential exclusion from a surface of a protein.

[0102] Next, with regard to the effect of an amino acid on the stability of MGM-IL-7-hyFc fusion protein under a stress condition, the protective effect against agitation was very low when 50 mM arginine, glutamate, glycine and histidine amino acids were respectively added to sodium citrate buffers. The monomer content decreased by 18.16% in the control group based on the reference value not exposed to a stress, while the monomer contents decreased by 49.71%, 64.75%, 3.96% and 46.75% when arginine, glutamate, glycine and histidine were, respectively, added. Meanwhile, arginine and glutamate were found to make a fusion protein of the present invention unstable when histidine-acetate buffer was used.

[0103] Meanwhile, when glycine was added to a histidine-acetate buffer, the monomer content of the MGM-IL-7-hyFc fusion protein did not decrease as compared to the reference value not exposed to a stress. The result was conflicting to the result from a sodium citrate buffer, suggesting that the stability of MGM-IL-7-hyFc fusion protein can change depending on the environmental factors including a buffer and an excipient.

[0104] Moreover, the protective effect against oxidation stress was observed when arginine and glycine were respectively added to sodium citrate buffers, but such effect was not observed when they were added to histidine-acetate buffers. However, with regard to the oxidation stress, when arginine and glycine were respectively added to sodium citrate buffers, the samples showed decreases of 30.66% and 3.21% respectively in the monomer contents based on the reference value not exposed to a stress, which were smaller than in the control group (decreased by 48.18%). Meanwhile, when arginine and glycine were added to histidine-acetate buffers, the monomer contents decreased significantly by 39.34% and 37.90% respectively based on the reference value not exposed to a stress, compared to the control group (decreased by 35.36%). Especially, in both types of buffers, the addition of glycine showed a protective effect on the fusion protein of the

present invention against agitation and oxidation stresses depending on buffer types, and the addition of glutamate resulted in an excellent protective effect on the fusion protein against oxidation stress.

TABLE-US-00004 TABLE 4 Sodium citrate pH 5.0 [MGM-IL-7-hyFc concentration: 9.5 mg/ml (Final concentration 3.0 mg/ml)] Aggregate (%) Monomer (%) Fragment (%) Excipient Reference* Agitation Oxidation Reference* Agitation Oxidation Reference* Agitation Oxidation Control group 0.81 20.58 47.98 96.34 78.18 48.16 2.85 1.24 3.86 Sugar & Sucrose 0.69 41.78 0.65 95.65 56.36 93.10 3.66 1.86 6.25 Sugar Sorbitol 0.57 55.79 0.98 96.08 42.57 93.29 3.34 1.64 5.74 alcohol Surfactant Tween 20 0.57 0.84 48.13 95.74 95.77 48.02 3.69 3.39 3.85 Tween 80 0.69 0.64 48.02 95.76 95.80 47.34 3.55 3.56 4.63 Poloxamer 0.69 0.74 50.17 95.71 95.68 46.45 3.60 3.58 3.38 188 Amino Arginine 0.75 52.65 31.39 95.62 45.91 64.96 3.36 1.44 3.65 acid Glutamate 0.55 67.66 0.81 95.60 30.85 93.73 3.85 1.48 5.46 Glycine 0.69 34.09 2.44 95.76 64.80 92.55 3.55 1.11 5.01 Histidine 0.77 50.34 22.37 95.70 48.95 73.53 3.53 0.72 4.10 *Note: Not exposed to a stress

TABLE-US-00005 TABLE 5 Histidine-acetate pH 5.0 [MGM-IL-7-hyFc concentration: 9.5 mg/ml (Final concentration 3.0 mg/ml)] Aggregate (%) Monomer (%) Fragment (%) Excipient Reference* Agitation Oxidation Reference* Agitation Oxidation Reference* Agitation Oxidation Control group 0.45 4.52 35.03 95.39 93.16 60.03 4.16 2.33 4.94 Sugar & Sucrose 0.42 13.71 0.52 95.62 84.09 93.94 3.96 2.20 5.54 Sugar Sorbitol 0.46 15.80 0.37 95.74 81.97 94.27 3.80 2.23 5.36 alcohol Surfactant Tween 20 0.43 0.55 39.36 95.32 95.73 56.59 4.25 3.72 4.06 Tween 80 0.36 0.60 40.93 96.00 95.30 55.29 3.64 4.10 3.79 Poloxamer 0.50 0.64 41.89 95.77 95.87 54.37 3.73 3.49 3.75 188 Amino Arginine 0.51 29.11 39.35 96.04 69.26 56.70 3.45 1.63 3.95 acid Glutamate 0.42 56.11 0.77 96.22 39.40 93.87 3.36 1.50 5.36 Glycine 0.53 2.93 38.42 95.55 94.83 57.65 3.92 2.24 3.93 *Note: Not exposed to a stress

2.2. Selection by DLS Analysis

[0105] To examine the effect of various excipients including a surfactant, a sugar and an amino acid on the stability of a modified IL-7 fusion protein, DLS analysis was conducted by the method described in Experimental Example 1.2. The measurements of a particle size and zeta potential were carried out three times, respectively.

[0106] Z-average size of MGM-IL-7-hyFc fusion proteins which were exposed to the agitation and oxidation stress in sodium citrate buffer with pH 5.0 was shown in FIG. 4 and Table 6.

TABLE-US-00006 TABLE 6 Sodium citrate pH 5.0 Reference* Agitation Oxidation Z-average Z-average Z-average Excipient size (d .Math. nm) PDI size (d .Math. nm) PDI size (d .Math. nm) PDI) Control group 11.29 0.07 19.40 0.19 177.10 0.85 Sugar & Sucrose 11.80 0.17 — — 13.07 0.28 Sugar Sorbitol 11.89 0.14 — — 14.11 0.32 alcohol Surfactant Tween 20 10.61 0.07 10.77 0.11 — — Tween 80 10.69 0.03 10.78 0.07 — — Poloxamer 11.01 0.05 11.32 0.11 — — 188 Amino acid Glutamate 11.09 0.06 — — 11.26 0.06 Glycine 11.65 0.13 — — 50.71 0.58 *Note: Not exposed to a stress

[0107] As shown in FIG. 4a, the control group showed only a single narrow peak in the particle size measurement which was conducted three times. However, the agitation control group, which was exposed to an agitation stress, showed another peak at the size of 100 nm (see arrow in the middle graph of FIG. 4a). In this control group, Z-average size increased from 11.29 nm to 19.40 nm, and PDI also increased from 0.07 to 0.19, indicating that an aggregation formation took place. Meanwhile, when Twin 20, Twin 80 and poloxamer 188 were added, Z-average size increased by 0.16, 0.09, 0.31 nm, respectively, and PDI increased by 0.04, 0.04 and 0.06, respectively. The increase in Z-average size and PDI implies that the aggregation is induced by agitation. A smaller increase was observed by the addition of a surfactant, indicating that the surfactants had a protein protective effect from aggregates formation. As shown in FIG. 4b, an aggregate peak was not observed when a surfactant was added.

[0108] The oxidation control group, which was exposed to an oxidation stress, showed an additional peak at the size of 1,000 nm (see arrow in the bottom graph of FIG. 4a), while Z-average

size and PDI increased from 11.29 nm to 177.10 nm, and 0.07 to 0.85 nm, respectively. This result indicates that an aggregate is formed by oxidation, and the formed aggregate is bigger than that formed by agitation. As shown in FIG. 4c, an aggregate peak was also observed when sucrose, sorbitol and glycine were respectively added (see arrows), but aggregates were not formed when glutamate was added. In the case of glutamate-added sample, Z-average size increased by 0.17 nm and PDI did not change. This result indicates that when glutamate is added to sodium citrate buffer with pH 5.0, it shows an excellent protective effect on MGM-IL-7-hyFc fusion protein against oxidation stress. Meanwhile, when sucrose, sorbitol and glycine were added, Z-average size increased by 1.27 nm, 2.22 nm and 39.06 nm, respectively, and PDI increased by 0.11, 0.18 and 0.45, respectively, which were considered to be small when compared to the samples in which other excipients were added. This result indicates that sucrose, sorbitol and glycine have a protective effect on the fusion protein of the present invention against oxidation stress, which is consistent with the result of Experimental Example 2.1.

[0109] Meanwhile, the Z-average size of MGM-IL-7-hyFc fusion proteins which were exposed to agitation and oxidation stress in histidine-acetate buffer with pH 5.0 was shown in FIG. 5 and Table 7.

TABLE-US-00007 TABLE 7 Histidine-acetate pH 5.0 Reference* Agitation Oxidation Z-average Z-average Z-average Excipient size (d .Math. nm) PDI size (d .Math. nm) PDI size (d .Math. nm) PDI Control group 10.50 0.02 11.70 0.10 17.43 0.37 Sugar & Sucrose 11.21 0.17 — — 12.47 0.22 Sugar Sorbitol 12.02 0.13 — — 11.61 0.09 alcohol Surfactant Tween 20 10.66 0.10 10.79 0.12 — — Tween 80 10.40 0.03 10.50 0.03 — — Poloxamer 188 10.83 0.04 10.74 0.07 — — Amino acid Glutamate 10.90 0.04 — — 11.31 0.06 Glycine 10.68 0.03 15.66 0.41 — — *Note: Not exposed to a stress

[0110] As shown in FIG. 5a, the control group and the agitation control group showed a single narrow peak, which is a different result from the agitation control group with the sodium citrate buffer (see arrow in bottom graph). However, as shown in FIG. 5b, an aggregate peak was observed when glycine was added (see arrows in bottom graph), but an aggregate peak was not observed and Z-average size and PDI did not change when surfactants such as Twin 20, Twin 80 and poloxamer 188 were added. This result indicates that an aggregates formation can take place by agitation when glycine is added to a MGM-IL-7-hyFc fusion protein formulation.

[0111] As shown in FIG. 5c, when sucrose was added, Z-average size and PDI, respectively, increased from 11.21 nm to 12.47 nm, and 0.17 to 0.22 by oxidation (see arrows in top graph). Both of the changes were smaller than the control group, indicating that sucrose has a protein protective effect against oxidation stress, but cause the problem of aggregates formation. Meanwhile, when sorbitol was added, Z-average size and PDI respectively decreased from 12.02 nm to 11.61 nm, and 0.13 to 0.09, while no aggregate peak was found. The decreases in Z-average size and PDI suggest the increase of fragments, even if not significant. When glutamate was added, Z-average size and PDI increased slightly, while neither aggregates nor fragments were found.

[0112] Therefore, it was found that the surfactants such as Twin 20, Twin 80 and poloxamer 188 have a protective effect on the fusion protein of the present invention against oxidation or agitation stress; a sugar or a sugar alcohol such as sucrose and sorbitol exhibits a protective effect against oxidation stress; and amino acids show different effects according to the basal buffer, but overall, glutamate shows a high protective effect.

Experimental Example 3: Combination of Excipients

[0113] From the results of Experimental Example 2, Tween 80 and sucrose were selected as the excipients showing a protective effect on the modified IL-7 fusion protein against agitation and oxidation stresses, respectively. For experiment for the combination of these two excipients, Design of Experiment software (Design-Expert, Stat-Ease Inc., U.S.A.) was used. The two excipients were combined as shown in Table 8, and the effect thereof on stability of the modified IL-7 fusion protein was examined. Stress condition in the present Experimental Example was the same as

Experimental Example 2.1., and the final concentration of the modified IL-7 fusion protein in the formulation was adjusted to 3 to 100 mg/ml.

TABLE-US-00008 TABLE 8 Excipient (Addition concentration w/v %) Tween 80 Sucrose

Sample	Excipient	Concentration (w/v %)	Basal	Final	SE-HPLC (%)
1	Tween 80	0.00	0	95.65	74.77
2	Sucrose	0.05	0	95.06	95.39
3	Tween 80	0.10	0	95.39	95.28
4	Sucrose	0.00	2.5	94.42	71.26
5	Tween 80	0.05	2.5	95.08	95.19
6	Sucrose	0.10	2.5	95.06	95.85
7	Tween 80	0.00	5	95.29	71.70
8	Sucrose	0.05	5	95.51	95.02
9	Tween 80	0.10	5	95.42	95.27
10	Sucrose	0.05	2.5	94.96	95.30

[0114] To examine the protective effect against a stress condition, size-exclusion chromatography was carried out [at 214 nm of UV wavelength of absorption spectrum using high performance liquid chromatography (HPLC) system (Waters e2695, U.S.A.)], under the same HPLC condition as Experimental Example 2.1.

[0115] First, an experiment was conducted regarding a formulation comprising the modified IL-7 fusion protein in a concentration of 3 mg/ml. As a result, as shown in Table 9, the control group with sodium citrate buffer showed a monomer content of 95.65%, which decreased to 74.77% by agitation, and to 49.81% by oxidation. The samples in which Tween 80 was added showed a protective effect against agitation stress, while those in which sucrose was added showed a protective effect against oxidation stress. The samples in which Tween 80 and sucrose were added in combination showed the protective effect against agitation stress and oxidation stress.

TABLE-US-00009 TABLE 9 Basal Final concentration SE-HPLC (%) Buffer Sample Excipient (w/v %) Reference* Agitation Oxidation Sodium

Sample	Excipient	Concentration (w/v %)	Basal	Final	SE-HPLC (%)
1	Tween 80	0.00	0	95.65	74.77
2	Sucrose	0.05	0	95.06	95.39
3	Tween 80	0.10	0	95.39	95.28
4	Sucrose	0.00	2.5	94.42	71.26
5	Tween 80	0.05	2.5	95.08	95.19
6	Sucrose	0.10	2.5	95.06	95.85
7	Tween 80	0.00	5	95.29	71.70
8	Sucrose	0.05	5	95.51	95.02
9	Tween 80	0.10	5	95.42	95.27
10	Sucrose	0.05	2.5	94.96	95.30

*Note: Not exposed to a stress

[0116] Meanwhile, as shown in Table 10, the control group with histidine-acetate buffer showed a monomer content of 95.46%, which decreased to 93.60% by agitation, and to 50.63% by oxidation. As compared to the sodium citrate buffer, the histidine-acetate buffer showed more excellent protective effect against agitation stress, and the protective effects of the added excipients was similar to those in the samples with sodium citrate.

TABLE-US-00010 TABLE 10 Basal Final concentration SE-HPLC (%) buffer Sample Excipient (w/v %) Reference* Agitation Oxidation Histidine-

Sample	Excipient	Concentration (w/v %)	Basal	Final	SE-HPLC (%)
1	Tween 80	0.00	0	95.46	93.60
2	Sucrose	0.05	0	95.77	94.62
3	Tween 80	0.10	0	95.35	95.30
4	Sucrose	0.00	2.5	95.49	94.93
5	Tween 80	0.05	2.5	95.52	94.36
6	Sucrose	0.10	2.5	95.75	94.82
7	Tween 80	0.00	5	95.34	94.86
8	Sucrose	0.05	5	95.29	94.90
9	Tween 80	0.10	5	95.38	94.42
10	Sucrose	0.05	2.5	94.90	94.90

[0117] From the results, the condition of the excipients was selected. To acquire the proper osmotic strength of an injection, sorbitol and mannitol were further added, and agitation stress and oxidation stress experiments were conducted, and additional experiment under the condition of adding 1.5 w/v % sorbitol was also conducted.

[0118] As a candidate formulation for each of sodium citrate buffer and histidine-acetate buffer, 5 w/v % sucrose, 1.5 w/v % sorbitol, and 0.05 w/v % Tween 80 were added to the formulations for MGM-IL-7-hyFc fusion protein. Using the above formulations, freeze/thaw, temperature stress and oxidation stress experiments were conducted. The stress conditions are shown in Table 11, and the analysis result is shown in Table 12.

TABLE-US-00011 TABLE 11 Stress condition Condition Stress method Control group -80° C. Freeze/thaw -80° C..fwdarw.Room temperature (5 times) Temperature 37° C. (39 h, 3 day), 50° C. (20 h) Oxidation 0.1% or 1% H.sub.2O.sub.2, room temperature 20 h

TABLE-US-00012 TABLE 12 Osmotic pH pressure SE-HPLC (%) Before Before Before stress stress stress 37° C., 50° C., 0.1% 1% Basal buffer exposure exposure exposure Freeze/thaw 39 h 20 h H.sub.2O.sub.2 H.sub.2O.sub.2 20 mM Sodium citrate, 5.02 303 95.76 95.77 94.83 93.81 95.71 92.86 5 w/v % Sucrose, 1.5 w/v % Sorbitol, 0.05 w/v % Tween 80, pH 5.0 20 mM Histidine-

acetate, 4.98 300 95.59 95.64 95.41 95.47 95.95 93.83 5 w/v % Sucrose, 1.5 w/v % Sorbitol, 0.05 w/v % Tween 80, pH 5.0

[0119] As a result of the stress test using the candidate formulations, as shown in Table 12, in this case where a sodium citrate buffer is used, the control group showed a monomer content of 95.76%, and about 3% of change by stress was observed.

[0120] In the case where a histidine-acetate buffer is used, the control group showed a monomer content of 95.59%, and about 1% of change by stress was observed, showing a slightly better protective effect compared to the sodium citrate buffer.

[0121] The effect of the candidate formulations on the modified IL-7 fusion protein of various concentrations was examined. First, a formulation comprising 3 to 100 mg/ml modified IL-7 fusion protein, 20 mM sodium citrate, and 5 w/v % sucrose was prepared. Then, 1 to 2 w/v % sorbitol or mannitol as a sugar alcohol, and 0.05 w/v % Tween 80 or poloxamer as a surfactant were added thereto, and the final pH was adjusted to 5.0. The result indicated that, 3 to 100 mg/ml of modified IL-7 fusion protein was well protected under the various combination conditions.

Claims

1. A pharmaceutical formulation comprising: (a) an interleukin-7 (IL-7) protein, which is conjugated to a Fc region of an immunoglobulin (IL-7 fusion protein); (b) a basal buffer with a concentration of 10 to 50 mM; (c) a sugar with a concentration of 2.5 to 5 w/v %; and (d) a surfactant with a concentration of 0.05 to 6 w/v %.
2. The pharmaceutical formulation of claim 1, wherein the modified IL-7 fusion protein comprises an oligopeptide consisting of 1 to 10 amino acid residues.
3. The pharmaceutical formulation of claim 2, wherein the oligopeptide is selected from the group consisting of M, G, MM, MG, GM, GG, MMM, GMM, MGM, MMG, GGM, GMG, MGG, GGG, MMMM (SEQ ID NO:9), GMMM (SEQ ID NO:10), MGMM (SEQ ID NO:11), MMGM (SEQ ID NO:12), MMMG (SEQ ID NO:13), GGMM (SEQ ID NO:14), MGGM (SEQ ID NO:15), MMGG (SEQ ID NO:16), GMGM (SEQ ID NO:17), MGMG (SEQ ID NO:18), GMMG (SEQ ID NO:19), GGGM (SEQ ID NO:20), MGGG (SEQ ID NO:21), GMGG (SEQ ID NO:22), GGMG (SEQ ID NO:23), GGGG (SEQ ID NO:24), MMMMM (SEQ ID NO:25), GMMMM (SEQ ID NO:26), GGMMM (SEQ ID NO:27), GGGMM (SEQ ID NO:28), GGGGM (SEQ ID NO:29), MGMMM (SEQ ID NO:30), MGGMM (SEQ ID NO:31), MGGGM (SEQ ID NO:32), MGGGG (SEQ ID NO:33), MMGMM (SEQ ID NO:34), MMGGM (SEQ ID NO:35), MMGGG (SEQ ID NO:36), MMMGM (SEQ ID NO:37), MMMGG (SEQ ID NO:38), MMMMG (SEQ ID NO:39), MGGGM (SEQ ID NO:40), MGMGM (SEQ ID NO:41), GMGMG (SEQ ID NO:42), GMMMG (SEQ ID NO:43), GGMGM (SEQ ID NO:44), GGMMG (SEQ ID NO:45), MGGMG (SEQ ID NO:46), MGMGG (SEQ ID NO:47), GMMGM (SEQ ID NO:48), MGMMG (SEQ ID NO:49), GMGGM (SEQ ID NO:50), MMGMG (SEQ ID NO:51), GMMGG (SEQ ID NO:52), GMGGG (SEQ ID NO:53), GGMGG (SEQ ID NO:54), GGGMG (SEQ ID NO:55), and GGGGG (SEQ ID NO:56).
4. The pharmaceutical formulation of claim 3, wherein the oligopeptide is bound to the N-terminal of the IL-7 protein.
5. The pharmaceutical formulation of claim 2, wherein the IL-7 protein comprises: (i) amino acid residues 26-177 of SEQ ID NO: 1, (ii) amino acid residues 26-154 of SEQ ID NO: 2, (iii) amino acid residues 26-154 of SEQ ID NO: 3, (iv) amino acid residues 26-177 of SEQ ID NO: 4, (v) amino acid residues 26-176 of SEQ ID NO: 5, or (iv) amino acid residues 26-176 of SEQ ID NO: 6.
6. The pharmaceutical formulation of claim 2, wherein the Fc region of the immunoglobulin is bound to the C-terminal of the IL-7 protein.
7. The pharmaceutical formulation of claim 6, wherein the Fc region of the immunoglobulin has the amino acid sequence of SEQ ID NO:7.

8. The pharmaceutical formulation of claim 1, wherein the basal buffer is histidine-acetate or sodium citrate.
9. The pharmaceutical formulation of claim 1, wherein the sugar is selected from the group consisting of sucrose, trehalose, dextrose, and a mixture thereof.
10. The pharmaceutical formulation of claim 1, wherein the surfactant is selected from the group consisting of polysorbate, polyoxyethylene alkyl ether, polyoxyethylene stearate, alkyl sulfates, polyvinyl pyridone, poloxamer and a mixture thereof.
11. The pharmaceutical formulation of claim 1, further comprising any one amino acid selected from the group consisting of arginine, glutamate, glycine, histidine, and a mixture thereof.
12. The pharmaceutical formulation of claim 11, wherein the concentration of the amino acid ranges from 40 to 60 mM.
13. The pharmaceutical formulation of claim 12, wherein the concentration of the amino acid is 50 mM.
14. The pharmaceutical formulation of claim 1, further comprising a sugar alcohol of 1 to 2 w/v %.
15. The pharmaceutical formulation of claim 14, wherein the sugar alcohol is selected from the group consisting of sorbitol, xylitol, maltitol, mannitol, and a mixture thereof.
16. The pharmaceutical formulation of claim 1, wherein the pH of the formulation is 5.0.
17. The pharmaceutical formulation of claim 1, wherein the formulation is a liquid formulation.
- 18-19. (canceled)
20. A method of increasing the stability of an interleukin-7 (IL-7) fusion protein, comprising admixing the IL-7 fusion protein in a formulation comprising: (i) a basal buffer with a concentration of 10 to 50 mM, (ii) a sugar with a concentration of 2.5 to 5 w/v %, and (c) a surfactant with a concentration of 0.05 to 6 w/v %, wherein the IL-7 fusion protein comprises an IL-7 protein which is conjugated to a half-life extending moiety.
21. A method of reducing an aggregation in a pharmaceutical formulation comprising an interleukin-7 (IL-7) fusion protein, comprising admixing the IL-7 fusion protein in a basal buffer with a concentration of 10 to 50 mM, wherein the basal buffer is selected from a histidine-acetate or sodium citrate, and wherein the IL-7 fusion protein comprises an IL-7 protein which is conjugated to a half-life extending moiety.
22. A method of treating a disease or disorder in a subject in need thereof, comprising administering to the subject the pharmaceutical formulation of claim 1.
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