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Diketo-piperazin-sók gyógyszeradagolás céljára

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(54) **Diketopiperazine salts for drug delivery**

Diketopiperazinsalze zur Arzneimitteldarreichung

Sels de diketopiperazine permettant l'administration de médicaments

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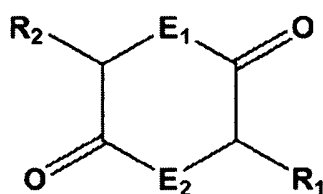
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Description**FIELD OF THE INVENTION**

[0001] This invention is generally in the field of drug delivery related to both small molecule and macromolecular drugs. More particularly it is related to 2,5-diketopiperazine salts, their use in the formulation of such drugs including therapeutic, prophylactic and diagnostic agents, stabilizing agents and systems for their delivery.

BACKGROUND TO THE INVENTION

[0002] Drug delivery has been a persistent challenge in the pharmaceutical arts, particularly when a drug is unstable and/or poorly absorbed at the locus in the body to which it is administered. One such class of drugs includes 2,5-diketopiperazines, which is represented by the compound of the general Formula 1 as shown below where E=NH.

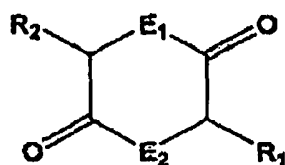
**Formula 1**

[0003] These 2,5 diketopiperazines have been shown to be useful in drug delivery, particularly those bearing acidic R groups (see for example U.S. Patent Nos. 5,352,461 entitled "Self Assembling Diketopiperazine Drug Delivery System;" 5,503,852 entitled "Method For Making Self-Assembling Diketopiperazine Drug Delivery System;" 6,071,497 entitled "Microparticles For Lung Delivery Comprising Diketopiperazine;" and 6,331,318 entitled "Carbon-Substituted Diketopiperazine Delivery System." Diketopiperazines can be formed into particles that incorporate a drug or particles onto which a drug can be adsorbed. The combination of a drug and a diketopiperazine can impart improved drug stability. These particles can be administered by various routes of administration. As dry powders these particles can be delivered by inhalation to specific areas of the respiratory system, depending on particle size. Additionally, the particles can be made small enough for incorporation into an intravenous suspension dosage form. Oral delivery is also possible with the particles incorporated into a suspension, tablets or capsules; or dissolved in an appropriate solvent Diketopiperazines may also facilitate absorption of an associated drug. Nonetheless difficulties can arise when diketopiperazines are diacids, or are in diacid form(s), due to the limited solubility of these diacids at non-basic pH (i.e., neutral or acid pH). Another difficulty arises because these diacid diketopiperazines may form disadvantageous association(s) with some drugs.

[0004] Therefore there is a need for diketopiperazine compositions having greater solubility at a neutral and/or acidic pH and methods for their use in the manufacture of therapeutic compositions.

SUMMARY OF THE INVENTION

[0005] A first aspect of the invention provides a pharmaceutically-acceptable salt of a heterocyclic compound according to Formula 1:

**Formula 1**

wherein R₁ and R₂ comprise 4-X-aminobutyl and wherein X is selected from the group consisting of succinyl, glutaryl, maleyl, citraconyl, malonyl, oxalyl and fumaryl;
E₁ and E₂ are NH; and

said salt further comprises at least one cation: and
 wherein said salt is not a lithium salt of 2,5-diaspartyl-3,8-diketopiperazine or 2,5-diglutamyl-3,8-diketopiperazine
 and said salt is in a solid form;
 wherein the heterocyclic compound comprises a diketopiperazine.

[0006] A second aspect of the invention provides microparticles comprising the pharmaceutical acceptable salt of the first aspect of the invention and a biologically active agent. The latter may be provided as a solid dosage form or dry powder medicament.

[0007] A third aspect of the invention provides a solid dosage form comprising a precipitate comprising the pharmaceutical acceptable salt of the first aspect of the invention and a biologically active agent.

[0008] A fourth aspect of the invention provides a composition according to the first, second or third aspect of the invention for use in medicine.

[0009] A fifth aspect of the invention provides a method of making the solid dosage form of the second or third aspect of the invention comprising forming by precipitation or by lyophilization.

[0010] The present invention provides improved drug delivery systems comprising carboxylate salts of heterocyclic compounds in combination with one or more drugs. In one embodiment of the present invention the heterocyclic compounds form microparticles that incorporate the drug or drugs to be delivered. These microparticles include microcapsules, which have an outer shell composed of either the heterocyclic compound alone or in combination with one or more drugs. The heterocyclic compounds of the present invention comprise diketopiperazines. The heterocyclic compositions of the present invention comprise rigid hexagonal rings with opposing heteroatoms and unbonded electron pairs.

[0011] Specifically preferred embodiments include, without limitation, derivatives of 3,6-di(4-aminobutyl)-2,5-diketopiperazine, such as 3,6-di(succinyl-4-aminobutyl)-2,5-diketopiperazine, 3,6-di(maleyl-4-aminobutyl)-2,5-diketopiperazine, 3,6-di(citraconyl-4-aminobutyl)-2,5-diketopiperazine, 3,6-di(glutaryl-4-aminobutyl)-2,5-diketopiperazine, 3,6-di(malonyl-4-aminobutyl)-2,5-diketopiperazine, 3,6-di(oxalyl-4-aminobutyl)-2,5-diketopiperazine, and 3,6-di(fumaryl-4-aminobutyl)-2,5-diketopiperazine (hereinafter fumaryl diketopiperazine or FDKP). Additionally, nonsymmetrical derivatives of the aforementioned are also contemplated. However, it is specifically noted herein that lithium salts of 2,5-diaspartyl-3,6-diketopiperazine and 2,5-diglutamyl-3,6-diketopiperazine (as defined further below) are not considered within the scope of the present invention and as such are hereby specifically disclaimed.

[0012] Representative drugs useful with the drug delivery systems of the present invention include, without limitation, insulin and other hormones, peptides, proteins, polysaccharides, such as heparin, nucleic acids (such as plasmids, oligonucleotides, antisense, or siRNA), lipids and lipopolysaccharides, anticoagulants, cytotoxic agents, antigens and antibodies and organic molecules having biological activity such as many of the antibiotics, anti-inflammatories, antivirals, vaso- and neuroactive agents.

[0013] In one embodiment, the cation is selected from the group consisting of sodium, potassium, calcium, lithium, triethylamine, butylamine, diethanolamine and triethanolamine.

[0014] Biologically active agents suitable for inclusion in the compositions of the present invention include hormones, anticoagulants, immunomodulating agents, cytotoxic agents, antibiotics, antivirals, antisense, antigens, antibodies and active fragments and analogues thereof. In one embodiment the biologically active agent is insulin.

[0015] In one embodiment, the therapeutic composition of the present invention is a precipitate and the precipitate is formulated into a solid dosage form suitable for oral, buccal, rectal, or vaginal administration. The solid dosage form may be a capsule, a tablet, and a suppository.

[0016] In an embodiment, the therapeutic composition of the present invention is a dry powder and the particles of said dry powder have a diameter between about 0.5 μm (microns) and 10 μm (microns). In one aspect of the embodiment the dry powder is suitable for pulmonary administration.

[0017] Herein is disclosed a method of preparing a solid composition for drug delivery comprising: preparing a solution containing a biologically active agent and a pharmaceutically-acceptable salt of a heterocyclic compound in a solvent and removing the solvent by a method selected from the group consisting of distillation, evaporation, and lyophilization.

[0018] The method of preparing a solid composition for drug delivery may further comprise the step of micronizing the solid to form a dry powder.

[0019] Herein is disclosed a method of preparing a dry powder for drug delivery comprising spray drying a solution of a pharmaceutically acceptable salt of a heterocyclic compound and a biologically active agent to form a dry powder wherein the dry powder releases a biologically active agent. The particles of the dry powder may be suitable for pulmonary delivery. The particles of the dry powder may have a rugosity of less than 2.

[0020] Also disclosed is a composition for delivering biologically active agents wherein the composition comprises a pharmaceutically acceptable salt of a heterocyclic compound and a biologically active agent spray dried to form a dry powder such that the dry powder releases said biologically active agents. The particles of the dry powder may be suitable for pulmonary delivery. The particles of the dry powder may have a rugosity of less than 2.

[0021] Also disclosed is a microparticulate system for drug delivery comprising a composition of pharmaceutically

acceptable salt of a heterocyclic compound and a biologically active agent and wherein the composition releases a biologically active agent. The biologically active agent can include hormones, anticoagulants, immunomodulating agents, cytotoxic agents, antibiotics, antivirals, antisense, antigens, antibodies and active fragments and analogues thereof.

[0022] The composition of the microparticulate system may be a dry powder which releases a biologically active agent in the pulmonary system. The composition can further be delivered to the pulmonary system. The composition of the microparticulate system can be absorbed into the systemic blood circulation or act locally in the lung after delivery to the pulmonary system.

[0023] The composition of the microparticulate system may comprise a liquid for drug delivery wherein the absorption of the biologically active agent is facilitated by the diketopiperazine. The liquid may be administered orally.

[0024] The composition of the microparticulate system may comprise a precipitate wherein the absorption of the biologically active agent is facilitated by the diketopiperazine. The precipitate may be administered orally.

[0025] Herein is disclosed a method for delivery of particles to the pulmonary system comprising: administering via inhalation to a patient in need of treatment an effective amount of a biologically active agent in the form of a dry powder, the dry powder prepared by spray drying a solution comprising a composition of a pharmaceutically acceptable salt of a heterocyclic compound and a biologically active agent, wherein the dry powder releases the biologically active agent in the pulmonary system

BRIEF DESCRIPTION OF THE DRAWINGS

[0026]

Figure 1 depicts a laser diffraction particle size analysis of particles made using a fumaryl diketopiperazine (FDKP) disodium salt according to one aspect of the present invention. (A) preparation A; (B) preparation B.

Figure 2 depicts particle size determination by laser diffraction of a formulation of a FDKP disodium salt containing 25% insulin (w:w) made according to the teachings of the present invention.

Figure 3 depicts scanning electron microscopy (SEM) of a spray dried microparticle preparation of a FDKP disodium salt containing 25% insulin (w:w) made according to the teachings of the present invention.

Figure 4 depicts an accelerated stability study of spray dried microparticles of a FDKP disodium salt/insulin formulation containing 25% insulin made according to the teachings of the present invention (stippled) compared to control lyophilized powder (hatched).

Figure 5 depicts the effect of solution concentration on insulin stability of spray dried microparticles of a FDKP disodium salt/insulin formulation containing 25% insulin made according to the teachings of the present invention compared to control lyophilized powder.

Figure 6 depicts SEM analysis of the insulin/disodium FDKP salt microparticles formed by the solvent/anti-solvent precipitation according to the teachings of the present invention.

Figure 7 depicts particle size determination by laser diffraction of spray dried microparticles of a FDKP diammonium salt/insulin formulation containing 25% insulin (w:w) made according to the teachings of the present invention.

Figure 8 depicts particle size determination by laser diffraction of spray dried microparticles of a FDKP diammonium salt/insulin formulation containing 50% insulin (w:w) made according to the teachings of the present invention.

Figure 9 depicts particle size determination by laser diffraction of spray dried microparticles of a diammonium salt of succinyl diketopiperazine (SDKP) containing 25% insulin (w:w) made according to the teachings of the present invention.

Figure 10 depicts SEM of the FDKP ammonium salt formulated with 25% insulin according to the teachings of the present invention.

Figure 11 depicts SEM of the SDKP ammonium salt formulated with 25% insulin according to the teachings of the present invention.

Figure 12 depicts an accelerated stability study of the spray dried microparticles of a FDKP diammonium salt/insulin

formulation containing 25% or 50% insulin made according to the teachings of the present invention compared to control lyophilized powder.

Figure 13 depicts the generation of the A₂₁ degradant during an accelerated stability study of the spray dried microparticles of a FDKP diammonium salt/insulin formulation containing 25% or 50% insulin made according to the teachings of the present invention compared to control lyophilized powder.

Figure 14 depicts an accelerated stability study of the spray dried microparticles of a diammonium SDKP salt/insulin formulation containing 25% insulin made according to the teachings of the present invention compared to control lyophilized powder.

Figure 15 depicts the generation of the A₂₁ degradant during an accelerated stability study of the spray dried microparticles of a diammonium SDKP salt/insulin formulation containing 25% insulin made according to the teachings of the present invention compared to control lyophilized powder.

Figure 16 depicts the aerodynamic performance of spray dried FDKP disodium salt/insulin particles containing increasing insulin concentrations made according to the teachings of the present invention.

Figure 17 depicts the aerodynamic performance of spray dried FDKP diammonium salt/insulin particles containing increasing insulin concentrations made according to the teachings of the present invention.

DEFINITION OF TERMS

[0027] Prior to setting forth the invention, it may be helpful to provide an understanding of certain terms that will be used hereinafter:

[0028] Acidic: As used herein, "acidic" refers to a pH range of from 0, up to, but not including 6.

[0029] Basic: As used herein, "basic" refers to a pH range of from 8, up to and including 14.

[0030] Biological agents: See "Drug" below.

[0031] Cargo: See "Drug" below.

[0032] Diketopiperazine: As used herein, "diketopiperazines" or "DKP" includes diketopiperazines and derivatives and modifications thereof falling within the scope of Formula 1.

[0033] Drug: As used herein, "drug", "cargo" or "biological agent" refer to the pharmacologically active agent incorporated with the microparticles discussed herein. Examples include proteins and peptides (wherein protein is defined as consisting of 100 amino acid residues or more and a peptide is less than 100 amino acid residues), such as insulin and other hormones; polysaccharides, such as heparin; nucleic acids, such as plasmids, oligonucleotides, antisense, or siRNA; lipids and lipopolysaccharides; and organic molecules having biological activity such as many of the antibiotics, anti-inflammatories, antivirals, vaso- and neuroactive agents. Specific examples include hormones, anticoagulant, immunomodulating agents, cytotoxic agents, antibiotics, antivirals, antisense, antigens, and antibodies.

[0034] Dry powder: As used herein "dry powder" refers to a fine particulate composition that is not suspended or dissolved in a propellant, carrier, or other liquid. It is not meant to imply a complete absence of all water molecules.

[0035] Microparticles: As used herein, the term "microparticles" includes microcapsules having an outer shell composed of either a diketopiperazine alone or a combination of a diketopiperazine and one or more drugs. It also includes microspheres containing drug dispersed throughout the sphere; particles of irregular shape; and particles in which the drug is coated in the surface(s) of the particle or fills voids therein.

[0036] Neutral: As used herein, "neutral" refers to a pH range of from 6, up to, but not including 8.

[0037] Weakly alkaline: As used herein, "weakly alkaline" refers to a pH range of from 8, up to, but not including 10.

DETAILED DESCRIPTION OF THE INVENTION

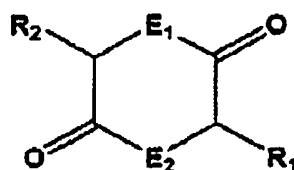
[0038] The present invention provides improved drug delivery systems comprising carboxylate salts of heterocyclic compounds in combination with one or more drugs. In one embodiment of the present invention the heterocyclic compounds form microparticles that incorporate the drug or drugs to be delivered. These microparticles include microcapsules, which have an outer shell composed of either the heterocyclic compound alone or in combination with one or more drugs. The heterocyclic compounds comprised in the compositions of the present invention are diketopiperazines. The heterocyclic compositions of the present invention comprise rigid hexagonal rings with opposing heteroatoms and unbonded electron pairs.

[0039] One aspect of the present invention includes a drug delivery system comprising the carboxylate salts of heterocyclic compounds in combination with one or more drugs. In one embodiment of the present invention the heterocyclic

compounds form microparticles that incorporate the drug or drugs to be delivered. These microparticles include microcapsules, which have an outer shell composed of either the heterocyclic compound alone or in combination with one or more drug(s). This outer shell may surround a core material. This outer shell may also surround or constitute microspheres that are either solid or hollow, or a combination thereof, which contain one or more drugs dispersed throughout the sphere and/or adsorbed onto the surface of the sphere. The outer shell also may surround microparticles having irregular shape, either alone or in combination with the aforementioned microspheres.

[0040] In a preferred embodiment for pulmonary delivery, the microparticles are from about 0.1 μm (microns) to about ten μm (microns) in diameter. Within drug delivery systems, these microparticles exhibit desirable size distributions as well as good cargo tolerance.

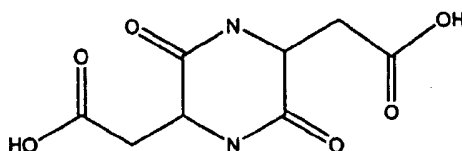
[0041] The heterocyclic compounds comprised in the compositions of the present invention are diketopiperazines. These heterocyclic compositions comprise rigid hexagonal rings with opposing heteroatoms and unbonded electron pairs. The general formula for diketopiperazine and its analogs is shown below in the compound of Formula 1.



Formula 1

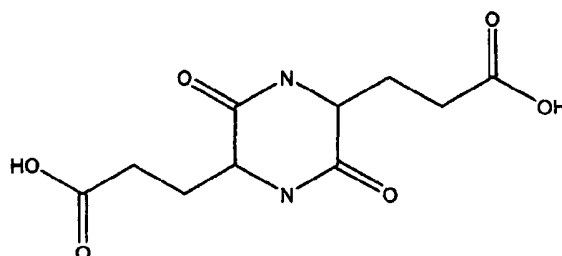
[0042] In the compound of Formula 1 the ring atoms E_1 and E_2 at positions 1 and 4 are NH. At least one of the side chains R_1 and R_2 located at positions 3 and 6 respectively contains a carboxylate group (i.e., OR).

[0043] General methods for the synthesis of diketopiperazines are known in the art and have been described in United States Patent Nos. 5,352,461, 5,503,852, and 6,331,318. In a preferred embodiment of the invention the diketopiperazine is a derivative of 3,6-di(4-aminobutyl)-2,5-diketopiperazine, which may be formed by condensation of the amino acid lysine. Exemplary derivatives include 3,6-di(succinyl-4-aminobutyl)-(succinyl diketopiperazine or SDKP), 3,6-di(maleyl-4-aminobutyl)-, 3,6-di(citraconyl-4-aminobutyl)-3,6-di(glutaryl-4-aminobutyl)-, 3,6-di(malonyl-4-aminobutyl)-, 3,6-di(oxalyl-4-aminobutyl)-, and 3,6-di(fumaryl-4-aminobutyl)-2,5-diketopiperazine (hereinafter fumaryl diketopiperazine or FD-KP). However, it is specifically noted herein that the lithium salts of 2,5-diaspartyl-3,6-diketopiperazine and 2,5-diglutamyl-3,6-diketopiperazine are not considered within the scope of the present invention and as such are hereby specifically disclaimed. The free acids of these disclaimed compounds are depicted below in Formula 2 and Formula 3 respectively.



Formula 2

(5-Carboxymethyl-3,6-dioxo-piperazin-2-yl)-acetic acid



Formula 3

3-[5-(2-carboxy-ethyl)-3,6-dioxo-piperazin-2-yl]-propionic acid

[0044] For convenience, the compound of Formula 2 will be referred to hereinafter as 2,5-diaspartyl-3,6-diketopiperazine. The compound of Formula 3 will be referred to hereinafter as 2,5-diglutamyl-3,6-diketopiperazine. It is understood that all other heterocyclic compounds based on Formula 1 are considered within the scope of the present invention.

[0045] For exemplary purposes, diketopiperazines salts and their derivatives will be described in detail. These compounds are the preferred embodiments of the present invention. However, this does not exclude other heterocyclic compounds based on the compound of Formula 1.

[0046] The use of DKP salts for the delivery of phosphodiesterase type 5-inhibitors is described in co-pending U.S. Patent Application No. 11/210,709 (US2006099269) filed August 23, 2005 and entitled "Pulmonary Delivery of Inhibitors of Phosphodiesterase Type 5". Pulmonary drug delivery using DKP microparticles is disclosed in U.S. Patent No. 6,428,771 entitled "Method For Drug Delivery To The Pulmonary System".

[0047] Diketopiperazine facilitate transcellular transport of biologically active agents across biological tissues however they are not penetration enhancers. Penetration enhancers are compounds that improve drug movement across biological tissues by disrupting cell membranes. Examples of penetration enhancers are surfactants and soaps. Diketopiperazines do not disrupt cell membranes either *in vitro* or *in vivo*. *In vitro* studies demonstrate that FDKP does not disrupt cell membranes or tight junctions and does not compromise cell viability. Diketopiperazine/insulin powder compositions are soluble at the physiological pH of the lung surface and dissolve rapidly after inhalation. Once dissolved, the DKP facilitates passive transcellular transport of the insulin.

[0048] Applicants have discovered improved diketopiperazine compositions having greater solubility at a neutral and/or acidic pH. Applicants have also discovered that therapeutic complexes between improved diketopiperazines and drug(s) of interest can be formed.

[0049] The salts of the present invention can be prepared by reacting the diketopiperazine free acid with a solution of an appropriate base as described in Examples 1 and 2 below. In a preferred embodiment, the salt is a pharmaceutically acceptable salt such as the sodium (Na), potassium (K), lithium (Li), magnesium (Mg), calcium (Ca), ammonium, or mono-, di- or tri-alkylammonium (as derived from triethylamine, butylamine, diethanolamine, triethanolamine, or pyridines, and the like) salts of diketopiperazine, for example. The salt may be a mono-, di-, or mixed salt. Higher order salts are also contemplated for diketopiperazines in which the R groups contain more than one acid group. In other aspects of the invention, a basic form of the agent may be mixed with the DKP in order to form a drug salt of the DKP, such that the drug is the counter cation of the DKP.

[0050] For drug delivery, biologically active agents or drugs having therapeutic, prophylactic, or diagnostic activities can be delivered using diketopiperazines. Essentially, the biologically active agent is associated with the diketopiperazine particles of the present invention. As used herein, "associated" means a biologically active agent-diketopiperazine composition formed by, among other methods, co-precipitation, spray drying or binding (complexation) of the diketopiperazine with the biologically active agent. The resulting diketopiperazine particles include those that have entrapped, encapsulated and/or been coated with the biologically active agent. While the exact mechanism of association has not been conclusively identified, it is believed that the association is a function of physical entrapment (molecular entanglement) in addition to electrostatic attraction including hydrogen bonding, van der Waal's forces and adsorption.

[0051] The biologically active agents that can be associated with the diketopiperazine particles of the present invention include, but are not limited to, organic or inorganic compounds, proteins, or a wide variety of other compounds, including nutritional agents such as vitamins, minerals, amino acids, carbohydrates, sugars, and fats. In preferred embodiments, the drugs include biologically active agents that are to be released in the circulatory system after transport from the GI tract following oral delivery. In other preferred embodiments the materials are biologically active agents that are to be released in the circulatory system following pulmonary or nasal delivery. In other preferred embodiments the materials are biologically active agents that are to be release in the central nervous system following nasal delivery. Additionally, the drug can be absorbed through mucosal tissue such as rectal, vaginal, and/or buccal tissue. Non-limiting examples of biologically active agents include proteins and peptides (wherein protein is defined as consisting of 100 amino acid residues or more and a peptide is less than 100 amino acid residues), such as insulin and other hormones, polysaccharides, such as heparin, nucleic acids (such as plasmids, oligonucleotides, antisense, or siRNA), lipids and lipopolysaccharides, and organic molecules having biological activity such as many of the antibiotics, anti-inflammatories, vasoactive agents (including agents used to treat erectile dysfunction) and neuroactive agents. Specific non-limiting examples include steroids, hormones, decongestants, anticoagulants, immunomodulating agents, cytotoxic agents, antibiotics, antivirals, anesthetics, sedatives, antidepressants, cannabinoids, anticoagulants, antisense agents, antigens, and antibodies. In some instances, the proteins may be antibodies or antigens which otherwise would have to be administered by injection to elicit an appropriate response. More particularly, compounds that can be associated with the diketopiperazine compositions of the present invention include insulin, heparins, calcitonin, felbamate, parathyroid hormone and fragments thereof, growth hormone, erythropoietin, glucagon-like peptide-1, somatotrophin-releasing hormone, follicle stimulating hormone, cromolyn, adiponectin, RNase, ghrelin, zidovudine, didanosine, tetrahydrocannabinol (i.e., cannabinoids), atropine, granulocytes colony stimulating factor, lamotrigine, chorionic gonadotropin releasing factor, luteinizing releasing hormone, beta-galactosidase and Argatroban. Compounds with a wide range of molecular weight can

be associated, for example, between 100 and 500,000 grams per mole.

[0052] Imaging agents including metals, radioactive isotopes, radiopaque agents, and radiolucent agents, can also be incorporated into diketopiperazine delivery systems. Radioisotopes and radiopaque agents include gallium, technetium, indium, strontium, iodine, barium, and phosphorus.

[0053] Additionally the drugs can be in various forms, such as uncharged molecules, metal or organic salts, or prodrugs. For acidic drugs, metal salts, amines or organic cations (e.g., quaternary ammonium) can in some cases be used.

[0054] In some embodiment, the drugs include biologically active agents that are to be released in the circulatory system after transport from the gastrointestinal tract following oral delivery. In other embodiments, the biologically active agents are to be released in the circulatory system following pulmonary or nasal delivery. In still other embodiments, the biologically active agents are to be released in the central nervous system following nasal delivery. Additionally, the drugs can be absorbed through mucosal tissue such as rectal, vaginal, and/or buccal tissue.

[0055] Some of these biological agents are unstable in gastric acid, diffuse slowly through gastrointestinal membranes, are poorly soluble at physiological pH, and/or are susceptible to enzymatic destruction in the gastrointestinal tract. The biological agents are combined with the diketopiperazine salts to protect them in the gastrointestinal tract prior to release in the blood stream. In a preferred embodiment the diketopiperazines are not biologically active and do not alter the pharmacologic properties of the therapeutic agents.

[0056] To associate one or more drugs with a DKP salt, the drug and the DKP salt are preferably mixed in solution or suspension and subsequently dried. Either component may be present as solute or suspendate. In different embodiments the mixture is spray dried or lyophilized.

[0057] Spray drying is a thermal processing method used to form, load or dry particulate solids from a variety of solutions or suspensions. The use of spray drying for the formation of dry particulate pharmaceuticals is known in the art however in the past its use had been limited by its incompatibility with biological macromolecular drugs, including protein, peptides and nucleic acids due to the nature of the spray drying process. During spray drying, a solution or suspension is formed into droplets through aerosolization and then passed through a heated gas stream having sufficient heat energy to evaporate water and solvents in the particles to a desired level before the particles are collected. The inlet temperature is the temperature of the gas stream leaving its source and its level is selected based upon the lability of the macromolecule being treated. The outlet temperature is a function of the inlet temperature, the heat load required to dry the product along with other factors.

[0058] The present inventors have unexpectedly determined that the particles of the present invention, have aerodynamic performance which improves with increasing content of a biologically active agent which has not been seen with other particles. The respirable fraction (%rf), the percentage of particles between 0.5 and 5.8 μm (microns) in diameter, of the spray dried particles of the present invention increases with increasing insulin content, rather than decreasing as was expected. Therefore using the methods of the present invention, diketopiperazine microparticles can be formed which have higher biologically active agent content that was previously achievable.

[0059] Additionally, the present inventors have surprisingly determined that spray dried FDKP disodium salt/insulin compositions have increased insulin stability as the concentration of the FDKP disodium salt in the starting solution increases. Stability was measured by insulin loss after 17 days at 40°C/75% relative humidity. For example, 8.5% insulin was lost from powder spray dried from a solution containing 37 mg/mL solids (total weight of FDKP disodium salt/insulin). By comparison, 4.5% insulin was lost from powder spray dried from a solution containing 45 mg/mL solids and 2.7% insulin was lost from powder spray dried from a solution containing 67 mg/mL solids.

[0060] In a further observation, inlet temperature was found to have surprising effects on insulin stability. The data indicate that insulin stability in the powder increases with increasing inlet temperature as measured by insulin loss after 17 days at 40°C/75% RH. For example, about 4% insulin was lost from powder spray dried at an inlet temperature of 180°C. By comparison, <1% insulin was lost from powder spray dried at an inlet temperature of 200°C.

[0061] In an embodiment of the present invention, microparticles suitable for delivery to the pulmonary system are provided wherein the microparticles have a rugosity of less than 2. Another aspect of the present invention influenced by spray drying is the particle morphology, measured by rugosity, which is the ratio of the specific area and the surface area calculated from the particle size distribution and particle density. The drying operation may be controlled to provide dried particles having particular characteristics, such as rugosity. Rugosity of spray dried particles is a measure of the morphology of the surface of the particles, such as the degree of folding or convolution.

[0062] It had previously been thought that a rugosity above 2 was needed in order to obtain particles with sufficient dispersability to form a free-flowing powder. Surprisingly, the present inventors have produced particles suitable for pulmonary delivery with a rugosity below 2.

[0063] The microparticle formulations of the present invention can be administered as a liquid or solid form. These can include solutions, suspensions, dry powders, tablets, capsules, suppositories, patches for transdermal delivery, and the like. These different forms offer distinct, but overlapping, advantages. The solid forms provide convenient bulk transport of drugs and can improve their stability. They can also be formed into microparticles enabling administration by inhalation specifically to the nasal mucosa or deep lung, depending on the size of the microparticle. Diketopiperazines

can also facilitate absorption of the associated drug even when delivered as a solution. Some of the DKP salts (for example, the sodium and potassium salts) offer improved solubility at neutral and acidic pH as compared to the free acid, which can lead to improved absorption in the stomach of orally administered solid forms.

[0064] Dikeopiperazine salt counter cations may be selected to produce salts having varying solubilities. These varying solubilities can be the result of differences in dissolution rate and/or Intrinsic solubility. By controlling the rate of DKP salt dissolution, the rate of drug absorption from the DKP salt/drug combination can also be controlled to provide formulations having immediate and/or sustained release profiles. For example, sodium salts of organic compounds are characteristically highly soluble in biological systems, while calcium salts are characteristically only slightly soluble in biological systems. Thus, a formulation comprised of a DKP sodium salt/drug combination would provide immediate drug absorption, while a formulation comprised of a DKP calcium salt/drug combination would provide slower drug absorption. A formulation containing a combination of both of the latter formulations could be used to provide immediate drug absorption followed by a period of sustained absorption.

[0065] Diketopiperazine salt formulations of biologically active agents may be administered orally. Microparticles, depending on the chemical nature and size, are absorbed through the epithelial lining of the gastrointestinal tract into the bloodstream or lymphatic system. Alternatively, the composition can be administered as a solution in which the DKP salt serves to facilitate the absorption of the drug. Additionally, the microparticles can be administered as a suspension or a solid dosage form that dissolves completely and is absorbed following dissolution.

[0066] For parenteral administration, microparticles of less than 5 μm (five microns) readily pass through a needle for intravenous administration. Suitable pharmaceutical carriers, for example, phosphate buffered saline, are known and commercially available. Similarly, microparticles can be injected or implanted subcutaneously. Intramuscularly, or intraperitoneally. Additionally, the microparticles can be placed in an implantable device to facilitate sustained and/or controlled delivery.

[0067] For topical or transdermal administration, microparticles can be suspended in a suitable pharmaceutical carrier for administration using methods appropriate for the carrier and site of administration. For example, microparticles are administered to the eye in a buffered saline solution, at a pH of approximately 7.4, or in an ointment such as mineral oil. The dosage will be dependent on the compound to be released as well as the rate of release. The microparticles, or aggregations of microparticles into films, disks, or tablets, with incorporated compound can be administered to the skin in an ointment, cream, or patch. Suitable pharmaceutical carriers are known to those skilled in the art and commercially available. Mucosal administration, including buccal, vaginal, rectal, nasal administration is also contemplated.

[0068] Pulmonary delivery can be very effectively accomplished using dry powders comprising the microparticles of the invention and can lead to rapid absorption into the circulation (bloodstream). Dry powder Inhalers are known in the art and particularly suitable inhaler systems are described in U.S. patent application nos. 09/621,092 and 10/655,153, both entitled "Unit Dose Capsules and Dry Powder Inhaler". Information on pulmonary delivery using microparticles comprising diketopiperazine can be found in U.S. Patent No. 6,428,771 entitled "Method for Drug Delivery to the Pulmonary System". The following examples are meant to illustrate one or more embodiments of the invention and are not meant to limit the invention to that which is described below.

EXAMPLES

Example 1. Preparation A of FDKP Disodium Salt

[0069] Thirteen grams of fumaryl diketopiperazine (FDKP) (28.73 mmol, 1 equiv.) were placed into a 250 mL 3-neck round bottom flask equipped with a reflux condenser, magnetic stir bar, and thermometer. The reaction was run under a nitrogen atmosphere. Water (150 mL) and 50% sodium hydroxide (4.48 g, 1.95 equiv.) were added sequentially to the flask. The resulting yellow solution was heated to 50°C and held for 2 hours. The solution was then hot filtered to remove any insoluble material. The water was removed from the sample via rotary evaporation. The recovered solids were dried in the vacuum oven (50°C, 30 inches of mercury) overnight. The salt was then assayed for moisture content (Karl Fischer) and sodium content (elemental analysis and titration). The yield of the salt was from about 90% to about 95%.

Molecular Formula: $\text{C}_{20}\text{H}_{26}\text{N}_4\text{Na}_2\text{O}_8 \cdot 1.4809 \text{ H}_2\text{O}$

% Water by Karl Fischer titration: 5.1 %

Elemental Analysis:

Calc	C 45.92	H 5.58	N 10.71	Na 8.79
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Found	C 45.05	H 5.23	N 10.34	Na 9.18
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Titration: 97% disodium salt (weight percent)

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Table 1. Laser diffraction particle size analysis (Preparation A particles):

Lot#	X ₁₀	X ₁₆	X ₅₀	X ₈₄	X ₉₀	X ₉₉	VMD	GSD
Preparation A	1.60 μm	1.44 μm	2.89 μm	4.60 μm	5.47 μm	19.20 μm	3.70 μm	1.59
Lot#	Particle Size							
	<3 μm	0.5 - 5 μm		Fine Particle Fraction (<5.8 μm)				
Preparation A	53.39%	87.91%		91.46%				
VMD = Volume median diameter; GSD = geometric standard deviation.								

Example 2. Preparation B of FDKP Disodium Salt

[0070] Thirteen grams of FDKP (28.73 mmol, 1 equiv.) and ethanol (150 mL) were placed into a 250 mL 3-neck round bottom flask equipped with a reflux condenser, magnetic stir bar, and thermometer. The reaction was run under a nitrogen atmosphere. The slurry was heated to 50°C. Sodium hydroxide, 50% w/w aqueous solution (4.71 g, 2.05 equiv.) was added in one portion. The resulting slurry was held at 50°C for 2 hours. The reaction contents were then cooled to ambient temperature (20-30°C) and the solids isolated by vacuum filtration. The recovered salt was washed with ethanol (300 mL) and acetone (150 mL) and dried in the vacuum oven (50°C, 30 inches of mercury) overnight. No further purification was required. The salt was then assayed for moisture content (Karl Fischer) and sodium content (elemental analysis and titration). The yield of the salt was from about 90% to about 95%.

Molecular Formula: C₂₀H₂₆N₄Na₂O₆·1.4503 H₂O

% Water by Karl Fischer titration: 5%

Elemental Analysis:

Calc	C 45.97	H 5.57	N 10.72	Na 8.8
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Found	C 46.28	H 5.26	N 10.60	Na 8.96
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Titration: 98.8% disodium salt (weight percent)

Table 2. Laser diffraction particle size analysis (Preparation B particles):

Lot#	X ₁₀	X ₁₆	X ₅₀	X ₈₄	X ₉₀	X ₉₉	VMD	GSD
Preparation A	1.55 μm	1.36 μm	3.11 μm	5.53 μm	6.64 μm	14.04 μm	3.76 μm	1.75
Lot#	Particle Size							
	<3 μm		0.5 - 5 μm		Fine Particle Fraction (<5.8 μm)			
	47.37%		80.13%		86.01%			
Preparation A	47.37%		80.13%		86.01%			
VMD = Volume median diameter; GSD = geometric standard deviation.								

Example 3. Preparation A of FDKP Dilithium Salt

[0071] Ten grams of FDKP (22.10 mmol, 1 equiv.) and 100 mL of water were placed into a 200 mL 3-neck round bottom flask equipped with a reflux condenser, magnetic stir bar, and thermometer. The reaction was run under a nitrogen atmosphere. In a separate flask, an aqueous solution of lithium hydroxide (1.81 g, 1.95 equiv.) in 40 mL of water was prepared. Once all of the lithium hydroxide had dissolved, this solution was added in one portion to the aqueous slurry of FDKP. The resulting solution was heated to 50°C and held for 1 hour. The reaction contents were then cooled to ambient temperature and filtered to remove any undissolved particles. The water was removed from the sample via rotary evaporation. The recovered solids were dried in a vacuum oven (50°C, 30 inches of mercury) overnight. The salt was then assayed for moisture content (Karl Fischer) and lithium content (elemental analysis and titration). The yield of the salt was about 98%.

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Molecular Formula: $C_{20}H_{26}N_4Li_2O_8 \cdot 0.0801 H_2O$

Karl Fischer: 0.31%

Elemental Analysis:

Calc C 51.57 H 5.66N 12.03 Li 2.98

Found C 50.98 H 5.74N 11.95 Li 2.91

Titration: 98.3% dilithium salt (weight percent)

Example 4. Preparation A of FDKP Dipotassium Salt

[0072] Twelve grams of FDKP (26.52 mmol, 1 equiv.) were placed into a 250 mL 3-neck round bottom flask equipped with a reflux condenser, magnetic stir bar, and thermometer. The reaction was run under a nitrogen atmosphere. Potassium hydroxide (0.5N, 105 g, 1.98 equiv.) was added to the flask. The resulting solution was heated to 50°C and held for 2 hours. The reactants were cooled to ambient temperature and the water was removed from the sample via rotary evaporation. The recovered solids were dried in the vacuum oven (50°C, 30 inches of mercury) overnight. The salt was then assayed for moisture content (Karl Fischer) and potassium content (elemental analysis and titration). The yield of the salt was from about 95% to about 98%.

Molecular Formula: $C_{20}H_{26}N_4K_2O_8 \cdot 0.4529 H_2O$

Karl Fischer: 4.98%

Elemental Analysis:

Calc C 44.75 H 5.05N 10.44 K 14.56

Found C 44.88 H 4.74N 10.36 K 14.34

Titration: 97.0% dipotassium salt (weight percent)

Example 5. Preparation B of FDKP Dipotassium Salt

[0073] Ten grams of FDKP (22.10 mmol, 1 equiv.) and ethanol (150 mL) were placed into a 250 mL 3-neck round bottom flask equipped with a reflux condenser, magnetic stir bar, and thermometer. The reaction was run under a nitrogen atmosphere. The slurry was heated to 50°C. Potassium hydroxide (10N, 4.64 g, 2.10 equiv.) was added in one portion. The resulting slurry was held at 50°C for a minimum of 3 hours. The reaction contents were cooled to ambient temperature (20-30°C) and the solids isolated by vacuum filtration. The recovered salt was washed with ethanol (100 mL) and acetone (200 mL) and dried in a vacuum oven (50°C, 30 inches of mercury) overnight. No further purification was required. The salt was then assayed for moisture content (Karl Fischer) and potassium content (elemental analysis and titration). The yield of the salt was from about 94% to about 98%.

Molecular Formula: $C_{20}H_{26}N_4K_2O_8 \cdot 0.6386 H_2O$

Karl Fischer: 2.13%

Elemental Analysis:

Calc C 44.47 H 5.09N 10.37 K 14.47

Found C 44.48 H 5.03N 10.31 K 13.92

Titration: 97% dipotassium salt (weight percent)

Example 6. Preparation A of Disodium FDKP-Insulin Microparticles

[0074] Two and a half grams of FDKP disodium salt (Preparation A) was placed in a 250 mL beaker with a magnetic stir bar. The material was suspended in 75 mL of deionized water. Insulin (0.84 g) was added to the FDKP salt suspension. The resulting slurry was titrated to a pH of 8.3 with NH_4OH to form a solution. The FDKP disodium salt and insulin solution was brought to a volume of 100 mL with deionized water and filtered through a 0.22 μm polyethersulfone membrane. The solution was spray-dried using a BUCHI® Mini Spray Dryer B-191 (Buchi Labortechnik AG, Switzerland) under the following conditions.

Inlet Temperature set at 170°C

Outlet Temperature = 75°C

Aspiration rate 80% of maximum

Atomization = 600 l/hr of dry nitrogen
 Feed pump rate 25% of maximum (8.5 ml/min)
 Nozzle chiller return water 22°C.

5 Example 7. Preparation B of Disodium FDKP-Insulin Microparticles

[0075] Five grams of FDKP disodium salt (Preparation B) was placed in a 250 mL beaker with a magnetic stir bar. The material was suspended in 75 mL of deionized water. Insulin (1.68 g) was added to the FDKP salt suspension. The resulting slurry was titrated to a pH of 8.3 with NH₄OH to form a solution. The FDKP disodium salt and insulin solution
 10 was brought to a volume of 100 mL with deionized water and filtered through a 0.22 µm polyethersulfone membrane. The solution was spray-dried using a BUCHI® Mini Spray Dryer B-191 (Buchi Labortechnik AG, Switzerland) under the following conditions.

Inlet Temperature set at 149°C
 15 Outlet Temperature = 75°C
 Aspiration rate 80% of maximum
 Atomization = 600 l/hr of dry nitrogen
 Feed pump rate 25% of maximum (8.5 mL/min)
 Nozzle chiller return water 23°C.

20 Example 8. Characterization of Disodium FDKP-Insulin Microparticle

[0076] The microparticles described in Examples 6 and 7 were subjected to laser diffraction particle size analysis (Sympatec GmbH, Germany) (FIGs. 1A and 1B). The particles of Example 6 displayed an average respirable fraction (according to the USP definition of 0.5 to 5.8 microns) of 87.93% with a standard deviation of 1.60 and a %CV (coefficient of variation) of 1.82. The particles of Example 7 displayed an average respirable fraction of 81.36% with a standard deviation of 4.20 and a %CV of 5.16.

30 Example 9. Pulmonary Administration of Disodium FDKP-Insulin

[0077] A dry powder containing the disodium FDKP salt and insulin is inhaled at the beginning of meal. The particles that comprise the dry powder are preferably in the range of approximately 0.5 - 5.8 microns in size. The exact dosage is patient-specific, but generally on the order of 5-150 Units of insulin per dose. The insulin absorption from this dosage regimen mimics physiologic first-phase insulin release, and attenuates post-prandial blood glucose excursions.

35 Example 10. Preparation of an Oral Dosage Form

[0078] Spray-dried disodium FDKP/insulin powder as described in Examples 6 or 7 is packed into hard gelatin capsules. The capsules can contain approximately 50-100 mg of powder. The FDKP salt/insulin powders prepared in Examples
 40 6 and 7 were 25% insulin by weight and insulin activity was about 26 units/mg. Thus, 50 mg would be on the order of 1300 units, significantly larger than a typical dose. About 2-30 mg of the FDKP salt/insulin powder is mixed with methyl cellulose (other bulking agents are well known in the art) to make up the balance of the desired mass.

45 Example 11. Oral Administration of Disodium FDKP-Insulin

[0079] Capsules containing the FDKP salt and insulin are taken before a meal. The exact dosage is patient-specific, but generally on the order of approximately 10-150 units of insulin is administered per dose. The subsequent insulin absorption attenuates post-prandial blood glucose excursions. This oral insulin formulation is used to replace pre-meal insulin injections in patients with diabetes. Additionally, insulin absorbed through the gastrointestinal tract mimics endogenous insulin secretion. Endogenous insulin is secreted by the pancreas into the portal circulation. Insulin absorbed
 50 following oral administration also goes directly to the portal circulation. Thus, the oral route of insulin administration delivers insulin to its site of action in the liver, offering the potential to control glucose levels while limiting systemic exposure to insulin. Oral insulin delivery using a combination of insulin and the diacid form of FDKP is hindered by the poor solubility of the FDKP diacid in the low pH environment of the gastrointestinal tract. The FDKP salts, however,
 55 provide a local buffering effect that facilitates their dissolution in low pH.

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Example 12. Preparation C of FDKP Di-Sodium Salt

[0080] Fifty grams of fumaryl diketopiperazine (FDKP, 221.01 mmol, 1 equiv.), water (200 mL), and 10 N sodium hydroxide (21.9 mL, 437.61 mmol, 1.98 equiv.) were charged to a 1-liter, 4-neck, round bottom flask equipped with a reflux condenser, overhead stirrer, nitrogen inlet, and thermometer. The mixture was heated to 50°C to achieve a yellow solution and ethanol (650 mL) was added over 15 minutes. When the addition was complete, the slurry was held at 50°C for 30 - 60 minutes. The reaction mixture was vacuum filtered and the isolated solids were washed with ethanol (150 mL) and acetone (150 mL x 2) then dried in a vacuum oven (50°C, 30 inches of mercury) overnight. No further purification was required. The salt was assayed for moisture content (Karl Fischer) and sodium content (elemental analysis and titration). The yield of the salt was from about 90% to about 95%.

Karl Fischer: 7.19 %

Elemental Analysis:

Calc	C 44.91	H 5.70	N 10.47	Na 8.6
Found	C 45.29	H 5.47	N 10.59	Na 8.24

Titration: 98.8 % disodium salt (weight percent)

[0081] The following are various processes described with regard to various formulations of the present invention.

Example 13: FDKP salt/insulin powder prepared by spray drying

[0082] The disodium salt of FDKP (5 g) was dissolved in deionized water (150 mL) and insulin (1.69 g) was added. The pH of the suspension was adjusted to 8.3 with ammonium hydroxide (NH₄OH) to give a solution that was subsequently diluted to 200 mL with deionized water and filtered. The solution was spray dried using the following conditions:

Inlet temperature - 200°C

Outlet temperature - 80°C

Atomization gas - 600 liter N₂/hr

Process gas - 80% of maximum

The spray nozzle was cooled to 28°C

[0083] The resultant particles were analyzed for their aerodynamic properties and the data are reported in Table 3.

Table 3. Aerodynamic properties of spray dried disodium FDKP/insulin.

Sample	%rf	%empty	%rf fill	mmad	gsd	inlet °C	%load	LOD
FDKP disodium salt with 25% insulin (w:w)	44.5	85.6	38.1	3.1	1.9	200	25.00	5.4

[0084] Table 3 shows the respirable fraction (%rf), which is the percentage of particles between 0.5 and 5.8 microns in diameter, the percentage of powder that empties from the cartridge upon discharge (% empty), the percentage of respirable fraction per fill (% rf fill, %rf X %empty - this measures the % of the respirable particles in the powder emptied from the cartridge, the mass median aerodynamic diameter (mmad), the inlet °C (the inlet temperature in degrees Celsius), the percentage of load (%load - the insulin content of particles in weight %), and the loss on drying (LOD), a measure of the residual water in the powder expressed as the % volatile material removed when the powder is dried in an oven overnight.

[0085] Particle size measured by laser diffraction demonstrated a size range of approximately 2µm - 15µm and the data are displayed in Table 4 and in Figure 2.

Table 4

Lot#	Run	X ₁₀	X ₅₀	X ₉₀	VMD	GSD	Fine Particle Fraction (<5.8 µm)
FDKP disodium salt with 25% insulin (w:w)	168	2.14 µm	5.88 µm	15.16 µm	7.76 µm	2.10	49.21 %

[0086] Scanning electron microscopy (SEM) was utilized to study particle morphology. A representative SEM is shown

in Figure 3. The particle morphology is consistent with a collapsed hollow sphere.

[0087] The stability of the disodium salt/insulin particles was evaluated under accelerated room temperature conditions (40°/75% relative humidity [RH]). Compared to a control formulation prepared by lyophilization, the spray-dried particles demonstrated superior insulin stability as measured by insulin degradation (Figure 4).

[0088] The starting concentration of the FDKP disodium salt/25%insulin solution prior to spray drying was evaluated for its effect on final particle stability. The data (Figure 5) shows that insulin stability on the particle increases with increasing solution concentrations as measured by insulin loss after 17 days at 40°/75% RH.

Example 14. Solvent/anti-solvent precipitation of a solution of FDKP salt/insulin with an organic solvent

[0089] The precipitation was controlled using harmonic ultrasonic atomization. Alternate cavitation methods as well as high shear mixing and homogenization are also applicable.

[0090] The disodium salt of FDKP (5 g) was dissolved in deionized water (80 mL). Insulin (0.65 g) was added to the solution to produce a suspension. The pH of the suspension was adjusted to 8.3 with NH₄OH to obtain a solution that was diluted to 100 mL with deionized water and filtered. The particles were precipitated by pumping the insulin/disodium salt of FDKP solution and ethanol in a 1:5 ratio through a dual inlet atomization hom vibrating at a frequency between 20 kHz and 40 kHz. The precipitate was collected in a media bottle containing ethanol (200 mL). Post-precipitation the material was washed with ethanol and dried via rotary evaporation or by bubbling nitrogen through the suspension. The particles contained 12.5% insulin by weight. Particle morphology was evaluated by SEM (Figure 6).

[0091] The particles illustrated in FIG. 6A (10k x) and FIG. 6B (20K x) are in the 1 to 5 micron range while at lower magnification (FIG. 6C, 2.5k x and FIG. 6D, 1.0k x) particles in the 10 to 40 micron range are seen. It is the non-binding hypothesis of the present inventors that the drying methods utilized in this study resulted in recrystallization of the primary particles into much larger secondary particles and that the use of a method that maintains a constant ratio of organic to aqueous components throughout the drying process, such as spray drying, can preserve the primary particles to the exclusion of the formation of a significant number of secondary particles.

Example 15. *In situ* diammonium salt formation and formulation.

[0092] FDKP or SDKP (succinyl DKP) diammonium salt/insulin particles were formed by spray drying. A representative procedure is given for the FDKP ammonium salt/insulin formulation containing 25% insulin.

[0093] FDKP (5 g) was suspended in deionized water (150 mL) and titrated to a pH of 7.5 to 8.0 with ammonium hydroxide (NH₄OH). Insulin (1.69 g) was added to the resulting solution (FDKP) to give a suspension. The pH of the suspension was adjusted to 8.3 with ammonium hydroxide (NH₄OH) to give a solution that was diluted to 200 mL with deionized water and filtered. The powder was produced by spray drying the solution under the following conditions.

Inlet temperature - 200°C
 Outlet temperature - 80°C
 Atomization gas - 600 liter N₂/hr
 Process gas - 80% of maximum
 The spray nozzle was cooled to 28°C

[0094] The %rf of the diammonium salts is about 10% higher than the %rf of the disodium salt. The counter cation has a large effect on particle performance. Also, the 50% FDKP ammonium salt/Insulin powder has a %rf comparable to that of the corresponding 25% FDKP ammonium salt/insulin powder. This is surprising because with the powders prepared by lyophilization from the FDKP free acid, the %rf decreases as the insulin content increases.

[0095] The resultant particles were analyzed for their aerodynamic properties and the data are reported in Table 5.

Table 5. Aerodynamic properties of spray dried diammonium FDKP/insulin and diammonium SDKP/insulin

Sample	%rf	%empty	%rf fill	mmad	gsd	inlet °C	%load	LOD
FDKP diammonium salt with 25% insulin (w:w)	52.1	88.7	46.2	2.9	1.9	200	25.00	6.6
FDKP diammonium salt with 50% insulin (w:w)	55.7	85.4	47.5	2.9	1.8	200	50.00	6.2
SDKP diammonium salt with 25% insulin (w:w)	56.0	90.1	55.7	3.0	2.0	200	25.00	3.8

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[0096] Particle size measured by laser diffraction and the data are displayed Table 6 and in Figures 7-9.

Table 6

Lot#	Run	X ₁₀	X ₅₀	X ₉₀	VMD	GSD	Fine Particle Fraction (<5.8 μm)
FDKP diammonium salt with 25% insulin (w:w)	078	1.70 μm	4.10 μm	8.40 μm	4.68 μm	1.86	72.13%

[0097] Particle size of a preparation of the diammonium salt of FDKP containing 25% insulin (w:w) was determined by laser diffraction and demonstrated a size range of approximately 1.7 μm - 8.4 μm for the FDKP ammonium salt formulated with 25% insulin (Figure 7 and Table 7).

Table 7

Lot#	Run	X ₁₀	X ₅₀	X ₉₀	VMD	GSD	Fine Particle Fraction (<5.8 μm)
FDKP diammonium salt with 50% insulin (w:w)	076	1.57 μm	4.51 μm	8.79 μm	4.97 μm	1.91	66.95%

[0098] Particle size of a preparation of the diammonium salt of FDKP containing 50% Insulin (w:w) was determined by laser diffraction and demonstrated a size range of approximately 1.6 μm - 8.8 μm for the FDKP ammonium salt formulated with 50% insulin (Table 8).

Table 8

Lot#	Run	X ₁₀	X ₅₀	X ₉₀	VMD	GSD	Fine Particle Fraction (<5.8 μm)
SDKP diammonium salt with 25% insulin (w:w)	084	1.66 μm	4.64 μm	927 μm	5.17 μm	1.82	64.69%

[0099] Particle size for the SDKP diammonium salt formulated with 25% insulin (w:w) was determined by laser diffraction and demonstrated a size range of approximately 1.7 μm - 9.3 μm for the SDKP diammonium salt formulated with 25% insulin.

[0100] Scanning electron microscopy was utilized to study particle morphology. Representative SEMs are shown in the Figure 10 (FDKP) and Figure 11 (SDKP). The particle morphology is consistent with a collapsed hollow sphere.

[0101] The stability of the *In situ* salt formation and formulation of the diammonium salt/insulin particles was evaluated under accelerated room temperature conditions (40°/75% RH). Compared to a control formulation prepared by lyophilization, the spray dried particles demonstrated superior Insulin stability as measured by Insulin degradation (FDKP, Figure 12 and SDKP, Figure 14) and formation of the desamino degradant (A₂₁) (FDKP, Figure 13 and SDKP, Figure 15).

Example 16. Characteristics of Spray Dried Microparticles.

[0102] Spray dried FDKP salt/insulin particles demonstrate a surprising and unexpected trend in aerodynamic performance. Previously observed insulin-containing microparticles, which had been formed from DKP free acid microparticles onto which insulin had been loaded and the solvent removed by lyophilization, demonstrated decreased aerodynamic performance with increasing insulin content. For example, the %rf (respirable fraction) for 25% loaded particles was significantly lower than the %rf for 5% loaded particles. For spray dried FDKP salt microparticles containing insulin, the opposite trend is observed. As insulin load increases, %rf increases.

[0103] Spray dried powders of the FDKP disodium salt were prepared with insulin contents of 11.4%, 50.0%, 70.0%, or 90.0% (w:w). Figure 16 shows that %rf increases with increasing insulin load.

[0104] A similar trend was also observed in spray dried FDKP diammonium salt/insulin powders having insulin contents of 11.4%, 50.0%, 70.0%, or 90.0% (w:w). The %rf increased with insulin load (Figure 17).

[0105] The starting concentration of the FDKP disodium salt solution prior to spray drying was evaluated for its effect on final particle insulin stability. The data indicate that insulin stability in the powder increases with increasing solution concentrations as measured by insulin loss after 17 days at 40°/75% RH. For example, 8.5% insulin was lost from powder spray dried from a solution containing 37 mg/mL solids. By comparison, 4.5% insulin was lost from powder spray dried

from a solution containing 45 mg/mL solids and 2.7% insulin was lost from powder spray dried from a solution containing 67 mg/mL solids.

[0106] The inlet temperatures used to spray dry solutions of the FDKP disodium salt and insulin to form particles containing 50% insulin was evaluated for its effect on final particle insulin stability. The data indicate that insulin stability in the powder increases with increasing inlet temperature as measured by insulin loss after 17 days at 40°/75% RH. For example, about 4% insulin was lost from powder spray dried at an inlet temperature of 180°C. By comparison, <1% insulin was lost from powder spray dried at an inlet temperature of 200°C.

[0107] Additionally, the present inventors have unexpectedly found that these particles, which are suitable for pulmonary delivery, have a rugosity of approximately 1.

[0108] Unless otherwise indicated, all numbers expressing quantities of ingredients, properties such as molecular weight, reaction conditions, and so forth used in the specification and claims are to be understood as being modified in all instances by the term "about". Accordingly, unless indicated to the contrary, the numerical parameters set forth in the following specification and attached claims are approximations that may vary depending upon the desired properties sought to be obtained by the present invention.

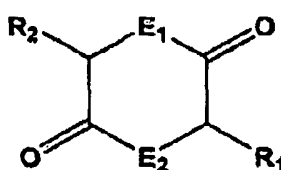
[0109] The terms "a" and "an" and "the" and similar references used in the context of describing the invention (especially in the context of the following claims) are to be construed to cover both the singular and the plural, unless otherwise indicated herein or clearly contradicted by context. All methods described herein can be performed in any suitable order unless otherwise indicated herein or otherwise clearly contradicted by context. The use of any and all examples, or exemplary language (e.g. "such as") provided herein is intended merely to better illuminate the invention and does not pose a limitation on the scope of the invention otherwise claimed. No language in the specification should be construed as indicating any non-daimed element essential to the practice of the invention.

[0110] Groupings of alternative elements or embodiments of the invention disclosed herein are not to be construed as limitations. Each group member may be referred to and claimed individual or in any combination with other members of the group or other elements found herein. It is anticipated that one or more members of a group may be included in, or deleted from, a group for reasons of convenience and/or patentability. When any such inclusion or deletion occurs, the specification is herein deemed to contain the group as modified thus fulfilling the written description of any and all Markush groups used in the appended claims.

[0111] Preferred embodiments of this invention are described herein, including the best mode known to the inventors for carrying out the invention. Of course, variations on those preferred embodiments will become apparent to those of ordinary skill in the art upon reading the foregoing description. The inventors expect skilled artisans to employ such variations as appropriate, and the inventors intend for the invention to be practiced otherwise than specifically described herein.

Claims

1. A pharmaceutically-acceptable salt of a heterocyclic compound according to Formula 1:



Formula 1

wherein R₁ and R₂ comprise 4-X-aminobutyl and wherein X is selected from the group consisting of succinyl, glutaryl, methyl, citraconyl, malonyl, oxalyl and fumaryl;

E₁ and E₂ are NH; and

said salt further comprises at least one cation; and

wherein said salt is not a lithium salt of 2,5-diaspartyl-3,6-diketopiperazine or 2,5-diglutamyl-3,6-diketopiperazine and said salt is in a solid form;

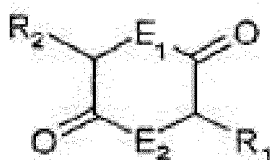
wherein the heterocyclic compound comprises a diketopiperazine.

2. The pharmaceutically-acceptable salt of claim 1 wherein said cation is selected from the group consisting of sodium, magnesium, potassium, calcium, lithium, ammonium, triethylamine, butylamine, diethanolamine, triethanolamine and pyridines.

3. Microparticles comprising the pharmaceutically acceptable salt of claim 1 or 2 and a biologically active agent.
4. The microparticles of claim 3 having a diameter between about 0.5 microns and about ten microns.
5. A solid dosage form comprising a precipitate comprising the pharmaceutically acceptable salt of claim 1 or 2 and a biologically active agent
6. A solid dosage form or dry powder medicament comprising the microparticles of claim 3 or 4.
7. A composition comprising the pharmaceutical acceptable salt of claim 1 or 2 and a biologically active agent or the microparticles of claim 3 or 4, or the solid dosage form of claim 5 or the solid dosage form or dry powder medicament of claim 6, for use in medicine.
8. The composition or the microparticles for use according to claim 7, provided as a dry powder.
9. The composition or the microparticles or the solid dosage form or dry powder medicament for use according to claim 7 or 8 wherein said biologically active agent is selected from the group consisting of hormones, anticoagulants, immunomodulating agents, cytotoxic agents, antibiotics, antivirals, antisense, antigens, antibodies and active fragments and analogues thereof.
10. The composition or the microparticles or the solid dosage form or dry powder medicament for use according to claim 7 or 8 wherein said biologically active agent is selected from the group consisting of insulin, cannabinoids, heparins, calcitonin, felbamate, parathyroid hormone and fragments thereof, growth hormone, erythropoietin, glucagon-like peptide-1, somatotrophin-releasing hormone, follicle stimulating hormone, cromolyn, adiponectin, RNase, ghrelin, zidovudine, didanosine, tetrahydrocannabinol, atropine, granulocytes colony stimulating factor, lamotrigine, chorionic gonadotropin releasing factor, luteinizing releasing hormone, beta-galactosidase and Argatroban.
11. The composition or the microparticles for use according to claim 8 wherein the dry powder releases said biologically active agent in the pulmonary system after administration to a subject by inhalation.
12. The dry powder medicament or the microparticles for use according to claim 9, wherein the biologically active agent is absorbed into the systemic blood circulation.
13. The dry powder medicament or the microparticles for use according to claim 9 wherein the biologically active agent acts locally in the lung.
14. A method of making the solid dosage form of claim 5 or 6 comprising forming by precipitation or by lyophilization.

Patentansprüche

1. Pharmazeutisch unbedenkliches Salz einer heterocyclischen Verbindung gemäß Formel 1:



Formel 1

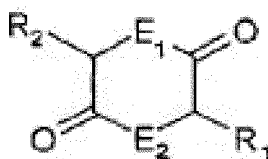
wobei R_1 und R_2 4-X-Aminobutyl beinhalten und wobei X aus der Gruppe ausgewählt ist, die aus Succinyl, Glutaryl, Maleyl, Citranconyl, Malonyl, Oxalyl und Fumaryl besteht;
 E_1 und E_2 NH sind; und
das Salz ferner mindestens ein Kation beinhaltet; und
wobei das Salz kein Lithiumsalz von 2,5-Diaspartyl-3,6-diketopiperazin oder 2,5-Diglutamyl-3,6-diketopiperazin ist und das Salz in fester Form vorliegt;

wobei die heterocyclische Verbindung ein Diketopiperazin beinhaltet.

2. Pharmazeutisch unbedenkliches Salz gemäß Anspruch 1, wobei das Kation aus der Gruppe ausgewählt ist, die aus Natrium, Magnesium, Kalium, Calcium, Lithium, Ammonium, Triethylamin, Butylamin, Diethanolamin, Triethanolamin und Pyridinen besteht.
3. Mikropartikel, die das pharmazeutisch unbedenkliche Salz gemäß Anspruch 1 oder 2 und einen biologischen Wirkstoff beinhalten.
4. Mikropartikel gemäß Anspruch 3 mit einem Durchmesser zwischen etwa 0,5 Mikrometer und etwa zehn Mikrometer.
5. Feste Arzneiform, die ein Präzipitat umfasst, welches das pharmazeutisch unbedenkliche Salz gemäß Anspruch 1 oder 2 und einen biologischen Wirkstoff umfasst.
6. Feste Arzneiform oder Trockenpulvermedikament, umfassend die Mikropartikel gemäß Anspruch 3 oder 4.
7. Zusammensetzung, umfassend das pharmazeutisch unbedenkliche Salz gemäß Anspruch 1 oder 2 und einen biologischen Wirkstoff, oder die Mikropartikel gemäß Anspruch 3 oder 4, oder die feste Arzneiform gemäß Anspruch 5 oder die feste Arzneiform oder das Trockenpulvermedikament gemäß Anspruch 6 zur Verwendung in der Medizin.
8. Zusammensetzung oder Mikropartikel zur Verwendung gemäß Anspruch 7, bereitgestellt als Trockenpulver.
9. Zusammensetzung oder Mikropartikel oder feste Arzneiform oder Trockenpulvermedikament zur Verwendung gemäß Anspruch 7 oder 8, wobei der biologische Wirkstoff aus der Gruppe ausgewählt ist, die aus Hormonen, Antikoagulanzen, Immunmodulatoren, Zytotoxika, Antibiotika, antiviralen Mitteln, Antisense, Antigenen, Antikörpern und aktiven Fragmenten und Analoga davon besteht.
10. Zusammensetzung oder Mikropartikel oder feste Arzneiform oder Trockenpulvermedikament zur Verwendung gemäß Anspruch 7 oder 8, wobei der biologische Wirkstoff aus der Gruppe ausgewählt ist, die aus Folgenden besteht: Insulin, Cannabinoiden, Heparinen, Calcitonin, Felbamat, Parathormon und Fragmenten davon, Wachstumshormon, Erythropoetin, Glucagon-ähnlichem Peptid-1, Somatotropin-Releasing-Hormon, follikelstimulierendem Hormon, Cromolyn, Adiponectin, RNase, Ghrelin, Zidovudin, Didanosin, Tetrahydrocannabinol, Atropin, Granulozytenkolonie-stimulierendem Faktor, Lamotrigin, Choriongonadotropin-Releasing-Faktor, Luteinisierendes-Releasing-Hormon, Beta-Galactosidase und Argatroban.
11. Zusammensetzung oder Mikropartikel zur Verwendung gemäß Anspruch 8, wobei das Trockenpulver den biologischen Wirkstoff nach dem Verabreichen in einem Subjekt durch Inhalieren im Lungensystem freisetzt.
12. Trockenpulvermedikament oder Mikropartikel zur Verwendung gemäß Anspruch 9, wobei der biologische Wirkstoff in den systemischen Blutkreislauf absorbiert wird.
13. Trockenpulvermedikament oder Mikropartikel zur Verwendung gemäß Anspruch 9, wobei der biologische Wirkstoff lokal in der Lunge wirkt.
14. Verfahren zum Herstellen der festen Arzneiform gemäß Anspruch 5 oder 6, umfassend das Ausbilden durch Präzipitation oder durch Gefriertrocknen.

Revendications

1. Sel pharmaceutiquement acceptable d'un composé hétérocyclique de Formule 1 :



Formule 1

- où R_1 et R_2 comprennent du 4-X-aminobutyle et où X est choisi dans le groupe constitué par le succinyle, le glutaryle, le maléyle, le citraconyle, le malonyl, l'oxalyle et le fumaryle ;
 E_1 et E_2 sont NH ; et
 ledit sel comprend en outre au moins un cation ; et
 où ledit sel n'est pas un sel de lithium de 2,5-diaspartyl-3,6-dicétopipérazine ou de 2,5-diglutamyl-3,6-dicétopipérazine et ledit sel est sous forme solide ;
 où le composé hétérocyclique comprend une dicétopipérazine.
2. Sel pharmaceutiquement acceptable selon la revendication 1, dans lequel ledit cation est choisi dans le groupe constitué par le sodium, le magnésium, le potassium, le calcium, le lithium, l'ammonium, la triéthylamine, la butylamine, la diéthanolamine, la triéthanolamine et les pyridines.
 3. Microparticules comprenant le sel pharmaceutiquement acceptable selon la revendication 1 ou 2 et un agent biologiquement actif.
 4. Microparticules selon la revendication 3, ayant un diamètre compris entre 0,5 micron et 10 microns environ.
 5. Forme pharmaceutique solide comprenant un précipité comprenant le sel pharmaceutiquement acceptable selon la revendication 1 ou 2 et un agent biologiquement actif.
 6. Forme pharmaceutique solide ou médicament en poudre sèche comprenant les microparticules selon la revendication 3 ou 4.
 7. Composition comprenant le sel pharmaceutiquement acceptable selon la revendication 1 ou 2 et un agent biologiquement actif, ou les microparticules selon la revendication 3 ou 4, ou forme pharmaceutique solide selon la revendication 5 ou forme pharmaceutique solide ou médicament en poudre sèche selon la revendication 6, pour une utilisation en médecine.
 8. Composition ou microparticules pour une utilisation selon la revendication 7, fournie(s) sous forme de poudre sèche.
 9. Composition ou microparticules ou forme pharmaceutique solide ou médicament en poudre sèche, pour une utilisation selon la revendication 7 ou 8, dans lesquels ledit agent biologiquement actif est choisi dans le groupe constitué par les hormones, les anticoagulants, les agents immunomodulateurs, les agents cytotoxiques, les antibiotiques, les antiviraux, les antisens, les antigènes, les anticorps, ainsi que leurs fragments et analogues actifs.
 10. Composition ou microparticules ou forme pharmaceutique solide ou médicament en poudre sèche, pour une utilisation selon la revendication 7 ou 8, dans lesquels ledit agent biologiquement actif est choisi dans le groupe constitué par l'insuline, les cannabinoïdes, les héparines, la calcitonine, le felbamate, l'hormone parathyroïdienne et ses fragments, l'hormone de croissance, l'érythropoïétine, le peptide-1 apparenté au glucagon, la somatolibérine, la folliculostimuline, l'acide cromoglycique, l'adiponectine, la RNase, la ghréline, la zidovudine, la didanosine, le tétrahydrocannabinol, l'atropine, le facteur de croissance des granulocytes et monocytes, la lamotrigine, le gonadolibérine chorionique, la lutéolibérine, la bêta-galactosidase et l'Argatroban.
 11. Composition ou microparticules pour une utilisation selon la revendication 8, dans lesquelles la poudre sèche libère ledit agent biologiquement actif dans l'appareil respiratoire après administration à un patient par inhalation.
 12. Médicament en poudre sèche ou microparticules pour une utilisation selon la revendication 9, dans lesquels l'agent biologiquement actif est absorbé dans la circulation sanguine systémique.

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13. Médicament en poudre sèche ou microparticules pour une utilisation selon la revendication 9, dans lesquels l'agent biologiquement actif agit localement dans les poumons.

5 14. Procédé de fabrication d'une forme pharmaceutique solide selon la revendication 5 ou 6, comprenant la formation par précipitation ou par lyophilisation.

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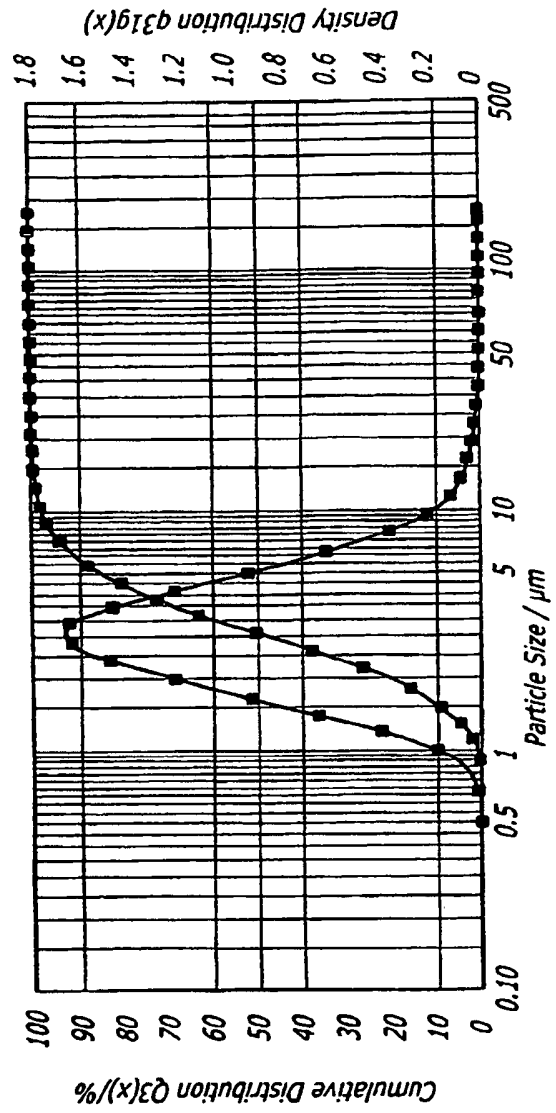
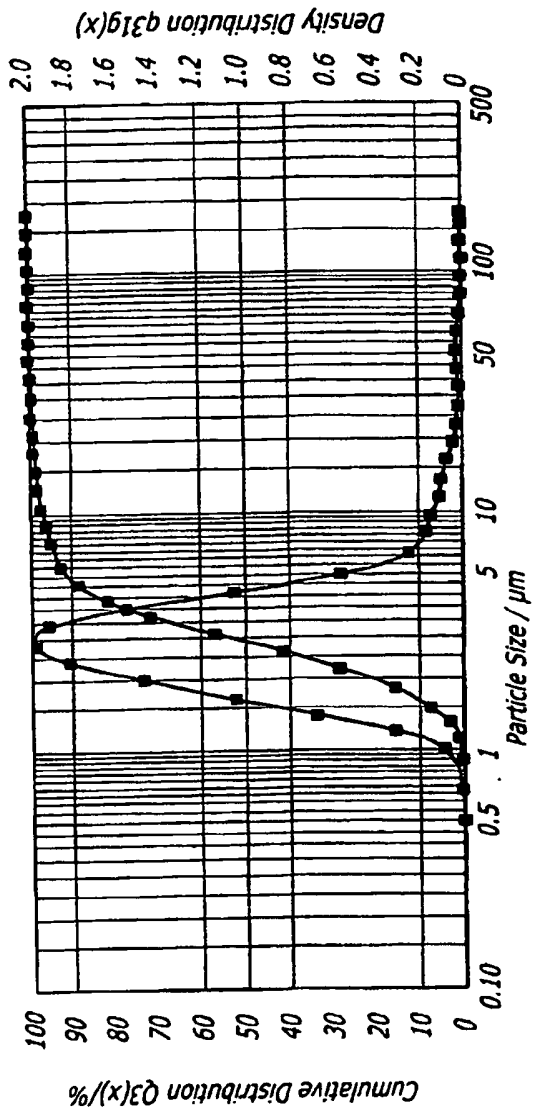
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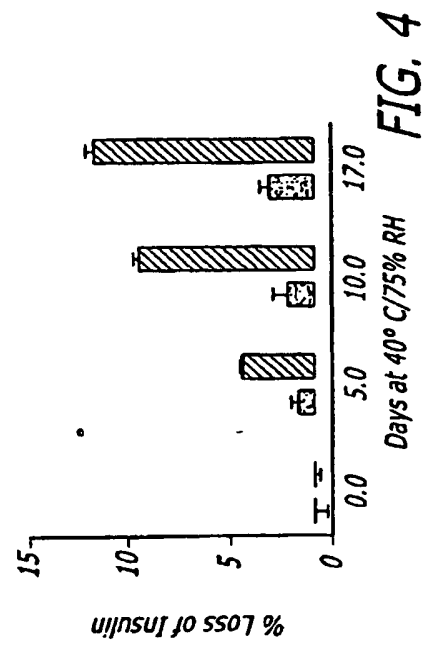
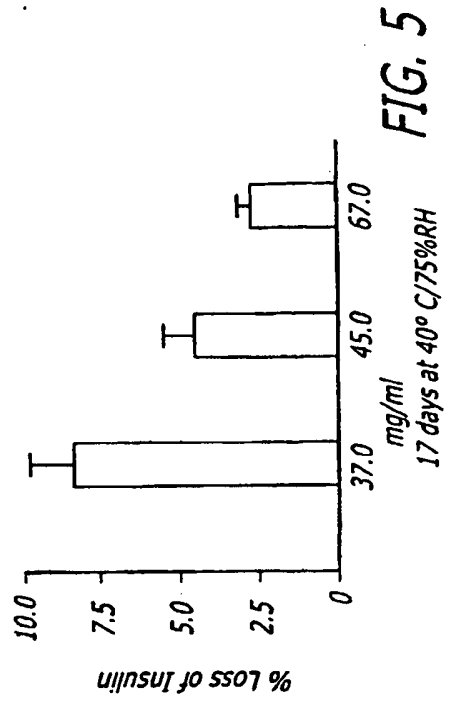
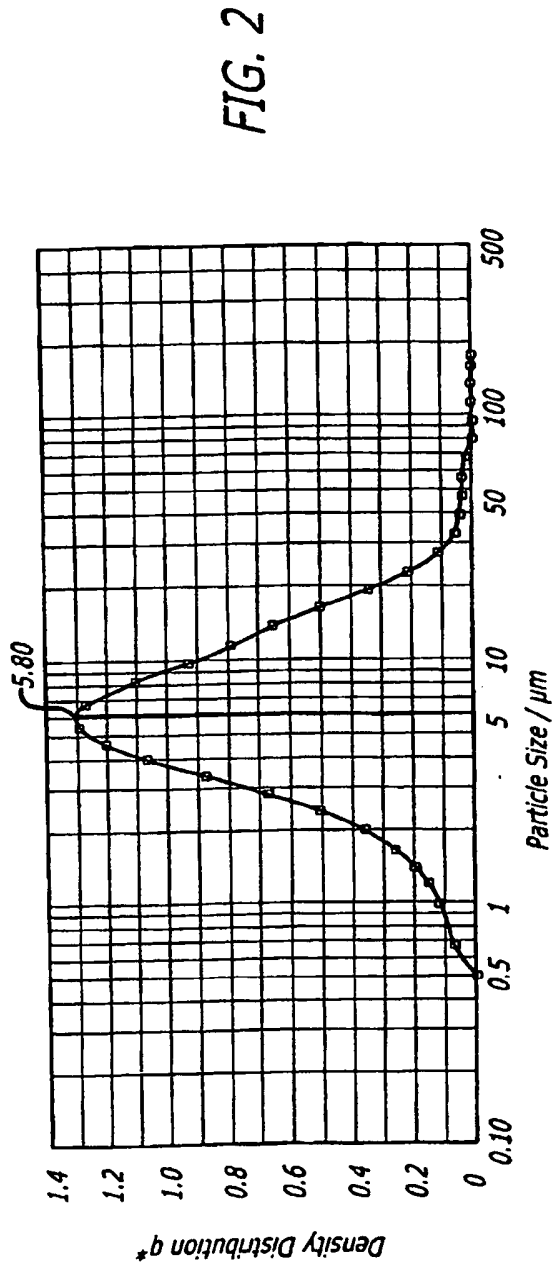
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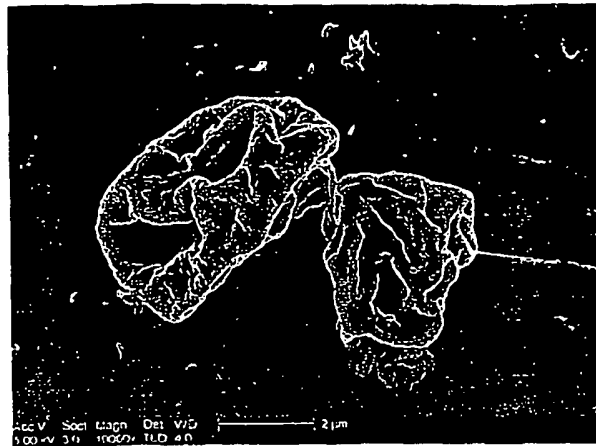


FIG. 3

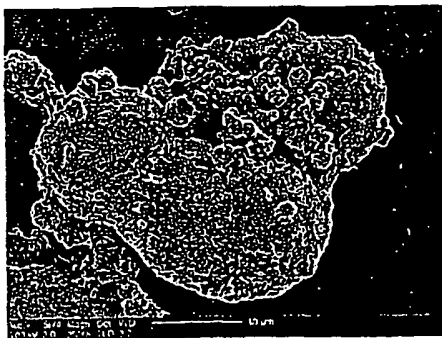
FIG. 6



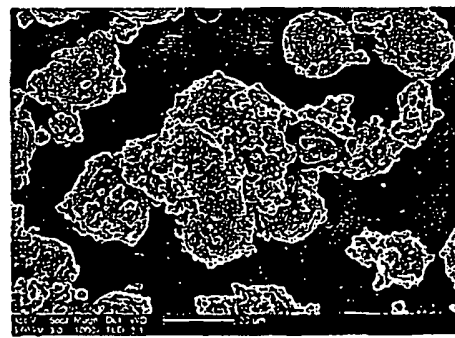
A



B



C



D

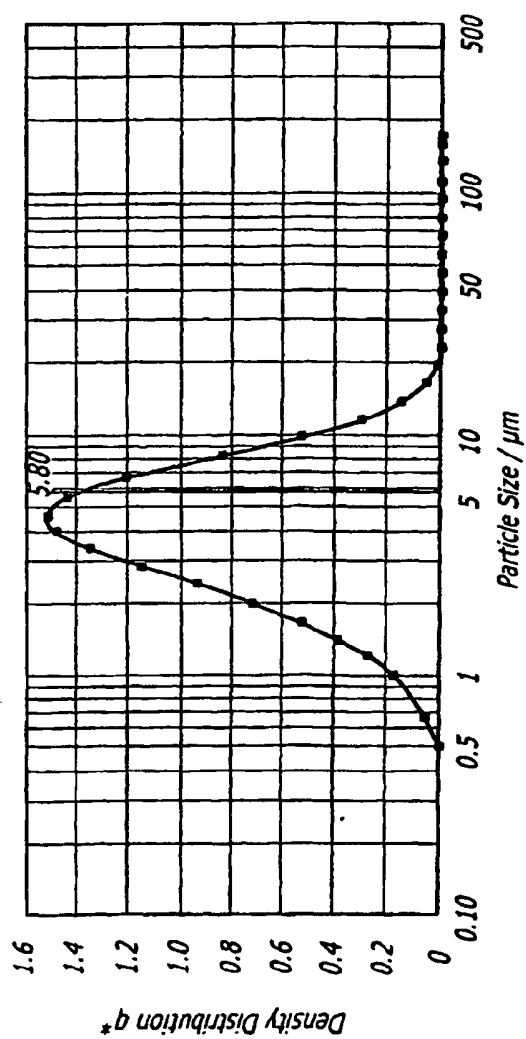


FIG. 7

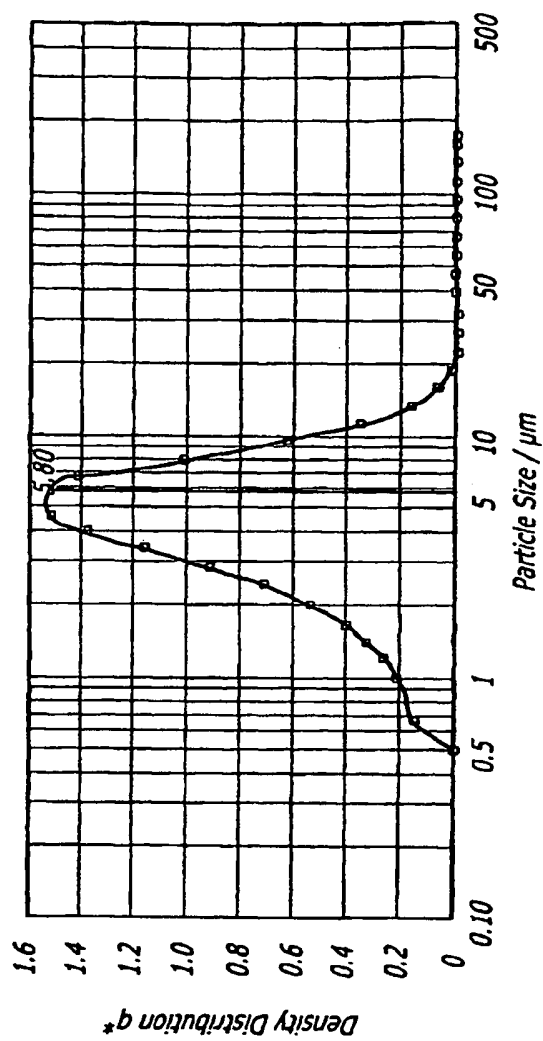
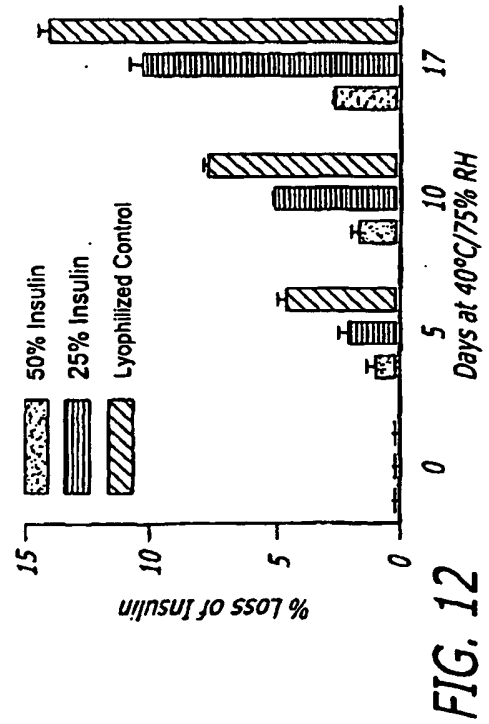
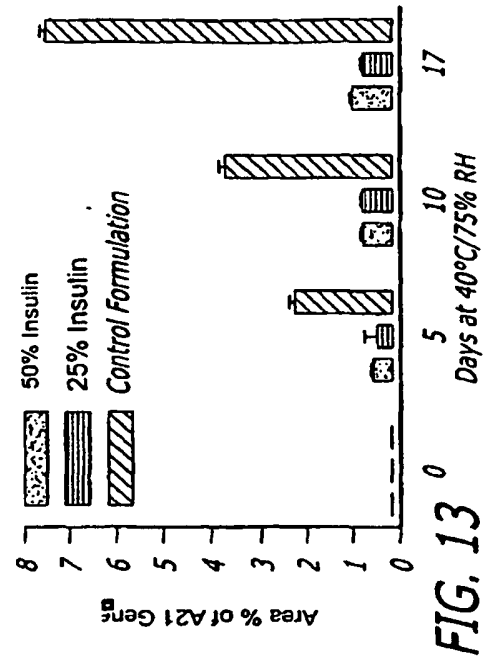
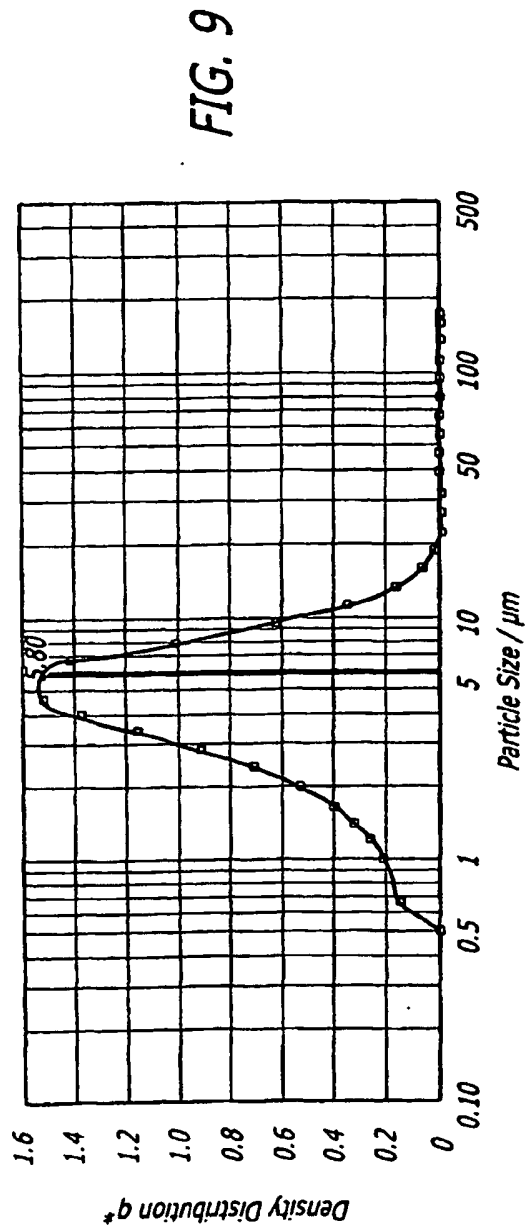


FIG. 8



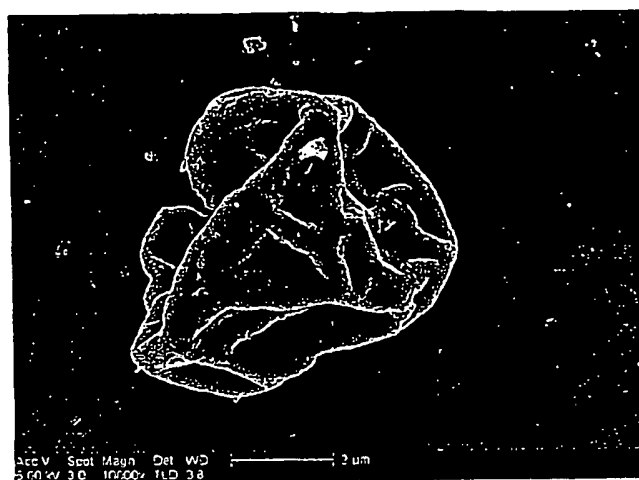


FIG. 10



FIG. 11

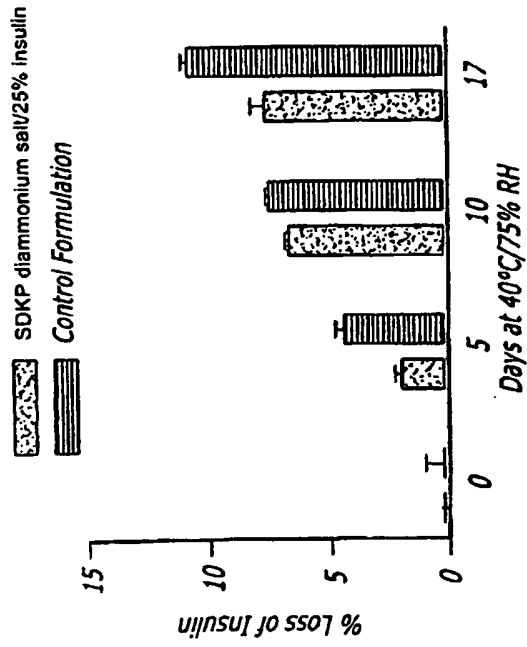


FIG. 14

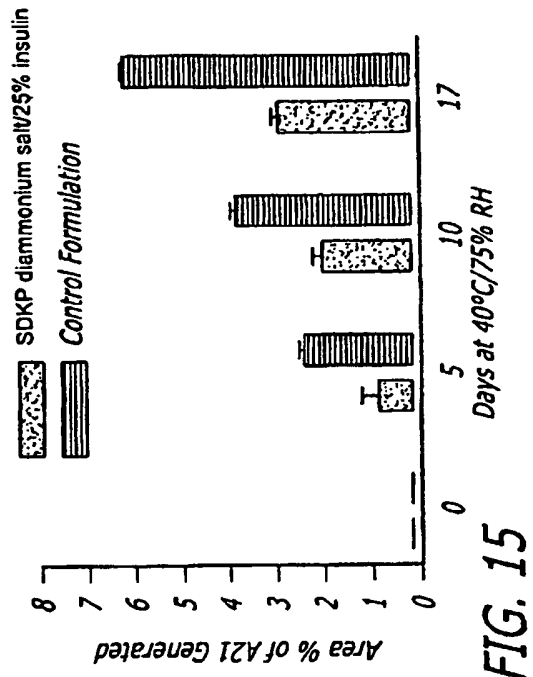


FIG. 15

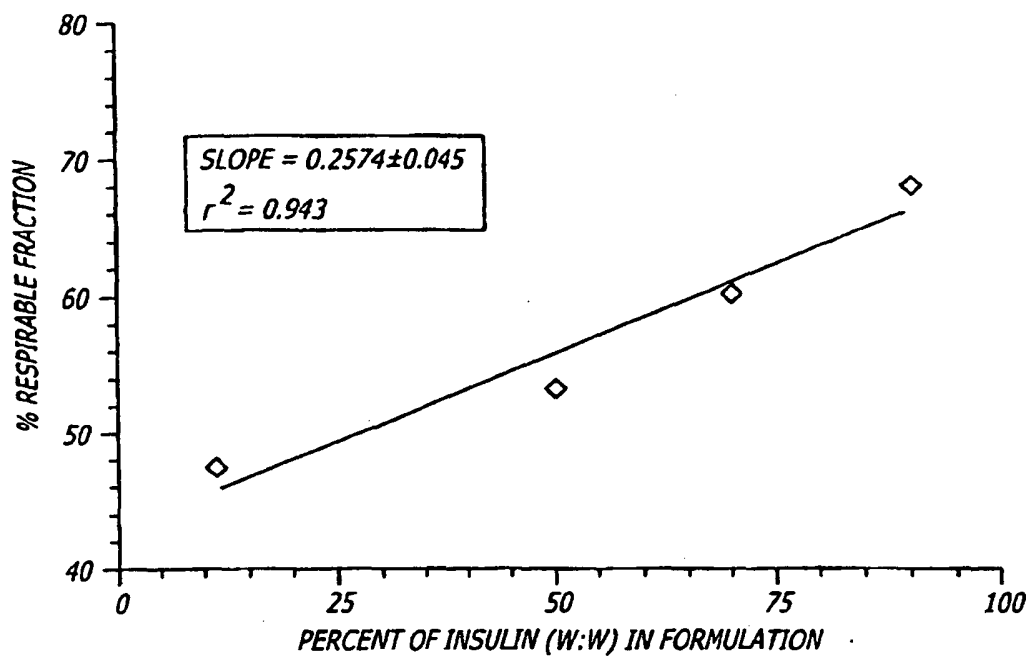


FIG. 16

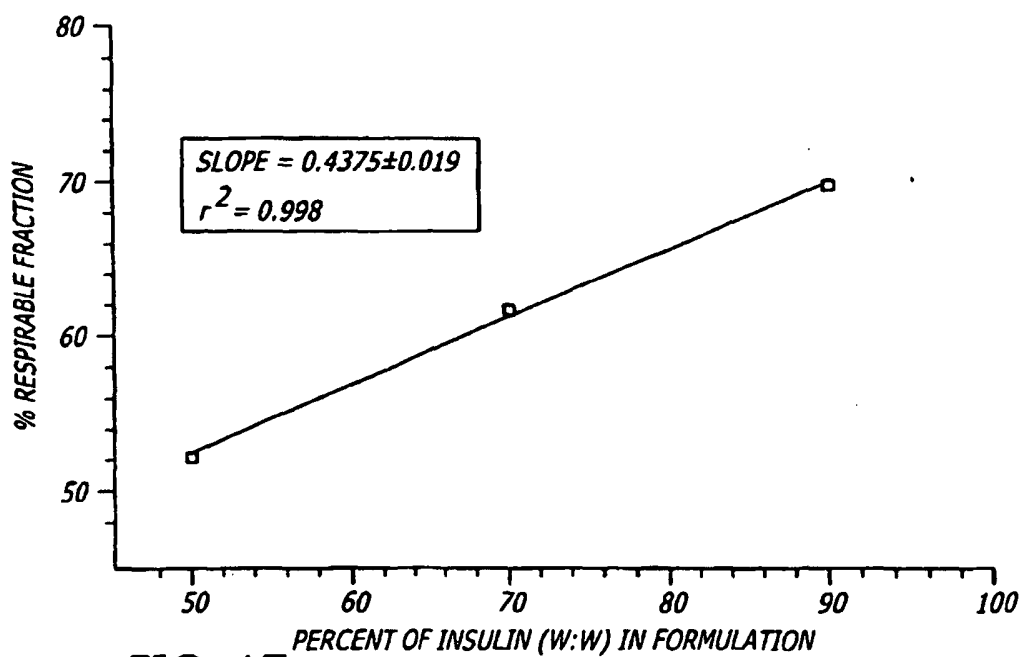


FIG. 17

REFERENCES CITED IN THE DESCRIPTION

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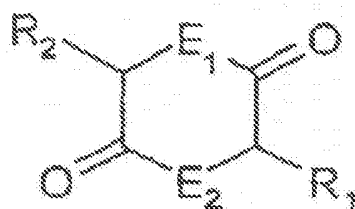
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Diketo-piperazin-sók gyógyszeradagolás céljára

Szabadalmi igénypontok

1. Az I általános képletű heterociklusos vegyület gyógyászatilag elfogadható sója



I

ahol

R₁ vagy R₂ 4-X-amino-butyl-csoportot tartalmaz és ahol X jelentése a szukcinil, glutaril, maleil, citrakonil, malonil, oxalil és fumaril közül kerül kiválasztásra;

E₁ és E₂ jelentése NH; és

a só továbbá legalább egy kationt is tartalmaz; és

ahol a só nem 2,5-diaszpartil-3,6-diketo-piperazin-lítiumsó vagy 2,5-diglutamil-3,6-diketo-piperazin-lítiumsó, és a só szilárd formájú;

ahol a heterociklusos vegyület egy diketo-piperazint tartalmaz.

2. Az 1. igénypont szerinti gyógyászatilag elfogadható só, ahol a kation a nátrium, magnézium, kálium, kalcium, lítium, ammónium, trietil-amin, butil-amin, dietanol-amin, trietanol-amin és piridinek közül kerül kiválasztásra.

3. Mikrorészecskék, amelyek az 1. vagy 2. igénypont szerinti gyógyászatilag elfogadható sót és biológiailag aktív szert tartalmaznak.

4. A 3. igénypont szerinti mikrorészecskék, amelyek átmérője körülbelül 0,5 mikron és körülbelül tíz mikron közötti.

5. Szilárd gyógyszeradagolási forma, amely az 1. vagy 2. igénypont szerinti gyógyászatilag elfogadható sót és biológiailag aktív szert tartalmazó kicsapott anyagot tartalmaz.

6. Szilárd gyógyszeradagolási forma vagy száraz por formájú gyógyszer, amely a 3. vagy 4. igénypont szerinti mikrorészecskéket tartalmazza.

7. Készítmény, amely az 1. vagy 2. igénypont szerinti gyógyászatiilag elfogadható só és biológiailag aktív szert vagy a 3. vagy 4. igénypont szerinti mikrorészecskéket vagy az 5. igénypont szerinti szilárd gyógyszeradagolási formát vagy a 6. igénypont szerinti szilárd gyógyszeradagolási formát vagy száraz por formájú gyógyszert tartalmazza a gyógyászatban történő felhasználásra.

8. A készítmény vagy a mikrorészecskék a 7. igénypont szerinti felhasználásra száraz por formájában.

9. A készítmény vagy a mikrorészecskék vagy a szilárd gyógyszerforma vagy a száraz por formájú gyógyszer a 7. vagy 8. igénypont szerinti felhasználásra, ahol a biológiailag aktív szer a hormonok, antikoagulánsok, immunmoduláló szerek, citotoxikus szerek, antibiotikumok, vírusellenes szerek, antiszenz, antigének, antitestek és aktív fragmentumok és analógjaik közül kerül kiválasztásra.

10. A készítmény vagy a mikrorészecskék vagy a szilárd gyógyszerforma vagy a száraz por formájú gyógyszer a 7. vagy 8. igénypont szerinti felhasználásra, ahol a biológiailag aktív szer az inzulin, kannabinoidok, heparinok, kalcitonin, felbamát, paratiroid hormon és fragmentumai, növekedési hormon, eritropoietin, glukagonszerű peptid-1, szomatotrofint felszabadító hormon, follikulus stimuláló hormon, kromolin, adiponektin, RNáz, grelin, zidovudin, didanozin, tetrahydro-kannabinol, atropin, granulociták tenyésztését stimuláló tényező, lamotrigin, korion-gonadotropint felszabadító faktor, luteinizáló hormont felszabadító hormon, béta-galaktoszidáz és Argatroban közül kerül kiválasztásra.

11. A készítmény vagy a mikrorészecskék a 8. igénypont szerinti felhasználásra, ahol a száraz por leadja a biológiailag aktív szert a pulmonális rendszerben inhalálással való bevitelt követően.

12. A készítmény vagy a mikrorészecskék a 9. igénypont szerinti felhasználásra, ahol a biológiailag aktív szer felszívódik a szisztémás vérkeringésbe.

13. A készítmény vagy a mikrorészecskék a 9. igénypont szerinti felhasználásra, ahol a biológiailag aktív szer helyileg hat a tüdőben.

14. Eljárás az 5. vagy 6. igénypont szerinti szilárd gyógyszerforma előállítására, amelynél a formulálást kicsapással vagy liofilizálással végezzük.

A meghatalmazott:

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1255 Budapest, Pf.: 80.