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(54) **MICROPARTICULES SECHEES PAR PULVERISATION, SERVANT DE VEHICULES THERAPEUTIQUES EN THERAPIE GENETIQUE**

(54) **SPRAY-DRIED MICROPARTICLES AS THERAPEUTIC VEHICLES FOR USE IN GENE THERAPY**

(57) La présente invention concerne des microparticules lisses et sphériques, dont au moins 90 % présente un diamètre de particule moyen en volume de 1 à 10 .mu.m, constituées d'un mélange sensiblement homogène d'un agent de thérapie génétique et d'un excipient. Par exemple, un gène nu ou encapsulé peut ainsi être administré au moyen d'un inhalateur à poudre.

(57) Microparticles, which are smooth and spherical, and at least 90 % of which have a volume median particle size of 1 to 10 .mu.m, comprise a substantially uniform mixture of an agent for gene therapy and an excipient. For example, a naked or encapsulated gene can thus be administered, using a dry powder inhaler.

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(54) Title: SPRAY-DRIED MICROPARTICLES AS THERAPEUTIC VEHICLES FOR USE IN GENE THERAPY

(57) Abstract

Microparticles, which are smooth and spherical, and at least 90 % of which have a volume median particle size of 1 to 10 μm, comprise a substantially uniform mixture of an agent for gene therapy and an excipient. For example, a naked or encapsulated gene can thus be administered, using a dry powder inhaler.

SPRAY-DRIED MICROPARTICLES AS THERAPEUTIC VEHICLES FOR USE IN GENE THERAPY

Field of the Invention

This invention relates to spray-dried microparticles
and their use as therapeutic vehicles. In particular, the
invention relates to means for delivery of agents for gene
therapy.

Background of the Invention

Current methods of aerosolising drugs for inhalation
include nebulisation, metered dose inhalers and dry powder
inhaler systems. Nebulisation of aqueous solutions and
suspensions requires large volumes of drugs and involves
the use of bulky and non-portable devices.

The most common method of administration of
therapeutic agents to the lung is by the use of volatile
propellant-based devices, commonly termed metered dose
inhalers. A solution of propellant, commonly CFC 11, 12 or
114, is used, containing either dissolved drug or a
suspension of the drug in a pressurised canister. Dosing
is achieved by depressing an actuator which releases a
propellant aerosol of drug suspension or solution which is
carried on the airways. During its passage to the lung,
the propellant evaporates to yield microscopic precipitates
from solution or free particles from suspension. The
dosing is fairly reproducible and cheap, but there is
growing environmental pressure to reduce the use of CFCs.
CFCs will shortly be replaced by non-ozone-depleting
propellants, so-called hydrofluoroalkanes. Furthermore,
the use of CFC solvents remains largely incompatible with
many drugs, because of their susceptibility to denaturation
and low stability.

Concurrently, there is a move towards using dry powder
inhalers (DPIs) containing drugs, usually admixed with a
carrier, such as mannitol, lactose or glucose, which
facilitates the aerosolisation and dispersion of the drug
particles. The energy for disaggregation is often supplied

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by the breath or inspiration of air through the device by the patient.

Drugs are currently micronised, to reduce particle size. This approach is not applicable for biotechnology products. In general, biotechnology products are available in low quantity and, furthermore, are susceptible to the methods currently employed to dry and micronise drugs, prior to mixing with carrier. Further, it is particularly difficult to provide blends of drug and carrier which are sufficiently free-flowing that they flow and dose reproducibly in modern multiple dose inhalers such as the Turbohaler (Astra) and Diskhaler (Glaxo).

WO-A-9116882 describes the use of spray-drying to produce lipid/drug powders of conventional asthma drugs which purportedly showed a sustained release effect due to the presence of the lipid. The conditions given for spray-drying produce dry powders which are bimodally distributed, with peaks at 1 μm and 7-10 μm.

WO-A-8806441 describes the use of spray-drying in the production of stable liposome formulations. A preserving additive such as a sugar is spray-dried with the lipid material. The liposomes retain their size distribution and integrity, on reconstitution. The stabilised liposomes may contain a therapeutic agent, apparently for use after storage of the liposomes and the removal of the preserving additive.

1-5 μm is the optimal particle size for penetration to the alveolar regions. Adsorption in the alveolar region offers the largest surface area for access to the bloodstream, and therefore it is desirable that a dry powder formulation should contain a large preponderance of particles with these dimensions. It has proved difficult to produce particles in this range reproducibly by micronisation techniques. Whilst there have been preliminary reports on the use of spray-drying to produce respirable powders, the first description of the precise process conditions leading to particles for therapeutic

use, having satisfactory narrow interquartile ranges, in the 1-5 μm range, is in WO-A-9609814 (Andaris).

While microparticles in the 1-5 μm range are of optimal size for alveolar penetration when aerosolised as discrete particles, they do not always possess the necessary bulk powder flow properties which are required for DPI device filling and dose dispensing. Particles of this size are highly cohesive and will generally require coarser-sized particles to impart the bulk flow properties required; larger particles flow better. For drugs that have low potency, dispensing and blending with a conventional carrier, e.g. lactose, present few problems in terms of reproducibility and accuracy. For highly potent drugs, however, accurate blending and dispensing becomes a real problem. When the required drug dose may be as small as 1 μg, contained in an emitted mass of 12-25 mg, the problem of accurate dosing becomes a real impediment to the development of such highly potent drugs.

Summary of the Invention

Microparticles according to the present invention, of which at least 90% can have a size of 1-10 μm, are intended for use in gene therapy, and comprise a substantially uniform mixture of an agent for gene therapy and an excipient.

Microparticles of this invention may be formulated with a carrier ("carrier" is used herein for distinction from the "excipient" in the microparticles). The formulation may be administered, e.g. from a dry powder inhaler. On administration, the carrier disperses in the airstream; because of its large size, it is deposited in the mouth and throat. The smaller microparticles go to the peripheral airways; on reaching the lung, the excipient dissolves and its diluting effect is lost. The dispersed therapeutic agent is reproducibly dosed to the desired locus of administration.

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Description of the Invention

Microparticles of this invention may be obtained by spray-drying under conditions described in WO-A-9609814, the contents of which are incorporated herein by reference.

5   Formulations containing them overcome some of the fundamental problems associated with dry powders for DPIs, specifically, poor powder mixing and limited accuracy of blending and dosing for highly potent drugs.

Particles of the same size are also suitable for

10   intravenous administration, e.g. for targeted delivery to the liver, and for gene therapy. Soluble microparticles prepared by spray-drying may be cross-linked, by known technology, for this purpose. By comparison with, say, iv administration of adenovirus, targeted administration is

15   achieved by means of the invention.

More generally, a process for preparing microcapsules of the invention comprises atomising a solution (or dispersion) of excipient and a therapeutic agent that is itself, or is used together with, a wall-forming material.

20   In particular, it has been found that, under the conditions stated herein, and more generally described in WO-A-9218164, remarkably smooth spherical microparticles of various materials may be produced. The particle size and size distribution of the product can be reproducibly

25   controlled within a tight range. For example, by Coulter analysis, 98% of the particles can be smaller than 6 μm on a number basis, within an interquartile range of 2 μm, and with less than 0.5 μm mean size variation between batches. Furthermore, when tested in a dry powder inhaler,

30   reproducible dosing was achieved, and subsequent aerosolisation under normal but low flow conditions (30 l/min) resulted in excellent separation of microparticles from carrier.

The therapeutic agent that is used in this invention

35   is intended for gene therapy. It may be a naked or encapsulated gene. Alternatively, it may be a virus particle that is stabilised by the presence of the

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excipient. Agents suitable for use in gene therapy are
receptor-targeted complexes, adenovirus, adeno-associated
virus, retrovirus, parvovirus, vaccinia virus, herpes virus
and sindbis virus. Cationic lipid-DNA complexes may be
5   used.

Antisense oligonucleotides may also be administered by
means of this invention. For example, the oligonucleotide
may be chosen to degenerate the template activity of
template RNAs. It is demonstrated by Nyce and Metzger,
10   Nature 385:721-5 (1997), that aerosolised phosphorothioate
antisense oligonucleotide targeting the adenosine A_1
receptor desensitised rabbits to subsequent challenge with
adenosine or dust-mite allergen, and is therefore of
potential utility in the treatment of asthma.

15   DNA being anionic, microparticles of the invention may
comprise a cationic material. In particular, cationic
lipids capable of forming positively-charged liposomes have
the ability to bind DNA and can be used for gene delivery
to mammalian cells both *in vitro* and *in vivo*. By way of
20   example, DNA may be encapsulated by binding to lipid
domains in a lactose or other carrier. Ideally, the lipid
domains should be <1 μm and the microcapsules of the order
of 2-5 μm, for deposition in the peripheral airways. The
principle has been tested using control lipids and control
25   DNA and the methodology transferred to produce an active
plasmid DNA/lipid/lactose microcapsule using the lipids DC-
Chol and DOPE and marker plasmid.

Other suitable cationic materials are chitosan and
lysine. Lysine has a very polar side-chain which is
30   positively charged at neutral pH and highly hydrophilic.
These properties make it suitable for co-spray-drying with
HSA (human serum albumin), to form microcapsules.
Successful DNA binding has been achieved on microparticles
containing 5% w/w of 25% lysine:75% HSA.

35   Chitosan is a natural polysaccharide suitable for
biomedical applications. Lysozyme will biodegrade chitosan
in vivo. By cross-linking the polymer matrix using heat or

chemical methods, e.g. glycolaldehyde, chitosan becomes less susceptible to lysosomal degradation and thus suitable for prolonged delivery of drugs. The cationic property of the chitosan has proved to bind DNA.

The excipient will usually be chosen on account of its ability to dilute/stabilise the therapeutic agent, and because of its wall-forming properties. It should be capable of formulation for spray-drying. Suitable excipients include carbohydrates, e.g. sugars such as glucose, lactose and mannitol. α-Lactose and mannitol may have the particularly desirable property of resisting moisture uptake. See also EP-A-0372777 (incorporated herein by reference).

The amount of excipient in the mixture of the microparticles will usually be chosen having regard to the desired degree of, say, dilution or stabilisation. It may be at least 50% by weight of the mixture, and often at least 70% or 80%.

Another inert wall-forming, material may also be used, if desired, to form the microparticles. Suitable such materials are water-soluble.

The wall-forming material and process conditions should be so chosen that the product is sufficiently non-toxic and non-immunogenic in the conditions of use, which will clearly depend on the dose administered and duration of treatment. The wall-forming material may be a starch derivative, a synthetic polymer such as tert-butyloxy-carbonylmethyl polyglutamate (US-A-4888398) or a polysaccharide such as polydextrose. Generally, the wall-forming material can be selected from most hydrophilic, biodegradable physiologically compatible polymers, as described in more detail in WO-A-9218164.

Especially for pulmonary delivery, the microparticles are preferably soluble, and preferably comprise a carbohydrate excipient. A sugar may facilitate dissolution of the microparticles on the lung surface.

For parenteral administration in particular, the microparticles are preferably stabilised, e.g. cross-linked by chemical means or heat. In this case, a wall-forming material used for gene therapy may be proteinaceous. For example, it may be collagen, gelatin or (serum) albumin, in each case preferably of human origin (i.e. derived from humans or corresponding in structure to the human protein). Most preferably, it is HSA derived from blood donations or, ideally, derived recombinantly from the fermentation of microorganisms (including cell lines) which have been transformed or transfected to express recombinant serum albumin. Further detail is given in WO-A-9218164.

For use in a DPI, microparticles of the invention may be mixed with a conventional carrier such as mannitol, lactose or glucose (which may be the same as the excipient). The carrier will have a particle size, and be in an amount, chosen in accordance with approved usage of the device. By way of example, 3 μm microparticles of the invention may be mixed with lactose particles 80 μm in size.

There are various considerations, in determining an appropriate percentage of the agent to total solids in the spray-drying medium. These criteria include the dose of the agent from one "shot" of the device; the dispensed weight of the formulation, i.e. the amount of powder in one "shot"; and the microparticle:carrier ratio, i.e. the proportion of microcapsules to carrier.

The spray-drying process can be controlled in order to obtain microspheres with desired characteristics. Thus, the pressure at which the seed solution is supplied to the spray nozzle may be varied, for example from 1.0-10.0 x 10^5 Pa, preferably 2-8 x 10^5 Pa, and most preferably about 7.5 x 10^5 Pa. Other parameters may be varied as disclosed below.

More than 30%, preferably more than 40%, 50%, or 60%, of the microspheres have a diameter within a 2 μm range and at least 90%, preferably at least 95% or 99%, have a

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diameter within the range 1.0-8.0 μm. The interquartile range may be 2 μm, with a median diameter of 3.5, 4.0, 4.5, 5.0, 5.5, 6.0 or 6.5 μm.

5 Thus, at least 30%, 40%, 50% or 60% of the microspheres may have diameters within the range 1.5-3.5 μm, 2.0-4.0 μm, 3.0-5.0 μm, 4.0-6.0 μm, 5.0-7.0 μm or 6.0-8.0 μm. Preferably, a said percentage of the microspheres have diameters within a 1.0 μm range, such as 1.5-2.5 μm, 2.0-3.0 μm, 3.0-4.0 μm, 4.0-5.0 μm, 5.0-6.0 μm, 6.0-7.0 μm

10 or 7.0-8.0 μm.

A product of the invention may be in the form of hollow microspheres, in which at least 90%, preferably at least 95% or 99%, of the microspheres have a diameter in the range 1.0-8.0 μm.

15 For spray-drying, a solution, suspension, emulsion or dispersion preferably contains 0.1 to 50% w/v, more preferably about 5.0-25.0%, of the materials of microcapsules. About 20% is optimal.

The solution or dispersion (preferably solution),
20 referred to herein as the "seed solution", is atomised and spray-dried by any suitable technique which results in discrete microspheres or microcapsules of 1 to 10 μm diameter. These figures refer to at least 90% of the population of microcapsules, the diameter being measured
25 with a Coulter Multisizer II. The term "microcapsules" means hollow particles enclosing a space, which space is filled with a gas or vapour but not with any solid materials.

The atomising comprises forming an aerosol of the
30 preparation by, for example, forcing the preparation through at least one orifice under pressure into, or by using a centrifugal atomiser in a chamber of warm air or other inert gas. The chamber should be big enough for the largest ejected drops not to strike the walls before
35 drying. The gas or vapour in the chamber is clean (i.e. preferably sterile and pyrogen-free) and non-toxic when administered into the bloodstream in the amounts

concomitant with administration of the microcapsules in
use.   The rate of evaporation of the liquid from the
preparation should be sufficiently high to form hollow
microcapsules but not so high as to burst the
5   microcapsules. The rate of evaporation may be controlled
by varying the gas flow rate, concentration of drug and
excipient in the seed solution, nature of liquid carrier,
feed rate of the solution and, more importantly, the
temperature of the gas encountered by the aerosol. With an
10   albumin concentration of 15-25% in water, an inlet gas
temperature of at least about 100°C, preferably at least
110°C, is generally sufficient to ensure hollowness and the
temperature may be as high as 250°C without the capsules
bursting. About 180-240°C, preferably about 210-230°C and
15   most preferably about 220°C, is optimal, at least for
albumin. Since the temperature of the gas encountered by
the aerosol will depend also on the rate at which the
aerosol is delivered and on the liquid content of the
protein preparation, the outlet temperature may be
20   monitored to ensure an adequate temperature in the chamber.
An outlet temperature of 40-150°C has been found to be
suitable. Controlling the flow rate has been found to be
useful in controlling the other parameters such as the
number of intact hollow particles.
25   Due to the heat-sensitive nature of DNA, it would be
advantageous to spray-dry using a low outlet temperature.
An increase in size has been observed, on reducing the
temperature.   This size increase may be due to the
microcapsules having a higher moisture content which in
30   turn results in particle agglomeration. To avoid this
agglomeration, an outlet temperature of 70°C is favoured.
More particularly, microparticles of the invention
preferably have a maximum interquartile range of 3 μm, more
preferably 2 μm, and most preferably 1.5 μm, with respect
35   to their volume median particle size. The volume median
particle diameter is determined by Coulter counter. This
is achieved by spray-drying in which there is a combination

of low feed stock flow rate with high levels of atomisation and drying air. The effect is to produce microcapsules of very defined size and tight size distribution.

Several workers have designed equations to define the mean droplet size of pneumatic nozzles; a simple version of the various parameters which affect mean droplet size is as follows:

$$D = A/(V^2.d)^a + B.(M_{air}/M_{liq})^{-b}$$

Where   D = Mean droplet size

            A = Constant related to nozzle design

            B = Constant related to liquid viscosity

            V = Relative air velocity between liquid and
                nozzle

            d = Air density

M_{air} and M_{liq} = Mass of air and liquid flow

            a and b = Constants related to nozzle design

Clearly, for any given nozzle design, the droplet size is most affected by the relative velocity at the nozzle and concurrently the mass ratio of air to liquid. For most common drying use, the air to liquid ratio is in the range of 0.1-10 and at these ratios it appears that the average droplet size is 15-20 μm. For the production of microparticles in the size range described herein, the air to liquid ratio is preferably from 20-1000:1. The effect is to produce particles at the high ratios which are exceedingly small by comparative standards, with very narrow size distributions. For microparticles, produced at the lower ratios of air to liquid, slightly larger particles are produced, but they still nevertheless have tight size distributions which are superior to microparticles produced by emulsion techniques.

The invention allows for the nature of the dry microcapsules to be manipulated, in order to optimise the flow or vehicle properties, by changing and reducing the

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forces of cohesion and adhesion within the microcapsule preparation. For example, if so required, the microcapsules may be made predominantly positive or negative by the use of highly-charged monomeric or polymeric materials, e.g. lysine or poly-lysine and glutamate or poly-glutamate in systems without HSA or heterogeneous systems including HSA and active principles.

In the Examples, dry powder formulations of gene vectors suitable for pulmonary delivery using a dry powder inhaler device were prepared by spray-drying. The resulting microparticles comprised the gene vector (cationic lipids:DNA complexes or adenovirus) dispersed within the sugar excipient shell. The sugar excipient component of the microparticles enables their dissolution on the lung surface, thereby releasing the gene vector at the optimal site within the lung.

The gene vectors carried reporter genes encoding luciferase of β-galactosidase to investigate DNA activity through the spray drying process. Gene transfection assays were performed on the dry powder formulations of both types of vectors, using *in vitro* cell culture methods.

Example 1

Plasmid pCMVLuc (CMV promoter:luciferase gene) was reconstituted with the lipids DC-Chol/DOPE at a charge ratio of 5:1. DC-Chol is 3β-[N-(N',N'-dimethylaminoethane)-carbamoyl]cholesterol; DOPE is dioleoylphosphatidylethanol-amine.

A feedstock for spray-drying was prepared containing 1 mg plasmid DNA, 8 mg DC-Chol, 8 mg DOPE and 700 mg lactose as excipient, in 7 ml water. The feed solution was spray-dried using a laboratory bench top spray-dryer, under the following conditions:

Inlet temperature	130°C
Outlet temperature	70.1°C
Atomisation pressure	3.0 bar
Feed rate	0.75 g/min

200 mg of spray-dried product was obtained, equivalent to a recovery of 35%. Analysis of the finished product using a Coulter LS230 Laser Sizer demonstrated that the lactose:cationic lipid:DNA microparticles had a median particle diameter (volume) of 3.8 μm. Typically, the cationic lipid:DNA domains were less than 200 nm in size.

Restriction analysis of plasmid DNA extracted from the microparticles using standard procedures demonstrated that the restriction pattern of the spray-dried DNA was identical to that of the control plasmid.

Two such samples (1A and 1B) were prepared. Microcapsules were also prepared comprising lactose only (1C) and lactose:cationic lipid (1D).

A549 cells (derived from a human lung tumour) were grown in 60 mm diameter cell culture dishes. Cells were seeded at 5×10^5 the day before transfection. To mimic delivery to the lung, i.e. the use of intra-tracheal powder aerosolisation, the microparticles/microcapsules were sprayed as a dry powder onto the surface of the cells using a syringe device after aspiration of the tissue culture medium. It was estimated that less than 10 μg pCMVLuc DNA was added to each dish. After 30 min, 2 ml fresh cell culture medium was added. At 48 hr post-transfection, cells were lysed in 500 μl lysis buffer (Promega), and the luciferase activity was measured in 20 μl aliquots of cell lysate.

The positive controls were DC-Chol/DOPE (40 μg):pCMVLuc (10 μg) in Experiment 1 and Spermidine-Chol/DOPE (40 μg):pCMVLuc (10 μg) in Experiment 2. Gene transfection data, expressed in terms of luciferase activity (RLU/min), are given in Table 1. Two values are given, where luciferase activity was measured in two independent dishes.

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Table 1

SAMPLE	EXPERIMENT 1 (RLU/min)	EXPERIMENT 2 (RLU/min)
Control Cells	826 776	792
Lactose	830 736	
pCMVLuc	760 810	
+ve Control	8636412 9896528	4192473 3490039
1A	125984 180776	32084 241805
1B	2051592	7505 31604
1C	1392 1394	
1D	878	771

Example 2

A plasmid carrying the β-galactosidase gene (CMV promoter) as reporter was reconstituted with the lipid DOTAP, i.e. N-[1-(2,3-dioleoyloxy)propyl]-N,N,N-trimethylammonium methylsulphate, and spray-dried with mannitol as excipient. The feedstock contained 2 mg plasmid DNA, 10 mg DOPE and 388 mg mannitol, in 4 ml diluent. The feed solution was spray-dried using a bench top spray dryer under the following conditions:

Inlet temperature $130^{\circ}C$
Outlet temperature $78^{\circ}C$
Atomisation pressure   3.0 bar
Feed rate   0.72 g/min

175 mg of spray-dried product was obtained, equivalent to a recovery of 43.8%. Analyses of the spray-dried product on a Coulter LS230 Laser Sizer demonstrated that the microparticles had a median diameter (volume) of 3-6 μm.

Standard procedures were used to assess gene transfection efficiency. B16 mouse melanoma cells were

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seeded at 10^5 cells per well and incubated for 24 hr. Samples of cationic lipid:plasmid DNA:mannitol microparticles containing 2 µg plasmid DNA were dissolved in water and used to transfect the confluent cells. The

5   controls were 2 µg plasmid DNA only and DOTAP/DNA at 2 µg DNA. The cells were incubated in the transfection solution for 4 hr and then grown for a further 48 hr before harvesting. Protein extracts were prepared and these were analysed for β-galactosidase activity. Transfection

10   efficiencies in terms of β-galactosidase activity are given in Table 2.

<div align="center">Table 2</div>

Sample	β-galactosidase activity (mU/mg protein)	Mean	Standard Deviation
Plasmid DNA only	3.41 2.32 2.08	2.60	0.71
DOTAP/DNA	10000.18 9803.90 10142.92	9982.33	170.21
DOTAP/DNA/ mannitol microparticles	6530.99 5043.20 5063.85	5546.01	853.08

Example 3

A feedstock was prepared containing an adenoviral vector (luciferase gene as reporter molecule) at a titre of

25   5.3×10^{10} IU/ml (7.8×10^{10} PFU/ml) in 10 mM Tris, 1 mM $MgCl_2$, 150 mM, Tween 80 (54 mg/ml) and 1M sucrose, at pH 8.5. The feedstock (50 ml), at a concentration of 0.352 g/ml, was spray-dried using a laboratory bench top spray-dryer under the following conditions:

30   Inlet temperature $56^{\circ}C$
         Outlet temperature $39.9^{\circ}C$
         Atomisation pressure   3.0 bar
         Feed rate   0.75 g/min

15

The recovery of product after spray drying is summarised in Table 3.

Table 3

	Before Spray-Drying	Post-Spray-Drying	Recovery
Mass (g)	17.62	6.46	36.6%
Titre (IU/ml)	5.3×10^{10}	2.6×10^{10}	49.0%

The dry powder microparticle formulation of adenovirus was investigated for gene transfection activity in an *in vitro* cell culture assay using A549 cells. The microparticles were dissolved in water, and samples equivalent to 0.5 mg, 2 mg, 5 mg and 20 mg dry powder were used to transfect the cells. Adenovirus at a MOI (multiplicity of infection) of 100 PFU/cell was used as control. All samples were assayed in duplicate. The efficiency of gene transfection measured in luciferase activity is given in Table 4.

Table 4

Sample	Luciferase Activity (RLU/min)	
Adenovirus control	17473732	17949356
Adenovirus control	26138380	27845016
+20 mg dry powder	42104272	42878048
+2 mg dry powder	13697072	13401940
+0.5 mg dry powder	5214460	5816246
+5 mg dry powder	24963076	24989568
Control	760	574
Control	203	183

These data demonstrate that the adenovirus retains significant gene transfection activity.

<u>CLAIMS</u>:

1.   Microparticles, which are smooth and spherical, at
least 90% of which have a particle size of 1 to 10 μm, and
which comprise a substantially uniform mixture of an agent
5   for gene therapy and an excipient.

2.   Microparticles according to claim 1, wherein the
excipient constitutes a major proportion of the mixture.

3.   Microparticles according to claim 1 or claim 2,
obtainable by spray-drying a solution of the mixture.

10   4.   Microparticles according to any preceding claim,
wherein said particle size is 1 to 5 μm.

5.   Microparticles according to any preceding claim, which
have a maximum interquartile range of 3 μm.

6.   Microparticles according to claim 5, which have a
15   maximum interquartile range of 2 μm.

7.   Microparticles according to any preceding claim,
obtainable by an aseptic process.

8.   Microparticles according to any preceding claim,
wherein the excipient comprises a carbohydrate.

20   9.   Microparticles according to any preceding claim,
wherein said agent is a naked or encapsulated gene, or a
virus particle.

10.   Microparticles according to any preceding claim, which
additionally comprises a cationic material, and wherein
25   said agent comprises DNA.

11.   Microparticles according to claim 10, wherein the
cationic material comprises lipid domains.

12.   A free-flowing particulate composition comprising
microparticles according to any preceding claim and
30   relatively large particles of a carrier material which may
be the same as or different from the excipient.

13.   An inhaler device adapted to deliver a therapeutic
agent via the pulmonary airways, which comprises a
composition according to claim 12.

35   14.   Use of an agent for gene therapy, for the manufacture
of a medicament for treatment of a complaint on which the
said agent acts on administration via the pulmonary

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airways, characterised in that the medicament is in the
form of microparticles according to any of claims 1 to 11
or a composition according to claim 12.

15.   In a method of treating a complaint by administration
to the patient of an effective amount of an agent for gene
therapy that acts via pulmonary airways to treat the
complaint, the improvement comprising administration of
said agent in the form of microparticles according to
claim 1.