

PHARMACEUTICAL COMPOSITION COMPRISING RECOMBINANT HEMOGLOBIN PROTEIN OR SUBUNIT-BASED THERAPEUTIC AGENT FOR CANCER TARGETING TREATMENT

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CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] The present application claims priority from the US provisional patent application serial number 62/019,925 filed July 2, 2014 and US non-provisional patent application serial number 14/752,999 filed June 28, 2015, and the disclosure of which is incorporated herein by reference in its entirety.

TECHNICAL FIELD

[0002] The present invention provides recombinant hemoglobin protein, recombinant hemoglobin tetramer, dimer and subunit including α , β , $\gamma 1$, $\gamma 2$ monomer (with or without heme molecule) with oxygen carrying capacity to replace the natural hemoglobin protein. The invention also provides method of using the recombinant hemoglobin protein, the recombinant hemoglobin tetramer, dimer or subunit for use in oxygenation of *in vivo* and *ex vivo* tissue. The

present invention further provides recombinant hemoglobin protein-, the recombinant hemoglobin tetramer-, dimer-, or subunit-based therapeutic agent that at least one of the hemoglobin subunit has been chemically modified to create a material having the ability to target and/or kill cancer cells. The present invention additionally provides a method for construction and expression of different recombinant hemoglobin subunits including α , β , $\gamma 1$, $\gamma 2$ monomer (with or without heme molecule), and the dimer and tetramer thereof and for chemical modification or engineering thereof. The present invention further describes a design for chemical engineering for creating a recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit-based therapeutic agent. The present invention further relates to recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit-based therapeutic agent-containing pharmaceutical compositions for cancer targeting treatment in humans and other animals, in particular, for liver cancer, breast cancer, pancreatic cancer, and tumor induced or associated with respective progenitor cells.

BACKGROUND OF INVENTION

[0003] Hemoglobin-based oxygen carriers (HBOCs) were initially developed as blood substitutes. Human placenta hemoglobin is a particularly promising HBOC. It allows transporting more oxygen to hypoxic tissues owing to its higher oxygen affinity, lower viscosity, and smaller mean diameter than human red blood cells. However, the major problems posed by the use of placenta hemoglobin on a large scale are caused by the conditions under which it is extracted from the placentas and purified. Placentas are kept in frozen storage and to extract hemoglobin they have to be crushed before thawing. Then, tissues and fibrinogen are removed by filtration and several steps of chromatography, precipitation and diafiltration are carried out on the resulting filtrate. Also, the sources of human placenta hemoglobin and human blood are

limited. Therefore, the construction and expression of recombinant hemoglobin protein or recombinant hemoglobin tetramer or dimer or subunits are important to solve this issue.

[0004] Hypoxia is common in cancers. Hypoxia and anemia (which contributes to tumor hypoxia) can lead to ionizing radiation and chemotherapy resistance by depriving tumor cells of the oxygen essential for the cytotoxic activities of these agents. Hypoxia may also reduce tumor sensitivity to radiation therapy and chemotherapy through one or more indirect mechanisms that include proteomic and genomic changes. Therefore, there is a need for improved cancer-treatment compositions, particularly, improved cancer-treatment compositions that facilitate targeting cancer cells and enhance the efficacy of cytotoxic agents.

Summary of the Invention

[0005] Accordingly, in the present invention, a recombinant hemoglobin protein, recombinant hemoglobin tetramer, dimer, and/or subunit is/are constructed with oxygen carrying capacity. The present invention provides recombinant hemoglobin protein, recombinant hemoglobin tetramer, dimer and subunit including α , β , $\gamma 1$, $\gamma 2$ monomer (with or without heme molecule) with oxygen carrying capacity to replace the natural hemoglobin protein. In addition, a therapeutic agent based on any of the recombinant hemoglobin protein, recombinant hemoglobin tetramer, dimer, and/or subunit of the present invention capable of targeting cancer cells in order to efficiently kill cancer cells by a conjugated therapeutic drug (e.g. chemotherapeutic agent, radiotherapeutic agent, anti-cancer protein drug) is provided. Common chemotherapeutic agents, radiotherapeutic agents and anti-cancer protein drugs which are widely used in different patients, however, have many side-effects are found. These problems can be overcome by chemically modifying any of the recombinant hemoglobin protein, tetramer, dimer, or subunit(s) of the present invention and linking thereof to one or more therapeutic drugs. Therefore, the present

invention further provides recombinant hemoglobin protein-, the recombinant hemoglobin tetramer-, dimer-, or subunit-based therapeutic agent that has been chemically modified to create a material having the ability to target and/or kill cancer cells. The present invention additionally provides a method for construction and expression of different recombinant hemoglobin subunits including α , β , $\gamma 1$, $\gamma 2$ monomer (with or without heme molecule), and the dimer and tetramer thereof and for chemical modification or engineering thereof. The present invention further describes a design for chemical engineering for creating a recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit-based therapeutic agent. The present invention further relates to recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit-based therapeutic agent-containing pharmaceutical compositions for cancer targeting treatment in humans and other animals, in particular, for liver cancer, breast cancer, pancreatic cancer, and tumor induced or associated with respective progenitor cells. When compared to the conventional therapeutic drugs alone for treating cancer (e.g. chemotherapeutic drug including 5-Fluorouracil, Temozolomide, Cisplatin), the recombinant hemoglobin protein-, tetramer-, dimer- or subunit-based therapeutic agents of the present invention not only can target cancer cells, but are much more efficacious in the treatment of cancer and/or tumors since the killing effect on the cancer cells/tumor is further enhanced. Further, since the cancer-targeting recombinant hemoglobin protein or recombinant hemoglobin tetramer or dimer or subunit-based therapeutic agents can be used in a relatively lower dose than other known hemoglobin-based oxygen carriers, the adverse side effect from the conventional therapeutic drug for cancers is greatly attenuated.

[0006] Most conventional therapeutic drugs are very expensive. The treatment cost can be cut down significantly for each patient if the therapeutically effective dose is lowered. Recombinant hemoglobin protein or recombinant hemoglobin subunit-based therapeutic agents of the present

invention are a good candidate for lowering the therapeutically effective dose as the modified recombinant hemoglobin protein or recombinant hemoglobin subunit can target cancer cells. In one embodiment, the therapeutically effective dose of the recombinant hemoglobin protein or recombinant hemoglobin tetramer or dimer or subunit-based therapeutic agent for treating cancer in human can be 0.0024 mg/kg (for a single dose) of the subject's body weight or lower. It can be used in treating cancer in multiple doses (once per week). The invention also provides method of using the recombinant hemoglobin protein, the recombinant hemoglobin tetramer, dimer or subunit for use in oxygenation of *in vivo* and *ex vivo* tissue.

[0007] Optionally, the presently claimed recombinant hemoglobin protein or recombinant hemoglobin tetramer or dimer or subunit-based therapeutic agent can also be linked to fluorescent probe(s) to facilitate the live-cell imaging. The recombinant hemoglobin protein or tetramer or dimer or subunit-based therapeutic agent conjugated with fluorescein can be taken up by liver cancer cells. The uptake of freshly fluorescein conjugated recombinant hemoglobin protein- or tetramer- or dimer- or subunit-based therapeutic agents by cells is verified by immediately employing the same to the cells in a series of live cell uptake studies as described hereinafter. In one example, the fluorescein conjugated recombinant hemoglobin protein- or tetramer- or dimer- or subunit-based therapeutic agent is shown to be taken up by liver cancer cells (e.g. HepG2 cell line).

[0008] Therefore, in the first aspect of the present invention, different recombinant hemoglobin proteins, recombinant tetramer, recombinant dimer, and/or recombinant hemoglobin subunits (e.g. α , β , $\gamma 1$, $\gamma 2$; with or without heme molecule), and the dimer and tetramer being formed thereof are provided, and said recombinant hemoglobin proteins, tetramer, dimer, and subunits are also provided for further chemical engineering or modification. A method for constructing

and expressing said recombinant hemoglobin proteins or subunit(s), and the dimer and tetramer thereof in a host is also provided.

[0009] The second aspect of the present invention is to construct a chemically modified recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit with one or more functional groups that can be used as a linkage or linker (cleavable or non-cleavable) to at least one type of therapeutic drug for targeting the cancer cells. In one embodiment, the chemical reactions involved in chemically modifying the recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit of the present invention include but not limited to amine reactions, thiol reactions, carboxylate reactions, hydroxyl reactions, native chemical ligations using thioesters, incorporation of bioorthogonal functionalities, photochemical reactions, and metal-mediated reactions. One or more hemoglobin subunits of the recombinant hemoglobin protein can be chemically modified.

[0010] The third aspect of the present invention is to chemically link the recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit of the present invention to at least one type of therapeutic drug (active agent) *via* said cleavable or non-cleavable linkage or linker in order to kill the cancer cells. The therapeutic drug or active agent which can be linked to the recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit of the present invention includes but is not limited to a chemotherapeutic drug (e.g., 5-Fluorouracil, Temozolomide, Cisplatin), a radiotherapeutic drug (e.g., Rhodium-105 complex, Samarium-153 complex and other related complex), an anti-cancer protein drug (e.g. arginase, arginine deiminase), any other therapeutic drug and/or compound which is proven to be effective for treating and/or alleviating cancer and capable of being readily linked to the recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit of the present invention, through the cleavable or non-cleavable linkage or linker to the recombinant

hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit or to the chemically modified recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit in the second aspect of the present invention.

[0011] The present invention further relates to recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit-based therapeutic agent-containing pharmaceutical composition for cancer targeting treatment in humans and other animals. The composition includes a therapeutically effective amount of said therapeutic agent and a pharmaceutically acceptable carrier, salt, buffer, water, or a combination thereof, in order for targeting and treating cancer. In one embodiment, the pharmaceutical composition has a pH in a range of 5.5 to 9.5. In another embodiment, the pharmaceutical composition has a pH in a range of 7.2 to 8.0.

[0012] In the fourth aspect of the present invention, it is provided a method for targeting and/or treating cancer comprising administering a pharmaceutical composition comprising a therapeutically effective amount of the recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit-based therapeutic agent of the present invention to a subject in need thereof suffering from various tumors and/or cancers. The composition can be administered to the subject by various routes including but not limited to intravenous injection, intraperitoneal injection, and subcutaneous injections. Both cleavable and non-cleavable forms of the recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit-based therapeutic agent containing an active agent such as chemotherapeutic agent (e.g. 5-Fluorouracil, 5FU) can be prepared for cancer targeting treatment in the present invention.

[0013] The recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit-based therapeutic agent of the present invention is also chemically modified to facilitate the targeting of the therapeutic agent to cancer cells such that it is more efficient to kill the cancer cells. The recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or

subunit can be chemically modified and linked to different therapeutic agents (e.g. 5FU, Temozolomide, Cisplatin, etc). The recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit can target cancer cells. This targeting property facilitates killing cancerous cells, cancer stem cells and/or cancer progenitor cells efficiently *via* inducing apoptosis of these cells. As such, a dose of the therapeutic agent can be lowered.

[0014] The recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit-based therapeutic agent provided in the present invention can be used in the treatment of various cancers such as pancreatic cancer, leukemia, head and neck cancer, colorectal cancer, lung cancer, breast cancer, liver cancer, nasopharyngeal cancer, esophageal cancer and brain cancer. The present invention is directed to recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit-based therapeutic agent, to methods of treating cancer, and to methods of treating and/or inhibiting metastasis of cancerous tissue and recurrence of cancerous tissue, including but not limited to liver cancer.

[0015] Cells within a tumor are heterogeneous in nature. A tumor is generally thought to be made up of (1) a majority of cancer cells with limited ability to divide, and (2) a rare population of cancer stem-like cells (CSCs), also known as progenitor cells, which can form new tumor cells and are highly metastatic in nature. Due to their inherent properties of being chemoresistant and metastatic, CSCs have been postulated to be responsible for recurrence in cancer patients.

[0016] Since the recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit moiety of the present recombinant hemoglobin protein or tetramer, dimer or subunit-based therapeutic agent can bring oxygen to CSCs or progenitor cells to facilitate killing of the cancer stem cells or the progenitor cells while the active agent moiety of the present recombinant hemoglobin protein or subunit-based therapeutic agent can kill the cancer cells, the present

recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit-based therapeutic agent of the present invention has a synergistic effect in cancer treatment.

BRIEF DESCRIPTION OF THE DRAWINGS

[0017] **FIG. 1** shows the design approach for construction of recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit-based therapeutic agent. One or more therapeutic drugs can be linked to recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit (e.g. α , β , $\gamma 1$, $\gamma 2$) to form the recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit-based therapeutic agent. The recombinant hemoglobin protein or recombinant hemoglobin tetramer, dimer or subunit can be chemically linked to therapeutic agent *via* cleavable (**A**) or non-cleavable linker/linkage (**B**). Cleavable linkers can be but not limited to carbinolamine, disulfide, carbamide, aminal, carbonate, ester, carbamate, phosphate, amide, acetal, imine, ether, and sulfonamide groups; non-cleavable linkers can be but not limited to alkyl and aryl groups.

[0018] **FIG. 2** depicts the amino acid sequence of different hemoglobin subunits (α , β , $\gamma 1$, $\gamma 2$) of human origin (*Homo sapiens*) and the design for recombinant hemoglobin monomer (i.e., monomer α , monomer β , monomer $\gamma 1$, and monomer $\gamma 2$), dimer (i.e., dimer $\alpha\beta$, dimer $\alpha\gamma 1$, dimer $\alpha\gamma 2$, dimer $\beta\gamma 1$, and dimer $\beta\gamma 2$) and tetramer (i.e., tetramer $2\alpha\beta_2$, tetramer $2\alpha\gamma 1_2$, tetramer $2\alpha\gamma 2_2$, tetramer $2\beta\gamma 1_2$, tetramer $2\beta\gamma 2_2$).

[0019] **FIG. 3** is a flow chart depicting an overview of the method for preparing the recombinant hemoglobin protein or tetramer or dimer or subunit of the present invention from cells transformed with different constructs expressing the proteins of different hemoglobin tetramer, dimer or subunits with the amino acid sequence described herein.

[0020] **FIG. 4** shows the growth curve of *E. coli* cells JM109(DE3) for the expression of recombinant hemoglobin subunit gamma-1 ($\gamma 1$).

[0021] **FIG. 5** shows molecular size of (A) purified recombinant hemoglobin subunits (α , β , $\gamma 1$, $\gamma 2$) and (B) purified recombinant hemoglobin dimer and tetramer in SDS-PAGE.

[0022] **FIG. 6** shows the purified recombinant hemoglobin subunits (α , β , $\gamma 1$, $\gamma 2$) with or without heme, labeled with fluorescent dyes.

[0023] **FIG. 7** shows the oxygen equilibrium curve for (A) one of the recombinant hemoglobin tetramer ($2\alpha\beta_2$) and another one of the recombinant hemoglobin tetramer ($2\alpha\gamma 1_2$) of the present invention.

[0024] **FIG. 8** depicts electrospray ionization mass spectrometry (ESI-MS) analysis for one of the recombinant hemoglobin dimers of the present invention ($\alpha\beta$).

[0025] **FIG. 9** depicts electrospray ionization mass spectrometry (ESI-MS) analysis for one of the recombinant hemoglobin tetramers of the present invention ($2\alpha\beta_2$).

[0026] **FIG. 10** illustrates that the fluorescent labeled recombinant hemoglobin subunits (α , β , $\gamma 1$, $\gamma 2$) with heme (+) or without heme (-) can enter into liver cancer cells (HepG2) successfully.

[0027] **FIG. 11** shows the liquid chromatography–mass spectrometry (LC-MS) results for (A) 5FU-linked recombinant hemoglobin tetramer ($2\alpha\beta_2$) and (B) 5FU-linked recombinant hemoglobin tetramer ($2\alpha\gamma 1_2$).

[0028] **FIG. 12** shows (A) the cleavage of *N*-(2-mercaptoethyl)benzamide (model compound) with disulfide bond under reducing condition (i.e., GSH (0.4-1,000 equiv.); PBS (pH 7.4); 37°C, 0-2 hrs) and (B) the conversion rate of the model compound at different GSH levels: GSH (1 equiv.) is similar to that in blood stream while GSH (1,000 equiv.) is similar to that in cytoplasm.

[0029] **FIG. 13** shows the efficacy of native bovine crosslinked hemoglobin (Hb) tetramer and recombinant hemoglobin tetramer of the present invention ($2\alpha\beta_2$) with or without Gemcitabine

(GEM) in reducing tumor size in terms of the tumor weight (g) in a pancreatic cancer Capan-1 animal model as compared to a control and a group of animal model treated with GEM only; Group 1: PBS buffer (Control); Group 2: Gemcitabine (GEM), 100 mg/kg; Group 3: Native bovine crosslinked hemoglobin (Hb) tetramer, 30 mg/kg; Group 4: Native bovine crosslinked hemoglobin tetramer + Gemcitabine (Hb + GEM); Group 5: Recombinant hemoglobin tetramer ($2\alpha\beta_2$), 30 mg/kg; Group 6: Recombinant hemoglobin tetramer ($2\alpha\beta_2$) + Gemcitabine.

[0030] **FIG. 14** shows the efficacy of native bovine crosslinked hemoglobin (Hb) tetramer and recombinant hemoglobin tetramer of the present invention ($2\alpha\gamma_1_2$) with or without Gemcitabine (GEM) in reducing tumor size in terms of the tumor weight (g) in a pancreatic cancer Capan-1 animal model as compared to a control and a group of animal model treated with GEM only; Group 1: PBS buffer (Control); Group 2: Gemcitabine (GEM), 100 mg/kg; Group 3: Native bovine crosslinked hemoglobin (Hb) tetramer, 30 mg/kg; Group 4: Native bovine crosslinked hemoglobin tetramer + Gemcitabine (Hb + GEM); Group 5: Recombinant hemoglobin tetramer ($2\alpha\gamma_1_2$), 30 mg/kg; Group 6: Recombinant hemoglobin tetramer ($2\alpha\gamma_1_2$) + Gemcitabine.

DEFINITIONS

[0031] The term “cancer stem cell” refers to a biologically distinct cell within a neoplastic clone that is capable of initiating and sustaining tumor growth *in vivo* (i.e. the cancer-initiating cell).

[0032] The term “cleavable conjugate” refers to a conjugate with at least one cleavable linker or linkage that can easily release a linked therapeutic drug/ active agent by hydrolysis or redox reaction.

[0033] The term “non-cleavable conjugate” refers to a conjugate with at least one non-cleavable linker or linkage and it cannot easily release a linked therapeutic drug/ active agent by hydrolysis or redox reaction.

[0034] The term “recombinant hemoglobin protein/ proteins” refers to hemoglobin molecule in a molecular size of at least approximately 65 KDa and synthesized by any standard molecular chemistry techniques and not being isolated or purified from any animal or human source. In the present invention, the recombinant hemoglobin protein/ proteins can include but not limited to the recombinant hemoglobin tetramer, dimer, and subunit (monomer) of the present invention (with or without heme) or be used interchangeably with the recombinant hemoglobin tetramer.

[0035] The term “recombinant hemoglobin tetramer” refers to hemoglobin molecule in a molecular size of at least approximately 64 KDa and synthesized by any standard molecular chemistry technique and not being isolated or purified from any animal or human source. In the present invention, the recombinant hemoglobin tetramer can comprise any four of the recombinant hemoglobin subunits described herein (with or without heme) or any four of the hemoglobin subunits synthesized by any standard molecular chemistry technique and not being isolated or purified from any animal or human source.

[0036] The term “recombinant hemoglobin dimer” refers to hemoglobin molecule in a molecular size of at least approximately 32 KDa and synthesized by any standard molecular chemistry techniques and not being isolated or purified from any animal or human source. In the present invention, the recombinant hemoglobin dimer can comprise any two of the recombinant hemoglobin subunits described herein or any two of the hemoglobin subunits synthesized by any standard molecular chemistry technique and not being isolated or purified from any animal or human source.

[0037] The term “recombinant hemoglobin subunit/subunits” refers to hemoglobin subunit or a fragment thereof in a molecular size of approximately 16 KDa or less and synthesized by standard molecular chemistry techniques and not being isolated or purified from animal or human source, which can be modified with or without heme group.

DETAILED DESCRIPTION OF THE INVENTION

[0038] Since most cancerous tissues, such as cancerous tumors, are hypoxic, they can become resistant to conventional chemotherapeutic/radiotherapeutic agent. Recombinant hemoglobin proteins or recombinant hemoglobin tetramer or dimer or subunits of the present invention can be used to alleviate this hypoxic condition. It allows transporting more oxygen to hypoxic tissues owing to its higher oxygen affinity, lower viscosity, and smaller mean diameter than human red blood cells, leading to reduction of chemotherapeutic/radiotherapeutic/other anti-cancer drug-resistance in cancer cells.

[0039] In one embodiment of the present invention, the recombinant hemoglobin protein or tetramer or dimer or subunit is produced. The recombinant hemoglobin protein or tetramer or dimer or subunit has the oxygen transport feature and can target cancerous cells or tissues in a human or animal body. The recombinant hemoglobin protein or recombinant hemoglobin tetramer or dimer or subunit of the present invention as an oxygen carrier is chemically modified and linked to at least one type of therapeutic drug, e.g., the chemotherapeutic agent, capable for triggering a receptor-mediated mechanism and leading a combined/conjugated chemotherapeutic agent to localize together with the recombinant hemoglobin protein or recombinant hemoglobin tetramer or dimer or subunit in at least the cytoplasm of the cancerous cells in order to increase the efficacy of both recombinant hemoglobin protein or tetramer or dimer or subunit and the chemotherapeutic agent in reducing tumor size leading to treatment of cancer. Preferably, said cancer is resistant to conventional therapeutic drugs such as chemotherapeutic and/or radiotherapeutic agents when administered solely.

[0040] A design for construction of a recombinant hemoglobin protein or recombinant hemoglobin tetramer or dimer or subunit-based therapeutic drug is shown in FIG. 1A and FIG.

1B. One or more active agents (or “therapeutic drug” are used interchangeably herein) are linked to the recombinant hemoglobin protein or recombinant hemoglobin tetramer or dimer or subunit to form the presently claimed recombinant hemoglobin protein- or subunit-based therapeutic agent. The selection of one or more particular active agent(s) can be made depending upon the type of cancer tissue to be targeted and the desired molecular size of the resulting chemically modified material. Furthermore, the selected active agents may be the same or different in the case of more than one active agents. That is, an active agent (or “therapeutic drug”), etc., as long as the resultant molecule retains the efficacy and is also able to link with the recombinant hemoglobin protein or tetramer or dimer or subunit of the present invention for targeting the cancer cells. The recombinant hemoglobin protein or tetramer or dimer or subunit of the present invention can be chemically linked to therapeutic drug/ active agent *via* cleavable (FIG. 1A) or non-cleavable linkage or link (FIG. 1B) or, alternatively, can be directly conjugated with the active agent in the absence of said linkage or link. Different chemical groups can be used for chemical modification of the recombinant hemoglobin protein or tetramer or dimer or subunit in the present invention and the recombinant hemoglobin protein or tetramer or dimer or subunit can be linked to the therapeutic drug/ active agent *via* these chemical groups. In one embodiment, the cleavable linkers can be but are not limited to carbinolamine, disulfide, carbamide, amination, carbonate, ester, carbamate, phosphate, amide, acetal, imine, oxime, ether, and sulfonamide groups; the non-cleavable linkers can be but not limited to alkyl and aryl groups.

[0041] FIG. 2 shows the amino acid sequences alignment of different hemoglobin subunits (α , β , $\gamma 1$, $\gamma 2$) and the design of recombinant hemoglobin monomer, dimer and tetramer. The recombinant hemoglobin can be produced using sequences from various species including, but not limited to, human, bovine, porcine, equine, and canine hemoglobin. In this example, different *Homo sapiens* hemoglobin subunit amino acid sequences with about 141-146 amino acids,

namely alpha, beta, gamma 1, and gamma 2 (SEQ ID Nos. 1 to 4), are selected for design of the DNA construct to be inserted into an expression vector and optionally tagged with His-sumo/Poly-His sequence for separation and purification after being expressed in a host (e.g., bacteria) which is transformed with the expression vector carrying the corresponding DNA construct. The process for producing the recombinant hemoglobin protein / subunits is outlined in the flow chart of FIG. 3. Any two of the subunits, e.g. alpha and beta subunits or alpha and gamma 1 subunits, can be expressed by using another expression vector carrying corresponding DNA sequences expressing alpha and beta subunits or alpha and gamma 1 subunits. After purification following the process illustrated in the flow chart of FIG. 3, dimers $\alpha\beta$, $\alpha\gamma_1$, $\alpha\gamma_2$, $\beta\gamma_1$, and $\beta\gamma_2$ can be formed, respectively. Further recombination of more than two subunits, e.g. two alpha subunits and two beta subunits, or two alpha subunits and two gamma-1 subunits, or two alpha subunits and two gamma-2 subunits, to form a tetramer, i.e. $2\alpha\beta_2$, $2\alpha\gamma_1_2$, $2\alpha\gamma_2_2$, $2\beta\gamma_1_2$, $2\beta\gamma_2_2$ respectively, can also be formed using similar approach. Further modification such as cross-linking between two hemoglobin subunits within the tetramer can be performed. Schematic diagram of the design of amino acid sequence(s) of the recombinant hemoglobin subunit, dimer, and tetramer is shown in FIG. 2A, 2B, and 2C, respectively.

[0042] Process depicted in FIG. 3 is based on a standard molecular chemistry technique for expressing a recombinant protein in a host system transformed with an expression vector design for carrying the corresponding DNA construct in order for expressing the corresponding hemoglobin subunit, dimer, or tetramer of the present invention. Other possible method which can express the recombinant protein can also be used for expressing the present recombinant hemoglobin protein/tetramer/dimer/subunit. Initially, a seed culture with transformed host cells carrying the DNA construct for expressing the corresponding hemoglobin subunit is prepared (301) for fermentation (302). The cells are harvested by centrifugation (302). The harvested cells

are then lysed to isolate the crude protein (303). The crude protein is then purified using affinity chromatography (304). The salt in the eluate from the affinity chromatography is removed by a desalting column and/or ultrafiltration (305). The semi-purified protein is further purified by an ion-exchange column chromatography (306) in order to remove the protein impurities and endotoxin (307). The process of the present invention is applicable to large-scale industrial production of recombinant monomeric, dimeric and tetrameric hemoglobin. In addition, the recombinant hemoglobin protein/tetramer/dimer/subunit in combination with a pharmaceutical carrier (e.g. water, physiological buffer, in capsule) is suitable for mammalian use.

[0043] The construction and expression of different recombinant hemoglobin proteins or recombinant hemoglobin subunits are carried out in *E. coli*. For example, *E. coli* JM109 (DE3) transformed with expression vector carrying the DNA construct for expression of Gamma-1 ($\gamma 1$) subunit are seeded and cultured for about 16 hours in order to harvest the cells for cell lysis and protein extraction. The growth curve of *E. coli* JM109 (DE3) for expressing Gamma-1 ($\gamma 1$) subunit is shown in FIG. 4. The recombinant hemoglobin proteins/tetramer/dimer/subunits are then purified by column chromatography methods. The recombinant hemoglobin subunits (α , β , $\gamma 1$, $\gamma 2$) expressed and purified according to the process of the present invention are shown in FIG. 5A and the recombinant hemoglobin dimer and tetramer expressed and purified according to the process of the present invention are shown in FIG. 5B. FIG. 6 shows that the unpurified but fluorescent labeled subunit with or without heme.

[0044] A p_{50} value of a test compound, meaning the oxygen partial pressure necessary to produce 50 percent saturation of hemoglobin in the present invention, is measured and the result is shown in FIG. 7. A native human hemoglobin has a p_{50} value on the order of approximately 23-30 mm Hg. FIG. 7 shows the oxygen equilibrium curve for (A) the recombinant hemoglobin tetramer ($2\alpha\beta_2$) and (B) the recombinant hemoglobin tetramer ($2\alpha\gamma 1_2$). As shown in FIG. 7, the

recombinant hemoglobin tetramer $2\alpha\beta_2$ has a p50 value on the order of ~15 mm Hg and the recombinant hemoglobin tetramer $2\alpha\gamma_1$ has a p50 value on the order of ~21 mm Hg. Recombinant hemoglobin with a relatively higher oxygen affinity and a lower p50 value of less than approximately 23 mm Hg is formed, compared with native human hemoglobin which has a p50 value on the order of 23-30 mm Hg. Compositions containing hemoglobin-based oxygen carrier with lower oxygen affinity are used when rapid oxygenation is desired in cases of tissue hypoxia resulting from extensive blood loss (e.g., hemorrhagic shock). Lower oxygen affinity means that the material can “offload” oxygen to a target more easily than a material with a higher oxygen affinity. Compositions with higher oxygen affinity are useful as oxygenation adjunct therapies in cancer treatment where a slower delivery rate of oxygen is desired in that case.

[0045] For use in the treatment of oxygen-deprivation disorders and for heart preservation, the hemoglobin-based oxygen carrier-containing pharmaceutical composition with a lower oxygen affinity of the present invention provides oxygen to a target organ or subject. The present recombinant hemoglobin protein or tetramer or dimer or subunit with lower oxygen affinity is useful for applications requiring rapid tissue oxygenation (e.g. hemorrhagic shock and *ex vivo* organ preservation).

[0046] For applications in cancer treatment, the hemoglobin-based oxygen carrier-containing pharmaceutical composition with a higher oxygen affinity of the present invention serves as a tissue oxygenation agent to improve the oxygenation in tumor tissues, thereby enhancing chemo- and radiation sensitivity. The present recombinant hemoglobin with a relatively higher oxygen affinity is useful for applications requiring a slower rate of oxygenation (e.g. cancer adjunct therapy).

[0047] ESI-MS allows the analysis of very large molecules. It is an ionization technique that analyzes the high molecular weight compound by ionizing the protein, and then separating the ionized protein based on mass/charge ratio. Therefore, the molecular weight and the protein interactions can be determined accurately. An electrospray ionization mass spectrometry (ESI-MS) is used to analyze and characterize the recombinant hemoglobin monomer, dimer and tetramer in this invention. FIG. 8 depicts electrospray ionization mass spectrometry (ESI-MS) analysis for the recombinant hemoglobin dimer ($\alpha\beta$). The size of α subunit is 15,189 Da and the size of β subunit is 15,899 Da. FIG. 9 depicts electrospray ionization mass spectrometry (ESI-MS) analysis for the recombinant hemoglobin tetramer ($2\alpha\beta_2$). The size of β subunit is 15,899 Da and the size of 2α is 30,356 Da. From this analysis, the estimated total molecular size of $2\alpha\beta_2$ is about 46 KDa. ESI-MS also performs for the recombinant hemoglobin tetramer ($2\alpha\gamma_1_2$), other recombinant hemoglobin dimer and monomer.

[0048] The recombinant hemoglobin protein/tetramer/dimer/subunit of the present invention is used to produce medicaments for tissue oxygenation, for cancer treatment, for the treatment of an oxygen-deprivation disorder such as hemorrhagic shock, and in heart preservation under a low oxygen content environment (e.g. heart transplant). The dosage of recombinant hemoglobin tetramer of the present invention is selected at a concentration range of approximately 0.03g/kg or less of an animal's body weight or 0.0024g/kg or less of a human's body weight (human dose is determined by dividing the dose for mouse by a coefficient of 12.3 according to a FDA guidance for industry and reviewers: Estimating the Safety Starting Dose in Clinical Trials for Therapeutics in Adult Healthy Volunteers, November 18, 2002).

[0049] For a medicament useful in cancer treatment, the hemoglobin-based oxygen-carrier-containing pharmaceutical composition of the present invention serves as a tissue oxygenation

agent to improve the oxygenation in tumor tissues, thereby enhancing chemo-sensitivity (e.g., sensitivity to chemotherapy) and radiation sensitivity.

[0050] The pharmaceutical composition of the present invention comprises recombinant hemoglobin protein or recombinant hemoglobin tetramer or dimer or subunit-based therapeutic agent capable of both targeting and killing the cancer cells. FIG. 6 shows the purified recombinant hemoglobin subunits (α , β , $\gamma 1$, $\gamma 2$) with or without heme, labeled with fluorescent dyes. The fluorescent labeled recombinant hemoglobin subunits can also enter into the cancer cells (e.g., liver cancer cells, HepG2) and localize in there, and the result is illustrated in FIG. 10. It is expected that the chemically modified recombinant hemoglobin protein or tetramer or dimer or subunit-based therapeutic agent can also kill the cancer cells and progenitor cells / cancer stem cells effectively.

[0051] Some conventional therapeutic drugs (e.g. chemotherapeutic drug, 5FU) cannot be used in high dose because of high toxicity. In the present invention, the chemotherapeutic agent, e.g., 5FU, is chemically linked to the recombinant hemoglobin protein or recombinant hemoglobin tetramer or dimer or any of the subunits thereof or the recombinant hemoglobin subunit itself. The chemotherapeutic agent linked to the presently claimed recombinant hemoglobin protein or tetramer or dimer or the subunit thereof can be lower in dose than other conventional method/agents for cancer being administered alone because the presently claimed recombinant hemoglobin protein or tetramer or dimer or subunit facilitates localization of the chemotherapeutic agent in the cytoplasm of the cancerous cells in order to increase the efficacy of both the recombinant hemoglobin protein or tetramer or dimer or subunit of the present invention and the chemotherapeutic agent. The recombinant hemoglobin protein or tetramer or dimer or subunit of the present invention can also improve the efficacy of the radiotherapeutic

agent and/or other anti-cancer drugs on cancer cells or tumor, especially on those which is/are hypoxic that is more resistant to these conventional therapeutic drug for cancer treatment. FIG. 11 shows the liquid chromatography–mass spectrometry (LC-MS) results for (A) 5FU-linked recombinant hemoglobin tetramer ($2\alpha\beta_2$) and (B) 5FU-linked recombinant hemoglobin tetramer ($2\alpha\gamma_1\beta_2$). 5FU is chemically linked successfully to the recombinant hemoglobin tetramer. It also applies to the other chemically linked recombinant hemoglobin dimer and recombinant hemoglobin subunit (or monomer).

[0052] The recombinant hemoglobin protein or tetramer or dimer or its subunit can be modified chemically by different functional groups before linking to the therapeutic drug. The recombinant hemoglobin protein or tetramer or dimer or its subunit can be modified by one or more of the following compound(s) or reaction(s): (1) for amine reactions: anhydride, ketene, NHS ester, isothiocyanates, isocyanates, activated esters which include fluorophenyl esters and carbonyl azides, sulfonyl chlorides, carbonyls followed by reductive amination, epoxides, carbonates, fluorobenzenes, imidoesters, hydroxymethyl phosphine derivatives, mannich condensation, diazonium derivatives, 4-sulfo-2,3,5,6-tetrafluorophenol, carbonyl diimidazole, sulfo-NHS, and N-terminal modification by pyridoxal-5-phosphate-based biomimetic transamination; (2) for thiol reactions: maleimides, alkyl halides, haloacetamides, disulfides, thiosulfates, aziridine-containing reagents, acryloyl derivatives, arylating agents, and vinylsulfone derivatives; (3) for carboxylate reactions: diazoalkanes and diazoacetyl compounds, carbonyldiimidazoles, and carbodiimides; (4) for hydroxyl reactions: epoxides and oxiranes, carbonyldiimidazoles, carbonates, chloroformates, chemical and enzymatic oxidations, alkyl halogens, and isocyanates; (5) for native chemical ligations using thioesters; (6) N-terminal modification using periodate oxidation of N-terminal serine or threonine to generate aldehydes for coupling with hydroxylamines, hydrazines, or hydrazides, carbodiimides; (7) for

incorporation of bioorthogonal functionalities: including alkynes and azides with subsequent bioorthogonal conjugation reactions which include dipolar addition Huisgen 1,3-dipolar additions of alkynes and azides, Staudinger ligation of azides and triarylphosphines, Diels-alder reaction of alkenes and tetrazines, and coupling of alkenes and tetrazoles; (8) for photochemical reactions: photoaffinity labeling agents, diazirine derivatives, benzophenones and anthraquinones; (9) for metal-mediated reactions: metal carbenoids and palladium-activated allyl reagents.

[0053] The glutathione (GSH) level is much higher in the cytoplasm of cancer cells (~1000 equiv) when compared to blood stream (~1 equiv). A therapeutic drug with a disulfide bond can be cleaved much more easily in the cytoplasm of cancer cells with high concentration of GSH. A high-performance liquid chromatography (HPLC) analysis for the cleavage of model compound (N-(2-mercaptoethyl)benzamide) with disulfide bond under reducing condition is shown in FIG. 12. The cleavage of model compound is observed in PBS buffer (pH 7.4) at 37°C with different concentration of glutathione (GSH) (0.4-1000 equiv). The cleavage of the model compound is better at high concentration of GSH (>50 equiv). In the present invention, the recombinant hemoglobin protein or recombinant hemoglobin tetramer or dimer or subunit linked with a therapeutic drug (e.g. chemotherapeutic agent, radiotherapeutic agent, anti-cancer protein drug) by a disulfide bond is prepared for cancer treatment.

[0054] No recombinant hemoglobin protein or recombinant hemoglobin tetramer or dimer or subunit-based therapeutic agent is available in the market. The chemically modified recombinant hemoglobin protein or recombinant hemoglobin tetramer or dimer or subunit-based therapeutic agent-containing pharmaceutical composition provided in the present invention can target cancer cells with therapeutic effect. For uses in cancer treatment, the chemically modified recombinant hemoglobin protein or recombinant hemoglobin tetramer or dimer or subunit-based therapeutic agent-containing pharmaceutical composition of the present invention serves as an anti-cancer

agent to kill cancer cells. The chemically modified recombinant hemoglobin protein or recombinant hemoglobin tetramer or dimer or subunit-based therapeutic agent is a good candidate to be used in lower dose and can be combined with other molecular targeting or cytotoxic agents.

EXAMPLES

[0055] The following examples are provided by way of describing specific embodiments of this invention without intending to limit the scope of this invention in any way.

[0056] Example 1

[0057] (a) Construction of expression vector with recombinant hemoglobin protein and/ or tetramer and/ or dimer and/or its subunit

[0058] The DNA sequences of $\alpha/\beta/\gamma 1/\gamma 2$ chain of human origin are synthesized and cloned in PUC 57 vector. The $\alpha/\beta/\gamma 1/\gamma 2$ chains are amplified by PCR method. The primers used in PCR are designed based on both $\alpha/\beta/\gamma 1/\gamma 2$ chains. A PCR is conducted in the Gradient Thermal Cycler (Life sci). The reaction is performed in a 25 ul reaction mixture containing Taq Buffer, 10 mM dNTP, 25 mM $MgCl_2$, 10 uM of primer, 2.5 U of Taq DNA polymerase and 100ng of template DNA. The gradient PCR condition is comprised of an initial denaturation step of 3 min at 95°C followed by 25 cycles of amplification consisting of denaturation at 95 °C for 30 s, primers annealing at 50–55 °C for 30 s and extension at 72 °C for 45 s prior to a final extension of 5 min at 72 °C. The PCR products are separated by 1.5 % agarose gel electrophoresis along with standard marker and stained with gel red. The product band of the expected size is excised and DNA fragment is extracted with the PCR clean-up Gel extraction kit following the manufacturer's instructions. The purified DNA fragment is cloned into a pSumo vector or PUC 19 vector. Ligation mixture is transformed into *E. coli* DH5a strain, and the transformants are

selected on LB agar containing 100 ng/ml ampicillin. The putative clones are verified by nucleotide sequencing. Analysis of cDNA sequences is performed using BLAST algorithm program.

[0059] (b) Different recombinant hemoglobin protein and/ or tetramer and/ or dimer and/ or subunits

[0060] Seven recombinant hemoglobin protein and/ or tetramer and/ or dimer and/or subunits are generated as shown in FIG. 2 and Table 1. The protein sequences of $\alpha/\beta/\gamma1/\gamma2$ chains are shown in FIG. 2. For recombinant hemoglobin monomers, $\alpha/\beta/\gamma1/\gamma2$ chains are cloned into pSumo vector and the fusion protein expression are controlled by T7 promoter. For recombinant hemoglobin heterodimer, the α chain and $\beta/\gamma1$ chain are co-expressed in pUC19 vector and under the control of pTac promoter. For recombinant hemoglobin tetramer, the two α chain are linked by a glycine linker and co-expressed with $\beta/\gamma1$ chain in pUC19 vector and under the control of pTac promoter.

[0061] Table 1: Different recombinant hemoglobin protein and/ or tetramer and/ or dimer and/or subunits

Numbering in FIG. 2	Recombinant	Expressed hemoglobin subunit/chain/dimer/tetramer
#1	Monomer	α (141 amino acids)
#2		β (146 amino acids)
#3		$\gamma1$ (146 amino acids)
#4		$\gamma2$ (146 amino acids)
#5	Dimer	$\alpha\beta$ (287 amino acids)
#6		$\alpha\gamma1$ (287 amino acids)
#7		$\alpha\gamma2$ (287 amino acids)
#8		$\beta\gamma1$ (292 amino acids)
#9		$\beta\gamma2$ (292 amino acids)

#10	Tetramer	$2\alpha\beta_2$ (575 amino acids)
#11		$2\alpha\gamma_1$ (575 amino acids)
#12		$2\alpha\gamma_2$ (575 amino acids)
#13		$2\beta\gamma_1$ (585 amino acids)
#14		$2\beta\gamma_2$ (585 amino acids)

[0062] **Example 2**

[0063] **Recombinant Protein Expression**

[0064] The plasmids for expression of recombinant hemoglobin protein/ tetramer/ dimer/ subunit are transformed into *E.coli* JM109(DE3) or *E.coli* BL21(DE3). *E.coli* cells are grown in LB medium supplemented with ampicillin (100 mg/ml) at 37°C. For shake flask expression, overnight cultures are 1:100 added into LB medium contained 100mg/ml ampicillin. The cells are grown in a shake flask at 37°C until the optical density at 600 nm reached 0.6. Expression of recombinant hemoglobin protein/ tetramer/ dimer/ subunit is induced by adding isopropyl b-thiogalactopyranoside to 0.4 mM. The culture is then supplemented with hemin (20mg/L) and the growth is continued for at least 16 h at 28°C. For fermenter cultures, medium used for the expression of recombinant hemoglobin protein/ tetramer/ dimer/ subunit contained 1.28% bactotryptone, 0.8% bacto yeast extract, 1% glycerol, 25 mM KH_2PO_4 , 25mM Na_2HPO_4 , 45mM NH_4Cl , 5mM $(\text{NH}_4)_2\text{SO}_4$ and 100 mg/L ampicillin. The cells are grown in a 2 L fermenter (Sartorius, BIOSTAT® B) at 30°C until the optical density at 600 nm reached 8. Expression of recombinant hemoglobin protein/ tetramer/ dimer/ subunit is induced by adding isopropyl b-thiogalactopyranoside (Sigma) to mM. The culture is then supplemented with hemin (50 mg/L) and the growth is continued for at least 16 h at 25°C. The cells are harvested by centrifugation and stored frozen at –80°C until needed for purification.

[0065] **Example 3**

[0066] **Protein yield for recombinant hemoglobin in shake flasks and fermentor**

[0067] The protein yield of recombinant hemoglobin protein and/or tetramer and/or dimer and/or subunit is summarized in Table 2. The protein yield from fermentation method is much higher, can be up to 180 mg/L.

[0068] **Table 2: Protein yield of recombinant human hemoglobin by shaking flask and fermentation.**

Protein	Culture method	Protein yield
Monomer - α	Shake flask	20-40 mg/L
Monomer - β	Shake flask	20-40 mg/L
Monomer - $\gamma 1$	Shake flask	20-40 mg/L
Monomer - $\gamma 2$	Shake flask	20-40 mg/L
Dimer - $\alpha\beta$	Shake flask	5-10 mg/L
Dimer - $\alpha\gamma 1$	Shake flask	5-10 mg/L
Tetramer - $2\alpha\beta_2$	Shake flask	5-10 mg/L
Monomer - $\gamma 1$	Fermentation	180 mg/ L
Tetramer - $2\alpha\beta_2$	Fermentation	30 – 40mg/L

[0069] **Example 4**

[0070] **Culture and reagents for cancer cell line**

[0071] Cancer cells (HepG2) are cultured in DMEM (Invitrogen) with 10% Fetal bovine serum (FBS), 100 U/mL penicillin and 100 μ g/mL streptomycin at 37 °C. For normoxic condition, cells are incubated with ambient O₂ concentration and 5% CO₂; for hypoxic condition, cells are incubated with 0.1- 0.5% O₂ (Quorum FC-7 automatic CO₂/O₂/N₂ gas mixer) and 5% CO₂.

[0072] **Example 5**

[0073] **Live cell time-lapse microscopy in cancer cells**

[0074] Cancer cells (e.g. HepG2 liver cancer cells) are seeded onto glass bottom microwell dishes (MatTek Corporation). Live cells at defined zooms (63x, 20x) are acquired using Zeiss Observer Z1 widefield microscope, equipped with atmospheric/temperature-controlled chamber and motorized stage for multi-positional acquisition. The incubation is performed in an enclosed live cell imaging system purged with 0.1% O₂ and 5% CO₂ (premixed). Cells are exposed to the fluorescent labeled recombinant hemoglobin subunits (α , β , γ_1 , γ_2); for 15 min prior to the acquisition of images every 3 min for a period of 2 h. Images are deconvolved and compacted into time-lapse movies using the MetaMorph software (Molecular Device). The images are shown in FIG. 10.

[0075] **Example 6**

[0076] **(a) *In vivo* efficacy of recombinant hemoglobin tetramer ($2\alpha\beta_2$) on cancer cell xenografts**

[0077] Nude balb/c mice (5-7 weeks) are used in this example and they are allowed to acclimatize for a week before the experiment. Mice are inoculated subcutaneously with 2×10^6 cancer cells in 100 μ l of fresh culture medium. Ten days later, the mice are randomly separated into control and treatment group. Control group receives 100 μ l PBS and treatment group (Group 2-6) receives the drug intraperitoneally weekly (twice per week, for 4 weeks). Gemcitabine is an anti-cancer ("antineoplastic" or "cytotoxic") chemotherapy drug, and it is used in Group 2, Group 4 & Group 6. Tumor size is measured by caliper and tumor volume is calculated using formula: (length x width²)/2. The result is shown in FIG. 13. The result shows that the recombinant hemoglobin tetramer ($2\alpha\beta_2$) can inhibit the growth of tumor cells.

[0078] **(b) *In vivo* efficacy of recombinant hemoglobin tetramer ($2\alpha\gamma_1\gamma_2$) on cancer cell xenografts**

[0079] Nude balb/c mice (5-7 weeks) are used in this example and they are allowed to acclimatize for a week before the experiment. Mice are inoculated subcutaneously with 2×10^6 cancer cells in 100 μ l of fresh culture medium. Ten days later, the mice are randomly separated into control and treatment group. Control group receives 100 μ l PBS and treatment group (Group 2-6) receives the drug intraperitoneally weekly (twice per week, for 4 weeks). Gemcitabine is an anti-cancer ("antineoplastic" or "cytotoxic") chemotherapy drug, and it is used in Group 2, Group 4 & Group 6. Tumor size is measured by caliper and tumor volume is calculated using formula: $(\text{length} \times \text{width}^2)/2$. The result is shown in FIG. 14. The result shows that the recombinant hemoglobin tetramer ($2\alpha\gamma 1_2$) can inhibit the growth of tumor cells. It also gives a better result when combined with the other chemotherapy drug (gemcitabine). A synergistic effect on cancer treatment, inhibiting metastasis and/or reducing recurrence is observed.

INDUSTRIAL APPLICABILITY

[0080] The recombinant hemoglobin protein and/or tetramer and/or dimer and/or subunit of the present invention are useful for oxygenation, in targeting cancer cells or tissue or tumor which might be resistant to conventional therapeutic drug for cancer. The chemically modified recombinant hemoglobin protein tetramer and/or dimer and/or subunit of the present invention is also capable of being cleaved or degradable *in vivo* such that no cytotoxic effect is generated to the subject being administered. The recombinant hemoglobin protein and/or tetramer and/or dimer and/or subunit configured to link with one or more of the conventional therapeutic drug can overcome the resistant problems in some types of cancers or at certain stage of the cancer in patients. The recombinant hemoglobin protein and/or tetramer and/or dimer and/or subunit is readily used for being formulated into pharmaceutical composition and said pharmaceutical composition comprising a therapeutically effective amount of the recombinant hemoglobin

protein and/or tetramer and/or dimer and/or subunit with or without linking to one or more therapeutic drug can be used in various applications including oxygenating tissues, treating hemorrhagic shock and oxygen-deprivation disease, and/or targeting drug-resistant cancer cells/tissue.

CLAIMS

1. A recombinant hemoglobin protein, with or without heme, comprising recombinant hemoglobin tetramer, dimer, and/or subunit as an oxygen carrier and supply replacing natural or native hemoglobin molecule isolated and/or purified from an animal or human source.
2. The recombinant hemoglobin protein of claim 1, wherein said recombinant hemoglobin subunit is a recombinant hemoglobin monomer being constructed and expressed based on a sequence of 141-146 amino acids originated from *Homo sapiens*.
3. The recombinant hemoglobin protein of claim 1, wherein said recombinant hemoglobin dimer comprises any two of the recombinant hemoglobin subunits, each of said recombinant hemoglobin subunits being a recombinant hemoglobin monomer and having a sequence of 141-146 amino acids originated from *Homo sapiens*.
4. The recombinant hemoglobin protein of claim 1, wherein said recombinant hemoglobin tetramer comprises any four of the recombinant hemoglobin subunits, each of said recombinant hemoglobin subunits being a recombinant hemoglobin monomer and having a sequence of 141-146 amino acids originated from *Homo sapiens*.
5. The recombinant hemoglobin protein of claim 2, wherein said recombinant hemoglobin subunit comprises alpha subunit, beta subunit, gamma-1 subunit, and gamma-2 subunit having SEQ ID Nos. 1 to 4, respectively.
6. The recombinant hemoglobin protein of claim 3, wherein said recombinant hemoglobin dimer comprises alpha-beta heterodimer, alpha-gamma-1 heterodimer, alpha-gamma-2 heterodimer, beta-gamma-1 heterodimer, and beta-gamma-2 heterodimer, each of said recombinant hemoglobin subunits having an amino acid sequence of any of SEQ ID Nos. 1 to 4.

7. The recombinant hemoglobin protein of claim 4, wherein said recombinant hemoglobin tetramer comprises $2\alpha\beta_2$, $2\alpha\gamma_1$, $2\alpha\gamma_2$, $2\beta\gamma_1$, and $2\beta\gamma_2$ tetramer, each of said recombinant hemoglobin subunits having an amino acid sequence of any of SEQ ID Nos. 1 to 4.
8. The recombinant hemoglobin protein of claim 1, wherein said recombinant hemoglobin subunit comprises with at least one of or without a Poly-N or a His tag.
9. A method of making the recombinant hemoglobin molecule of claim 1 comprising constructing a fusion gene coding for the recombinant hemoglobin protein, tetramer, dimer and/or subunit, inserting the fusion gene into a vector, inserting the vector into a host organism and expressing said protein, tetramer, dimer or subunit of claim 1.
10. A pharmaceutical composition comprising the recombinant hemoglobin protein of claim 1 in a pharmaceutically-acceptable carrier.
11. A recombinant hemoglobin protein, tetramer, dimer or subunit-based therapeutic agent comprising the recombinant hemoglobin protein, tetramer, dimer, and/or subunit of claim 1 and an active agent, the active agent selected from one or more of a chemotherapeutic agent, a radiotherapeutic agent, a cell labeling agent, or a fluorescent labeling agent, the recombinant hemoglobin protein, tetramer, dimer and/or subunit being either directly conjugated to the active agent or indirectly conjugated to the active agent through a cleavable or non-cleavable linker.
12. The therapeutic agent of claim 11, wherein said recombinant hemoglobin subunit is a recombinant hemoglobin monomer being constructed and expressed based on a sequence of 141-146 amino acids originated from *Homo sapiens*.
13. The therapeutic agent of claim 11, wherein said recombinant hemoglobin dimer comprises any two of the recombinant hemoglobin subunits, each of said recombinant hemoglobin subunits being a recombinant hemoglobin monomer and having a sequence of 141-146 amino acids originated from *Homo sapiens*.

14. The therapeutic agent of claim 11, wherein said recombinant hemoglobin tetramer comprises any four of the recombinant hemoglobin subunits, each of said recombinant hemoglobin subunits being a recombinant hemoglobin monomer and having a sequence of 141-146 amino acids originated from *Homo sapiens*.
15. The therapeutic agent of claim 11, wherein said recombinant hemoglobin protein is linked with one or more of said linker molecule(s) at one or more subunits of said recombinant hemoglobin protein.
16. The therapeutic agent of claim 11, wherein said recombinant hemoglobin protein comprises with at least one of or without a Poly-N or a His tag.
17. The therapeutic agent of claim 12, wherein said recombinant hemoglobin subunit comprises alpha subunit, beta subunit, gamma-1 subunit, and gamma-2 subunit having SEQ ID Nos. 1 to 4, respectively.
18. The therapeutic agent of claim 13, wherein said recombinant hemoglobin dimer comprises alpha-beta heterodimer, alpha-gamma-1 heterodimer, alpha-gamma-2 heterodimer, beta-gamma-1 heterodimer, and beta-gamma-2 heterodimer, each of said recombinant hemoglobin subunits having an amino acid sequence of any of SEQ ID Nos. 1 to 4.
19. The therapeutic agent of claim 14, wherein said recombinant hemoglobin tetramer comprises $2\alpha\beta_2$, $2\alpha\gamma_1$, $2\alpha\gamma_2$, $2\beta\gamma_1$, and $2\beta\gamma_2$ tetramer, each of said recombinant hemoglobin subunits having an amino acid sequence of any of SEQ ID Nos. 1 to 4.
20. The therapeutic agent of claim 11, wherein said recombinant hemoglobin protein, tetramer, dimer, or subunit is chemically modified by one or more functional groups before being conjugated to said active agent, said chemical modification comprising reacting the recombinant hemoglobin molecule with one or more of the following compound(s) in respect of different chemical reactions for said chemical modification:

a) for amine reactions, the compounds involved comprising anhydride, ketene, NHS ester, isothiocyanates, isocyanates, activated esters which include fluorophenyl esters and carbonyl azides, sulfonyl chlorides, carbonyls followed by reductive amination, epoxides, carbonates, fluorobenzenes, imidoesters, hydroxymethyl phosphine derivatives, mannich condensation, diazonium derivatives, 4-sulfo-2,3,5,6-tetrafluorophenol, carbonyl diimidazole, sulfo-NHS, and N-terminal modification by pyridoxal-5-phosphate-based biomimetic transamination;

b) for thiol reactions, the compounds involved comprising maleimides, alkyl halides, haloacetamides, disulfides, thiosulfates, aziridine-containing reagents, acryloyl derivatives, arylating agents, and vinylsulfone derivatives;

c) for carboxylate reactions, the compounds involved comprising diazoalkanes and diazoacetyl compounds, carbonyldiimidazoles, and carbodiimides;

d) for hydroxyl reactions, the compounds involved comprising epoxides and oxiranes, carbonyldiimidazoles, carbonates, chloroformates, chemical and enzymatic oxidations, alkyl halogens, and isocyanates;

e) for native chemical ligations, the compounds involved comprising thioesters;

f) for N-terminal modification, the compounds involved comprising N-terminal serine or threonine using periodate oxidation thereof to generate aldehydes for coupling with hydroxylamines, hydrazines, or hydrazides, carbodiimides;

g) for incorporation of bioorthogonal functionalities, the compounds involved comprising alkynes and azides with subsequent bioorthogonal conjugation reactions which include dipolar addition Huisgen 1,3-dipolar additions of alkynes and azides, Staudinger ligation of azides and triarylphosphines, Diels-alder reaction of alkenes and tetrazines, and coupling of alkenes and tetrazoles;

h) for photochemical reactions, the compounds involved comprising photoaffinity labeling agents, diazirine derivatives, benzophenones and anthraquinones;

i) for metal-mediated reactions, the compounds involved comprising metal carbenoids and palladium-activated allyl reagents.

21. The therapeutic agent of claim 11, wherein said cleavable linker comprises carbinolamine, disulfide, carbamide, aminal, carbonate, ester, carbamate, phosphate, amide, acetal, imine, oxime, ether and sulfonamide group linker.

22. The therapeutic agent of claim 11, wherein said non-cleavable linker comprises alkyl and aryl group linker.

23. The therapeutic agent of claim 11, wherein said chemotherapeutic agent comprises 5-Fluorouracil, temozolomide, cisplatin and other anti-cancer drugs.

24. The therapeutic agent of claim 11, wherein said radiotherapeutic agent comprises Rhodium-105 complex, Samarium-153 complex and other related complexes.

25. The therapeutic agent of claim 11, wherein said cell or fluorescent labeling agent comprises one or more fluorescent proteins, non-protein organic fluorophores, fluorescent nanoparticles and metal-based luminescent dyes.

26. The therapeutic agent of claim 11, wherein said cancer comprises pancreatic cancer, leukemia, head and neck cancer, colorectal cancer, lung cancer, breast cancer, liver cancer, nasopharyngeal cancer, esophageal cancer, prostate cancer, stomach cancer and brain cancer.

27. A pharmaceutical composition comprising a therapeutically effective amount of the recombinant hemoglobin protein, tetramer, dimer or subunit-based therapeutic agent of claim 11, said composition being administered to a subject in need thereof who suffers from cancer.

28. The composition of claim 27, wherein the therapeutically effective amount of said therapeutic agent is at ≤ 0.0024 g/kg of said subject and said subject is human.

29. The composition of claim 27, wherein the therapeutically effective amount of said therapeutic agent is at ≤ 0.03 g/kg of said subject and said subject is animal.
30. The composition of claim 27 further comprising a pharmaceutically acceptable carrier, salt, buffer, water, or a combination thereof.
31. The composition of claim 27, wherein route of said administering comprises intravenous injection, intraperitoneal injection, and subcutaneous injection, and said administering of the composition can be in single dose or in multiple doses for once a week.
32. The composition of claim 27, wherein said cancer comprises pancreatic cancer, leukemia, head and neck cancer, colorectal cancer, lung cancer, breast cancer, liver cancer, nasopharyngeal cancer, esophageal cancer or brain cancer.
33. The composition of claim 27, wherein said cancer is tumor comprising hepatocellular carcinoma, liver cancer progenitor cells-induced tumor, glioblastoma, or triple negative progenitor cells-induced tumor.
34. The pharmaceutical composition of claim 27 wherein the composition has a pH in a range of 5.5 to 9.5.
35. The pharmaceutical composition of claim 27 wherein the composition has a pH of 7.2 to 8.0.
36. A method for preparing the therapeutic agent of claim 11, said method comprising:
- a) providing a recombinant hemoglobin protein, tetramer, dimer or subunit;
 - b) purifying the recombinant hemoglobin protein, tetramer, dimer or subunit using different column chromatography methods;
 - c) stabilizing the recombinant hemoglobin protein or tetramer by a cross-linker to form a cross-linked recombinant hemoglobin protein or tetramer;

d) chemically modifying the recombinant hemoglobin protein, tetramer, dimer or subunit comprising reacting with one or more functional groups followed by conjugating said recombinant hemoglobin protein, tetramer, dimer or subunit with an active agent, or conjugating an active agent with a cleavable or non-cleavable linker to form a linker-active agent conjugate prior to linking said conjugate to said recombinant hemoglobin protein, tetramer, dimer or subunit.

37. The method of claim 36, wherein said chemically modifying comprises one or more of the following reactions or modifications:

a) amine reactions, wherein the compounds involved comprises anhydride, ketene, NHS ester, isothiocyanates, isocyanates, activated esters which include fluorophenyl esters and carbonyl azides, sulfonyl chlorides, carbonyls followed by reductive amination, epoxides, carbonates, fluorobenzenes, imidoesters, hydroxymethyl phosphine derivatives, mannich condensation, diazonium derivatives, 4-sulfo-2,3,5,6-tetrafluorophenol, carbonyl diimidazole, sulfo-NHS, and N-terminal modification by pyridoxal-5-phosphate-based biomimetic transamination;

b) thiol reactions, wherein the compounds involved comprises maleimides, alkyl halides, haloacetamides, disulfides, thiosulfates, aziridine-containing reagents, acryloyl derivatives, arylating agents, and vinylsulfone derivatives;

c) carboxylate reactions, wherein the compounds involved comprises diazoalkanes and diazoacetyl compounds, carbonyldiimidazoles, and carbodiimides;

d) hydroxyl reactions, wherein the compounds involved comprises epoxides and oxiranes, carbonyldiimidazoles, carbonates, chloroformates, chemical and enzymatic oxidations, alkyl halogens, and isocyanates;

e) native chemical ligations, wherein the compounds involved comprises thioesters;

f) N-terminal modification, wherein the compounds involved comprises N-terminal serine or threonine using periodate oxidation thereof to generate aldehydes for coupling with hydroxylamines, hydrazines, or hydrazides, carbodiimides;

g) incorporation of bioorthogonal functionalities, wherein the compounds involved comprises alkynes and azides with subsequent bioorthogonal conjugation reactions which include dipolar addition Huisgen 1,3-dipolar additions of alkynes and azides, Staudinger ligation of azides and triarylphosphines, Diels-alder reaction of alkenes and tetrazines, and coupling of alkenes and tetrazoles;

h) photochemical reactions, wherein the compounds involved comprises photoaffinity labeling agents, diazirine derivatives, benzophenones and anthraquinones;

i) metal-mediated reactions, wherein the compounds involved comprises metal carbenoids and palladium-activated allyl reagents.

38. The method of claim 36, wherein said cleavable linker comprises carbinolamine, disulfide, carbamide, aминаl, carbonate, ester, carbamate, phosphate, amide, acetal, imine, oxime, ether or sulfonamide group linker.

39. The method of claim 36, wherein said non-cleavable linker comprises alkyl or aryl group linker.

40. The method of claim 36, wherein said active agent is a chemotherapeutic agent comprising 5-Fluorouracil, temozolomide, cisplatin and other anti-cancer drug.

41. The method of claim 36, wherein said active agent is a cell/fluorescent labeling agent comprising one or more fluorescent proteins, non-protein organic fluorophores, fluorescent nanoparticles and metal-based luminescent dye.

42. Use of the composition of claim 10 for locally oxygenating tissue from or in an animal or human subject.

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43. The use of claim 42, wherein the tissue is cancerous tumor tissue.
 44. The use of claim 42, wherein the tissue is hypoxic.
 45. The use of claim 42, wherein the tissue is *ex vivo*.
 46. The use of claim 42, wherein the tissue is normal organ tissue.
 47. Use of the composition of claim 10 in manufacturing a medicament for treating oxygen-deprivation disorder in an animal or human subject in needs thereof.
 48. Use of the composition of claim 10 in manufacturing a medicament for treating hemorrhagic shock in an animal or human subject in needs thereof.
 49. Use of the therapeutic agent of claim 11 in manufacturing a medicament for targeting and inducing apoptosis of cancer cells in a subject in needs thereof, wherein said medicament comprises a therapeutically effective amount of the therapeutic agent of claim 11.
 50. Use of the therapeutic agent of claim 11 in manufacturing a medicament for inhibiting tumor metastasis and reducing recurrence in a subject in needs thereof, wherein said medicament comprises a therapeutically effective amount of the therapeutic agent of claim 11.
 51. Use of the therapeutic agent of claim 11 for live-cell imaging and diagnostic imaging of cancer cells or tumor cells or progenitor cells or cancer stem cells.