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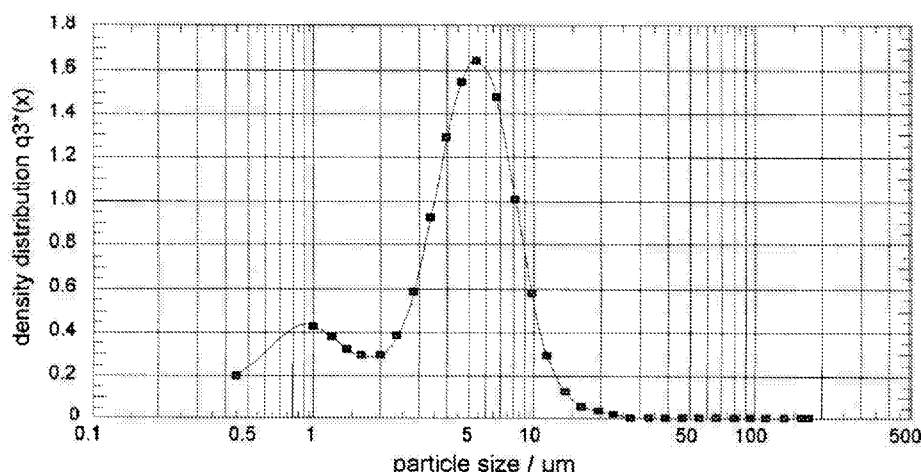
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## (54) Title: MICROPARTICLES COMPRISING A SULPHUR-CONTAINING COMPOUND

Figure 1



(57) Abstract: The present invention provides microparticles comprising a sulphur-containing compound, such as cysteamine, or a pharmaceutically acceptable salt, hydrate or ester thereof. Also provided is a composition comprising the microparticles and a stabilizing agent.

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## MICROPARTICLES COMPRISING A SULPHUR-CONTAINING COMPOUND

**Field of the Invention**

The present invention relates to microparticles comprising a sulphur-containing compound, such as cysteamine or cystamine, or a pharmaceutically acceptable salt, hydrate or ester thereof.

**5 Background to the Invention**

Cystic fibrosis (CF) is a multisystem disorder caused by mutations in the cystic fibrosis transmembrane conductance regulator (CFTR) gene, located on chromosome.

Lung disease remains the leading cause of morbidity and mortality in patients with CF [Davis PB, Drumm M, Konstan MW. Cystic fibrosis. Am J Respir Crit Care Med 1996; 154:1229; Goss CH, 10 Rosenfeld M. Update on cystic fibrosis epidemiology; Curr Opin Pulm Med 2004; 10:510; Brennan AL, Geddes DM. Cystic fibrosis. Curr Opin Infect Dis 2002; 15:175; Gibson RL, Burns JL, Ramsey BW. Pathophysiology and management of pulmonary infections in cystic fibrosis. Am J Respir Crit Care Med 2003; 168:918.

One of the major drivers of CF lung disease is infection (Sagel SD, Gibson RL, Emerson J, et al. Impact 15 of Pseudomonas and Staphylococcus infection on inflammation and clinical status in young children with cystic fibrosis. J Pediatr 2009; 154:183; Cystic Fibrosis Foundation Annual Patient Registry 2013. Available at: <http://www.cff.org/research/ClinicalResearch/PatientRegistryReport/> (Accessed on August 07, 2015).

The approach to treating infection in CF is multifaceted, involving antibiotics, chest physiotherapy, 20 inhaled medications to promote secretion clearance, and anti-inflammatory agents. Undoubtedly, improved use of antibiotics is responsible for a substantial portion of the increased survival that has occurred in patients with CF (Brennan AL, Geddes DM. Cystic fibrosis. Curr Opin Infect Dis 2002; 15:175; Sagel SD, Gibson RL, Emerson J, et al. Impact of Pseudomonas and Staphylococcus infection on inflammation and clinical status in young children with cystic fibrosis. J Pediatr 2009; 154:183).

25 There remains a need for better therapies for treating and preventing lung diseases/conditions in particular those associated with mucous-rich environments such as the CF lung. In addition there remains a need to limit the amount or doses of antibiotics used with the introduction of novel, replacement therapies or adjunct treatments that can improve the effectiveness of currently available treatments in the treatment or prevention of bacterial infections, in particular in the CF 30 lung.

Surprisingly, we have shown that microparticles provide a useful mode of delivery for cysteamine to patients with lung disease.

**Statements of the Invention**

35 According to a first aspect of the present invention there is provided a microparticle or microparticles comprising a sulphur-containing compound, or a pharmaceutically acceptable salt, hydrate or ester thereof.

As used herein "sulphur-containing compound" is intended to cover cysteamine, cystamine or a derivative thereof. The sulphur-containing compound may be an aminothiols. Examples of aminothiols include cysteamine and derivatives thereof. The term "derivative thereof" may encompass 2-methylthio ethylamine (cinnamate), 2-methyl thio ethylurea, N-(2-methylthio ethyl) p-acetamido benzamide, 2-aminoethanethiol, N-(2-methylthio ethyl)p-acetamido benzenesulfonamide, N-(2-propylthioethyl)-p-methoxy benzamide, N-(butylthio ethyl) nicotinamide, N-(2-dodecylthio ethyl) p-butoxybenzamide, N-(2-methylthio ethyl) p-toluenesulfonamide, N-(2-isopropylthio ethyl) propionamide, N-(2-octylthio ethyl) acetamide, N-(2-butylthio ethyl) methanesulfonamide, N-(2-isopentylthioethyl)butane, bis 1,4-(2-acetamido ethylthio), 2,3-butanediol, 2-hexadecylthio ethylamine hydrochloride, 2-allylthio ethylamine malate, 9-octadecene 2-ylthio ethylamine hydrochloride, 2-dodecylthio ethylamine hydrochloride, 2-isopentylthio ethylamine mandelate, 2-octadecylthio ethylamine salicylate, 2-β-hydroxyethyl thio ethylurea, 2-β-hydroxyethylthio ethylamine hydrochloride, 2-(2,3-dihydroxy propylthio)ethylamine p-toluenesulfonate, 2-(2-hydroxypropylthio)ethylamineoxalate, N-(2-methylthio ethyl)phenylacetamide, 2-(2,2-dimethoxy ethylthio) ethylamine hydrochloride, 2-(2,2-dimethoxy ethylthio) ethylamineundecylenate, 2-(2,2-diethoxy ethylthio) ethylamine undecylenate, 2-(2,2-diethoxy ethylthio)ethylamine acetate, 2-undecenylthio ethylamine, 2-β-ureidoethylthio ethylamine hydrochloride, 2-β-acetamidoethylthio ethylamine tropate, 2,2'-thio diethylamine fumarate, 2,2'-thio diethylurea, 3-β-aminoethylthio propylamine hydrochloride, S-β-ureidoethyl thiocarbamate, 2-ethoxycarbonylthio ethylamine hydrochloride, 2-dimethylamino carbonylthio ethylamine sulfate, 2-butoxycarbonyl methylthio ethylurea, 2-ethyloxycarbonylmethylthio ethylamine hydrochloride, 6-β-aminoethylthio hexanoate of methyl hydrochloride, 5-β-aminoethylthio pentanoic acid, 2-phenylthio ethylamine dihydrogen phosphate, 2-p-t-butylphenylthio ethylamine trichloroacetate, 2-p-methoxyphenylthio ethylamine ditartrate, 2-tolylthio ethylamine hydrobromide, 2-(1-biphenyl thio) ethylamine hydrochloride, 2-N-pentachlorophenylthio ethyl acetamide, 2-benzylthio ethylamine malate, 2-benzylthio ethylamine nicotinate, 2-benzylthio 2-methyl propylamine hydrochloride, 2-benzylthio propylamine lactate, N-(2-benzylthio ethyl)nicotinamide hydrochloride, N-(2-benzylthio ethyl) 10-undecene amide, N-(2-benzylthio ethyl) hexadecanamide, S-β-aminoethyl mercaptobutyric acid, N-(2-benzylthio ethyl)formamide, N-(2-benzylthio ethyl)phenylacetamide, N-[2-(2,6-dimethyl phenyl)ethyl] hexanamide, 2-o-aminophenylthio ethylamine succinate, N-(2-benzylthio ethyl) glutamine, S-β-aminoethyl mercapto acetic acid (3-S-β-aminoethyl) mercapto propionic acid, (3-S-γ-amino propyl) mercapto acetic acid, S(2-p-methoxybenzamido ethyl) mercapto 2-(2-naphthyl methylthio) ethylamine hydrochloride, 2-(2-naphthyl methylthio) ethylamine disuccinate, (2-thenyl) 2-thio ethylamine hydrobromide, 2-N-acetyl (2-thenylthio- ethylamine, 2-o-chlorobenzylthio ethylamine hydrochloride, 2-p-chlorobenzylthio ethylamine glycolate, 2-o-fluorobenzylthio ethylamine hydrochloride, 2-furfurylthio ethylamine hydrochloride, 2-tetrahydrofurfurylthio ethylamine p-amino-benzoate, 2-β-phenylethylthio ethylamine glutamate, 2-diphenylmethylthio ethylamine hydrochloride, 2-triphenyl methylthio ethylamine hydrochloride hemihydrate, 2-(2-pyridyl ethylthio)ethylamine hydrochloride, 2-(2-p-toluene sulfonamido ethylthio) pyridine N-oxide, 2-β-aminoethylthiomethyl pyridine N-oxide dihydrochloride, 2-β-aminoethylthio pyridine N-oxide hydrochloride, 2,4-dichloro 2-benzylthio ethylamine aspartate, N-[2-(3,4-dichloro benzylthio)ethyl] butyramide, N-[2-(2,6-dichloro benzylthio)ethyl] dodecanamide, N-[2-(3,5-dichloro benzylthio)ethyl] trifluoroacetamide

hydrochloride, 2-p-ethoxybenzylthio ethylamine hydrochloride, N-[2-m-fluorobenzylthio ethyl] chloroacetamide, 2-p-bromobenzylthio ethylamine succinate, 2-(3,4-dimethoxy benzylthio)ethylamine malate, 2-(3,4-methylenedioxy benzylthio)ethylamine hydrochloride, 2-(2,4-dichloro cetylthio)ethylamine, 2 (3,4,5-trimethoxy benzylthio)ethylamine hydrocinnamate, 2-p-methoxy benzylthio ethylamine salicylate, 2-o-methylbenzylthio ethylamine phenyl-acetate, N-[2-p-dimethylaminobenzylthio ethyl] methane-sulfonamide, 2-p-phenoxybenzylthio ethylamine hydrochloride, 2-.beta.-aminoethylthio pyridine hydrochloride, 2-benzylthio ethylamine citrate, N-[2-benzylthio ethyl] 2,4-dihydroxy 3,3-dimethyl butyramide, N-(2-benzylthio ethyl) 6,8-dihydroxy 7,7-dimethyl 5-oxo 4-aza octanamide, N-[2-(2-pyridyl thio)ethyl] propionamide, 2-(2-pyridyl methylthio)ethylamine dihydrochloride, 2-benzylthio ethylamine pantothenate, S-(.beta.-acetamidoethyl)mercaptoacetate of beta.-morpholinoethyl, S-(.beta.-phenylacetamidoethyl)mercaptoacetate N'-methyl 2-piperazino ethyl, S-(.beta.-ureidoethyl)mercaptoacetate of beta.-pyrrolidino-ethyl, S-(.beta.-trifluoroacetamidoethyl)-.beta.mercapto-propionate of .beta.-dimethylaminoethyl, 2-p-nitrobenzylthio ethylamine crotonate, 2-.beta.-morpholinocarbonyl ethylthio ethylamine hydrochloride, N,N-di(hydroxyethyl)S-(.beta.-benzamido-ethyl) mercapto-acetamido, N[2-N'-methyl piperazino carbonylthio ethyl] acetamide, 2-(1-naphthyl thio)ethylamine hydrochloride, N-(3-.beta.-ureidoethylthio propyl) succinamic acid, 3-allylthio propylamine, 3-(2,2'-dimethoxy ethylthio)propylamine, 3-(2,2'-dimethoxy ethylthio)propylamine sulfate, S-.beta.-aminoethylmercapto acetic acid, the hydrochloride of S-.beta.-aminoethyl mercapto acetic acid, N-(2-benzylthioethyl)acetamide, N-(2-benzylthioethyl)propionamide, N-(2-benzylthioethyl)butyramide, N-(2-benzylthioethyl)methanesulfonamide, N-(2-benzylthioethyl)ethanesulfonamide, N-(2-benzylthioethyl-propanesulfonamide, N-(2-benzylthioethyl)butanesulfonamide, S-(2-p-acetamidobenzenesulfonamido ethyl) mercapto acetic acid, S-(2-p-acetamidobenzamido ethyl) mercapto acetic acid, N-(2-thenylthioethyl)acetamide, 2-benzylthio propylamine, 2-benzylthio 2-methyl propylamine, 2-(2-p-toluenesulfonamido ethylthio) pyridine N-oxide, S-(2-p-butoxybenzamidoethyl)mercapto acetic acid, 2-t-butylthio ethylamine hydrochloride, 2-methoxycarbonyl methylthio ethylamine hydrochloride, 2-ethoxycarbonylmethylthio ethylamine hydrochloride, 2-propoxycarbonylmethyl thio ethylamine hydrochloride, 2-butoxycarbonylmethylthio ethylamine hydrochloride, 2,2'-thio diethylamine dihydrochloride, 3-(2-aminoethylthio)alanine hydrochloride, 2-benzylthio ethylammonium diacid phosphate, 2-methylthio ethylamine, N-(methylthioethyl) p-acetamidobenzamide, N-(2-methylthioethyl)nicotinamide, N-(2-methylthioethyl)benzamide, N-(2-methylthioethyl) p-butoxybenzamide, N-(2-methylthioethyl) butyramide, N-(2-methylthioethyl) propionamide, N-(2-methylthioethyl) acetamide, N-(2-methylthioethyl) butanesulfonamide, N-(2-octylthioethyl) methanesulfonamide, 2-cetylthio ethylamine hydrochloride, 2-(2-hydroxyethylthio) ethylamine hydrochloride, 2-methylthio ethylamine phenylacetatesnd 2-methylthio ethylamine undecylenate.

Alternatively, the sulphur-containing compound may be an organic disulphide, such as cystamine.

The sulphur-containing compound of the invention may be administered in the form of pharmaceutically acceptable salts. The pharmaceutically acceptable salts of the present invention can be synthesized from the parent compound which contains a basic or acidic moiety by conventional chemical methods. Generally, such salts can be prepared by reacting the free acid or base forms of these compounds with a stoichiometric amount of the appropriate base or acid in water or in an organic solvent, or in a mixture of the two; generally, nonaqueous media like ether, ethyl acetate, ethanol, isopropanol, or acetonitrile are preferred. Lists of suitable salts are found in *Remington's Pharmaceutical Sciences*, 17th ed., Mack Publishing Company, Easton, Pa., US, 1985, p. 1418, the disclosure of which is hereby incorporated by reference; see also Stahl et al, Eds, *"Handbook of Pharmaceutical Salts Properties Selection and Use"*, Verlag Helvetica Chimica Acta and Wiley-VCH, 2002. The phrase "pharmaceutically acceptable" is employed herein to refer to those compounds, materials, compositions, and/or dosage forms which are, within the scope of sound medical judgment, suitable for use in contact with the tissues of human beings or, as the case may be, an animal without excessive toxicity, irritation, allergic response, or other problem or complication, commensurate with a reasonable benefit/risk ratio.

The invention thus includes pharmaceutically-acceptable salts of the disclosed compounds wherein the parent compound is modified by making acid or base salts thereof for example the conventional non-toxic salts or the quaternary ammonium salts which are formed, e.g., from inorganic or organic acids or bases. Examples of such acid addition salts include acetate, adipate, alginate, aspartate, benzoate, benzenesulfonate, bisulfate, butyrate, citrate, camphorate, camphorsulfonate, cyclopentanepropionate, digluconate, dodecylsulfate, ethanesulfonate, fumarate, glucoheptanoate, glycerophosphate, hemisulfate, heptanoate, hexanoate, hydrochloride, hydrobromide, hydroiodide, 2-hydroxyethanesulfonate, lactate, maleate, methanesulfonate, 2-naphthalenesulfonate, nicotinate, oxalate, pamoate, pectinate, persulfate, 3-phenylpropionate, picrate, pivalate, propionate, succinate, tartrate, thiocyanate, tosylate, and undecanoate. Base salts include ammonium salts, alkali metal salts such as sodium and potassium salts, alkaline earth metal salts such as calcium and magnesium salts, salts with organic bases such as dicyclohexylamine salts, N-methyl-D-glucamine, and salts with amino acids such as arginine, lysine, and so forth. Also, the basic nitrogen-containing groups may be quaternized with such agents as lower alkyl halides, such as methyl, ethyl, propyl, and butyl chloride, bromides and iodides; dialkyl sulfates like dimethyl, diethyl, dibutyl; and diamyl sulfates, long chain halides such as decyl, lauryl, myristyl and stearyl chlorides, bromides and iodides, aralkyl halides like benzyl and phenethyl bromides and others.

In a preferred aspect of the invention, the microparticles have particle size of about 0.5 to 15 microns, for example 1 to 13 microns, including 4 to 8 microns. Particle size may be defined as "volume mean diameter" and as such the microparticles may have a volume mean diameter of about 0.5 to 15 microns, for example 1 to 13 microns, including 4 to 8 microns. The microparticles may have a volume mean diameter of 2 to 4 microns/micrometers (2-4 $\mu$ m).

Mean is a calculated value similar to the concept of average. The various mean calculations are defined in several standard documents (ISO 9276-2:2001: Representation of results of particle size analysis – Part 2: Calculation of average particle sizes/diameters and moments from particle size distributions; ASTM E 799-03 Standard Practice for Determining Data Criteria and Processing for Liquid Drop Size Analysis). There are multiple definitions for mean because the mean value is

associated with the basis of the distribution calculation (number, surface, volume), see (TN154, Particle Size Result Interpretation: Number vs. Volume Distributions, available at [www.horiba.com/us/particle](http://www.horiba.com/us/particle)) for an explanation of number, surface, and volume distributions. The equation for defining the volume mean is shown below. The best way to think about this calculation is to think of a histogram table showing the upper and lower limits of n size channels along with the percent within this channel. The  $D_i$  value for each channel is the geometric mean, the square root of upper x lower diameters. For the numerator take the geometric  $D_i$  to the fourth power multiplied by the percent in that channel, summed over all channels. For the denominator take the geometric  $D_i$  to the third power multiplied by the percent in that channel, summed over all channels.

$$D[4,3] = \frac{\sum_{i=1}^n D_i^4 p_i}{\sum_{i=1}^n D_i^3 p_i}$$

The volume mean diameter has several names including  $D_{4,3}$  or  $D_{50}/D_{90}$ .

As used herein, the terms "diameter" or "d" in reference to particles refers to the number average particle size, unless otherwise specified. An example of an equation that can be used to describe the number average particle size is shown below:

$$d = \frac{\sum_{i=1}^p n_i d_i}{\sum_{i=1}^p n_i}$$

where n=number of particles of a given diameter (d).

As used herein, the terms "geometric size", "geometric diameter", "volume average size", "volume average diameter" or " $d_g$ " refers to the volume weighted diameter average. An example of equations that can be used to describe the volume average diameter is shown below:

$$d_g = \left[ \frac{\sum_{i=1}^p n_i d_i^3}{\sum_{i=1}^p n_i} \right]^{1/3}$$

Where n=number of particles of a given diameter (d).

As used herein, the term “volume median” refers to the median diameter value of the “volume-weighted” distribution. The median is the diameter for which 50% of the total are smaller and 50% are larger and corresponds to a cumulative fraction of 50%.

Geometric particle size analysis can be performed on a Coulter counter, by light scattering, by light microscopy, scanning electron microscopy, or transmittance electron microscopy, as known in the art. It is a generally held belief that the ideal scenario for delivery to the lung is to have an aerodynamic diameter < 5 micrometers. See, e.g., Edwards et al., J Appl. Physiol. 85(2):379-85 (1998); Suarez & Hickey, Respir. Care. 45(6):652-66 (2000).

As used herein, the term “aerodynamic diameter” refers to the equivalent diameter of a sphere with density of 1g/mL were it to fall under gravity with the same velocity as the particle analysed. The aerodynamic diameter ( $d_a$ ) of a microparticle is related to the geometric diameter ( $d_g$ ) and the envelope density ( $\rho_e$ ) by the following:

$$d_a = d_g \sqrt{\rho_e}$$

Porosity affects envelope density which in turn affects aerodynamic diameter. Thus porosity can be used to affect both where the microparticles go in the lung and the rate at which the microparticles release the pharmaceutical agent in the lung. Gravitational settling (sedimentation), inertial impaction, Brownian diffusion, interception and electrostatic affect particle deposition in the lungs.

The microparticles may have an aerodynamic diameter of about 0.5 to 15 microns, for example 1 to 13 microns, including 4 to 8 microns. The microparticles may have an aerodynamic diameter of 2 to 4 microns/micrometers (2-4 $\mu$ m).

In a further aspect, the invention provides a composition comprising microparticles according to the first aspect of the invention and a stabilizing agent. In some instances the stabilizing agent is selected from the group consisting of monosaccharides, disaccharides, trisaccharides, oligosaccharides and their corresponding sugar alcohols, polysaccharides and chemically modified carbohydrates.

In a yet further aspect, the invention provides a composition comprising a sulphur-containing compound, or a pharmaceutically acceptable salt, hydrate or ester thereof, and a stabilizing agent as defined herein.

The stabilizing agent may be a sugar such as trehalose.

The stabilising agent may be a sugar alcohol selected from the group consisting of lactose, erytritol, ribitol, xylitol, galactitol, glucitol and mannitol. Preferably the stabilising agent is mannitol.

In a preferred composition of the invention, the composition comprises up to 20%w/w sulphur-containing compound, for example between 1 and 15% such as between about 5 and 10% w/w Sulphur-containing compound. Typically, the composition comprises about 5 or 10% Sulphur-containing compound.



As used herein, the term "about" is intended to vary the specified amount to allow for minor fluctuations of between + or – 10% of the specified amount.

In a preferred composition of the invention, the composition comprises up to 85% stabilising agent. The composition may comprise between 80 and 95% w/w stabilizing agent, for example between 85  
5 and 90% w/w stabilizing agent. Typically, the composition comprises about 90% stabilizing agent.

It has been shown that in compositions according to the invention which comprise trehalose or mannitol, cysteamine has increased formulation stability.

In a preferred embodiment of the invention the sulphur-containing compound is cysteamine or cystamine, preferably cysteamine. In a further embodiment of the invention, the sulphur-containing  
10 compound is cysteamine bitartrate.

In a preferred embodiment the composition is provided as an aqueous composition.

The composition of the invention may further comprise leucine. Leucine has surprisingly been shown to improve the stability of the formulation. In one embodiment of the invention, the composition comprises between 1 and 10% leucine, preferably about 5% leucine.

15 The composition may be in a solid dose form selected from the group consisting of microparticles, microspheres, and powders. Preferably the composition is provided as a dry powder. The powder may contain particles having a geometric diameter of about 3 to 8 microns, including 4 to 8 microns, such as 3 to 7 microns. In one embodiment, the powder contains particles having a geometric diameter of up to about 5 microns, for example 2 to 4 microns.

20 A further aspect of the invention provides microparticles according to the first aspect of the invention, or a composition according to the invention, for use in the treatment or prevention of lung disease.

A yet further aspect of the present invention relates to a method of treating or preventing lung disease comprising administering microparticles according to the first aspect of the invention, or a  
25 composition according to the invention, to a subject suffering, or having previously suffered from, from lung disease.

As used herein the term "lung disease" includes any disease or condition of the lung including cystic fibrosis, specifically lung infections associated with cystic fibrosis, and chronic obstructive pulmonary disease (COPD). COPD is the name for a collection of lung diseases including chronic  
30 bronchitis, bronchiectasis, emphysema and chronic obstructive airways disease. The term lung disease is also intended to include any respiratory disease which has a mucous or infectious element, for example a chronic cough, common cold, influenza, hantavirus, pneumonia and pleurisy.

A further aspect of the invention provides a therapeutic composition (or combination) which may be  
35 useful in the treatment of prevention of lung disease, which comprises microparticles according to the first aspect of the invention, or a composition according to the invention, and at least one additional pharmaceutical agent. The additional pharmaceutical agent may be selected from the

group consisting of antimicrobial agents such as antiviral, antifungal or antibacterial agents e.g. antibiotics, mucolytic agents, vasodilators such as bronchodilators, antihypertensive agents, cardiovascular drugs and calcium channel blockers. Preferably the additional pharmaceutical agent is an antibiotic.

- 5 The term “antibiotic” is used to refer to antibacterial agents that may be derived from bacterial sources. Antibiotic agents may be bactericidal and/or bacteriostatic.

The antibiotic agent may contain a  $\beta$ -lactam ring. The  $\beta$ -lactam ring is part of the core structure of several antibiotic families, the principal ones being the penicillins, cephalosporins, carbapenems, and monobactams. These antibiotic agent are called  $\beta$ -lactam antibiotics.

- 10 Generally the antibiotic agent is of the group consisting of aminoglycosides, ansamycins, carbacephem,  $\beta$ -lactams carbapenems, cephalosporins, (including first, second, third, fourth and fifth generation cephalosporins), penicillin, monobactams), glycylicyclines, lincosamides, lipopeptides, macrolides, nitrofurans, oxazolidinones, quinolones, sulfonamides, polypeptides and tetracyclins.

- 15 The antibiotic agent may be of the group consisting of aminoglycosides, ansamycins, carbacephem, carbapenems, cephalosporins (including first, second, third, fourth and fifth generation cephalosporins), lincosamides, macrolides, monobactams, nitrofurans, quinolones, penicillin, sulfonamides, polypeptides and tetracyclins. Alternatively or additionally the antibiotic agent may be effective against mycobacteria.

- 20 The antibiotic agent may be an aminoglycoside such as Amikacin, Gentamicin, Kanamycin, Neomycin, Netilmicin, Tobramycin or Paromomycin.

The antibiotic agent may be an such as Geldanamycin and Herbimycin

Alternatively the antibiotic agent may be a carbacephem such as Loracarbef.

- 25 The antibiotic agent is a carbapenem such as Ertapenem, Doripenem, Imipenem/Cilastatin or Meropenem.

- Alternatively the antibiotic agent may be a cephalosporins (first generation) such as Cefadroxil, Cefazolin, Cefalexin, Cefalotin or Cefalothin, or alternatively a Cephalosporins (second generation) such as Cefaclor, Cefamandole, Cefoxitin, Cefprozil or Cefuroxime. Alternatively the antibiotic agent may be a Cephalosporins (third generation) such as Cefixime, Cefdinir, Cefditoren, Cefoperazone, 30 Cefotaxime, Cefpodoxime, Ceftibuten, Ceftizoxime and Ceftriaxone or a Cephalosporins (fourth generation) such as Cefepime and Ceftobiprole.

The antibiotic agent may be a lincosamides such as Clindamycin and Azithromycin, or a macrolide such as Azithromycin, Clarithromycin, Dirithromycin, Erythromycin, Roxithromycin, Troleandomycin, Telithromycin and Spectinomycin.

- 35 Alternatively the antibiotic agent may be a monobactams such as Aztreonam, or a nitrofuran such as Furazolidone or Nitrofurantoin.

The antibiotic agent may be a penicillin such as Amoxicillin, Ampicillin, Azlocillin, Carbenicillin, Cloxacillin, Dicloxacillin, Flucloxacillin, Mezlocillin, Nafcillin, Oxacillin, Penicillin G or V, Piperacillin, Temocillin and Ticarcillin.

The antibiotic agent may be an oxazolidinone such as linezolid or tedizolid.

- 5 The antibiotic agent may be a sulfonamide such as Mafenide, Sulfonamidochrysoidine, Sulfacetamide, Sulfadiazine, Silver sulfadiazine, Sulfamethizole, Sulfamethoxazole, Sulfanilimide, Sulfasalazine, Sulfisoxazole, Trimethoprim, and Trimethoprim-Sulfamethoxazole (Co-trimoxazole) (TMP-SMX).

- 10 The antibiotic agent may be a quinolone such as Ciprofloxacin, Enoxacin, Gatifloxacin, Levofloxacin, Lomefloxacin, Moxifloxacin, Nalidixic acid, Norfloxacin, Ofloxacin, Trovafloxacin, Grepafloxacin, Sparfloxacin and Temafloxacin.

The antibiotic agent may be a polypeptide. Examples of such polypeptides include Bacitracin, Colistin and Polymyxin B. In one embodiment, the antibiotic agent is not a polypeptide.

- 15 The antibiotic agent may be a lipopeptide. Examples of such lipopeptides include Daptomycin and Surfactin.

Alternatively, the antibiotic agent may be a tetracycline such as Demeclocycline, Doxycycline, Minocycline and Oxytetracycline

Alternatively the antibiotic agent may be a glycylicycline. Examples of such glycylicyclines include tigecycline.

- 20 Alternatively or additionally the antibiotic agent may be effective against mycobacteria. In particular the antibiotic agent may be Clofazimine, Lamprene, Dapsone, Capreomycin, Cycloserine, Ethambutol, Ethionamide, Isoniazid, Pyrazinamide, Rifampicin, Rifabutin, Rifapentine or Streptomycin.

- 25 In one embodiment, the antibiotic agent is a macrolide and/or an aminoglycoside and/or sulphonamides.

In one embodiment, the antibiotic is selected from tobramycin, azithromycin, telithromycin, ciprofloxacin, ceftazidime.

In one embodiment, the antibiotic agent is not ciprofloxacin. In another embodiment the antibiotic is not tobramycin.

- 30 The antibiotic agent may be active in the treatment or prophylaxis of infections caused by Enterobacteriaceae (e.g. E.coli or Klebsiella spp., such as K. pneumoniae) or non-Enterobacteriaceae bacteria such as Burkholderia spp.

Generally the antibiotic agent is active in the treatment or prophylaxis of infections caused by gram-negative or gram-positive bacteria, such as Pseudomonas spp.

In one embodiment of the invention, the antibiotic is not a  $\beta$ -lactam antibiotic.

The active agents of the invention may be provided as pharmaceutical compositions additionally containing one or more pharmaceutically acceptable diluents, excipients and/or carriers. For example, the additional pharmaceutical agent may be provided as a composition comprising the agent and a carrier such as lactose or mannitol.

In a preferred aspect of the invention, the microparticles, or the composition, according to the invention, and the additional pharmaceutical agent may be administered simultaneously, sequentially or separately. The microparticles, or composition, and additional pharmaceutical agent may be provided as a combination package. The combination package may further instructions for simultaneous, separate or sequential administration of each of the microparticles, or composition, and additional pharmaceutical agent. For sequential administration, the microparticles, or composition, and additional pharmaceutical agent can be administered in any order.

The at least one additional pharmaceutical agent may be provided in microparticles distinct from said microparticles of the first aspect of the invention. Alternatively, the at least one additional pharmaceutical agent may be provided in a form other than microparticles.

In one embodiment of the invention, the microparticles of the first aspect of the invention, or composition according to the invention, comprise the at least one additional pharmaceutical agent,

In a further embodiment of the invention, the at least one additional pharmaceutical agent is administered in microparticles distinct from the microparticles of the first aspect, or composition, of the invention.

In a yet further embodiment of the invention, the at least one additional pharmaceutical agent is administered in a form other than microparticles.

In one embodiment, the microparticles or composition of the invention comprising a sulphur-containing compound, such as cysteamine, and/or an additional pharmaceutical agent to be administered in addition to the sulphur-containing compound have a volume average diameter between 0.1 and 5 micrometers (e.g., between 1 and 5 micrometers, between 2 and 5 micrometers, etc.). In another embodiment, the microparticles or composition of the invention, and/or an additional pharmaceutical agent, have a volume average diameter of up to 10 micrometers, for targeting delivery to the large bronchi. Particle size (geometric diameter and aerodynamic diameter) is selected to provide an easily dispersed powder that upon aerosolization and inhalation readily deposits at a targeted site in the respiratory tract (e.g., upper airway, deep lung, etc.), preferably while avoiding or minimizing excessive deposition of the particles in the oropharynx or nasal regions. In one preferred embodiment, the porous microparticles have a volume average diameter of between 2 and 5 micrometers, for example between 2 and 4 micrometers.

Considerable attention has been devoted to the design of therapeutic aerosol inhalers to improve the efficiency of inhalation therapies. Timsina et. al., Int. J Pharm., 101: 1-13 (1995); and Tansey, I. P., Spray Technol. Market, 4: 26-29 (1994). Attention has also been given to the design of dry powder aerosol surface texture, regarding particularly the need to avoid particle aggregation, a phenomenon which considerably diminishes the efficiency of inhalation therapies. French, D. L.,

Edwards, D. A. and Niven, R. W., *J. Aerosol Sci.*, 27: 769-783 (1996). Dry powder formulations ("DPFs") with large particle size have improved flowability characteristics, such as less aggregation (Visser, J., *Powder Technology* 58: 1-10 (1989)), easier aerosolization, and potentially less phagocytosis. Rudt, S. and R. H. Muller, *J. Controlled Release*, 22: 263-272 (1992); Tabata, Y. and Y. Bcada, *J. Biomed. Mater. Res.*, 22: 837-858 (1988). Dry powder aerosols for inhalation therapy are generally produced with mean geometric diameters primarily in the range of less than 5 micrometers. Ganderton, D., *J. Biopharmaceutical Sciences*, 3: 101-105 (1992); and Gonda, I. "Physico-Chemical Principles in Aerosol Delivery," in *Topics in Pharmaceutical Sciences* 1991, Crommelin, D. J. and K. K. Midha, Eds., Medpharm Scientific Publishers, Stuttgart, pp. 95-115, 1992. Large "carrier" particles (containing no drug) have been co-delivered with therapeutic aerosols to aid in achieving efficient aerosolization among other possible benefits. French, D. L., Edwards, D. A. and Niven, R. W., *J. Aerosol Sci.*, 27: 769- 783 (1996).

Drugs currently administered by inhalation come primarily as liquid aerosol formulations. However, many drugs and excipients, especially proteins, peptides (Liu, R., et al., *Biotechnol. Bioeng.*, 37: 177-184 (1991)), and biodegradable carriers such as poly(lactide-co-glycolides) (PLGA), are unstable in aqueous environments for extended periods of time. This can make storage as a liquid formulation problematic. In addition, protein denaturation can occur during aerosolization with liquid formulations. Considering these and other limitations, dry powder formulations (DPF's) are gaining increased interest as aerosol formulations for pulmonary delivery. Darnms, B. and W. Bains, *Nature Biotechnology* (1996); Kobayashi, S., et al, *Pharm. Res.*, 13(1): 80-83 (1996); and Timsina, M., et al., *Int. J. Pharm.*, 101: 1-13 (1994). However, among the disadvantages of DPF's is that powders of ultrafine particulates usually have poor flowability and aerosolization properties, leading to relatively low respirable fractions of aerosol, which are the fractions of inhaled aerosol that escape deposition in the mouth and throat. Gonda, I., in *Topics in Pharmaceutical Sciences* 1991, D. Crommelin and K. Midha, Editors, Stuttgart: Medpharm Scientific Publishers, 95-117 (1992). A primary concern with many aerosols is particulate aggregation caused by particle-particle interactions, such as hydrophobic, electrostatic, and capillary interactions. The present invention aims to address these issues.

Thus, in a further aspect, the invention provides an inhalation device comprising microparticles of the first aspect, or composition, of the invention. The device may be selected from a dry powder inhalation device and a metered dose inhaler.

In a further aspect of the invention the composition is obtained by preparing an aqueous solution of microparticles, or sulfhydryl (SH) compound, and stabilising agent and evaporating the water from the solution. Preferably the evaporating step is by spray drying.

Thus, a further aspect of the invention provides a process for preparing a composition according to the invention comprising preparing an aqueous solution of microparticles, or sulfhydryl (SH) compound, and stabilising agent and evaporating water from the aqueous solution. Preferably the evaporating step is by spray drying.

The microparticles according to the invention may be in the form of a dry powder. The microparticles may release an effective amount of a sulfhydryl (SH) compound, over a duration of at least two hours from inhalation of said microparticles by a human subject. In a preferred

embodiment, substantially all of the Sulphur-containing compound is released by 24 hours from inhalation of said microparticles by a human subject.

Microparticles are convenient to administer, thereby enhancing the extent of patient compliance. The microparticles, or composition, of the invention may be administered in a single puff.

- 5 Alternatively, the microparticles are formulated to provide sustained release of cysteamine. The microparticles may facilitate local delivery of cysteamine to the lungs or systemic delivery via the lungs.

- 10 The microparticles, or compositions, of the invention may also be administered intranasally or by inhalation and may be delivered in the form of a dry powder inhaler or an aerosol spray presentation from a pressurised container, pump, spray, atomiser, nebuliser, with or without the use of a suitable propellant. Preferably the microparticles, or compositions, of the invention are administered to the respiratory tract.

As used herein, the terms "comprise," "comprising," "include," and "including" are intended to be open, non-limiting terms, unless the contrary is expressly indicated.

- 15 The invention will now be described by way of example only with reference to the following figures:

Figure 1 is a graph showing Particle size distribution in batch 57#08a;

Figure 2 is a graph showing Particle size distribution in batch 57#08b;

Figure 3 is a graph showing Particle size distribution in batch 57#07 (placebo);

Figure 4. Lynovex/lactose study demonstrating reduction in *Pseudomonas* lung burden;

- 20 Figure 5. Lynovex (cysteamine) and Tobramycin combination resulting in reduced lung burden;

Figure 6. Mouse weight does not reduce in the presence of a combination of Lynovex and Tobramycin.

## **EXAMPLES**

EXAMPLE 1:

- 25 Spray drying as a potential formulation technique for the delivery of cysteamine bitartrate by oral inhalation

### **1 Materials**

Cysteamine Bitartrate: Manufactured by Recordati, batch number 140514-1 was supplied by Nova Biotics.

- 30 Oleic Acid: Fluka, 75096-1L, lot number BCBN9185V

Water: Deionized, Millipore, RiOs 5 system, serial number F8HN7 8491K

L-Leucine: Sigma, L-8000, lot number 91k0906

Trehalose: Sigma, T9449-1006, Lot number 011M7000N

## 2 Methods

### 2.1 Initial spray drying studies using solutions of cysteamine bitartrate formulated with oleic acid and trehalose

Several batches of cysteamine bitartrate were produced by spray drying solutions containing the active ingredient alone and with added trehalose and oleic acid (added as a potential taste masking agent).

Cysteamine bitartrate was allowed to warm to room temperature for 30 minutes before opening. For each batch to be spray dried, 100 mg cysteamine bitartrate powder was added to 10 ml deionised water, to give a total solids concentration of 1% w/v. This was stirred until fully dissolved.

Additional excipients (oleic acid and trehalose) were added to the cysteamine bitartrate solution to assess their impact on the powder properties after spray drying. The solutions were spray dried using a Buchi B290 spray dryer, fitted with a high-efficiency cyclone and a Buchi two-fluid nozzle. Full spray drying conditions are given in Table 1 below

Aspirator	100%
Liquid Feed Rate	2 ml/minute
Atomisation Pressure	5.5 bar
Inlet temperature	See Table 1
Outlet temperature	See Table 1

**Table 1 Spray drying conditions**

**Results from these initial studies confirmed that the presence of oleic acid in the formulation led to poor powder properties and low recoveries.**

A summary of the batches spray dried is described below in Table 2

Batch	Component A	Component B	Component C	Solvent	Result	Spray Dryer Temp
052#053	Cysteamine Bitartrate 95%	Oleic acid 5%	N/A	EtOH : Water, 2 :1	Waxy, glassy solid deposited on the walls of the cyclone	Inlet: 155 °C Outlet: 83 °C
052#055	Cysteamine Bitartrate 95%	Oleic acid 5%	N/A	EtOH : Water, 2 :1	Waxy, glassy solid deposited on the walls of the cyclone	Inlet: 78 °C Outlet: 48 °C
052#056	Cysteamine Bitartrate 70%	Oleic acid 5%	Trehalose 25%	EtOH : Water, 2 :1	Waxy, glassy solid deposited on the walls of the cyclone	Inlet: 75 °C Outlet: 46 °C
052#057	Cysteamine Bitartrate 32.6 %	Oleic acid 1.7%	Trehalose 65.7%	EtOH : Water, 2 :1	Waxy, glassy solid deposited on the walls of the cyclone	Inlet: 63 °C Outlet: 40 °C
052#058	Cysteamine Bitartrate 95%	Oleic acid 5%	N/A	Ethyl Acetate : Water, 5 :1	Waxy, glassy solid deposited on the walls of the cyclone	Inlet: 50 °C Outlet: 36 °C
052#059	Cysteamine Bitartrate 95%	Oleic acid 5%	N/A	Water : Ethyl acetate (added to crystals of API are formed)	Waxy, glassy solid deposited on the walls of the cyclone	Inlet: 50 °C Outlet: 38 °C

**Table 2 Production of initial feasibility batches containing oleic acid**

## **2.2 Initial spray drying studies using solutions of cysteamine bitartrate formulated with trehalose (no oleic acid)**

Based on the spray drying results obtained in 3.1 (above) it was decided to remove oleic acid from the formulation.

Cysteamine bitartrate was allowed to warm to room temperature for 30 minutes before opening. For each batch to be spray dried, 100 mg cysteamine bitartrate powder was added to 10 ml deionised water, to give a total solids concentration of 1% w/v. This was stirred until fully dissolved.



Trehalose was added to the cysteamine bitartrate solution to assess its impact on the properties of the spray dried powder. The solutions were spray dried using a Buchi B290 spray dryer, fitted with a high-efficiency cyclone and a Buchi two-fluid nozzle. Full spray drying conditions are given in Table 3 below

Aspirator	100%
Liquid Feed Rate	2 ml/minute
Atomisation Pressure	5.5 bar
Inlet temperature	See Table 4
Outlet temperature	See Table 4

5 **Table 3 Spray drying conditions**

A summary of the batches spray dried is described below in Table 4 below.

Batch	Component A	Component B	Component C	Solvent	Result	Spray Dryer Temp
052#060*	Cysteamine Bitartrate 10%	Trehalose 90%	N/A	Water	White powder.	Inlet: 81 °C Outlet: 42 °C
052#062	Cysteamine Bitartrate 50%	Trehalose 50%	N/A	Water	Waxy, glassy solid deposited on the walls of the cyclone	Inlet: 82 °C Outlet: 44 °C
052#063	Cysteamine Bitartrate 25%	Trehalose 75%	N/A	Water	Dry white powder.	Inlet: 114 °C Outlet: 61 °C
052#064	Cysteamine Bitartrate 25%	Trehalose 75%	N/A	Water	Dry white powder.	Inlet: 136 °C Outlet: 71 °C

052#65	Cysteamine Bitartrate 25%	Trehalose 75%	N/A	Water	Dry white powder.	Inlet: 162 °C Outlet: 79 °C
052#66	Cysteamine Bitartrate 35%	Trehalose 65%	N/A	Water	Wet looking powder. Not free-flowing.	Inlet: 148 °C Outlet: 70 °C
52#67	Cysteamine Bitartrate 30%	Trehalose 70%	N/A	Water	Damp looking powder. Forms aggregates.	Inlet: 147 °C Outlet: 72 °C
052#097*	Cysteamine Bitartrate 25%	Trehalose 75%	N/A	Water	Dry white powder.	Inlet: 121 °C Outlet: 71 °C Spray pressure 5.5 Bar

**Table 4 Spray drying conditions for trehalose formulations**

\*Used to generate additional data

**2.3 Spray drying cysteamine bitartrate formulated with Trehalose and L-Leucine (spray dried batch number 052#155, 052#140, 052#121 with Leucine 052#122 without Leucine)**

- 5 In order to further improve the properties of the spray dried powder, L-leucine was added to the formulation.

Cysteamine bitartrate was allowed to warm to room temperature for 30 minutes before opening. 100 mg Cysteamine Bitartrate powder, 50 mg of L-Leucine and 850 mg of Trehalose were added to 10 ml deionised water, to give a total solids concentration of 10% w/v. This was stirred until fully dissolved. Batches 052#140 and 052#155 were scaled to produce a 2 g batch size.

The solution was spray dried using a Buchi B290 spray dryer, fitted with a high-efficiency cyclone and a Buchi two-fluid nozzle. Full spray drying conditions are given in Table 5 below

Aspirator	100%
Liquid Feed Rate	2 ml/minute
Atomisation Pressure	5.5 bar
Inlet temperature	184°C
Outlet temperature	78°C

**Table 5 Spray drying conditions**

Following spray drying, the moisture content of the product was reduced further by secondary vacuum drying, at ambient temperature, overnight. The final product was then stored in a sealed glass vial prior to capsule filling. The solutions spray dried are summarised in Table 6 below.

Solution Number	Weight of Cysteamine Bitartrate	Weight of Trehalose	Weight of L-Leucine	Volume of deionised water	Spray dried powder reference
1	100 mg	850 mg	50 mg	10 ml	052#121
2	100 mg	900 mg	0 mg	10 ml	052#122
3	200mg	1700 mg	100 mg	20 ml	052#140
4	200 mg	1700 mg	100 mg	20 ml	052#155*

5 \* Collected as two batches of approximately 1 g

**Table 6 Spray drying of Cysteamine Bitartrate formulations containing L-leucine**

#### **2.4 Particle size analysis**

10 Particle size analysis was performed using a SympaTec HELOS particle size analyser with a RODOS disperser. Approximately 50mg of formulation was fed into the hopper. Dispersal was achieved using compressed air at a pressure of 2 bar. All instrument settings are detailed on the particle size analysis reports in appendix 1 (data not shown).

#### **2.5 Aerodynamic particle size analysis by Andersen Cascade Impactor**

15 The aerodynamic particle size of the spray dried powder was determined using a Copley Scientific 8 stage Andersen cascade impactor (ACI) fitted with a 60 l/minute pre-separator and stages -1 to 6. The method was as described in the US Pharmacopoeia 29 general chapter <601>, and the European Pharmacopoeia 5.1. 2.9.18 (procedure for dry powder inhalers).

The following parameters were used:

Dose: 2 x capsules

Capsules: Qualicaps HPMC standard size 3

Device: Plastiap, 3444, COQ, 23970000AA

Plate coating: None

Airflow: Approximately 60L/min (determined as a 4KPa pressure differential across the device).

Actuation time: Approximately 4seconds (determined by airflow to equate to a volume of 4 litres).

Plate washing: 0.1 M sodium phosphate buffer with EDTA, pH 8.

Detection: UV at 412 nm using Ellmans reagent to provide a suitable chromophore

10 Cysteamine bitartrate concentration in the washings was measured at 412 nm as described in section 3.6 below.

15 The mass of powder deposited at each stage was then calculated using the extinction coefficient determined in section 3.6. By analysing the amount of drug deposited on the various stages, it was then possible, using the dedicated Copley Scientific software, to calculate the Fine Particle Dose (FPD), the Fine Particle Fraction (FPF), the Mass Median Aerodynamic Distribution (MMAD) and Geometric Standard Deviation (GSD) of the peptide particles collected.

The Fine Particle Dose (FPD) was defined as the quantity of drug in the prescribed dose of an inhaled product that is generally considered to be of a size capable of penetrating the lung during inhalation i.e., respirable. This is usually considered to be about 5 microns or less.

20 The Fine Particle Fraction (FPF) was the FPD expressed as a percentage of the delivered dose.

## **2.6 Quantification of Cysteamine Bitartrate**

The quantification of Cysteamine Bitartrate was conducted using a Shimadzu UV-1650PC UV spectrometer. As Cysteamine Bitartrate as no UV chromophore Ellman's Reagent, 5,5-dithiobis(2-nitrobenzoic acid) was used.

### **25 2.6.1 Preparation of reagents**

Reaction Buffer: 0.1 M sodium phosphate, pH 8.0, containing 0.1 mM EDTA.

Ellman's Reagent Solution: Dissolve 40 mg Ellman's Reagent in 10 mL Reaction Buffer

Dissolve 34 mg of Cysteamine Bitartrate in 100 mL of Reaction Buffer to produce a 1.5 mM solution.

### **2.6.2 Preparation of Standard Curve**

30 Standards were prepared by dissolving Cysteamine Bitartrate in Reaction Buffer at the following concentrations:

Standard	Volume of Reaction Buffer mL	Amount of Cysteamine Bitartrate	Final Concentration
A	100	34 mg	1.5 mM
B	5	25 mL of Standard A	1.25 mM
C	10	20 mL of Standard A	1.0 mM
D	15	15 mL of Standard A	0.75 mM
E	20	10 mL of Standard A	0.5 mM
F	25	5 mL of Standard A	0.25 mM
G (Blank)	30	0 mL of Standard A	0.0 mM

**Table 2 Cysteamine Bitartrate standards**

A set of vials, each containing 50  $\mu$ L of Ellman's Reagent Solution and 2.5 mL of Reaction Buffer was prepared.

- 5 The assay solution or standard (250  $\mu$ L) was added to the vials prepared in the previous step. The reagents were mixed and analysed on the spectrophotometer immediately immediately. Absorbance was measured at 412 nm.

The values obtained from the standards were used to generate a standard curve. The experimental sample concentration of Cysteamine Bitartrate are determined from this curve.

## 10 3 Results

### **3.1 Initial studies on the spray drying cysteamine bitartrate with oleic acid and trehalose**

- Initial studies described in sections 3.1 confirmed that it was not possible to produce a suitable dry powder by spray drying solutions of cysteamine bitartrate containing oleic acid (with and without trehalose). Under all of the conditions used the resultant powder consisted of a glassy, solid material  
15 that stuck to the walls of the cyclone and collection jar.

Improved results were obtained when oleic acid was removed from the formulation (see section 3.2). Removal of oleic acid resulted in the production of a fine, white powder (rather than a waxy solid). However the powder was still cohesive and had relatively poor flow properties.

### **3.2 Spray drying of cysteamine bitartrate formulations containing trehalose and L-leucine**

- 20 Powder properties improved when L-leucine was added to the feed solution, resulting in fine white powders. Recoveries (yields) were high; in the range 50-83%. The spray dried powders had acceptable handling properties, and could be easily recovered from the collection vessel with

minimal static charge. Formulations containing L-Leucine had a higher % yield and improved flow characteristics over those without.

Yields obtained from spray dried solutions containing L-leucine are summarised in Table 7 below:

Sample Reference	Weight of powder recovered	% Yield**
052#122	0.5g	50
052#121	0.7 g	70
052#140*	1.6 g	80
052#155*	1.7 g	83

**Table 7 Spray drying yields from formulations containing L-leucine**

5 \*\* No residual moisture accounted for

\* 2 g batch size

### **3.3 Particle size analysis of spray dried cysteamine bitartrate formulations containing trehalose and L-leucine**

10 A summary of the particle size data for cysteamine bitartrate formulations containing trehalose and L-leucine are shown in Table 8.

Sample	X <sub>10</sub> * (µm)	X <sub>50</sub> ** (µm)	X <sub>90</sub> *** (µm)	VMD**** (µm)
052#122	0.88	2.28	4.61	2.56
052#121	1.46	2.65	4.59	2.89
052#140	0.93	2.75	6.39	3.34
052#155A	0.74	1.92	4.30	2.28
052#155B	1.06	2.84	6.19	3.42

**Table 8 Particle size analysis (summary)**

\* 10% of microparticles, by volume, below this figure

\*\* 50% of microparticles, by volume, below this figure

\*\*\* 90% of microparticles, by volume, below this figure

15 \*\*\*\* Volume mean diameter

### 3.4 Aerodynamic Particle Size Analysis by Anderson Cascade Impactor

A summary of the aerodynamic particle size data for spray dried batches of cysteamine bitartrate, formulated with trehalose and L-leucine is shown in Table 9. Full particle size analysis reports are detailed in appendix 2 (data not shown).

Batch	Capsule A fill wt (mg)	Capsule B fill wt (mg)	Gravimetric quantity of formulation released from device, Capsule A	Gravimetric quantity of formulation released from device, Capsule B	Mass of API recovered from the ACI	FPD (mg)	FPF (%)
<b>052#121</b> <b>Run 1</b>	132.8	130.7	120.7	121.7	11.3	N/A*	N/A*
<b>052#121</b> <b>Run 2</b>	114.8	119.6	105.0	109.5	18.5	6.9	37.7
<b>052#122</b>	84.0	75.2	54.8	54.9	11.2	3.0	27.0
<b>052#140</b> <b>Run 1</b>	105.9	114.1	105.0	109.5	23.1	4.5	19.6
<b>052#140</b> <b>Run 2</b>	96.9	100.9	90.2	93.4	25.9	5.6	21.5
<b>052#155</b> <b>Run 1</b>	82.7	84.9	76.6	41.7	14.0	6.06	43.2
<b>052#155</b> <b>Run 2</b>	96.2	94.3	89.5	87.8	15.3	3.6	23.7

5 **Table 9 Aerodynamic Particle Size**

\*Not included due to changes within the recovery process analytical method.

## 4 Conclusions

Cysteamine Bitartrate was successfully spray dried with trehalose and with, or without L-leucine. In these studies, a formulation containing cysteamine bitartrate (10%w/w), trehalose (85%w/w) and L-Leucine(5%w/w) were superior in terms of powder recoveries, handling properties and drug loading into the capsules.

The improved powder handling characteristics of the cysteamine bitartrate/trehalose/leucine formulations were translated into an increase in the Fine Particle Fraction (FPF), especially with formulations containing 5% Leucine.

- 5 Initial feasibility studies on DPI delivery confirm the spray dried powders can be delivered using commercially available DPI's without a lactose carrier. Providing an FPF between 20 % and 40 % and a FPM between 3 and 6.9 mg delivered from two capsules.

## EXAMPLE 2:

### Production of spray dried cysteamine bitartrate formulations for in vivo testing

## 10 5 Materials

Cysteamine bitartrate was supplied by NovaBiotics (Recordati 140514-1). All other reagents were analytical grade, supplied by Sigma.

## 6 Methods

### 6.1 *Spray drying of cysteamine bitartrate formulations*

#### 15 6.1.1 *Cysteamine bitartrate 5%(w/w), L-leucine 5%(w/w), mannitol 90%(w/w) (Batch 57#08a)*

The cysteamine bitartrate powder was warmed to room temperature for 30 minutes before opening. A solution containing 0.1g cysteamine bitartrate powder, 0.1g of L-Leucine and 1.8g of mannitol was prepared in 20 ml deionised water, to give a total solids concentration of 10% w/v. This was stirred until fully dissolved.

- 20 The solution was spray dried using a Buchi B290 spray dryer, fitted with a high-efficiency cyclone and a Buchi two-fluid nozzle. Full spray drying conditions are given in Table 1 below

Aspirator	100%
Liquid Feed Rate	2 ml/minute
Atomisation Pressure	5.5 bar
Inlet temperature	104°C
Outlet temperature	58°C

**Table 3 Spray drying conditions Batch 57#08a**

Following spray drying, the powder was collected and stored in a glass vial using laboratory film and foil overwrapped within a protective environment with a % RH < 10 %



6.1.2 Cysteamine bitartrate 10%(w/w), L-leucine 5%(w/w), mannitol 85%(w/w) (Batch 57#08b)

The cysteamine bitartrate powder was warmed to room temperature for 30 minutes before opening. A solution containing 0.2g cysteamine bitartrate powder, 0.1g of L-Leucine and 1.7g of mannitol was prepared in 20ml deionised water, to give a total solids concentration of 10% w/v. This was stirred until fully dissolved.

The solution was spray dried using a Buchi B290 spray dryer, fitted with a high-efficiency cyclone and a Buchi two-fluid nozzle. Full spray drying conditions are given in Table 2 below

Aspirator	100%
Liquid Feed Rate	2 ml/minute
Atomisation Pressure	5.5 bar
Inlet temperature	106°C
Outlet temperature	55°C

**Table 2 Spray drying conditions Batch 57#08b**

Following spray drying, the powder was collected and stored in a glass vial using laboratory film and foil overwrapped within a protective environment with a % RH < 10 %

6.1.3 Placebo batch containing L-leucine 5%(w/w), mannitol 95%(w/w) (Batch 57#07)

A solution containing 0.1g of L-Leucine and 1.9g of mannitol was prepared in 20ml deionised water, to give a total solids concentration of 10% w/v. This was stirred until fully dissolved.

The solution was spray dried using a Buchi B290 spray dryer, fitted with a high-efficiency cyclone and a Buchi two-fluid nozzle. Full spray drying conditions are given in Table 3 below

Aspirator	100%
Liquid Feed Rate	2 ml/minute
Atomisation Pressure	5.5 bar
Inlet temperature	100°C
Outlet temperature	64°C

**Table 3 Spray drying conditions Batch 57#007**

Following spray drying, the powder was collected and stored in a glass vial using laboratory film and foil overwrapped within a protective environment with a % RH < 10 %

## 6.2 Particle size analysis

Particle size analysis was performed using a SympaTec HELOS particle size analyser with a RODOS disperser. Approximately 50mg spray dried cysteamine bitartrate formulation was placed on the vibrating feeder and fed into the hopper. Dispersal was achieved using compressed air at a pressure of 2 bar.

## 6.3 Analysis of cysteamine bitartrate content in spray dried powders

The quantification of Cysteamine Bitartrate was conducted using a Shimadzu UV-1650PC UV spectrometer. As Cysteamine Bitartrate has no UV chromophore Ellman's Reagent, 5,5-dithiobis(2-nitrobenzoic acid) was used to measure the sulphhydryl group on the cysteamine.

### 6.3.1 Preparation of reagents

Reaction Buffer: 0.1 M sodium phosphate, pH 8.0, containing 0.1 mM EDTA.

Ellman's Reagent Solution: Dissolve 40 mg Ellman's Reagent in 10 mL Reaction Buffer

Dissolve 34 mg of Cysteamine Bitartrate in 100 mL of Reaction Buffer to produce a 1.5 mM solution.

### 6.3.2 Preparation of Standard Curve

Standards were prepared by dissolving Cysteamine Bitartrate in Reaction Buffer at the concentrations shown in Table 4:

Standard	Volume of Reaction Buffer mL	Amount of Cysteamine Bitartrate	Final Concentration
A	100	34 mg	1.5 mM
B	5	25 mL of Standard A	1.25 mM
C	10	20 mL of Standard A	1.0 mM
D	15	15 mL of Standard A	0.75 mM
E	20	10 mL of Standard A	0.5 mM
F	25	5 mL of Standard A	0.25 mM
G (Blank)	30	0 mL of Standard A	0.0 mM

**Table 4 Cysteamine bitartrate standards**

A set of vials, each containing 50 µL of Ellman's Reagent Solution and 2.5 mL of Reaction Buffer was prepared.

The assay solution or standard (250  $\mu$ L) was added to the vials prepared in the previous step. The reagents were mixed and analysed on the spectrophotometer immediately. Absorbance was measured at 412 nm.

The values obtained from the standards were used to generate a standard curve. The experimental sample concentration of cysteamine bitartrate are determined from this curve.

### 6.3.3 Analysis of cysteamine content in feed solutions and spray dried powders

The cysteamine bitartrate content was measured in each of the feed solutions used to produce the two spray dried batches. A 100 $\mu$ L aliquot of each solution was diluted into 10ml of DI water to produce a solution that fell within the linear region of the standard curve. The samples were analysed as described in section 3.3.2 and cysteamine bitartrate concentration determined.

The cysteamine bitartrate content was measured in the two spray dried formulations. A 50mg sample of each powder was diluted into 0.5ml DI water. A 100 $\mu$ L aliquot was diluted into 10ml of DI water to produce a solution that fell within the linear region of the standard curve. The samples were analysed as described in section 3.3.2 and cysteamine bitartrate concentration determined.

## 7 Results and Discussion

### 7.1 Spray drying of cysteamine bitartrate formulations

All feed solution was successfully spray dried, resulting in a fine white powder. Recoveries are summarised in Table 4 below:

Batch No	Amount spray dried (g)	Amount recovered (g)	Yield (%)
57#08a	2g	1.0	50
57#08b	2g	0.75	38
57#07 (placebo)	2g	1.1	55

**Table 5 Recovery of spray dried cysteamine formulations**

Recoveries for batches were lower than anticipated, however this is likely to be due to the small batch size (2g). All powders had good handling properties, however it was noticed that the 10% cysteamine formulation was slightly more cohesive than the 5% formulation.

### 7.2 Particle size analysis

A summary of the particle size data for all time points is shown in Table 5 and representative particle size distributions are shown in Figures 1-3.

**Table 6 Particle size analysis (summary)**

Batch	X <sub>10</sub> * (μm)	X <sub>50</sub> ** (μm)	X <sub>90</sub> *** (μm)	VMD**** (μm)
57#08a	0.85	4.51	8.58	4.73
	0.90	4.34	7.95	4.48
	0.90	4.41	8.27	4.75
57#08b	1.62	6.61	12.69	7.16
	1.83	6.89	13.28	7.49
	1.91	6.99	13.21	7.52
57#07(placebo)	1.29	4.2	7.99	4.66
	1.37	4.27	8.09	4.73
	2.68	4.49	7.38	6.72

\* 10% of microparticles, by volume, below this figure

\*\* 50% of microparticles, by volume, below this figure

\*\*\* 90% of microparticles, by volume, below this figure

\*\*\*\* Volume mean diameter

Examples of the size distributions obtained for each batch are shown in Figures 1-3.

### **7.3 Determination of cysteamine content in feed solution and in spray dried powders**

Both the spray dryer feed solution and the spray dried powders produced were analysed for cysteamine content. The results obtained are shown below in Table 6 below

Sample	Target concentration	Measured concentration
Batch 57#08a (feed solution)	5% (w/v)	5.9% (w/v)
Batch 57#08a (spray dried powder)	5% (w/w)	5.9% (w/w)
Batch 57#08b (feed solution)	10% (w/v)	11.5% (w/v)
Batch 57#08b (spray dried powder)	10% (w/w)	11.7% (w/w)

**Table 7 Cysteamine content in feed solutions and spray dried powders**

In all samples the measured concentration was higher than the expected concentration based on the theoretical content.

**EXAMPLE 3:**Assessment of efficacy of Lynovex (cysteamine) prep in a mouse IN neutropenic model of *Pseudomonas aeruginosa* ATCC 27853 (lung burden model)

## Chemicals

- 5 Animals were immunosuppressed/pre-conditioned with either 200 mg/kg or 150 mg/kg cyclophosphamide. Lynovex, chemical name cysteamine, and vehicle were either prepared either as Lynovex and lactose vehicle, or Lynovex and mannitol-based vehicle (both provided by Upperton (Upperton product)). These were prepared for treatment and vehicle-control alone respectively, and in combination. Tobramycin was prepared as an inhalation formulation in lactose. All
- 10 treatments were administered using a Penn Century device. Phosphate buffered saline (PBS) and *Pseudomonas* selective agar were required for bacterial tissue burden.

## Animals

- Male CD1 mice (n = 6 for treatment groups, plus five in pre-treatment group, totalling 35 mice) were used in this study. On day -4, the mice were immunosuppressed/pre-conditioned with 200 mg/kg
- 15 cyclophosphamide intraperitoneally; and with 150 mg/kg cyclophosphamide intraperitoneally on day -1. An infection was established with *P. aeruginosa* ATCC27853, with an inoculum of  $5 \times 10^6$  cfu/ml, administered intranasally in a volume of 40  $\mu$ l following anaesthetisation with a ketamine/xylazine anaesthetic cocktail for 15 minutes, for the Lynovex prepared in lactose study, and an inoculum of  $4 \times 10^6$  for the Upperton Lynovex product.

## 20 Treatment

All treatments were administered intratracheally using a Penn Century device.

- Lynovex (cysteamine) was administered at 1.5 mg alone, and in combination with lactose at the following concentrations: Lynovex 0.75 mg + 2.25 mg lactose powder, Lynovex 1.5 mg + 1.5 mg lactose powder, Lynovex 2.25 mg + 0.75 mg, along with a vehicle only control of 3 mg lactose. In
- 25 addition, Tobramycin at 188  $\mu$ g/dose was administered, as an inhaled formulation which was mixed with lactose to aid measurement. The treatments were administered approximately 5 minutes after infection.

- In a different study, Lynovex was administered at the following doses: 3 mg 5% Lynovex and 3 mg 10% Lynovex. Lynovex in combination with Tobramycin as follows: 3 mg 5% Lynovex + Tobramycin
- 30 0.188 mg in 1.5 mg vehicle, 3 mg 10% Lynovex + Tobramycin 0.188 mg in 1.5 mg Vehicle (mannitol-based, provided by Upperton) and a Tobramycin only control (0.188 mg/dose in lactose vehicle). The treatments were administered once approximately 10 minutes after infection.

## Bacterial burden in tissue

- The lung tissue burden of each animal, at the clinical end point of 24 h post-infection, was
- 35 determined. The lungs were homogenised in 2 ml PBS, serially diluted in PBS and plated onto *Pseudomonas* selective agar before quantification after 24-48 h at 37°C.

With the Lynovex/Lactose study, a variable infection was achieved in the lungs of the mice infected with *P. aeruginosa* ATCC27853. Intratracheal dosing with 0.188 mg of the inhalation formulation of Tobramycin resulted in a statistically significant reduction in lung burden when compared with vehicle-treated mice ( $P = 0.0097$  Kruskal Wallis test) and 5/6 animals cleared the infection to below the limit of detection. Intratracheal administration of 1.5 mg and 2.25 mg Lynovex also reduced the lung burden compared to vehicle ( $P = 0.0072$  and  $P = 0.0349$  respectively) with 5/6 and 4/6 mice respectively clearing the infection to below the detection limit (Figure 4).

In the Lynovex study with the Upperton product, a robust infection was achieved in the lungs of the mice infected with *P. aeruginosa* ATCC27853. Intratracheal dosing with 0.188 mg of the inhalation formulation of Tobramycin resulted in highly variable burdens with an average  $1.61 \log_{10}$  cfu/g reduction in lung burden when compared with vehicle-treated mice (Kruskal Wallis test). Intratracheal administration of 3 mg of 5% or 10% Lynovex as monotherapy did not reduce the lung burden compared to vehicle. However, combining 5% or 10% Lynovex with 0.188 mg Tobramycin resulted in a decrease in burden compared to vehicle mice ( $P < 0.0001$  and  $P < 0.0001$ , respectively). This reduction was compared to treatment with Tobramycin alone ( $P < 0.0001$  for 5% Lynovex+Tobramycin and  $P < 0.0015$  for 10% Lynovex+Tobramycin, Kruskal-Wallis test) (Figure 5).

Additionally, mouse weights were recorded before and after infection. Mice treated with vehicle, Lynovex monotherapy or Tobramycin monotherapy lost weight following infection. In contrast mice treated with the Lynovex + Tobramycin combinations maintained weight after infection indicating they remained relatively healthy post infection (Figure 6).

It should be noted that the Tobramycin for dry powder inhalation was suspended in lactose rather than mannitol. Suspension in lactose led to clumping of the powder resulting in some difficulties in delivery as many of the Penn Century devices blocked during dosing. The Lynovex suspensions were much easier to administer and all were delivered without issues due to the delivery device becoming blocked. Whilst the reductions in burden in the combination therapy arms are impressive and significantly superior to Tobramycin monotherapy, the data from some mice treated with Tobramycin monotherapy could be suspect due to the difficulty in delivery of the DPI. Even when animals with uncertain Tobramycin treatment are censored the greatly enhanced efficacy of the combination arms still remains.

**Claims:**

1. Microparticles comprising a sulphur-containing compound or a pharmaceutically acceptable salt, hydrate or ester thereof.
2. Microparticles as claimed in claim 1 wherein the compound is an aminothiols.
3. Microparticles as claimed in claim 2 wherein the aminothiols is cysteamine or a derivative thereof.
4. Microparticles as claimed in claim 1 wherein the compound is an organic disulphide.
5. Microparticles as claimed in claim 4 wherein as claimed in claim 7 wherein the organic disulphide is cystamine.
6. Microparticles as claimed in any one preceding claim wherein the particles have a particle size of about 0.5 to 15 microns.
7. Microparticles as claimed in claim 6 wherein the particles have a particle size of 1 to 13 microns.
8. Microparticles as claimed in claim 7 wherein the particles have a particle size of 4 to 8 microns.
9. Microparticles as claimed in claim 8 wherein the particles have a particle size of 2 to 4 microns.
10. A composition comprising microparticles as claimed in any one of claims 1 to 9 and a stabilizing agent.
11. The composition as claimed in claim 10 wherein the stabilizing agent is selected from the group consisting of monosaccharides, disaccharides, trisaccharides, oligosaccharides and corresponding sugar alcohols, polysaccharides and chemically modified carbohydrates.
12. The composition as claimed in any one preceding claim wherein the stabilizing agent is a sugar such as trehalose.
13. The composition as claimed in any one of claims 1 to 11 wherein the stabilizing agent is a sugar alcohol.
14. The composition as claimed in any preceding claim which comprises up to 20%w/w sulphur-containing compound.
15. The composition as claimed in claim 14 which comprises between about 5 and 10% w/w Sulphur-containing compound.
16. The composition as claimed in any preceding claim which comprises up to 85% stabilising agent.

17. The composition as claimed in any preceding claim which comprises between 80 and 95% w/w stabilizing agent.
18. The composition as claimed in any preceding claim which is an aqueous composition.
19. The composition as claimed in any preceding claim which further comprises leucine.
20. The composition as claimed in claim 19 which comprises  
between 1 and 10% leucine.
21. The composition as claimed in any preceding claim which is in a solid dose form selected from the group consisting of microparticles, microspheres, and powders.
22. The composition as claimed in claim 21 which is a dry powder.
23. Microparticles as claimed in any one of claims 1 to 9, or a composition as claimed in any one of claims 10 to 22, for use in the treatment or prevention of lung disease.
24. A method of treating or preventing lung disease comprising administering microparticles as claimed in any one of claims 1 to 9, or a composition as claimed in any one of claims 10 to 22, to a subject suffering, or having previously suffered from, from lung disease.
25. A therapeutic composition comprising microparticles microparticles as claimed in any one of claims 1 to 9, or a composition as claimed in any one of claims 10 to 22, and at least one additional pharmaceutical agent.
26. The therapeutic composition as claimed in claim 25 wherein the additional pharmaceutical agent is an antibacterial agent.
27. The therapeutic composition as claimed in claim 26 wherein the additional pharmaceutical agent is selected from the group consisting of antibiotics, mucolytic agents, vasodilators such as bronchodilators, antihypertensive agents, cardiovascular drugs and calcium channel blockers.
28. An inhalation device comprising microparticles microparticles as claimed in any one of claims 1 to 9, or a composition as claimed in any one of claims 10 to 22.



Figure 1

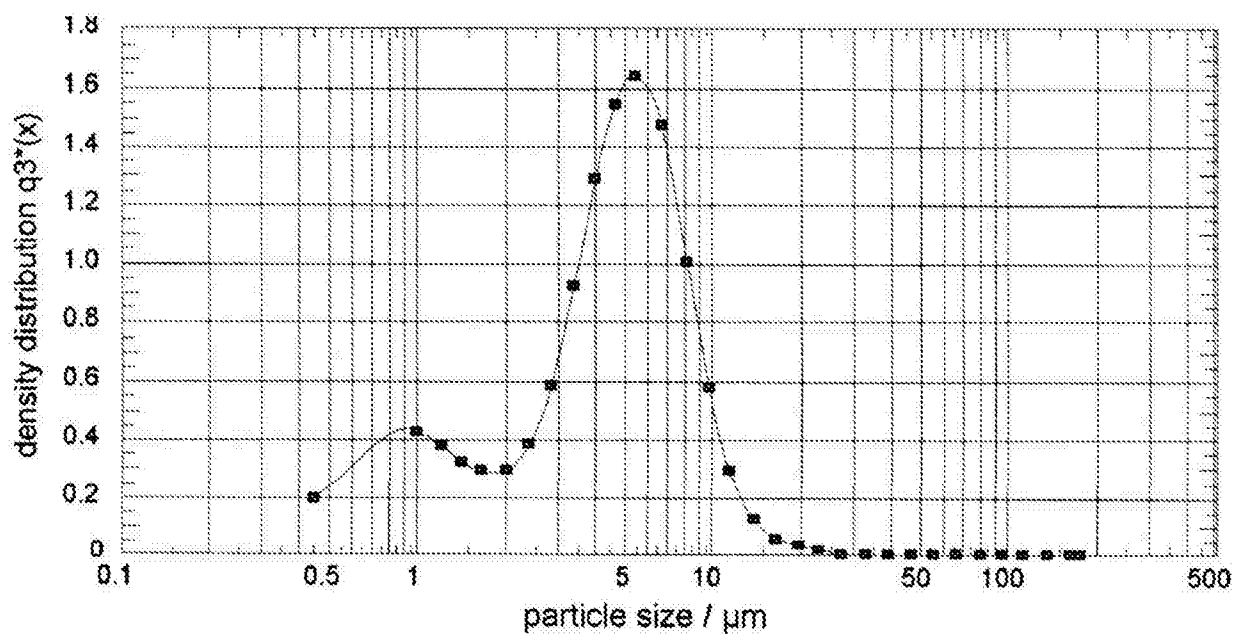


Figure 2

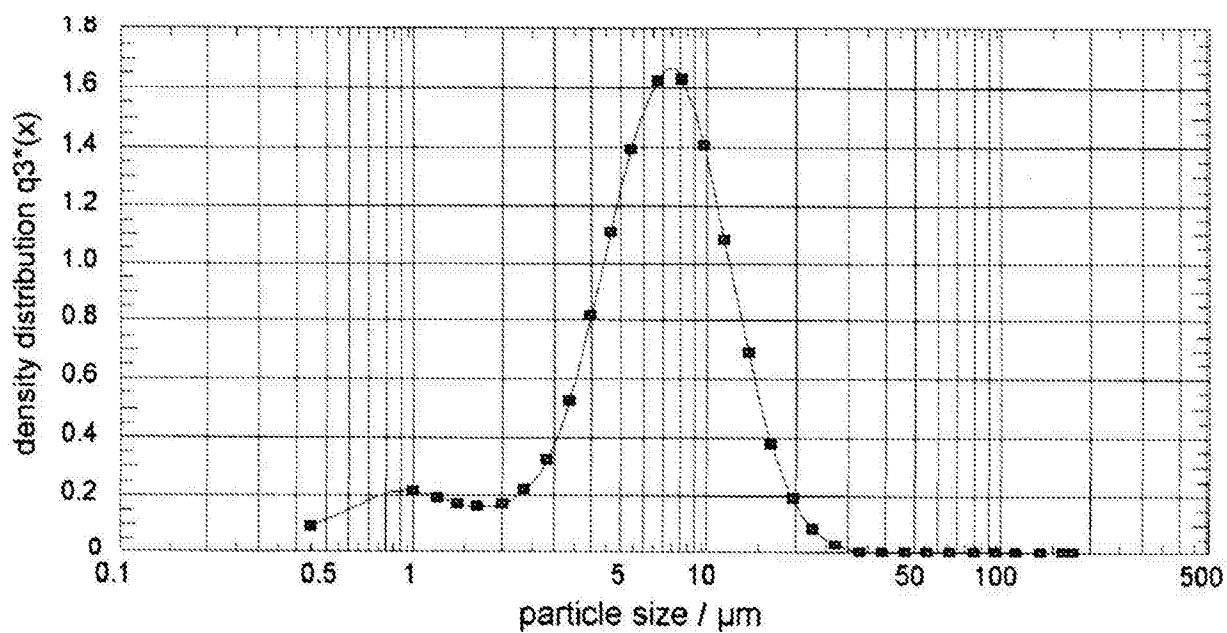


Figure 3

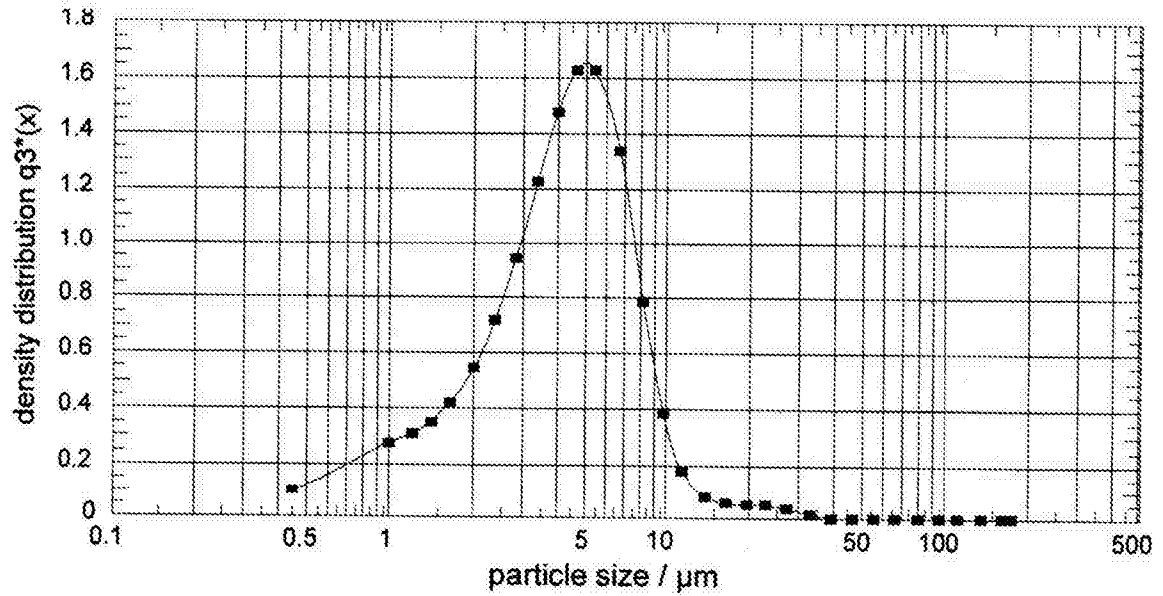


Figure 4

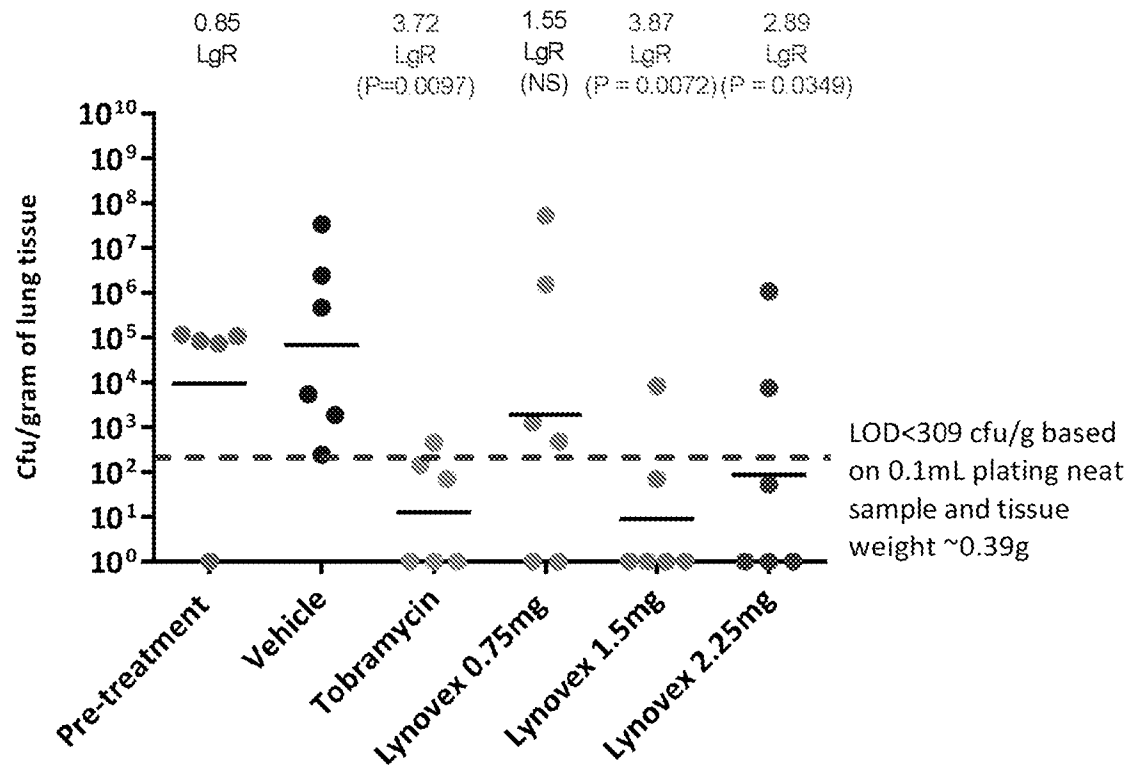


Figure 5

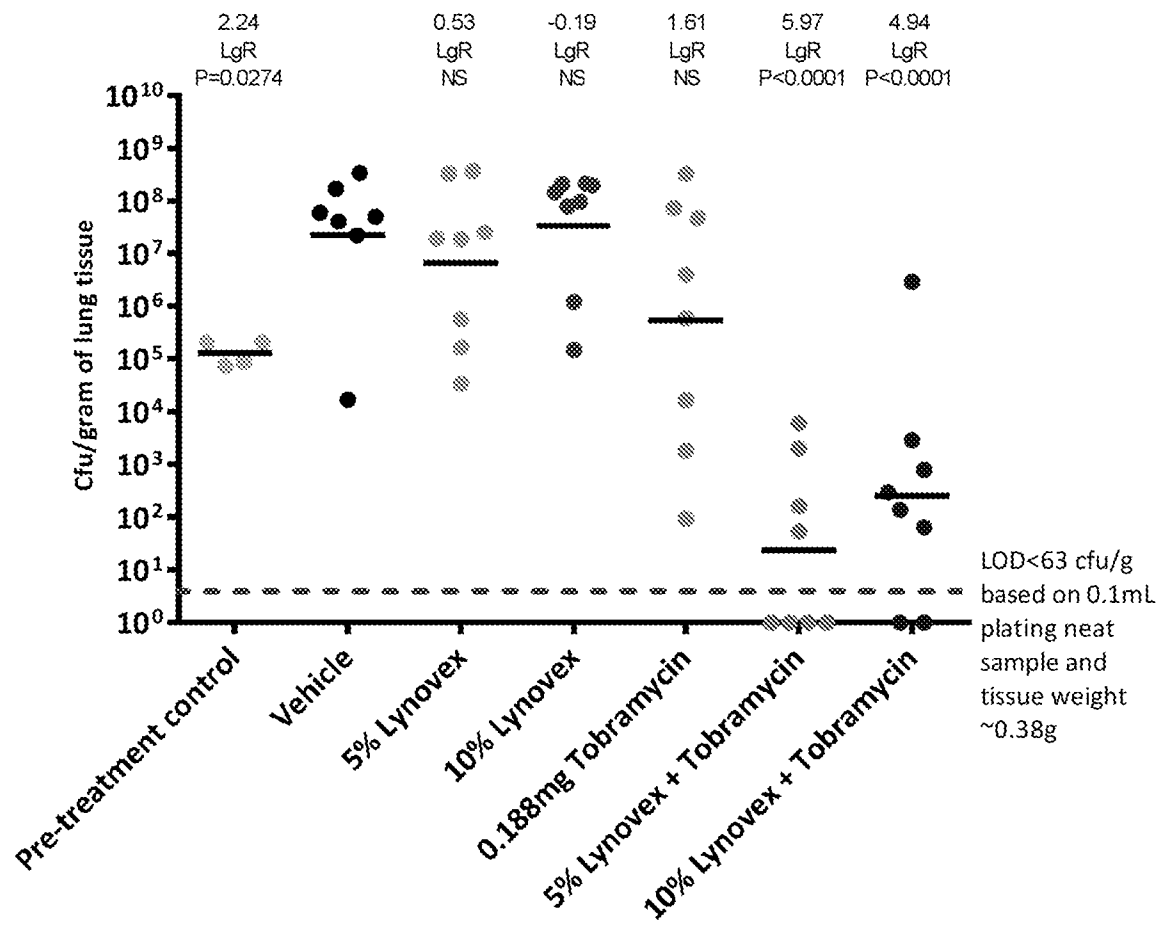
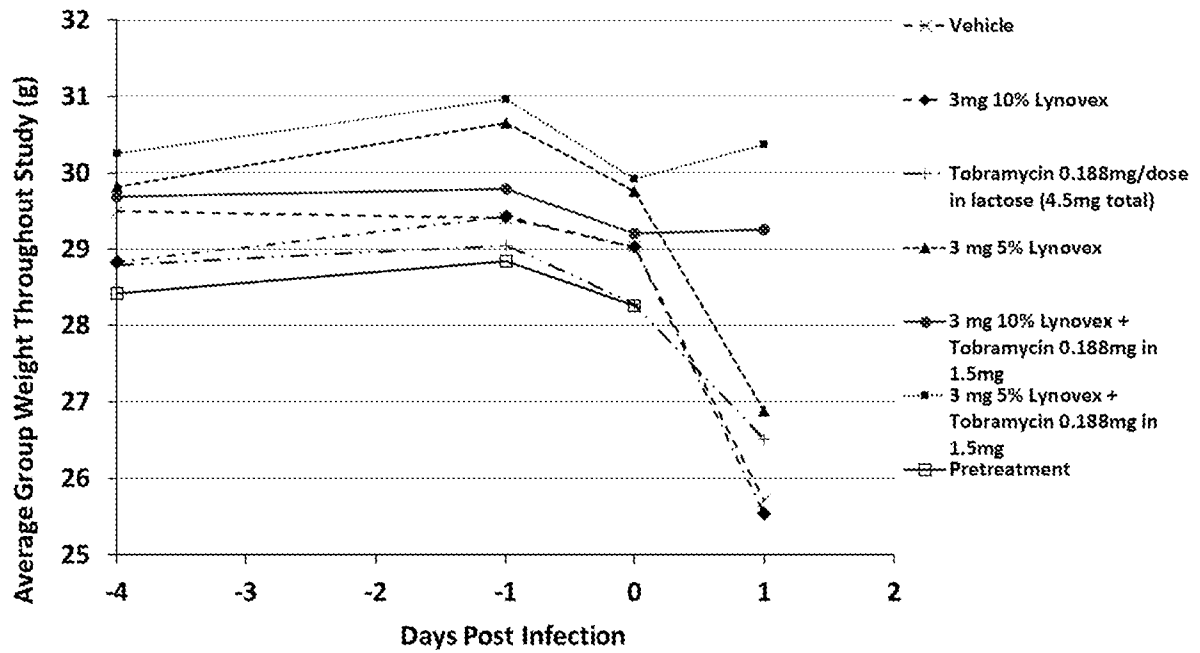


Figure 6



# INTERNATIONAL SEARCH REPORT

International application No  
PCT/GB2017/051637

<b>A. CLASSIFICATION OF SUBJECT MATTER</b> INV. A61K31/145 A61K9/00 A61K9/16 A61K45/06 A61P11/00 ADD.		
According to International Patent Classification (IPC) or to both national classification and IPC		
<b>B. FIELDS SEARCHED</b> Minimum documentation searched (classification system followed by classification symbols) A61K		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) EPO-Internal, CHEM ABS Data, BIOSIS, EMBASE, WPI Data		
<b>C. DOCUMENTS CONSIDERED TO BE RELEVANT</b>		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	B. E. Buchan: "FORMULATION STUDIES ON CYSTEAMINE FOR THE TREATMENT OF NEPHROPATHIC CYSTINOSIS", 1 August 2011 (2011-08-01), pages 1-223, XP55404192, Retrieved from the Internet: URL: <a href="https://core.ac.uk/download/pdf/1576950.pdf">https://core.ac.uk/download/pdf/1576950.pdf</a> [retrieved on 2017-09-06]	1-3, 6-11, 13-17, 19-25,28
Y	page 137, line 1 - page 138, paragraph 2 page 155, last paragraph - page 159, paragraph 2 tables 27,31 page 168, last paragraph page 173, last paragraph - page 176, last paragraph table 34 page 191, paragraph 2 - page 193, last -/--	4,5,26, 27
<div style="display: flex; justify-content: space-between;"> <span><input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C.</span> <span><input checked="" type="checkbox"/> See patent family annex.</span> </div>		
<div style="display: flex;"> <div style="flex: 1;"> <p>* Special categories of cited documents :</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier application or patent but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="flex: 1;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>"&amp;" document member of the same patent family</p> </div> </div>		
Date of the actual completion of the international search  <div style="text-align: center; font-size: 1.2em;">7 September 2017</div>		Date of mailing of the international search report  <div style="text-align: center; font-size: 1.2em;">15/09/2017</div>
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016		Authorized officer  <div style="text-align: center; font-size: 1.2em;">Hoff, Philippe</div>

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International application No

PCT/GB2017/051637

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Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>paragraph</p> <p>-----</p> <p>B.E. Buchan: "The formulation and evaluation of a dry powder for pulmonary delivery in cystinosis",</p> <p>1 January 2010 (2010-01-01), XP55403968, Retrieved from the Internet: URL: <a href="https://www.cystinosis.org.uk/wp-content/uploads/2010/09/2010_RGU_ResearchPoster.pdf">https://www.cystinosis.org.uk/wp-content/uploads/2010/09/2010_RGU_ResearchPoster.pdf</a> [retrieved on 2017-09-05] the whole document</p>	1-3,6-8, 10,16, 21,22, 25,28
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X	<p>-----</p> <p>WO 90/13538 A1 (DRACO AB [SE]) 15 November 1990 (1990-11-15)</p> <p>page 1, line 1 - line 14 page 2, line 21 - page 5, line 30 page 7, line 25 - line 29; claims; examples A,B</p>	1-3,6,7, 10-18, 21-24,28
X	<p>-----</p> <p>WO 2010/091124 A2 (MICROBION CORP [US]; BAKER BRETT HUGH JAMES [US]) 12 August 2010 (2010-08-12) abstract page 12, line 27 - page 13, line 27 page 15, line 21 - page 16, line 20 table 1 page 32, line 14 - line 27 page 35, line 6 - line 16 page 67, line 4 - line 18; claims 1,6,13,23-25</p> <p>-----</p> <p>-/--</p>	1-4, 6-18,21, 22,25-27

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International application No

PCT/GB2017/051637

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>WO 2013/120086 A1 (WHITEHEAD BIOMEDICAL INST [US]) 15 August 2013 (2013-08-15)</p> <p>abstract  page 68, paragraph 129 - paragraph 132  page 121, paragraph 245  page 123, paragraph 252 - page 124, line 1  page 127, paragraph 262; claims 1,9</p> <p>-----</p>	<p>1-7,10,  11,  13-18,  21-25,28</p>
Y	<p>CEDRIC CHARRIER ET AL: "Cysteamine (Lynovex?), a novel mucoactive antimicrobial &amp; antibiofilm agent for the treatment of cystic fibrosis", ORPHANET JOURNAL OF RARE DISEASES, BIOMED CENTRAL LTD, LO, vol. 9, no. 1, 30 November 2014 (2014-11-30), page 189, XP021206325, ISSN: 1750-1172, DOI: 10.1186/S13023-014-0189-2 the whole document</p> <p>-----</p>	<p>26,27</p>
Y	<p>WO 2016/046523 A1 (NOVABIOTICS LTD [GB]) 31 March 2016 (2016-03-31)  abstract  page 27, line 24 - page 28, line 31;  claims 1-3,10,11</p> <p>-----</p>	<p>4,5</p>

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Information on patent family members

International application No

PCT/GB2017/051637

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