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Referencia : Expediente No. 00.038.727
Solicitante : INHALE THERAPEUTIC SYSTEMS, INC.
Asunto : Solicitud de patente para: **APARATOS Y MÉTODOS MEJORADOS DE AEROSOLIZACIÓN**
Fecha de la solicitud : 25 de mayo de 2000

TRAMITE:002 - EVENTO:001 - ACTUACION:444

1. Yo, Emilio FERRERO, obrando como apoderado del solicitante en el asunto antes identificado, me refiero al auto número 1889, notificado por estado del 22 de junio de 2000.

2. Mediante el auto número 1889 se puso en conocimiento del solicitante el informe que contiene las observaciones hechas por ese Despacho sobre la presente solicitud de patente. Atiendo a las observaciones de la manera siguiente:

2.1. Título del invento. Con el objeto de corregir un error que aparece en el título originalmente presentado el solicitante desea presentar un nuevo título como sigue:

APARATOS Y METODOS MEJORADOS DE AEROSOLIZACION

Adicionalmente presento nuevo formato de solicitud en donde aparece el nuevo título que presento. Además presento nueva pagina 1 del texto de la solicitud con el título corregido.

RESPUESTA AUTO No. 1889 DEL ESTADO
DEL D.A. 22 de junio de 2000

2.2. En la descripción. Presento junto con este escrito nuevas hojas 7 y 34, correspondiente al capítulo reivindicatorio, en el cual se ha hecho la conversión a algunas unidades que no se encontraban en el Sistema Internacional.

2.3. En las reivindicaciones. Presento junto con este escrito nueva hoja 39, correspondiente al capítulo reivindicatorio, en el cual se ha hecho la conversión a algunas unidades que no se encontraban en el Sistema Internacional.

2.4. Extractos. Adjunto a este memorial presento nuevo extracto, nueva tarjeta temática y nueva tarjeta de archivo de propietarios (Formas P-103, P-104 y P-105), los cuales están en conformidad con las modificaciones hechas al pliego reivindicatorio y al título.

2.5. Arte Final. Adjunto a este memorial presento los artes finales de acuerdo con lo requerido.

2.6. Copia Certificada. Presento con este escrito copia certificada de la solicitud de patente Estadounidense 60/136,518 del 28 de mayo de 1999, junto con su respectiva traducción simple de sus reivindicaciones y carátula, en las cuales se basa la reivindicación de la prioridad.

3. Puesto que se han atendido a cabalidad las observaciones hechas por ese Despacho sobre la presente solicitud de patente, atentamente solicito que se continúe su tramitación.

Bogotá, ~8 AGO. 2000
Señora Jefe,



EMILIO FERRERO

T.P.A. No. 31.929



COLO 09101-801-010-7
RCB/MRH/86

78
76



SOLICITUD DE
PATENTE DE
INVENCION

TRAMITE: 02 EVENTO: 01 ACTUACION: 01

DATOS DEL SOLICITANTE

(71) Nombres y Apellidos o Razón Social **INHALE THERAPEUTIC SYSTEMS, INC.**

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Ciudad

DATOS DE LA PATENTE DE INVENCION

(54) Título de la Invención

APARATOS Y METODOS MEJORADOS DE AEROSOLIZACION

PRIORIDAD REIVINDICADA SI ☒ NO ☐ TIPO DE PRIORIDAD ANDINA ☐ UNIONISTA ☒

(33) País

Fecha

No. Solicitud

US.
Convención de París.

28 de Mayo de 1999

60/136.518

Para Publicar a los ☐ 6 meses ☐ 12 meses ☒ 18 meses a partir de la fecha de la presente solicitud

Colo: 9101-801-010-7

RC/mrm.Id-79

(51) CLASIFICACION INTERNACIONAL

RESUMEN CON EL OBJETO Y LA FINALIDAD DE LA INVENCION

La presente invención proporciona métodos y aparatos para aerosolizar una formulación farmacéutica contenida dentro de un receptáculo. En un método, se proporciona una cantidad medida de gas presurizado, estando el gas presurizado previamente en equilibrio con un líquido. El gas medido se libera para crear una corriente de gas de alta presión. La corriente de gas de alta presión fluye a través de un mecanismo de aerosolización para extraer la formulación farmacéutica proveniente del receptáculo y para dispersar la formulación farmacéutica dentro de la corriente de gas para formar un aerosol,

ANEXOS DE LA SOLICITUD

- | | |
|--|--------------------------|
| Certificado de Existencia y Representación Legal cuando el solicitante sea Persona Jurídica | ☐ |
| Los poderes debidamente otorgados, o mención de su protocolización ante la SIC | ☐ |
| Recibo de la tasa respectiva | ☐ |
| Recibo de la tasa por concepto de excedente de palabras en la publicación | ☐ |
| Si se reivindica prioridad Andina: | |
| Copia de la primera solicitud | ☐ |
| Traducción oficial | ☐ |
| Si se reivindica prioridad Unionista: | |
| Copia certificada de la primera solicitud: | ☐ |
| Nota: Debe presentarse dentro de los tres (3) meses, contados a partir de la fecha de radicación de la solicitud posterior | |
| Traducción simple | ☐ |
| Capítulo descriptivo | ☐ |
| Dibujos, planos necesarios | ☐ |
| Capítulo reivindicatorio | ☐ |
| Tarjeta archivo propietarios según formato | ☐ |
| Tarjeta archivo temático según formato | ☐ |
| Extracto para publicación según formato | ☐ |
| Arte final 12 x 12 cms Y 6x6 cms | ☐ |
| Otros Documentos | ☐ |

SOLICITO LA CONCESION DE LA PATENTE DE INVENCION Y DECLARO QUE LA INVENCION ES NUEVA Y DE MI PROPIEDAD

FIRMA

NOMBRE EMILIO FERRERO

CC. 438.019 de Usaquén

T.P.No. 31.929

propulsantes que se pueden utilizar incluyen CO₂ y CFC (clorofluorocarburo) 11,12 y 11. En razón de que los CFC se han identificado como sustancias que afectan la capa de ozono de la estratosfera, los HFC o los HFA son los preferidos. El gas presurizado es preferiblemente medido en volumen en un rango que va desde aproximadamente 0,5 ml hasta aproximadamente 2,0 ml. La presión del gas medido se encuentra preferiblemente en el rango de aproximadamente 276,2 kPa a aproximadamente 828,8 kPa; y más preferiblemente de aproximadamente 345,3 kPa a aproximadamente 552,5 kPa. Cuando se utiliza CO₂, la presión del gas medido se encuentra preferiblemente en el rango de aproximadamente 5525 kPa a aproximadamente 8287,6 kPa.

Para medir el gas, al gas se le permite preferiblemente fluir dentro de una cámara de medición que esta ubicada para recibir y medir una cantidad de la porción de fase de vapor del propulsante. Se proporciona una válvula de liberación para liberar el propulsante medido para formar una corriente de gas de alta presión. El aparato incluye además un mecanismo de aerosolización para recibir la corriente de gas de alta presión, para extraer la formulación farmacéutica del receptáculo y

Aunque mostrado con la unidad de reservorio 78, se apreciará que modalidades a alternativas de fuentes de propulsante se pueden proporcionar. Por ejemplo, la unidad de reservorio 78 se puede remplazar con una cánula u otro tanque reemplazable para facilitar el fácil reemplazo de la cánula o el tanque cuando el propulsante se ha agotado. En razón de que es deseable reemplazar periódicamente el mecanismo de aerosolización 50, la fuente del propulsante se puede configurar para acoplarse con el mecanismo de aerosolización de tal forma que ambos se pueden remover y reemplazar una vez el propulsante se ha agotado.

Cuando se utiliza HFA como propulsante, la regresión de la porción de la fase de vapor mantenida dentro de la cámara de medición 116 se encuentra preferiblemente en el rango de aproximadamente 276,3 kPa a aproximadamente 828,8 kPa y más preferiblemente aproximadamente 345,3 kPa a aproximadamente 552,5 kPa. Configurar el aparato 34 para operar utilizando un amplio rango de presiones es ventajoso por que la presión del HFC puede variar dependiendo de la temperatura ambiente. Como tal, el aparato 34 se puede utilizar en una amplia variedad de ambientes y climas.

8. Un método como en la reivindicación 6, en donde el propulsante comprende CO_2 .

9. Un método como en la reivindicación 7, en donde la cantidad medida tiene un volumen en el rango que va de 0,5 ml a aproximadamente 2 ml y se presuriza a una presión en el rango de aproximadamente 276,3 kPa a aproximadamente 828,8 kPa.

10. Un método como en la reivindicación 6, que comprende además abrir una válvula para liberar la cantidad medida.

11. Un método como en la reivindicación 6, en donde la etapa de medición comprende permitirle a la porción de fase de vapor fluir dentro de la cámara de medición y cerrar la cámara de medición después de que la cámara de medición se llena.

12. Un aparato para aerosolizar la formulación farmacéutica mantenida dentro del receptáculo, el aparato comprende:

FORMATO DE DIGITALIZACION
RELACION DE ANEXOS



80

Expediente No		00038727	No de Folio
Item		No de unidades	✓
1	CD		
2	DVD		
3	REVISTA		
4	LIBRO		
5	PERIODICO		
6	DISQUETES		
7	SOBRE DE MANILA ✓	1	✓
8	SOBRE BLANCO TAMAÑO CARTA		
9	SOBRE BLANCO TAMAÑO OFICIO		
10	OTROS:		

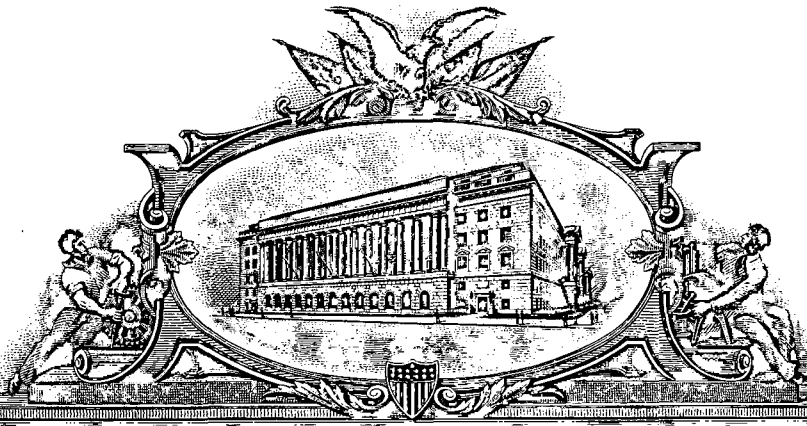
Observaciones:

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Arte final
Inhal Therapeutic Systems, Inc.

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PA 247400



THE UNITED STATES OF AMERICA

TO ALL TO WHOM THESE PRESENTS SHALL COME:

UNITED STATES DEPARTMENT OF COMMERCE

United States Patent and Trademark Office

May 11, 2000

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THE RECORDS OF THE UNITED STATES PATENT AND TRADEMARK
OFFICE OF THOSE PAPERS OF THE BELOW IDENTIFIED PATENT
APPLICATION THAT MET THE REQUIREMENTS TO BE GRANTED A
FILING DATE UNDER 35 USC 111.

APPLICATION NUMBER: 60/136,518

FILING DATE: May 28, 1999



By Authority of the
COMMISSIONER OF PATENTS AND TRADEMARKS

P. SWAIN
Certifying Officer

05/28/99
JCS53 U.S. PTO

PROVISIONAL APPLICATION

Attorney Docket No. 15225-004100

"Express Mail" Label No. EJ718885200US

Date of Deposit: May 28, 1999

I hereby certify that this is being deposited with the United States Postal Service "Express Mail Post Office to Addressee" service under 37 CFR 1.10 on the date indicated above, addressed to:

Assistant Commissioner for Patents
Box Provisional Patent Application
Washington, D.C. 20231

By: Janet Byrne

87
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JCS41 U.S. PTO
60/136518
05/28/99

ASSISTANT COMMISSIONER FOR PATENTS
BOX PROVISIONAL PATENT APPLICATION
Washington, D.C. 20231

Sir:

Transmitted herewith for filing is a provisional patent application under CFR 1.53(c) of:

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Bakshi	Ancesh	K.	Belmont, California
Tuttle	Derrick		San Mateo, California

Title: IMPROVED AEROSOLIZATION APPARATUS AND METHODS

Enclosed are:

- ☒ 17 pages of the specification (including description and claims).
☒ 12 sheet(s) of [] formal [X] informal drawing(s).
☒ Abstract.

Please charge Deposit Account No. 20-1430 as follows:

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PROVISIONAL PATENT APPLICATION

IMPROVED AEROSOLIZATION APPARATUS AND METHODS

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Entity: Large

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IMPROVED AEROSOLIZATION APPARATUS AND METHODS

BACKGROUND OF THE INVENTION

The present invention relates generally to the field of drug delivery, and in particular to the pulmonary delivery of drugs. More specifically, the invention relates to the aerosolization of pharmaceutical formulations for inhalation by a patient.

Effective drug delivery to a patient is a critical aspect of any successful drug therapy. Various routes of delivery exist, and each has its own advantages and disadvantages. Oral drug delivery of pills, capsules, elixirs, and the like, is perhaps the most convenient method, but many drugs are degraded in the digestive tract before they can be absorbed. Such degradation is a particular problem with modern protein drugs which are rapidly degraded by proteolytic enzymes in the digestive tract. Subcutaneous injection is frequently an effective route for systemic drug delivery, including the delivery of proteins, but enjoys a low patient acceptance. Since injection of drugs, such as insulin, one or more times a day can frequently be a source of poor patient compliance, a variety of alternative routes of administration have also been developed, including transdermal, intranasal, intrarectal, intravaginal, and pulmonary delivery.

Of particular interest to the present invention, pulmonary drug delivery relies on inhalation of a drug dispersion or aerosol by the patient so that the active drug within the dispersion can reach the distal (alveolar) regions of the lung. It has been found that certain drugs are readily absorbed through the alveolar region directly into blood circulation. Pulmonary delivery is particularly promising for the delivery of proteins and polypeptides which are difficult to deliver by other routes of administration. Such pulmonary delivery is effective both for systemic delivery and for localized delivery to treat diseases of the lungs.

Pulmonary drug delivery (including both systemic and local) can itself be achieved by different approaches, including liquid nebulizers, metered dose inhalers (MDI's) and dry powder dispersion devices. Dry powder dispersion devices are particularly promising for delivering protein and polypeptide drugs which may be readily formulated as dry powders. Many otherwise labile proteins and polypeptides may be stably stored as lyophilized or spray-dried powders by themselves or in combination with suitable powder carriers. The ability to deliver proteins and polypeptides as dry powders, however, is problematic in certain respects. The dosage of many protein and polypeptide

drugs is often critical so it is necessary that any dry powder delivery system be able to accurately, and precisely (repeatedly) deliver the intended amount of drug. Moreover, many proteins and polypeptides are quite expensive, typically being many times more costly than conventional drugs on a per-dose basis. Thus, the ability to efficiently deliver the dry powders to the target region of the lung with a minimal loss of drug is critical. It is further desirable that powder agglomerates present in the dry powder be sufficiently broken up prior to inhalation by the patient to assure effective systemic absorption or other pulmonary delivery.

U.S. Patent Nos. 5,785,049 and 5,740,794 describe exemplary apparatus and methods for aerosolizing a powdered medicament held within a receptacle. The complete disclosures of these patents are herein incorporated by reference. These apparatus and methods utilize a high pressure gas stream to extract a powdered medicament from a receptacle and to disperse the powder into the gas stream to produce an aerosol. Such apparatus and methods have proven to be generally effective in producing aerosolized medicaments for delivery to the lungs.

U.S. Patent No. 4,534,343 shows a typical metered dose inhaler where medication under pressure is stored in a canister. The drug is held in the pressurized liquid propellant that is held within the canister. The drug must be stable in this condition for the life of the system. This is undesirable for many labile medicaments such as proteins and peptides which may not be stable in this liquid for long periods of storage.

As described hereinafter, the invention provides alternative systems, devices and methods to aerosolize pharmaceutical formulations. In this way, alternative schemes are provided to assist in the delivery of pharmaceutical formulations to the lungs.

SUMMARY OF THE INVENTION

The invention provides exemplary systems, methods and apparatus for aerosolizing pharmaceutical formulations that are typically contained within a receptacle. In a broad sense, the invention utilizes a propellant's energy to aerosolize a pharmaceutical formulation. The pharmaceutical formulation may be stored separately from the propellant until aerosolized to prolong the life of the pharmaceutical formulation. The pharmaceutical formulation may be in powder or liquid form, with the energy of the propellant being employed to place the formulation in aerosolized form so that it may be inhaled and delivered to the lungs. In one aspect, a vapor phase portion of

phase portion of the propellant to enter into the metering chamber. Once the metering chamber is filled, the metering valve is preferably moved to a closed position. As one example, the metering valve may comprise a poppet valve which is pushed to move the valve to the open position. Once the chamber is filled, the poppet valve is released to
5 allow the valve to move to the closed position.

Conveniently, a lever may be provided which is movable to close the release valve and to open the metering valve. When the lever is moved back to its starting position, the metering valve closes while the release valve remains closed. In this way, the metered propellant remains held within the metering chamber. Preferably, a fire
10 button is provided to open the release valve and allow the metered gas to escape. In one particular aspect, a housing is provided to hold the reservoir, the metering chamber and the aerosolization mechanism. Preferably, a capture chamber is operably coupled to the housing. The capture chamber includes a mouthpiece through which the aerosolized powder may be withdrawn.

In one embodiment, a system is provided for aerosolizing a pharmaceutical
15 formulation. The system comprises a source of propellant which includes a vapor phase portion in equilibrium with a liquid phase portion. A pharmaceutical formulation is dispersed within at least a portion of the vapor phase portion of the propellant which has been separated from the liquid phase portion. Hence, with such a system a portion of a
20 vapor propellant may be used as an energy source to aerosolize a pharmaceutical formulation.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is a schematic diagram of an exemplary system for aerosolizing a
25 powder contained within a receptacle according to the invention.

Fig. 2 is a right side view of one embodiment of an aerosolization apparatus according to the invention.

Fig. 3 is a left side cross-sectional view of the apparatus of Fig. 2.

Fig. 3A is a magnified view of a bottom half of the apparatus of Fig. 3.

Fig. 4 is a bottom perspective view of a base unit of the apparatus of Fig.
30 2, with a cover being removed for convenience of illustration.

Fig. 5 is a partial left side cross-sectional view of the base unit of Fig. 4.

Fig. 6 illustrates the base unit of Fig. 4 with a lever moved to close the release valve.

Fig. 7 illustrates the base unit of Fig. 5 with the lever in the position shown in Fig. 6.

Fig. 8 illustrates the base unit of Fig. 7 with the lever moved to open a metering valve.

5 Fig. 9 is a top perspective view of a bottom portion of a reservoir unit of the apparatus of Fig. 3.

Fig. 10A is a bottom perspective view of a top portion of the reservoir unit.

Fig. 10B is a top perspective view of the top portion of Fig. 10A.

10 Fig. 11 is a top perspective view of a metering valve housing of the apparatus of Fig. 3.

Fig. 12 is a top perspective view of an actuator for the metering valve of the apparatus of Fig. 3.

Fig. 13A is a side view of a left arm of the base unit of Fig. 4.

Fig. 13B is a side view of a right arm of the base unit of Fig. 4.

15 Figs. 14A and 14B are rear and front views, respectively, of an actuator arm for the release valve of the base unit of Fig. 4.

Fig. 15 illustrates a filling unit to refill the reservoir unit of Fig. 3.

DETAILED DESCRIPTION OF THE SPECIFIC EMBODIMENTS

20 The invention provides exemplary apparatus and methods for aerosolizing pharmaceutical formulations. These formulations may be in dry powder form or they may be liquids. Examples of powdered medicaments that may be aerosolized according to the invention are described in PCT Publication No. WO96/32149, the complete disclosure of which is herein incorporated by reference. The powders aerosolized by the
25 invention preferably have a size in the range from about 0.5 μm to about 50 μm , and more preferably in the range from about 0.5 μm to about 5 μm . Alternatively, the pharmaceutical formulations may be in liquid form. The pharmaceutical formulations may be held in receptacles which have a puncturable access surface, and are sometimes referred to as "blister packages."

30 The invention preferably utilizes a high-pressure gas stream to extract the powder from the receptacle and to aerosolize the powder so that it may be inhaled by a patient. Examples of aerosolization mechanisms which may be utilized with the invention include those described in U.S. Patent Nos. 5,785,049 and 5,740,794,

previously incorporated by reference, and in copending U.S. provisional patent application serial number 60/087,929, filed June 4, 1998, the disclosure of which is herein incorporated by reference. At least one of the specific embodiments described hereinafter utilizes an aerosolization mechanism as well as other related components which are

5 constructed in a manner similar to those described in copending U.S. provisional patent application serial number 60/087,929, previously incorporated by reference. However, it will be appreciated that other types of aerosolization mechanisms which require the use of a high pressure gas stream to extract a powder from a receptacle and to entrain the powder within the gas stream may be utilized.

10 In one aspect, the invention utilizes energy from a propellant to aerosolize a pharmaceutical formulation. The propellant may include a vapor phase portion in equilibrium with a liquid phase portion. Some of the vapor phase portion may be extracted and employed as an energy source to aerosolize the pharmaceutical formulation. For example, the extracted vapor propellant may be employed to extract a powder from a

15 receptacle, to deagglomerate the powder and to place the powder in a gas stream in aerosolized form.

Referring now to Fig. 1, an exemplary system 10 for aerosolizing a powder contained within a receptacle 12 will be described. System 10 comprises a reservoir 14 which is filled with a propellant 16 having a liquid phase portion 18 and a vapor or gas

20 phase portion 20. Extending from a top end of reservoir 14 is a line 22 to which is coupled a metering valve 24. Also coupled to metering valve 24 is a metering chamber 26. Disposed at the other end of metering chamber 26 is a release valve 28 to which is coupled a line 30. Line 30 extends to an aerosolization mechanism 32 which interacts with receptacle 12 to extract and aerosolize the powder contained within receptacle 12.

25 Propellant 16 is selected so that vapor phase portion 20 is under pressure. Hence, when metering valve 24 is opened, the pressurized propellant enters into metering chamber 26. Release valve 28 is closed to prevent the pressurized propellant from proceeding into aerosolization mechanism 32. When metering chamber 26 is filled, metering valve 24 is closed to retain a metered amount of pressurized gas within metering

30 chamber 26. Hence, metering chamber 26 is filled with a metered amount of gas which is at a pressure determined by the type of propellant as well as the temperature of metering chamber 26.

When it is desired to aerosolize the powder, aerosolization mechanism 32 is inserted into receptacle 12 to gain access to the powder. Release valve 28 is then

rapidly opened to abruptly release the metered gas within metering chamber 26. The metered gas passes through line 30 and into aerosolization mechanism 32 which extracts the powder from receptacle 12 and entrains the powder in a high pressure gas stream to form an aerosol. The aerosol may be captured in a capture chamber so that it will be

5 available for inhalation by a patient.

Hence, system 10 is configured to operate simply by releasing a metered amount of pressurized gas which was obtained by extracting a vapor phase portion of a propellant that was previously in equilibrium with a liquid phase portion. System 10 is configured to meter a fixed volume of the gas at a fixed pressure. In this way,

10 aerosolization mechanism 32 is able to consistently receive a known volume of gas which is at a known pressure so that it may optimally extract and aerosolize the powder within receptacle 12. With such a configuration, a variety of advantages are provided. For example, system 10 is easy to operate because it only requires the opening of metering valve 24 when metering the gas, and the opening of release valve 28 to aerosolize the

15 powder. Another advantage is that the size of system 10 may be reduced by utilizing a propellant. Typically, only a small volume of propellant is needed so that the size of reservoir 14 may be made relatively small. Such a configuration reduces the overall size of system 10 so that it will be easy and convenient for a user to carry and store.

Typically, system 10 will have a longer life than comparable MDI devices because

20 system 10 employs only a vapor phase portion of the propellant to aerosolize the powder rather than the liquid phase as with MDI devices. Further, system 10 is more simple to manufacture and easier to use (because the patient is not required to shake the propellant) than comparable MDI devices since it does not require a drug to be suspended within the liquid phase portion of the propellant.

25 Preferable propellants that may be utilized with system 10 include HFA or HFC, such as hydrofluorocarbon 134a (HFC-134a, 1,1,2-tetrafluoroethane) and hydrofluorocarbon 227ea (HFC-227ea, 1,1,1,2,3,3,3-heptafluoropropane). These propellants are known in the art and are commercially available from the Minnesota Mining & Manufacturing Co. and Dupont. Such propellants are particularly useful in that

30 they do not utilize CFCs. Other propellants that may be used with the invention include CO₂, CFCs, and the like.

Referring now to Fig. 2, an exemplary embodiment of an apparatus 34 for aerosolizing a powder medicament will be described. Apparatus 34 comprises a base unit 36 and a capture chamber 38 which is removably attached to base unit 36. Capture

chamber 38 is configured to slide over base unit 36 to reduce the overall size of apparatus 34 during storage and to provide a degree of protection to the components within base unit 36. Capture chamber 38 also includes a mouthpiece 40 that is rotatable between an open position and a closed position. During aerosolization, mouthpiece 40 is in the closed position as illustrated in Fig. 2. When the patient is ready to inhale the aerosolized medicament, mouthpiece 40 is rotated 180 degrees to the open position where the patient may place his mouth over the mouthpiece and inhale the powdered medicament from capture chamber 38.

While their size is smaller, the overall shape and appearance of capture chamber 38 and base unit 36 are similar to those described in copending application number 60/087,929, previously incorporated by reference. By utilizing a propellant to aerosolize a medicament, base unit 36 does not require a cylinder to compress air from a large volume and does not require a lever which fits the patient's hand to provide the energy for compressing the gas. Hence, base unit 36 may have its size reduced. Merely by way of example, base unit 36 may have a height that is in the range from about 8 cm to 12 cm. Capture chamber 38 may have a height in the range from about 9 cm to about 13 cm.

Still referring to Fig. 2, base unit 36 includes an opening 42 into which a receptacle containing a powdered medicament is inserted. Base unit 36 also includes a fire button 44 which is pressed to pierce the receptacle and to release a volume of pressurized gas to aerosolize the medicament in the receptacle as described hereinafter. A release button 46 is provided to allow the receptacle to be removed from opening 42 after the medicament has been aerosolized. Base unit 36 further includes a lever 48 which may be operated to meter an amount of pressurized gas as described hereinafter.

Referring now to Figs. 3 and 3A, the components within base unit 36 will be described. Disposed within base unit 36 is an aerosolization mechanism 50 which is employed to extract a medicament from a receptacle 52, to entrain the medicament within a high pressure gas stream, and to deliver the aerosolized medicament into capture chamber 38. Base unit 36 also includes a carriage assembly 54 which moves receptacle 52 to engage aerosolization mechanism 50 when fire button 44 is pressed. Aerosolization mechanism 50, carriage assembly 54, and their related components are constructed essentially identical to those described in copending application number 60/087,929, previously incorporated by reference. As such, their manner of construction will only be briefly described in this application.

To operate apparatus 34, receptacle 52 is inserted into opening 42 (see Fig. 2) as previously described. Fire button 44 is then pushed to engage receptacle 52 with aerosolization mechanism 50. More specifically, when fire button 44 is pushed, gear teeth 56 are pivoted about a pivot pin 58, causing a lifter 60 of carriage assembly 54 to
 5 move receptacle 52 toward aerosolization mechanism 50. When fire button 44 is fully depressed, a pointed tip 62 and side punches 64 pierce through receptacle 52 and enter into the receptacle. Depression of fire button 44 also operates a valve 66 to allow for the release of a metered volume of high pressure gas into aerosolization mechanism 50 so that the powder within receptacle 52 may be aerosolized. More specifically, depression
 10 of fire button 44 causes a valve actuator 68 of an actuator arm 70 to be released from an over center position, thereby unlocking valve 66.

Once valve 66 is unlocked, the metered amount of high pressure gas within base unit 36 causes valve 66 to "pop open." More specifically, the release of valve actuator 68 causes the metered amount of high pressurized gas to come into contact with
 15 the underside of the diaphragm 72 causing a valve seat 74 to be lifted from a passage 76. In this manner, the propellant is allowed to flow through passage 76 and into aerosolization mechanism 50. The high pressurized gas then extracts the powdered medicament from receptacle 52, deagglomerates the powdered medicament and disperses the powdered medicament into capture chamber 38 as previously described. Hence, to
 20 operate apparatus 34, the user simply inserts receptacle 52 and then presses fire button 44 which will cause the aerosolized medicament to be introduced into capture chamber 38 where it may be withdrawn through mouthpiece 40.

Still referring to Figs. 3 and 3A, the manner in which the pressurized gas which is supplied to aerosolization mechanism 50 through valve 66 will be described.
 25 Base unit 36 includes a reservoir unit 78 which is constructed of a bottom member 80 and a top member 82. Bottom member 80 is also shown in Fig. 9, and top member 82 is shown in Figs. 10A and 10B. Bottom member 80 includes an opening 84 and top member 82 includes an opening 86. When top member 82 is secured to bottom member 80, openings 84 and 86 together form a cavity 88 which holds the propellant.
 30 Conveniently, top member 82 includes groove 90 for holding an O-ring seal 92 so that the propellant does not escape from cavity 88. Traveling through bottom member 80 is a passage 94. Cavity 88 is configured such that it may hold a liquid phase portion of the propellant in equilibrium with a vapor phase portion. The vapor phase portion of the

propellant extends into passage 94 where it is prevented from escaping by a metering valve 96.

As also shown in Fig. 11, metering valve 96 includes a housing 98 which is received into an opening 100 of bottom member 80 (see Fig. 9). Housing 98 has a central passage 102 which communicates with passage 94 of bottom member 80. As best shown in Figs. 3 and 3A, a valve poppet 104 is positioned within passage 102. A spring 106 biases valve poppet 104 upward so that an O-ring 108 disposed about valve poppet 104 engages housing 98. In this way, metering valve 96 is configured to be in a normally closed position where the vapor phase propellant is prevented from moving beyond valve poppet 104.

Housing 98 may conveniently be divided into a bottom portion 110 and a top portion 112. When metering valve 96 is in the closed position, the vapor phase propellant remains within bottom portion 110. When metering valve 96 is opened, the vapor phase propellant rushes into top portion 112 and into a metering tube 114. As such, the volume within top portion 112 and within metering tube 114 define a metering chamber 116. At the opposite end of metering chamber 116 is valve 66 whose function is to prevent the escape of the pressurized gas from metering chamber 116 until valve actuator 68 is released as previously described. When valve 66 is opened, the gas within metering chamber 116 rushes into passage 76 where it enters into aerosolization mechanism 50.

As best shown in Figs. 3, 4 and 12, a valve actuator 118 is disposed over housing 98. Valve actuator 118 has a passage 120 which allows for the passage of gases between an interior 122 and metering tube 114. As best shown in Figs. 3 and 3A, valve poppet 104 extends vertically above housing 98. In this way, when valve actuator 118 is moved downward, it will eventually engage valve poppet 104 to cause spring 106 to compress and to move O-ring 108 from housing 98. In this way, pressurized gas within bottom portion 110 may escape into metering chamber 116.

As best shown in Figs. 4 and 10A, top member 82 includes an opening 123 through which a bleed valve 124 is inserted. Bleed valve 124 may conveniently be constructed as a screw having an O-ring. Bleed valve 124 is opened when filling the cavity 88 with a liquid propellant. Preferably, cavity 88 is filled by inserting the propellant through passage 102 of housing 98 as described in greater detail with reference to Fig. 15.

Referring now to Figs. 4 and 5, construction of base unit 36 to facilitate opening and closing of valves 66 and 96 (See Fig. 3A) will be described. As best shown in Fig. 4, base unit 36 includes a chassis 126 to which a frame 128 is coupled. Although not shown, a similar frame is also coupled to chassis 126 on the other side of base unit 36.

5 Movably coupled to frames 128 are a left arm 130 and a right arm 132. Arms 130 and 132 are also shown in Figs. 13A and 13B, respectively. Arms 130 and 132 include a hole 134 through which a pin 136 is inserted as shown in Fig. 4. Pins 136 extend through slots (not shown) in frames 128 so that arms 130 and 132 may be moved vertically up and down relative to frames 128. Lever 48 is pivotally coupled between arms 130 and 132 by
10 a pin 137. Lever 48 includes an extension 138 which engages bottom member 80 when lever 48 is pivoted. In so doing, lever 48 is moved away from bottom member 80 to move arms 130 and 132 vertically downward relative to frames 128.

Also pivotally coupled to frame 128 is an actuator arm 140. As best shown in Fig. 4, a pin 142 is employed to pivotally couple actuator arm 140 to frame 128.
15 Actuator arm 140 is also shown in Figs. 14A and 14B. As best shown in Figs. 14A and 14B, actuator arm 140 includes a slot 144 through which a pin 146 (see Fig. 13B) on right arm 132 moves when right arm 132 is moved relative to frame 128. In this way, movement of arm 132 relative to frame 128 causes actuator arm 140 to pivot about pin 142.

20 As best shown in Fig. 4, actuator arm 140 engages a release valve set arm 148. Valve set arm 148 is configured to pivot with actuator arm 140. Further, valve set arm 148 is employed to move actuator arm 70 (see Fig. 3A) so that valve 66 may be held in a closed position until valve set arm 148 is moved to allow valve 66 to open. Conveniently, valve set arm 148 and actuator arm 70 may be configured to be essentially
25 identical to similar components described in copending application serial number 60/087,929, previously incorporated by reference.

Chassis 126 includes a boss 150 under which valve set arm 148 may be positioned to maintain valve 66 in the closed position. Valve set arm 148 is moved under boss 150 by moving lever 48, which causes valve set arm 148 to pivot until received
30 under boss 150. At this position, valve 66 is in an over center position where it will remain closed until valve set arm 148 is moved out from under boss 150.

Fire button 44 includes a tab 152 which engages a post 154 on valve set arm 148 when fire button 44 is depressed. As tab 152 pushes valve set arm 148 out from under boss 150, valve actuator arm 70 (see Fig. 3A) is allowed to move back away from

its over center position, unclamping diaphragm 72. In this way, the pressurized gas within metering chamber 116 is allowed to unseat valve actuator 68 and rush into passage 76 as previously described.

As best shown in Fig. 4, pin 136 also couples right arm 132 to valve actuator 118. Left arm 130 is coupled to valve actuator 118 in a similar manner. In this way, when lever 48 is moved, valve actuator 118 is moved downward to open valve 96 (see Fig. 3A). Hence, operation of lever 48 closes valve 66 and subsequently opens metering valve 96. In this way, the metering chamber may be filled with pressurized gas. As lever 48 is moved back, metering valve 96 is closed so that the pressurized gas remains held within the metering chamber.

As shown in Figs. 4 and 5, base unit 36 has lever 48 in a home position. As previously described, in this position metering valve 96 is in its normally closed position. Assuming that fire button 44 was previously pressed, valve 66 is in its normally open position. As shown in Figs. 6 and 7, lever 48 is beginning to be moved away from its home position. In so doing, extension 138 engages bottom member 80 to move arms 130 and 132 downward. In turn, valve set arm 148 is moved underneath boss 150 to close valve 66. At the same time, valve actuator 118 is moved downward until it engages valve poppet 104 (see Fig. 7). As shown in Fig. 5, valve actuator 118 is spaced apart from valve poppet 104 when lever 48 is in home position. Hence, when reaching the position shown in Fig. 7, actuator 118 has not yet pushed valve poppet 104 downward. As such, valve 66 is closed before metering valve 96 is opened.

As shown in Fig. 8, lever 48 is fully pivoted, causing valve actuator 118 to force valve poppet 104 downward. In so doing, metering valve 96 is opened to allow the pressurized gas within bottom portion 110 to escape into metering chamber 116. Lever 48 may then be moved back to the home position to close metering valve 96. As lever 48 is moved back to the home position, valve set arm 148 remains under boss 150 so that the pressurized gas is held within metering chamber 116. Fire button 44 may then be depressed to open valve 66 to allow the powdered medicament to be aerosolized as previously described. Although not shown, an interlock may be provided to ensure that fire button 44 may not be actuated unless a receptacle has been placed in apparatus 34. In this way, the propellant is prevented from being released unless a receptacle is inserted.

One advantage of utilizing reservoir unit 78 to hold a propellant is that the overall size of base unit 36 may be reduced. At the same time, reservoir unit 78 is able to hold a sufficient volume of propellant so that many doses may be supplied to the patient.

Merely by way of example, cavity 88 may be configured to hold about 2cc of HFC liquid. This will allow apparatus 34 to be actuated approximately 100 times using about 0.65cc of vapor per actuation. As such, metering chamber 116 will preferably be configured to hold about 0.65cc of the HFC vapor. However, it will be appreciated that reservoir unit 78 and metering chamber 116 may be configured to hold other volumes of propellant.

Although shown with reservoir unit 78, it will be appreciated that alternative embodiments of propellant sources may be provided. For example, reservoir unit 78 may be replaced with a canister or other replaceable tank to facilitate easy replacement of the canister or tank when the propellant has been depleted. Because it is desirable to periodically replace aerosolization mechanism 50, the source of propellant may be configured to be coupled to the aerosolization mechanism so that both can be removed and replaced once the propellant has been depleted.

When utilizing HFA as the propellant, the pressure of the vapor phase portion held within metering chamber 116 is preferably in the range from about 40 psi to about 120 psi, and more preferably about 50 psi to about 80 psi. Configuring apparatus 34 to operate utilizing a wide range of pressures is advantageous in that the pressure of the HFC may vary depending upon the ambient temperature. As such, apparatus 34 may be used in a wide variety of environments and climates.

Fig. 15 illustrates a filling unit 200 that may be employed to fill reservoir unit 78 (see Fig. 3) with a propellant. Filling unit 200 comprises a supply tank 202 to store the propellant. Coupled to tank 202 is a valve 204 that may be opened to allow the propellant to be transferred from tank 202. Tubing 206 extends from valve 204 and is configured to fit over metering valve housing 98 (see also Fig. 11).

In use, tubing 206 is placed over housing 98 and valve 204 is opened. The propellant passes through housing 98 and into cavity 88 (see Fig. 10A). Bleed valve 124 may be opened to provide a vent.

Optionally, apparatus 34 may include a visual display of the number of times the apparatus has been used and/or the number of times it may be used before requiring a propellant refill and/or replacement. Conveniently, a counting system may be coupled to the fire button 44 or lever 48 to count the number of actuations.

The invention has now been described in detail for purposes of clarity of understanding. However, it will be appreciated that certain changes and modifications may be practiced within the scope of the appended claims.

WHAT IS CLAIMED IS:

1 1. A method for aerosolizing a pharmaceutical formulation contained
2 within a receptacle, comprising:
3 providing a metered amount of a pressurized gas that was previously in
4 equilibrium with a liquid;
5 releasing the metered gas to create a high pressure gas stream; and
6 flowing the high pressure gas stream through an aerosolization mechanism to extract the
7 pharmaceutical formulation from the receptacle and to disperse the pharmaceutical
8 formulation within the gas stream to form an aerosol.

1 2. A method as in claim 1, wherein the pressurized gas comprises
2 HFC.

1 3. A method as in claim 1, wherein the pharmaceutical formulation is
2 in dry powder form.

1 4. A method as in claim 1, wherein the pharmaceutical formulation is
2 in liquid form.

1 5. A method as in claim 1, further comprising opening a valve to
2 release the pressurized gas.

1 6. A method for aerosolizing a pharmaceutical formulation contained
2 within a receptacle, the method comprising:
3 providing a propellant having a liquid phase portion and a vapor phase
4 portion;
5 metering an amount of the vapor phase portion;
6 releasing the metered amount to form a high pressure gas stream; and
7 flowing the high pressure gas stream through an aerosolization mechanism
8 to extract the pharmaceutical formulation from the receptacle and to disperse the
9 pharmaceutical formulation within the gas stream to form an aerosol.

1 7. A method as in claim 6, wherein the propellant comprises HFC.

1 8. A method as in claim 6, wherein the propellant comprises CO₂.

1 9. A method as in claim 7, wherein the metered amount has a volume
2 in the range from about 0.5 ml to about 2 ml and is pressurized to a pressure in the range
3 from about 40 psig to about 120 psig.

1 10. A method as in claim 6, further comprising opening a valve to
2 release the metered amount.

1 11. A method as in claim 6, wherein the metering step comprises
2 permitting the vapor phase portion to flow into a metering chamber and closing the
3 metering chamber after the metering chamber is filled.

1 12. An apparatus for aerosolizing a pharmaceutical formulation held
2 within a receptacle, the apparatus comprising:
3 a reservoir which is adapted to hold a propellant having in equilibrium a
4 liquid phase portion and a vapor phase portion;
5 a metering chamber which is adapted to receive an amount of the vapor
6 phase portion of the propellant;
7 a release valve which is adapted to release the metered propellant to form a
8 high pressure gas stream; and
9 an aerosolization mechanism which is adapted to receive the high pressure
10 gas stream, to extract the pharmaceutical formulation from the receptacle and to disperse
11 the powder within the gas stream to form an aerosol.

1 13. An apparatus as in claim 12, further comprising a metering valve
2 disposed between the reservoir and the metering chamber, wherein the metering valve is
3 movable to an open position to allow the vapor phase portion of the propellant to enter the
4 metering chamber.

1 14. An apparatus as in claim 13, further comprising a lever which is
2 moveable to close the release valve and to open the metering valve.

1 15. An apparatus as in claim 12, further comprising a fire button to
2 open the release valve.

1 16. An apparatus as in claim 12, further comprising a housing to hold
2 the reservoir, the metering chamber and the aerosolization mechanism.

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1 17. An apparatus as in claim 16, further comprising a capture chamber
2 operably coupled to the housing, wherein the capture chamber includes a mouthpiece.

1 18. An apparatus as in claim 13, wherein the metering valve comprises
2 a poppet valve which is pushed to move the valve to the open position.

1 19. A system to aerosolize a pharmaceutical formulation, the system
2 comprising:
3 a source of propellant which includes a vapor phase portion in equilibrium
4 with a liquid phase portion;
5 a metering chamber disposed to meter volumes of the vapor phase portion
6 of the propellant;
7 a release valve to release the metered propellant to form a high pressure
8 gas stream; and
9 an aerosolization mechanism disposed to receive the high pressure gas
10 stream, to extract the powder from the receptacle and to disperse the pharmaceutical
11 formulation within the gas stream to form an aerosol.

1 20. A system as in claim 19, further comprising a metering valve
2 disposed between the source of propellant and the metering chamber, wherein the
3 metering valve is movable to an open position to allow the vapor phase propellant to
4 move into the metering chamber.

1 21. A system as in claim 20, further comprising a lever which is
2 movable to close the release valve and to open the metering valve.

1 22. A system as in claim 19, further comprising a fire button which is
2 operable to open the release valve.

1 23. A system as in claim 19, further comprising a housing holding the
2 source of propellant, the metering chamber and the aerosolization mechanism, and a
3 capture chamber having a mouthpiece operably coupled to the housing.

1 24. A system to aerosolize a pharmaceutical formulation, said system
2 comprising:

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- 3 a source of propellant which includes a vapor phase portion in equilibrium
4 with a liquid phase portion; and
5 a pharmaceutical formulation dispersed within at least a portion of the
6 vapor phase portion of the propellant which has been separated from the liquid phase
7 portion.

[illegible]

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IMPROVED AEROSOLIZATION APPARATUS AND METHODS

ABSTRACT OF THE DISCLOSURE

The invention provides exemplary methods and apparatus for aerosolizing a pharmaceutical formulation contained within a receptacle. In one method, a metered amount of a pressurized gas is provided, with the pressurized gas previously being in equilibrium with a liquid. The metered gas is released to create a high pressure gas stream. The high pressure gas stream is flowed through an aerosolization mechanism to extract the pharmaceutical formulation from the receptacle and to disperse the pharmaceutical formulation within the gas stream to form an aerosol.

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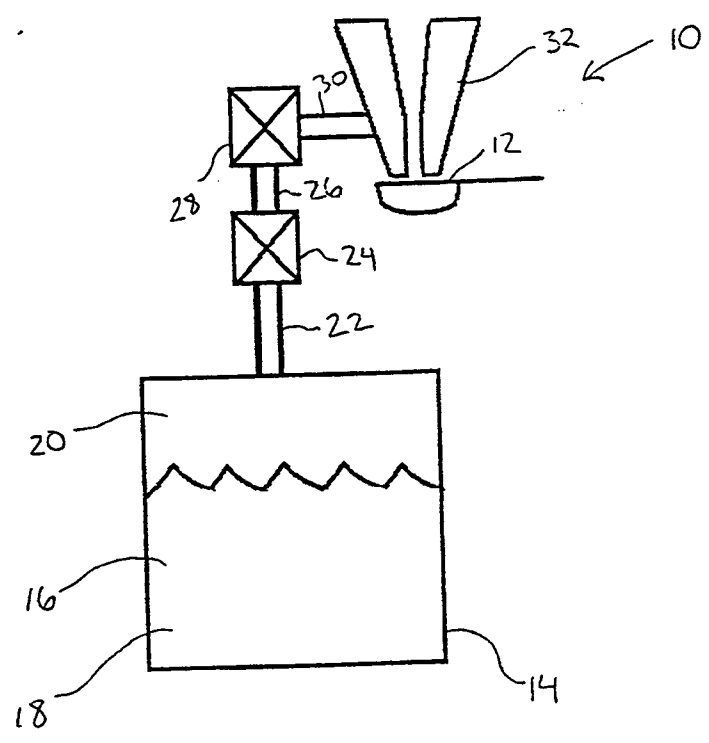
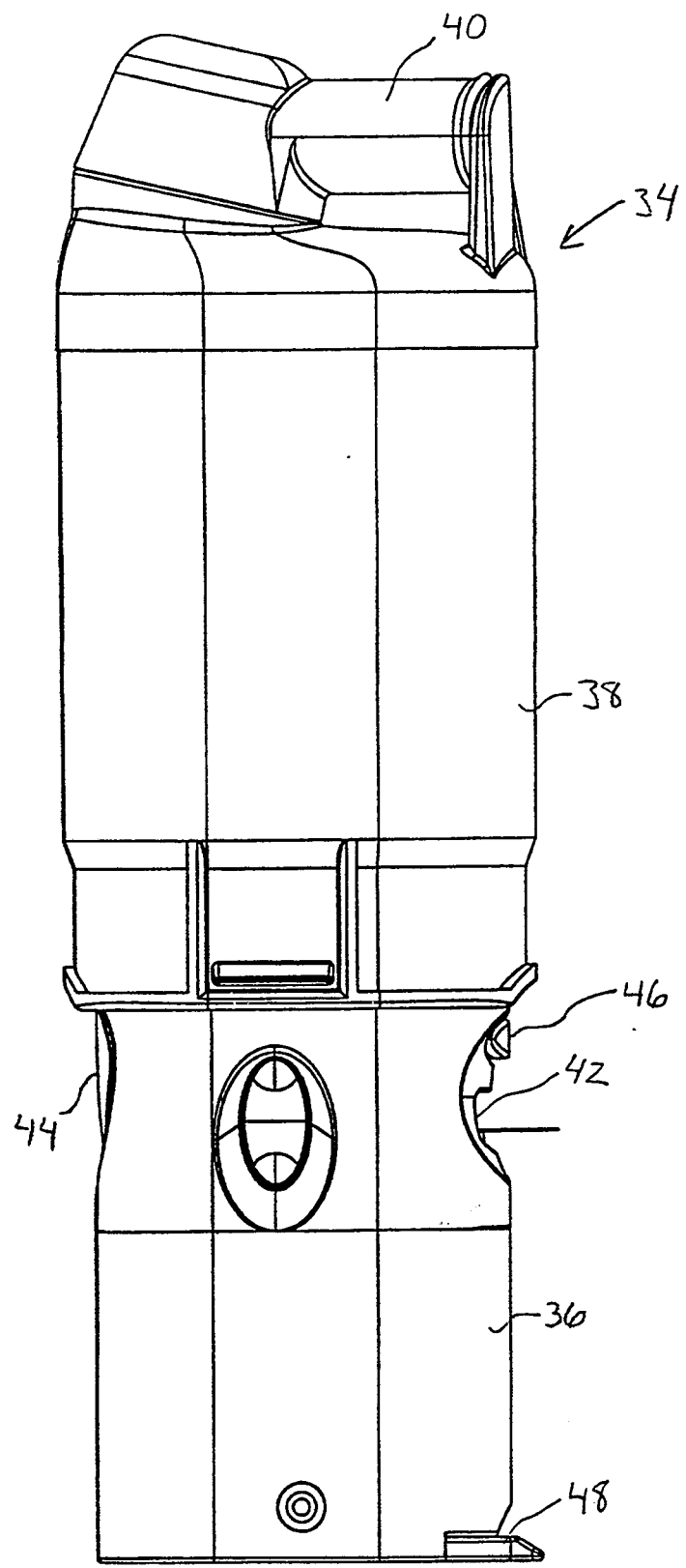


FIG. 1

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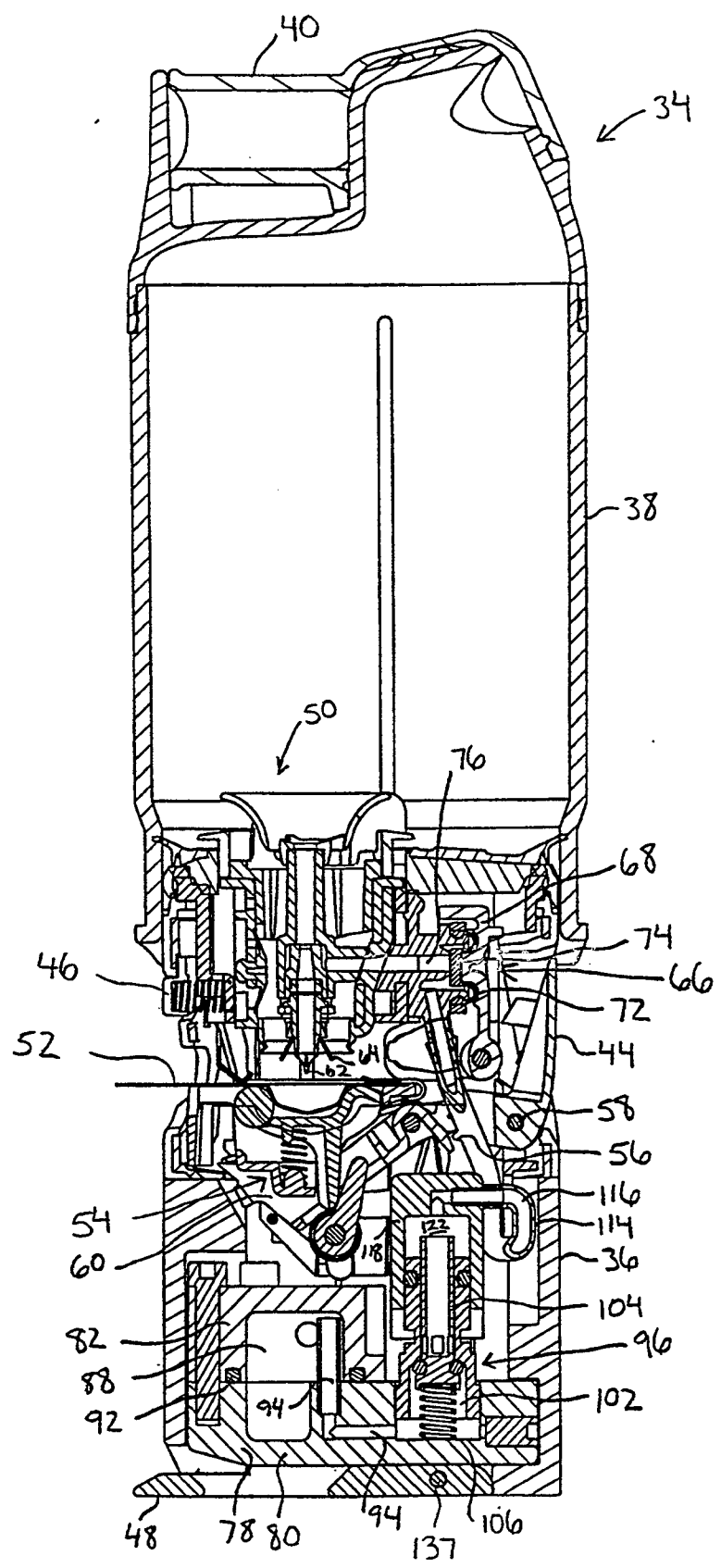
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FIG. 2

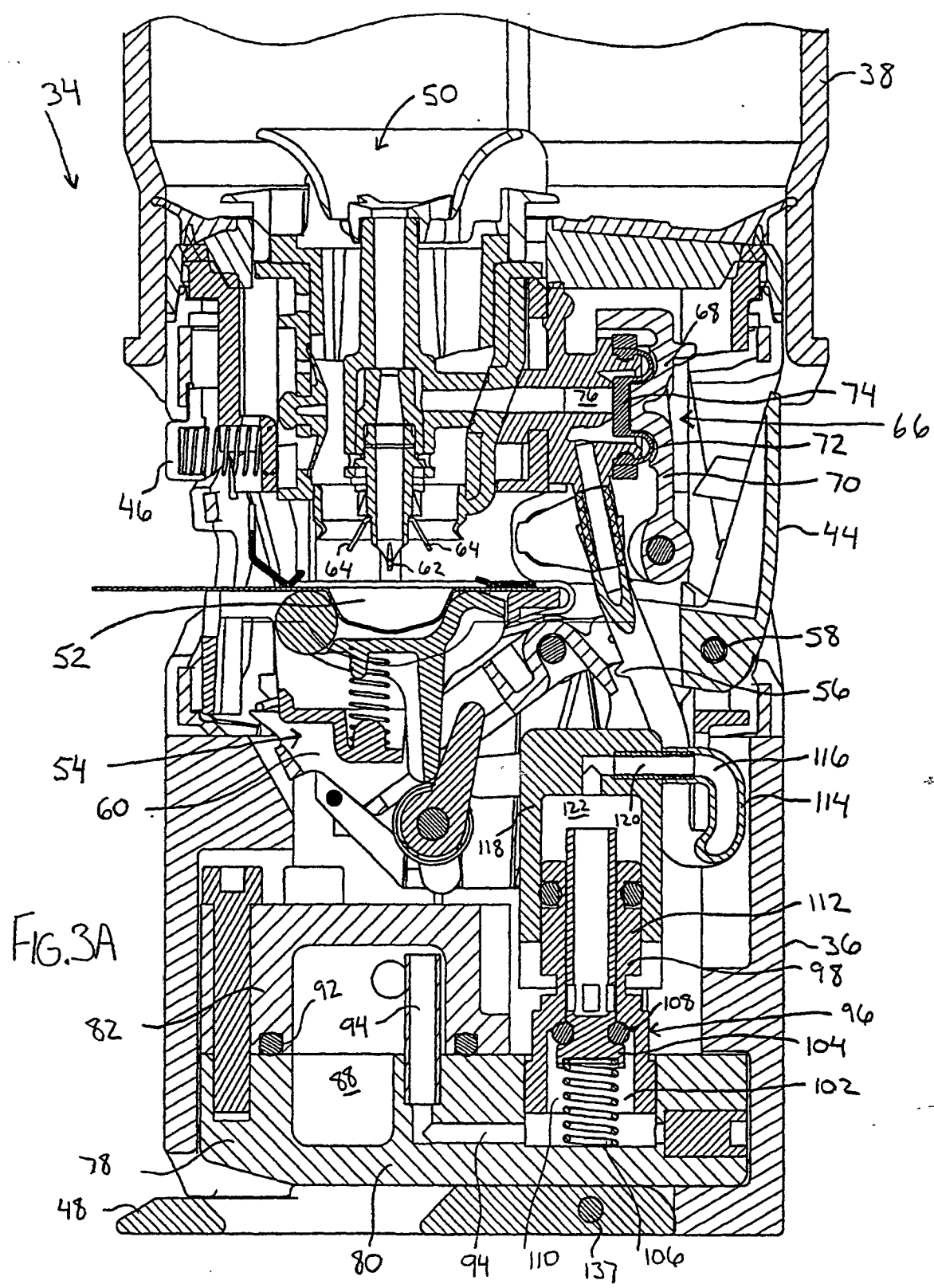


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FIG. 3



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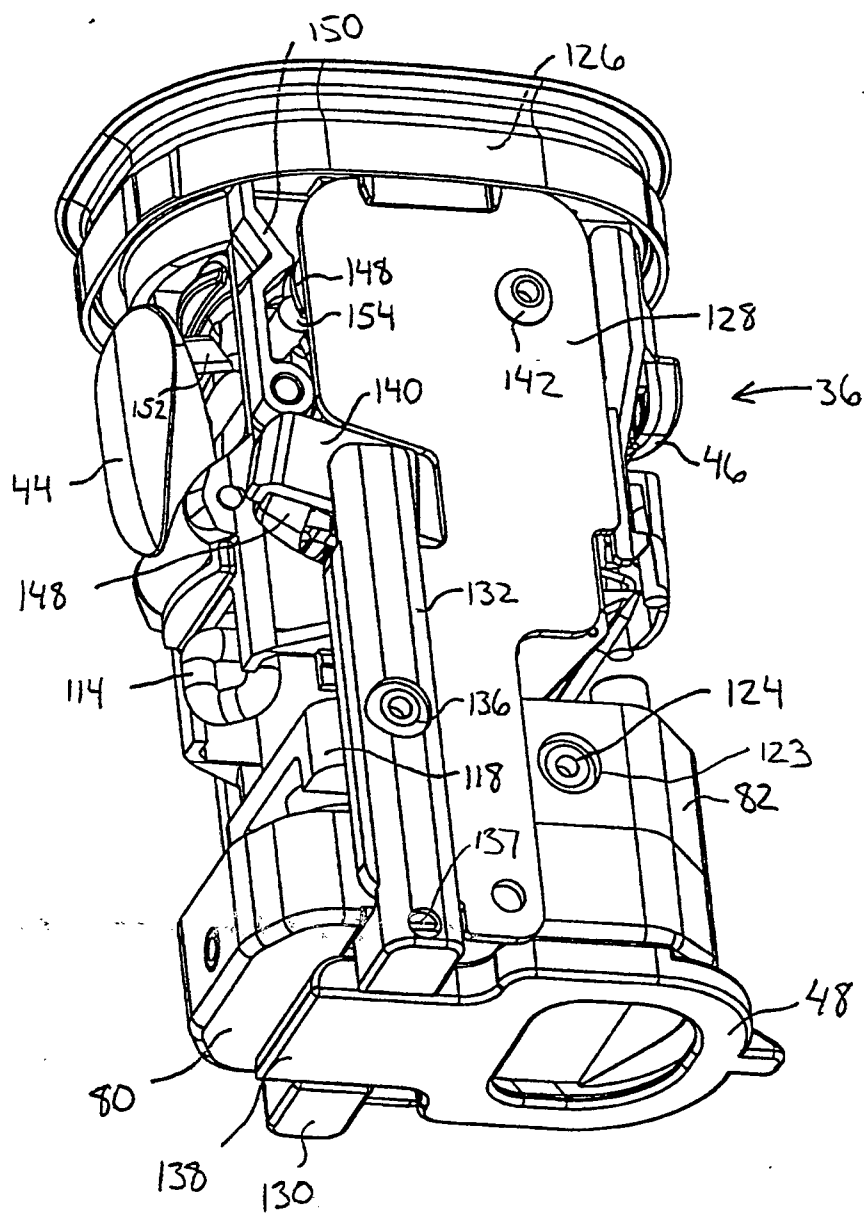
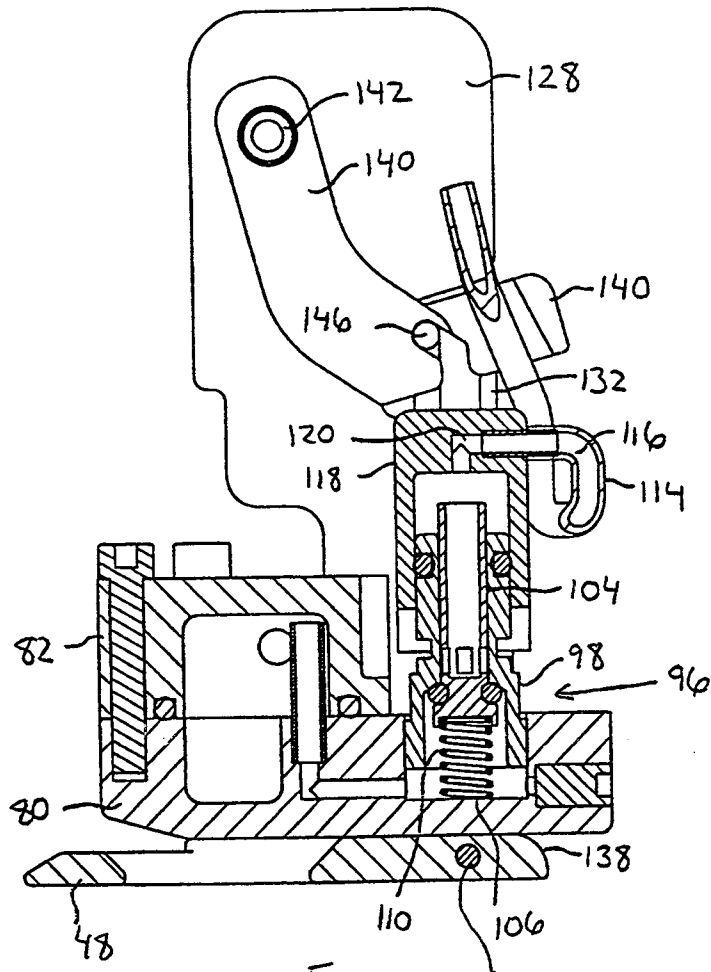
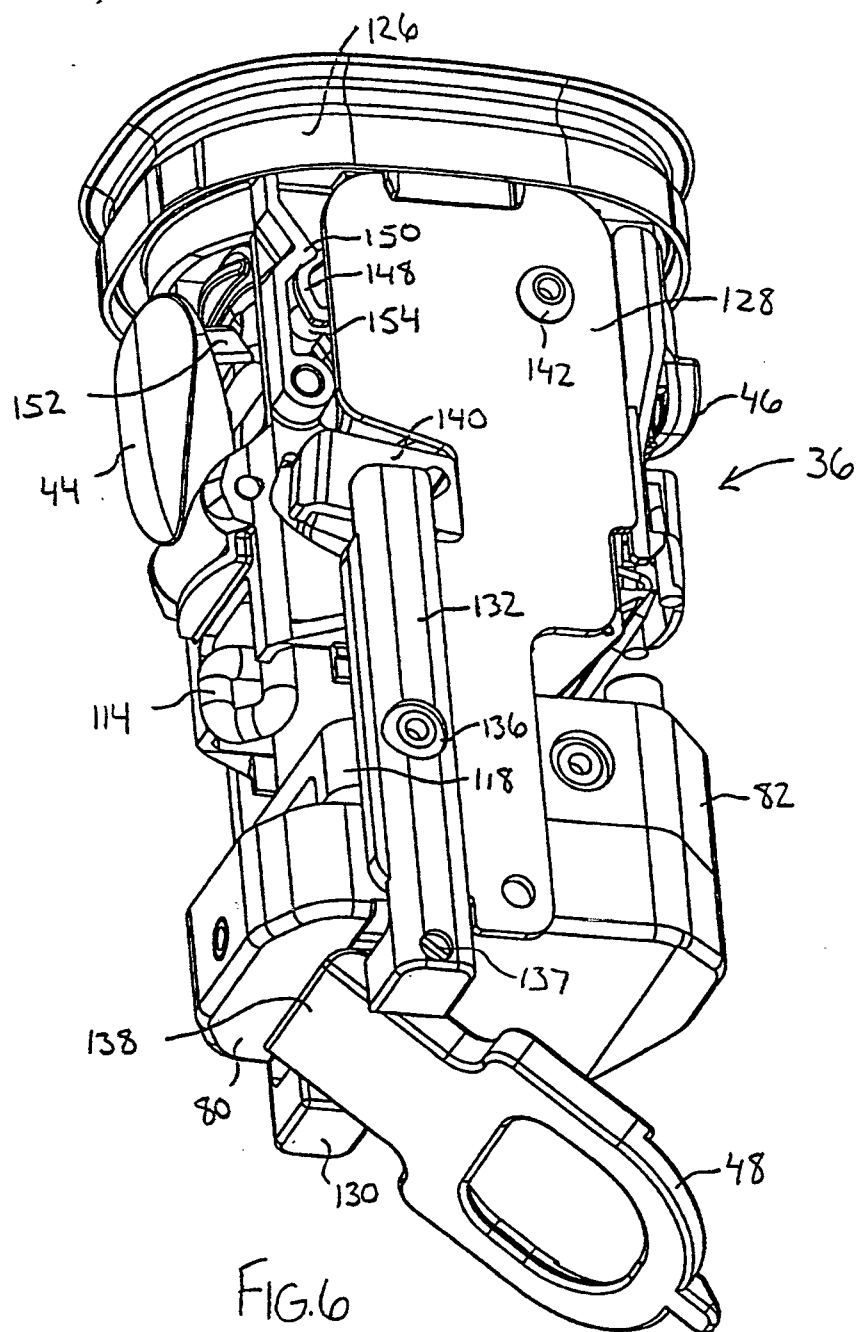


FIG. 4

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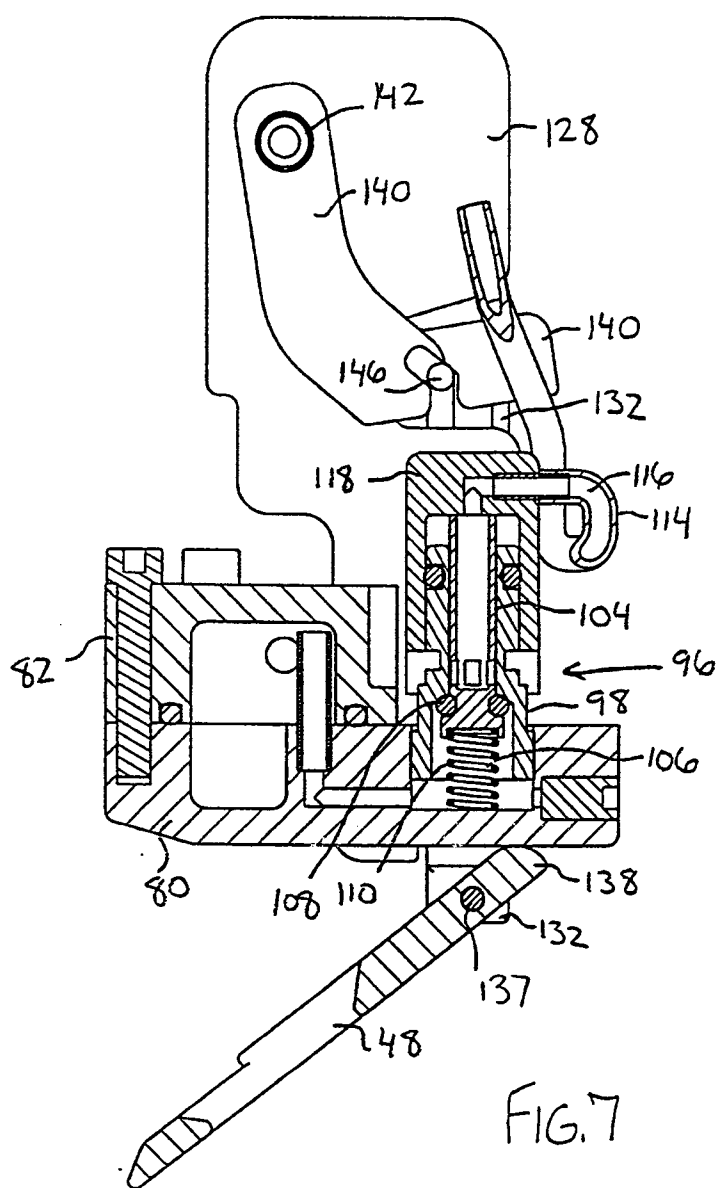


FIG. 8

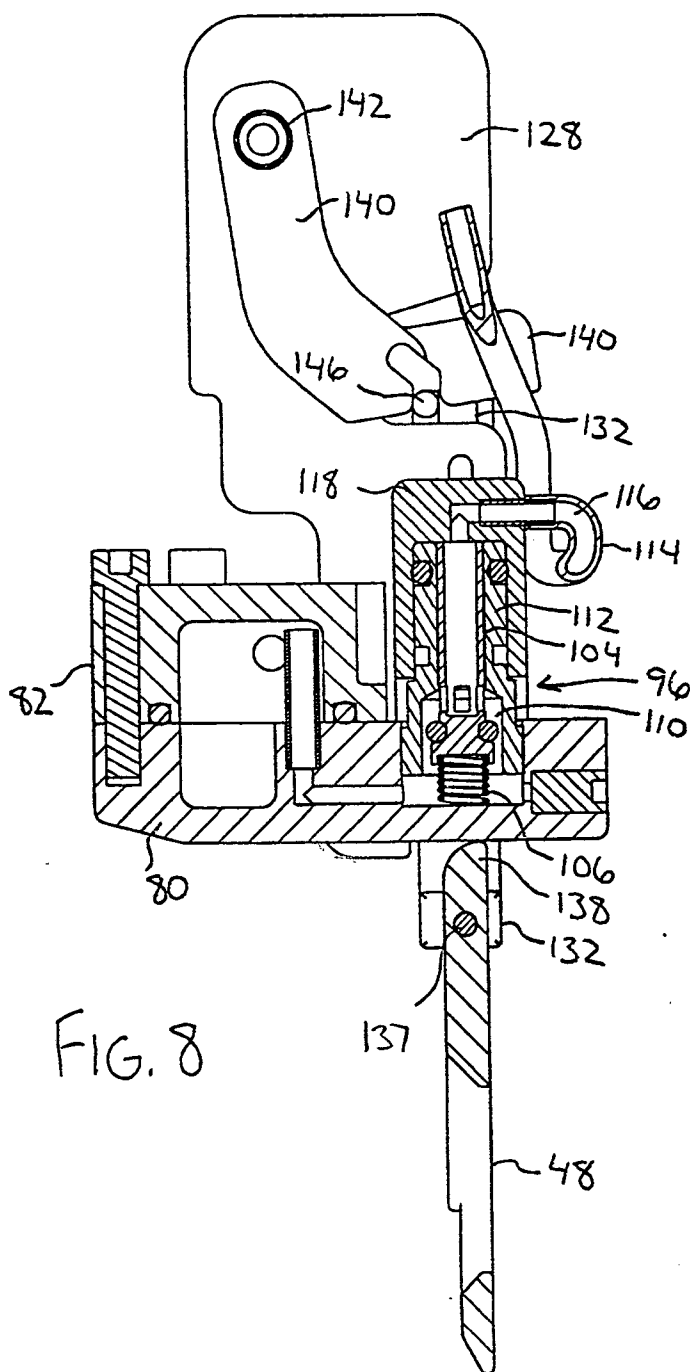


FIG. 8

FIG.10A

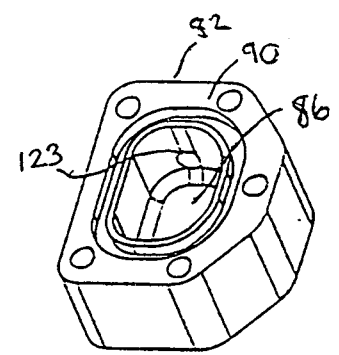
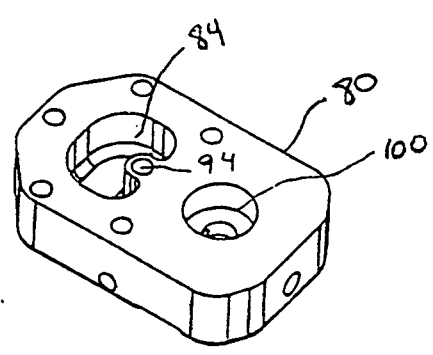


FIG.10B

FIG.9

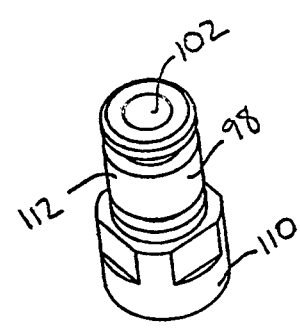


FIG.11

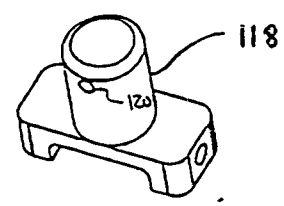


FIG.12

FIG. 10A

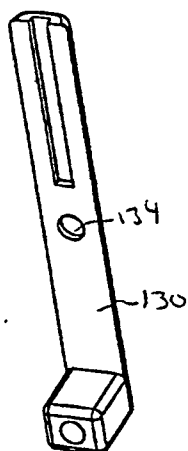


FIG. 13A

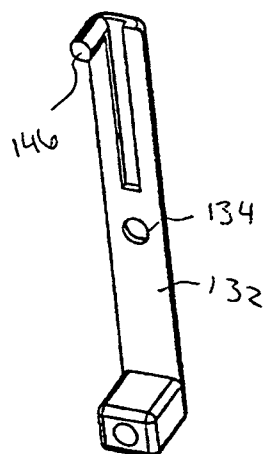


FIG. 13B

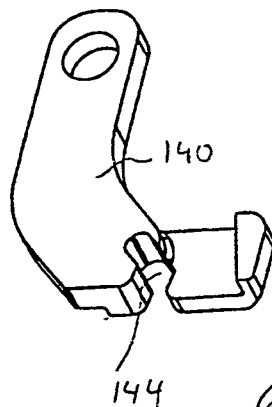


FIG. 14A

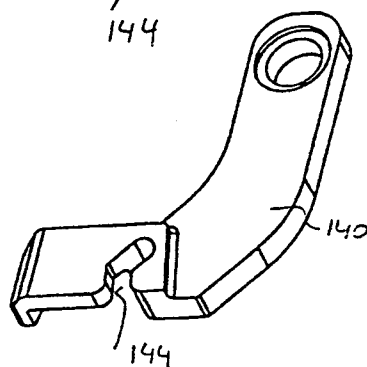
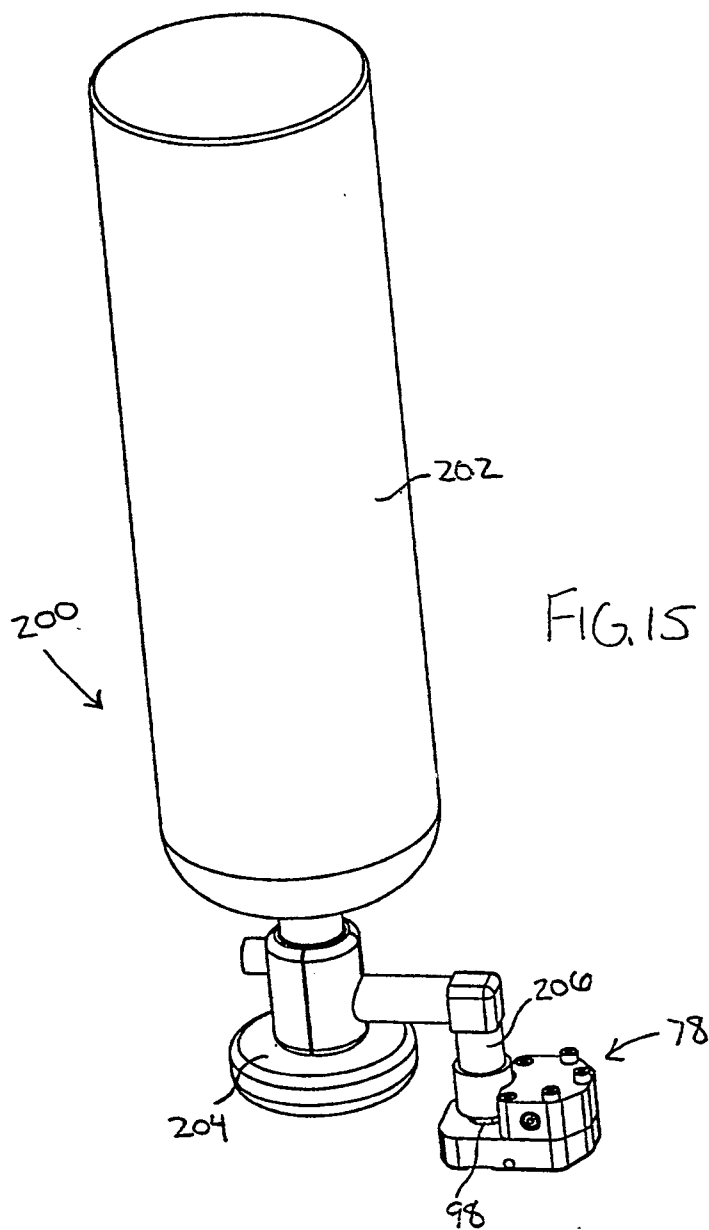


FIG. 14B

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TRADUCCION: Se traduce la carátula y las reivindicaciones anexas a una copia de la solicitud de patente estadounidense N° 60/136,518, escrita en inglés. Para el invento denominado: **APARATOS Y METODOS MEJORADOS DE AEROSOLIZACION**

LOS ESTADOS UNIDOS DE AMERICA
A TODOS AQUELLOS A QUIENES LA PRESENTE CONCIERNE
DEPARTAMENTO DE COMERCIO DE LOS ESTADOS UNIDOS

Oficina de Patentes y Marcas de los Estados Unidos

11 de mayo de 2000

LA PRESENTE ES PARA CERTIFICAR QUE LA ANEXA A ESTA ES UNA COPIA FIEL DE LOS REGISTROS DE LA OFICINA DE PATENTES Y MARCAS DE LOS ESTADOS UNIDOS DE LOS DOCUMENTOS DE LA SOLICITUD DE PATENTE IDENTIFICADA MAS ADELANTE QUE SATISFACEN LOS REQUISITOS PARA QUE SE OTORQUE UNA FECHA DE PRIORIDAD BAJO 35 USC 111.

SOLICITUD NUMERO: 60/136,518

FECHA DE PRESENTACION: 28 DE MAYO DE 1999

Por la Autoridad del
COMISIONADO DE PATENTES Y MARCAS

(RUBRICA)

P. SWAIN

Funcionario Certificador.

SELLO EN RELIEVE

REIVINDICACIONES

1. Un método para aerosolizar una formulación farmacéutica contenida dentro de un receptáculo, que comprende:

proporcionar una cantidad medida de un gas presurizado que estaba previamente en equilibrio con un líquido;

liberar el gas medido para crear una corriente de gas de alta presión; y

hacer fluir la corriente de gas de alta presión a través a través de un mecanismo de aerosolización para extraer la formulación farmacéutica desde el receptáculo y dispersar la formulación farmacéutica dentro de la corriente de gas para formar un aerosol.

2. Un método como en la reivindicación 1, en donde el gas presurizado comprende HFC.

3. Un método como en la reivindicación 1, en donde la formulación farmacéutica se encuentra en forma de polvo seco.

4. Un método como en la reivindicación 1, en donde la formulación farmacéutica se encuentra en forma líquida.

5. Un método como en la reivindicación 1, que comprende además abrir una válvula para liberar el gas presurizado.

6. Un método para aerosolizar una formulación farmacéutica contenida dentro de un réceptáculo, el método comprende:

proporcionar un propulsante que tiene una porción de fase líquida y una porción de fase de vapor;

medir una cantidad de la porción de fase de vapor;

liberar la cantidad medida para formar una corriente de gas de alta presión;

hacer fluir la corriente de gas de alta presión a través de un mecanismo de aerosolización para extraer la formulación farmacéutica del receptáculo y dispersar la formulación farmacéutica dentro de la corriente del gas para formar un aerosol.

7. Un método como en la reivindicación 6, en donde el propulsante comprende HFC.

8. Un método como en la reivindicación 6, en donde el propulsante comprende CO₂.

9. Un método como en la reivindicación 7, en donde la cantidad medida tiene un volumen en el rango que va de 0,5 ml a aproximadamente 2 ml y se presuriza a una presión en el rango de aproximadamente 40 psig a aproximadamente 120 psig.

10. Un método como en la reivindicación 6, que comprende además abrir una válvula para liberar la cantidad medida.

11. Un método como en la reivindicación 6, en donde la etapa de medición comprende permitirle a la porción de fase de vapor fluir dentro de la cámara de medición y cerrar la cámara de medición después de que la cámara de medición se llena.

12. Un aparato para aerosolizar la formulación farmacéutica mantenida dentro del receptáculo, el aparato comprende:

un reservorio que se adapta para mantener un propulsante que tiene en equilibrio una porción de fase líquida y una porción de fase de vapor;

una cámara de medición que se adapta para recibir una cantidad de la porción de fase de vapor del propulsante;

Una válvula de liberación que se ata para liberar el propulsante medido para formar una corriente de gas de alta presión;

Un mecanismo de aerosolización que se adapta para recibir una corriente de gas de alta presión, para extraer la formulación farmacéutica proveniente del receptáculo y para dispersar el polvo dentro de la corriente de gas para formar un aerosol.

13. Un aparato como la reivindicación 12 que comprende además una válvula de medición dispuesta entre reservorio y la cámara de medición, en donde la válvula de medición se mueve a una posición abierta para permitirle a la porción de fase de vapor del propulsante entrar a la cámara de medición.

14. Un aparato como la reivindicación 13, que comprende además una palanca que es movable para cerrar la válvula de liberación y abrir la válvula de medición.

15. Un aparato como en la reivindicación 12, que comprende además un botón de fuego para abrir la válvula de liberación.

16. Un aparato como en la reivindicación 12, que comprende además una carcasa para mantener el reservorio, la cámara de medición y el mecanismo de aerosolización.

17. Un aparato como en la reivindicación 16, que comprende además una cámara de captura operablemente acoplada a la carcasa, en donde la cámara de captura incluye una pieza de boca.

18. Un aparato como en la reivindicación 13, en donde la válvula de medición comprende una válvula de disco con movimiento vertical que se empuja para mover la válvula a la posición abierta.

19. Un sistema para aerosolizar una formulación farmacéutica, el sistema comprende:

una fuente de propulsante que incluye una porción de fase de vapor en equilibrio con una porción de fase líquida;

una cámara de medición dispuesta para medir volúmenes de la porción de fase de vapor del propulsante;

una válvula de liberación para liberar el propulsante medido para formar una corriente de gas de alta presión;

un mecanismo de aerosolización dispuesto para recibir la corriente del gas de alta presión, para extraer el polvo proveniente del receptáculo y para dispersar la formulación farmacéutica dentro de la corriente de gas para formar un aerosol.

20. Un sistema como la reivindicación 19, que comprende además una válvula de medición dispuesta entre la fuente del propulsante y la cámara de medición, en donde la válvula de medición se mueve a una posición abierta para permitirle al propulsante de la fase de vapor moverse hacia dentro de la cámara de medición.

21. Un sistema como en la reivindicación 20, que comprende además una palanca que es movible para cerrar

la válvula de liberación y para abrir la válvula de medición.

22. Un sistema como en la reivindicación 19, que comprende además un motor de fuego que es operable para abrir la válvula de liberación.

23. Un sistema como en la reivindicación 19, que comprende además una carcasa que mantiene la fuente de propulsante, la cámara de medición y el mecanismo de aerosolización, y una cámara de captura que tiene una pieza de boca operablemente acoplada a la carcasa.

24. Un sistema para aerosolizar una formulación farmacéutica, dicho sistema comprende:

una fuente de propulsante que incluye una porción de fase de vapor en equilibrio con una porción de fase líquida;

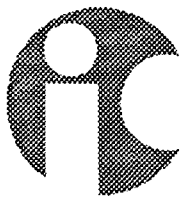
una formulación farmacéutica dispersada dentro de al menos una porción de la porción de fase de vapor del propulsante que ha sido separada de la porción de fase líquida.

APARATOS Y METODOS MEJORADOS DE AEROSOLIZACION

ANTECEDENTES DE LA INVENCION

La presente invención se relaciona en forma general con el campo del suministro de droga, y en particular con el suministro pulmonar de drogas. Más específicamente, la invención se relaciona con la aerosolización de formulaciones farmacéuticas para la inhalación por un paciente.

El suministro efectivo de drogas a un paciente es un aspecto crítico de cualquier terapia de drogas exitosa. Existen varias rutas de suministro, y cada una tiene sus propias ventajas y desventajas. El suministro oral de drogas en píldoras, cápsulas, elixires, y similares, es quizás el método más conveniente, pero muchas drogas se degradan en el tracto digestivo antes de que ellas puedan ser absorbidas. Tal degradación es un problema particular con las drogas modernas de proteína que se degradan rápidamente por las enzimas proteolíticas en el tracto digestivo. La inyección subcutánea es frecuentemente una ruta efectiva para el suministro sistémico de drogas que



2do. EXAMEN DE FORMA PATENTE DE INVENCION

Folio 129

Radicación no 00-38727

Concepto Técnico no 1710

Clasificación Internacional de Patentes: B65D 83/14, A61M 15/00

Practicado el examen de acuerdo a lo establecido en el Art. 38 de la Decisión 486 se determinó que esta solicitud de **Patente de Invención**:

- SI ☒ NO ☐ Contiene un resumen de la invención (Art.26-e).
- SI ☒ NO ☐ Contiene el título o nombre de la invención (Art.27-d).
- SI ☒ NO ☐ Contiene la descripción de la invención (Art.26-b).
- SI ☒ NO ☐ NO ES APLICABLE ☐ Contiene los dibujos necesarios para comprender la invención (Art.26-d)
- SI ☒ NO ☐ Contiene una o más reivindicaciones. (Art.26-c).
- SI ☐ NO ☐ NO ES APLICABLE ☒ Han sido desarrollados los productos y/o procedimientos a partir de recursos genéticos o de sus productos derivados de los que cualquiera de los Países Miembros es país de origen. (Art. 26- h).
- SI ☐ NO ☐ NO ES APLICABLE ☒ Los productos y/o procedimientos han sido obtenidos o desarrollados a partir de los conocimientos tradicionales de las comunidades indígenas, afroamericanas, o locales de los países miembros, de acuerdo a lo establecido en la Decisión 391 y sus modificaciones y reglamentaciones vigentes.(Art.26-i).
- SI ☐ NO ☒ La invención se refiere a un producto relativo a un material biológico. (Art.26-J).
- SI ☒ NO ☐ NO ES APLICABLE ☐ La prioridad coincide con la materia reivindicada.
- SI ☒ NO ☐ **CUMPLE LOS REQUISITOS TÉCNICOS**

OBSERVACIONES Y OTROS: SI ☐ NO ☐ ver hoja(s) anexa(s)

☐ NUEVO TITULO, FOLIO: 76

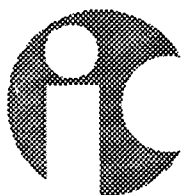
☐ NUEVO FORMATO DE SOLICITUD Folio:78

☐ NUEVA PAGINA 7 DESCRIPCION Folio: 79

☐ NUEVAS REIVINDICACIONES Folios: 121 - 127

EXAMINADOR TÉCNICO
202051

Bogotá D. C., noviembre 22 de 2002



2002-12-17
UIG

Folio 130

2020
Bogotá DC

Doctor(a)
EMILIO FERRERO
Apoderado(a)

REFERENCIA:	Radicación No.	00-38727
	Trámite	002
	Evento	001
	Actuación	416
	Oficio No.	2260
	Folios	001

Teniendo en cuenta que la solicitud de privilegio de patente que se tramita bajo el expediente indicado en la referencia, reúne los requisitos de forma establecidos en los artículos 26 y 27 de la Decisión 486 de la Comisión de la Comunidad Andina, y en las demás disposiciones vigentes sobre la materia, se envía el extracto de la solicitud a la Oficina de Comunicaciones para efecto de su publicación en la Gaceta de Propiedad Industrial, conforme lo establece el artículo 40 de la misma Decisión.

Cúmplase
Dado en Bogotá DC,

57.1.2003


ALIX CARMENZA CÉSPEDES DE VERGEL
Jefe de la División de Nuevas Creaciones

202051

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DIVISION DE NUEVAS CREACIONES

RADICACIÓN EXPEDIENTE No **00-38727**

EL EXTRACTO DE LA PRESENTE SOLICITUD DE **PATENTE DE INVENCION**
FUE PUBLICADO EN LA GACETA DE PROPIEDAD INDUSTRIAL No **527**, DE
FECHA **30 DE ABRIL DE 2003**, EL TERMINO HABIL SEÑALADO POR LA LEY
EXPIRA EL **29 DE JULIO DE 2003**.

NOTA: Dentro del plazo de los seis (6) meses siguientes a partir de la fecha de publicación en la gaceta de Propiedad Industrial, deberá solicitar y cancelar el examen de patentabilidad so pena de que la solicitud caiga en abandono, de acuerdo con lo establecido en el Artículo 44 de la Decisión 486/2000.

SE PRESENTARON OPOSICIONES POR PARTE DE TERCEROS.

SI _____

NO _____ **X** _____

Bogotá D.C., Octubre de 2003