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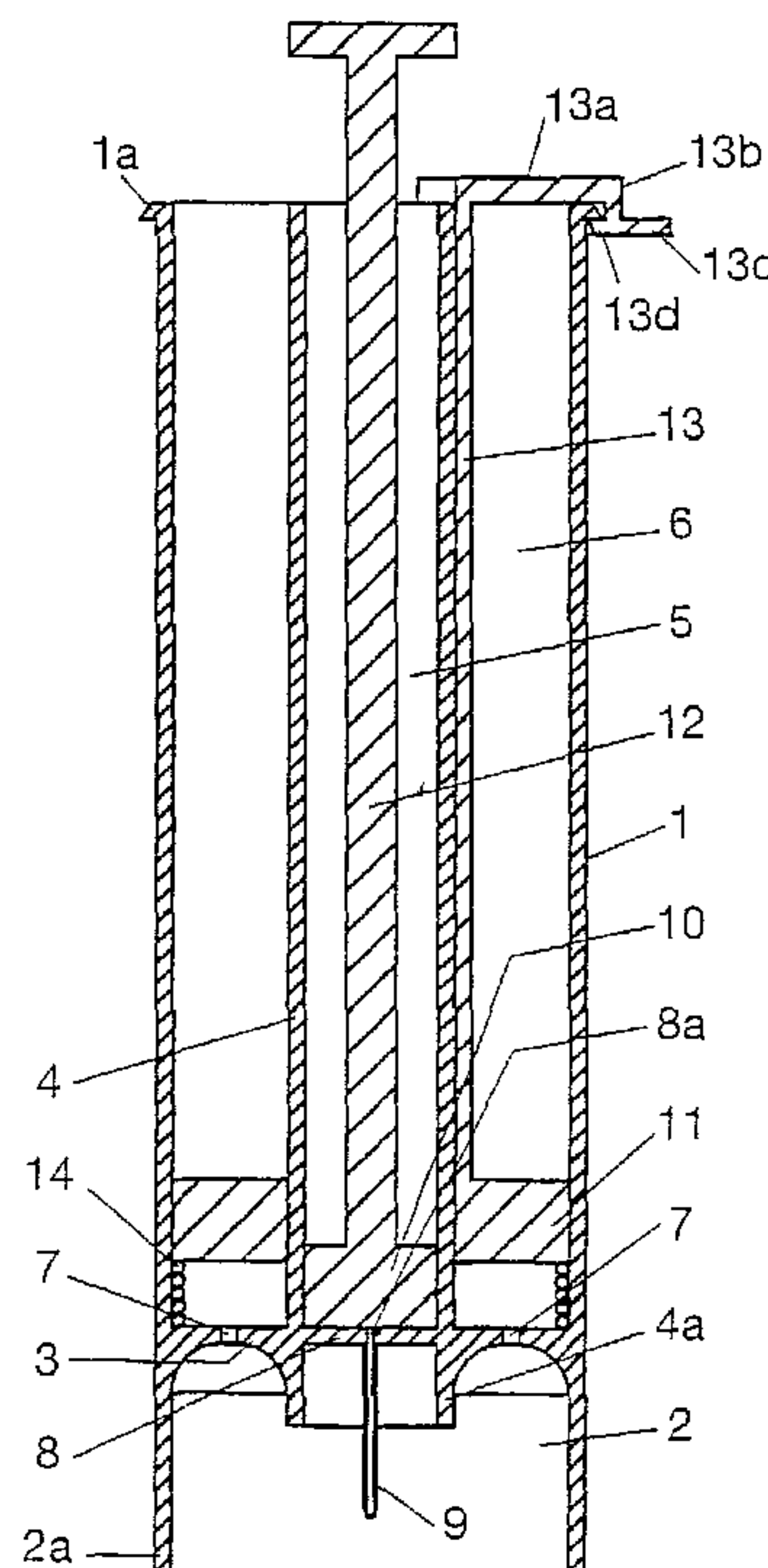
(71) Demandeur/Applicant:
ARES TRADING S.A., CH

(72) Inventeurs/Inventors:
BERGERON, LUC, CH;
BRIOSCHI, ELENA, CH;
JACQUET, REMY, CH;
MOULIN, JEROME, CH;
SOLDINI, LAURENT, CH

(74) Agent: KIRBY EADES GALE BAKER

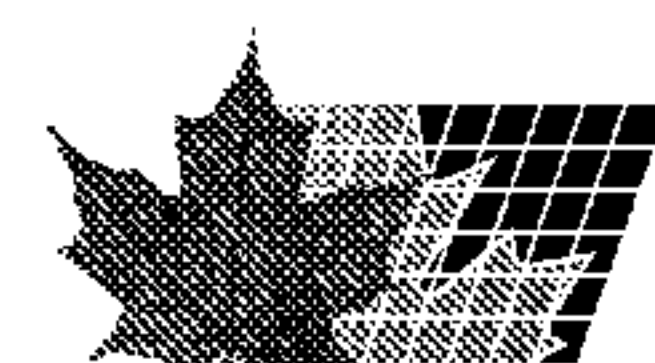
(54) Titre : DISPOSITIF D'INJECTION

(54) Title: INJECTION DEVICE



(57) **Abrégé/Abstract:**

The invention provides an injection device comprising an open cavity (2), an injection needle (9) projecting from the bottom of this cavity (2), a first leakproof compartment (6) delimited by a first section of a mobile barrier (11), first devices (13) for displacing the said first section of a mobile barrier (11) to vary the volume of the compartment, at least one passage (7) communicating between the said first leakproof compartment (6) and the said cavity (2), a second leakproof compartment (5) for a product to be injected, delimited by a second section of a mobile barrier (10), and second devices (12) for displacing the said second section of a mobile barrier (10) to vary the volume of the compartment. The said first and second leakproof compartments (5, 6) extend longitudinally within a tubular body (1). The cavity (2) is located at one end of this tubular body, while the said first and second devices (12, 13) for displacing the said first and second sections of a mobile barrier (10, 11) are accessible at the other end of the said tubular body (1).



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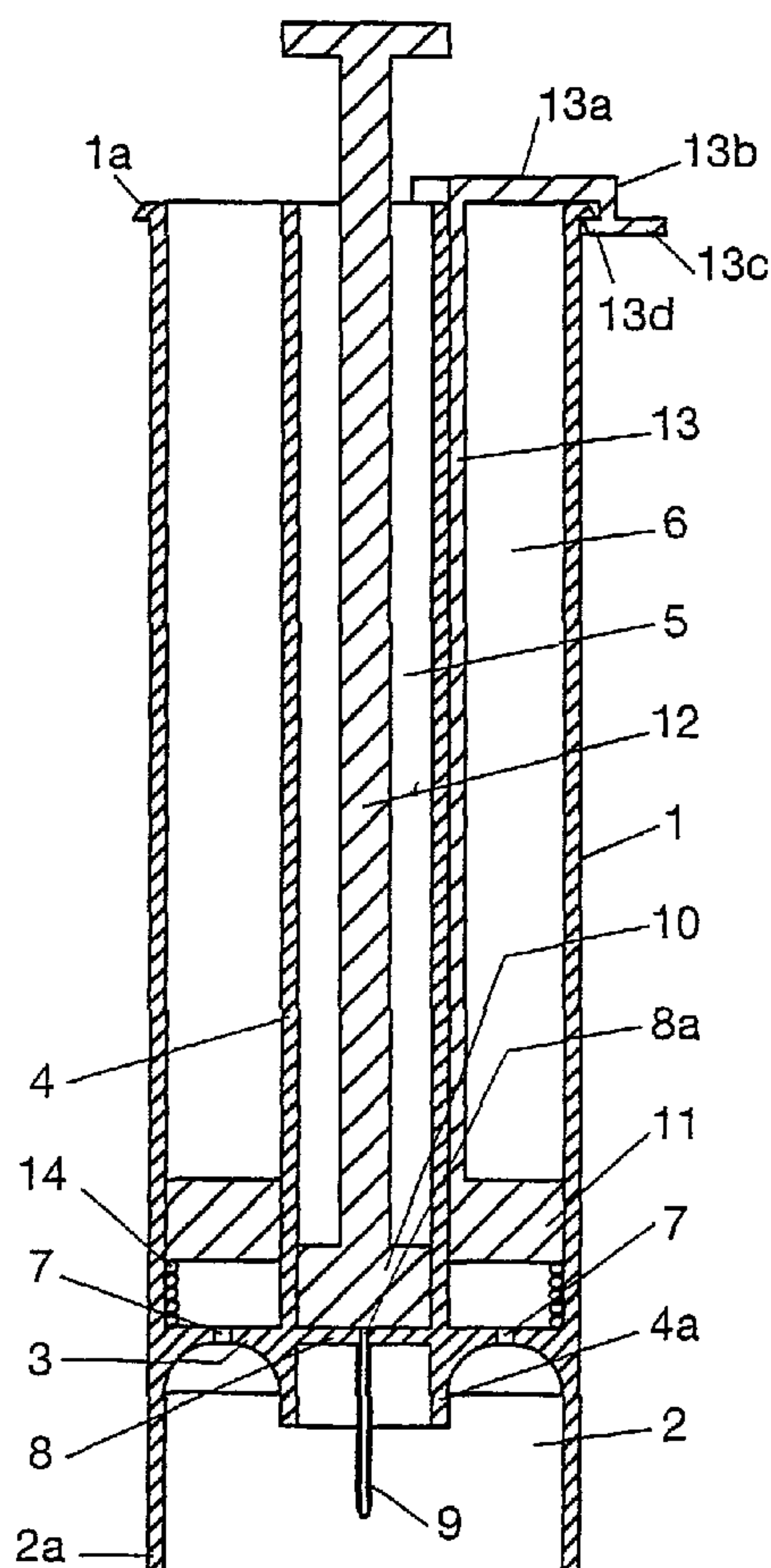
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00810414.3 15 May 2000 (15.05.2000) EP(71) Applicant (for all designated States except US): **ARES TRADING S.A.** [CH/CH]; Château de Vaumarcus, CH-2028 Vaumarcus (CH).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **BERGERON, Luc**[CH/CH]; Chemin de la Condémine, CH-1034 Boussens (CH). **BRIOSCHI, Elena** [CH/CH]; 34, Rue de Lyon, CH-1203 Geneva (CH). **JACQUET, Rémy** [CH/CH]; 7, Voisinage, CH-2400 Le Locle (CH). **MOULIN, Jérôme** [CH/CH]; 48, Grand-Rue, CH-1890 St.-maurice (CH). **SOLDINI, Laurent** [CH/CH]; 14bis Druey, CH-1018 Lausanne (CH).(74) Agents: **SAVOYE, Jean-Paul** et al.; Moinas & Savoye S.A., 42, Rue Plantamour, CH-1201 Geneva (CH).(81) Designated States (*national*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

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(54) Title: INJECTION DEVICE



(57) Abstract: The invention provides an injection device comprising an open cavity (2), an injection needle (9) projecting from the bottom of this cavity (2), a first leakproof compartment (6) delimited by a first section of a mobile barrier (11), first devices (13) for displacing the said first section of a mobile barrier (11) to vary the volume of the compartment, at least one passage (7) communicating between the said first leakproof compartment (6) and the said cavity (2), a second leakproof compartment (5) for a product to be injected, delimited by a second section of a mobile barrier (10), and second devices (12) for displacing the said second section of a mobile barrier (10) to vary the volume of the compartment. The said first and second leakproof compartments (5, 6) extend longitudinally within a tubular body (1). The cavity (2) is located at one end of this tubular body, while the said first and second devices (12, 13) for displacing the said first and second sections of a mobile barrier (10, 11) are accessible at the other end of the said tubular body (1).

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INJECTION DEVICE

FIELD OF THE INVENTION

The invention relates to an injection device.

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BACKGROUND OF THE INVENTION

An injection device using suction has been proposed in US 4'114'619. The proposed device has an open cavity with an injection needle projecting from the bottom of the cavity. The open cavity is applied to the skin. When the opening of the cavity is placed correctly sensors activate a suction piston which creates a vacuum in the cavity. This vacuum draws the skin towards the bottom of the cavity and then causes penetration by the injection needle projecting from the bottom of the cavity. A second detector, consisting of a pressure transducer, activates the piston of the injection syringe.

The purpose of the device of US 4,114,619 is to achieve complete automation of the injection process once the opening of the cavity is correctly placed in contact with this tissue. In consequence the design of the device for executing this automatic process is relatively complex.

SUMMARY OF THE INVENTION

It is an object of the present invention to provide an injection device of simplified design.

It is a further object of the invention to provide an injection device that can be simply manufactured, at low cost.

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It is a further object of the invention to provide an injection device that is easily portable.

The invention exploits the concept which consists of making the cutaneous tissue penetrate inside the needle rather than the inverse, as in traditional injection syringes. The invention provides a device of the simplest possible design so
5 that it can be manufactured at low cost and be simple and easy to use. This result is achieved by creating in the injection area, as in the device mentioned above, a vacuum by the use of an open cavity whose opening is applied to the skin so as to draw it up against the injection needle projecting from the
10 bottom of the cavity.

The invention provides an injection device having an open cavity whose opening is delimited by an annular rim intended for application to the skin of a subject under treatment, an injection needle projecting from the bottom of the cavity, its
15 forward end being set back in respect of the annular rim, a first, leakproof compartment delimited by a first mobile barrier, means for displacing the first mobile barrier so as to vary the volume of the first compartment, at least one communicating passage between the first leakproof compartment
20 and the cavity, a second leakproof compartment for a product to be injected, delimited by a second mobile barrier, a second means for displacing the said second section of a mobile barrier so as to vary the volume of the second leakproof compartment, and an opening to enable the second leakproof
25 compartment to communicate with the said injection needle.

DETAILED DESCRIPTION OF THE INVENTION

In the case of self-injection the person performing the injection does not himself have to make the needle penetrate
30 the cutaneous tissue, which is a notable improvement from the psychological standpoint. The drawing up of the skin reduces the perception of pain caused by the penetration of the needle.

The depth of needle penetration is perfectly regulated by the length to which the needle projects from the bottom of the cavity, a safety feature likely to be reassuring, particularly to the person who must perform self-injection.

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The attached drawings illustrate, schematically and by way of example, one embodiment of the injection device to which the invention refers.

Figure 1 is a section along the axis of this embodiment
10 showing the device in a first position;

Figure 2 is an enlarged view of a detail of Figure 1;

Figure 3 is a view similar to Figure 1 showing the device in a second position;

Figure 4 is a view similar to Figure 1 showing the device
15 in a third position;

Figure 5 is a view similar to Figure 1 showing the device in a fourth position.

The device illustrated has a tubular cylindrical body 1
20 having at one end an open cavity 2 the bottom of which is delimited by an annular wall 3 extending around a cylindrical wall 4 centred on the longitudinal axis of the tubular body 1. This cylindrical wall 4 extends from the open end of the tubular body 1, opposite to the open cavity 2, into the
25 interior of the latter, in the bottom of which a portion 4a of this cylindrical wall 4 projects.

The cylindrical wall 4 divides the interior space of the tubular body 1 adjoining the annular wall 3 opposite the open
30 cavity 2 into two elongated compartments, of which one, 5, has an annular section forming a cylinder concentric with the axis of the tubular body 1, the other, 6, having an annular section

concentric with the cylindrical compartment 5 which it surrounds.

Openings 7 are made through the wall 3 to place the compartment 6 in communication with the open cavity 2. The cylindrical compartment 5 is closed at the level of the wall 3 by a barrier 8 which forms one piece with an injection needle 9 extending inside the open cavity 2, its forward end being set back from the rim 2a bounding the opening of the cavity 2. If the barrier 8 is not directly formed by a single piece moulded together with the tubular body 1 it can be soldered or glued on. Its centre has a perforation 8a to place the channel of the injection needle in communication with the cylindrical compartment 5.

15

Each of the compartments 5, 6 encloses a piston, 10, 11 respectively. The piston 10 forms one piece with a plunger 12, while the piston 11 forms one piece with a plunger 13 which in section takes the form of an annular sector with its concave surface fitting tightly against the exterior surface of the cylindrical wall 4. The end of the plunger 13 located outside the tubular body 1 forms one piece with a circular sector 13a the exterior extremity of which terminates in a lip 13b which in turn forms one piece with a lug 13c (Figure 2).

25

A locking element 13d is located within the lip 13b and terminates in a bevel 13e. The end of the wall of the tubular body 1 consists of an annular lip 1a delimited externally by a bevel 1b. As is shown in Figure 2 the two bevels 1b and 13e are designed as detents, that is to say, they are designed to

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ensure, when locked together, that the plunger 13 and the piston 11 stay in position along the axis of the tubular body 1 when subjected to an axial force which tends to push the plunger 13 out of the tubular body 1. On the other hand, the bevels 1b and 13e allow the lip 13d to be locked onto the lip 1a of the tubular body 1 when the plunger 13 and the piston 11 are displaced along their axis from the open end of the compartment 6 towards its closed end, creating a component of centripetal force which can displace the lip 13b when the bevel 13e pushes against the bevel of the lip 1a to allow the lip 13d to pass the lip 1a of the tubular body 1, until the lip 13b locks under the lip 1a. This, of course, presupposes adequate elasticity in the circular sector 13a. For this purpose it is advantageous to use a material which can be deformed elastically.

A helicoidal spring 14 is placed in the compartment 6 between the bottom of the compartment delimited by the annular wall 3 and the piston 11. When the piston 11 compresses the spring 14 against the bottom 3 of the compartment 6 the circular sector 13a forming one piece with the plunger 13 of the piston 11 is locked to the lip 1a of the tubular body 1a [sic] as explained above. When not in use the various elements of the device described are in the position illustrated in Figure 1.

To perform an injection using this device the required amount of the product to be injected is drawn in, as in the case of a traditional syringe, by placing the forward end of the injection needle 9 in the appropriate product and drawing

in the desired dose of the product by displacing the piston 10 by means of the plunger 12, as is shown in Figure 3.

The opening 2a of the cavity 2 is then applied to the
5 surface of the skin (not shown) through which the injection is to be made. The piston 11 is freed by unlocking the lip 13d of the circular sector 13a from the lip 1a of the tubular body 1 by deforming the lug 13c upwards. The freed piston 11 allows the pressure developed by the release of the spring 14 to push
10 it towards the open end of the compartment 6. Since the cavity 2 is closed by the skin to which its rim 2a is applied the displacing of the piston 11 creates a vacuum within the compartment 6 and the cavity 2 which communicates with it via the openings 7. This vacuum has the effect of deforming the
15 portion of skin which is inside the rim 2a of the cavity 2 in the direction of the bottom of the cavity 2, thus forcing the end of the needle through the skin resting against the rim of the portion 4a of the cylindrical wall 4 projecting from the bottom of the cavity 2 concentrically with the injection needle
20 9. The various elements of the device are then in the position illustrated in Figure 4, ready for the injection proper.

For this purpose the plunger 12 is then pushed back within the compartment 5 until it rests against the barrier 8
25 delimiting the bottom of the compartment 5, as shown in Figure 5. The device is withdrawn simply by pushing back the plunger 13 forming one piece with the piston 11, thus once more raising the pressure and then making it possible to separate the opening 2a of the cavity 2 from the skin which is no longer
30 drawn in by the vacuum in the cavity 2. The locking element 13d

is then once again locked under the lip 1a of the cylindrical body 1 to hold the piston 11 in the position of Figure 1 against the pressure exerted upon it by the spring 14 which is once more compressed at the bottom of the annular compartment 5 6.

The device of the invention may be made of plastic, metal or glass. It is preferred that the pistons and cylindrical body be made of plastic.

10

In a preferred embodiment, the device is designed for a single use.

In a further preferred embodiment, the needle length can 15 be varied, thus allowing control of the depth of penetration.

CLAIMS

1. A handheld injection device comprising:
an open cavity (2) having a rim (2a), the rim being capable of
5 forming an airtight seal when applied to the skin of a patient;
a first piston (11) in communication with the open cavity (2)
for creating suction therein;
a needle (9) projecting into the open cavity (2), but not
projecting beyond the rim (2a);
10 a second piston (10) for containing a product to be injected,
the second piston being in communication with the needle (9);
wherein withdrawal of the first piston (11) creates suction in
the open cavity (2a), causing the patient's skin to be sucked
into the cavity (2) and to be pierced by the needle (9), and
15 actuation of the second piston (10) causes the product to be
injected;
characterised in that the first (11) and second (10) pistons
are located within a cylindrical body (1), the cylindrical body
having the needle (9) at one end and actuating means (13) for
20 the pistons (10, 11) at another end.

2. A device according to claim 1, wherein said cylindrical
body (1) comprises a first central cylinder (5) surrounded by a
second annular cylinder (6) in which are located respectively
25 said first 11, and second pistons.

3. An injection device comprising
an open cavity (2) the opening of which is delimited by an
annular rim (2a) intended to be applied to the skin of a
30 subject under treatment,

an injection needle (9) projecting from the bottom of the cavity (2) its forward end being set back in relation to the said annular rim (2a),
a first leakproof compartment (6) delimited by a first section
5 of a mobile barrier (11),
first means (13) for displacing the said first section of a mobile barrier (11) in order to vary the volume of the said first compartment (6),
at least one passage (7) communicating between the first
10 leakproof compartment (6) and the said cavity (2),
a second leakproof compartment (5) for a product to be injected, delimited by a second section of a mobile barrier (10),
second means (12) for displacing the said second section of a
15 mobile barrier (10) in order to vary the volume of the second leakproof compartment (5) and
an opening (8a) communicating between the second leakproof compartment (5) and the said injection needle (9),
characterized in that
20 the said first and second leakproof compartments (6, 5) extend longitudinally within a tubular body (1), that the said cavity (2) is located at one end of the tubular body (1), while the said first and second means (13, 12) for displacing the said first and second sections of a mobile barrier (11, 10)
25 respectively of the said first and second leakproof compartments (6, 5) are accessible at the other end of the said tubular body (1).

4. A Device according to Claim 3, characterized in that
30 the said first leakproof compartment (6) has an annular section

surrounding the said second leakproof compartment (5).

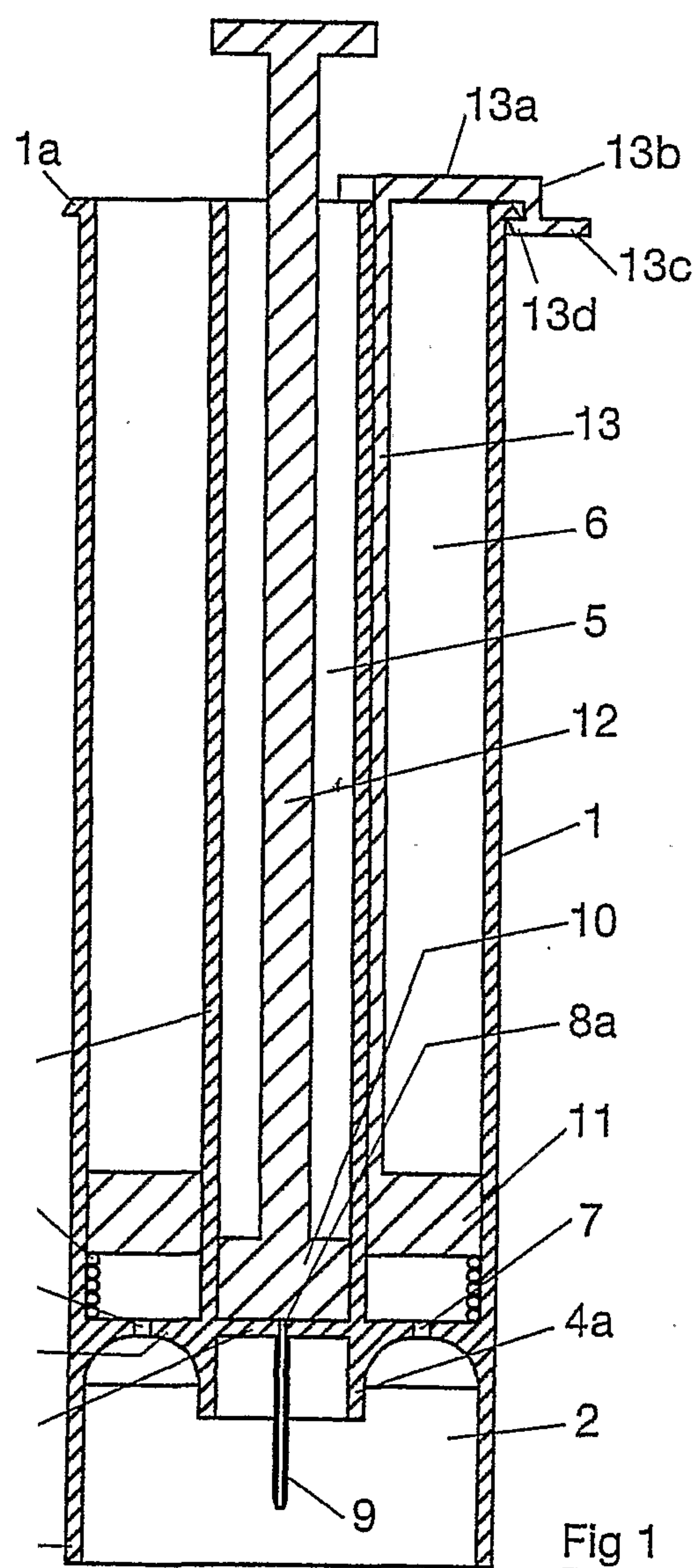
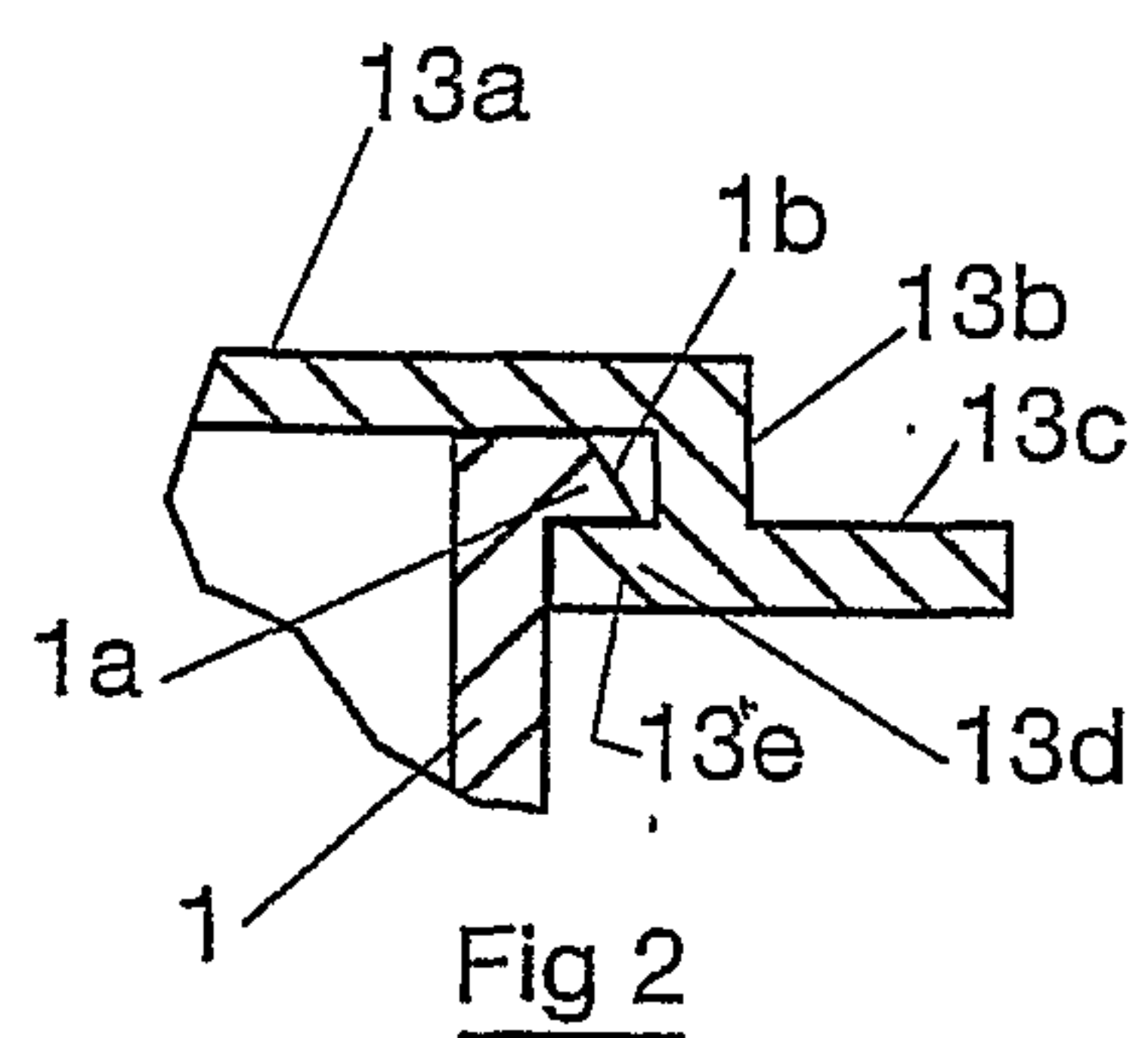
5. A Device according to claim 3 or 4, characterized in that it includes elastic pressure devices (14) acting on the said first section of a mobile barrier (11) to increase the volume of the said first compartment (6) and locking elements (1a, 13d) to hold the said first section of a mobile barrier (11) in a position in which the said elastic devices (14) are compressed, equivalent to the minimum volume of the said first compartment (6).

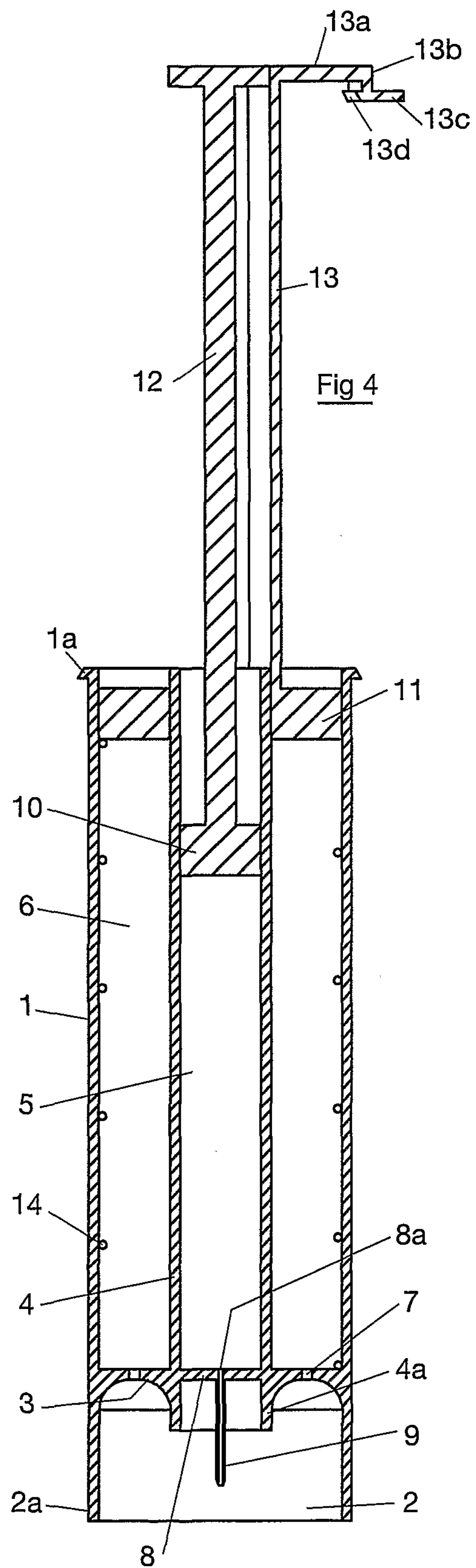
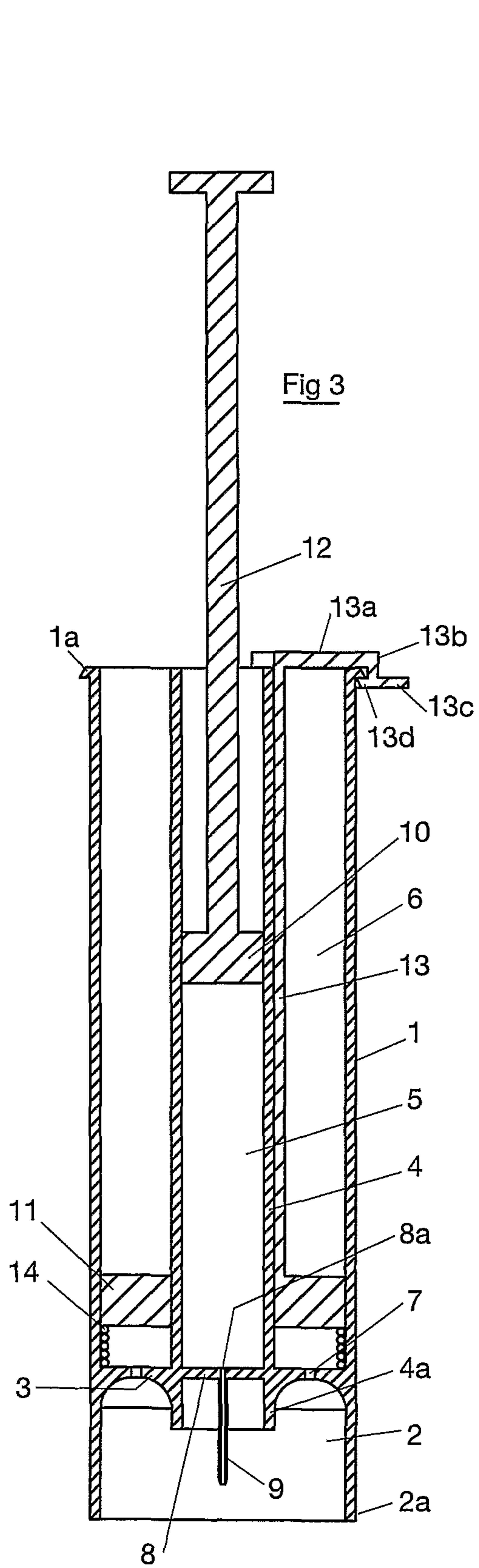
6. A Device according to Claim 5, characterized in that the said locking elements (1a, 13d) form one piece with both the said first devices (13) for displacing the said first section of a mobile barrier (11) and with the said tubular body (1).

7. A Device according to any one of claims 3 to 6, characterized in that each of the said first and second sections of a mobile barrier (11, 10) is formed by a piston sliding within its respective compartment (6, 5).

8. A Device according to any one of claims 3 to 7, characterized in that an annular wall (4a) extends concentrically around the portion of the said injection needle (9) which projects at the bottom of the said cavity (2), at approximately half the distance between the bottom and the forward end of the said injection needle (9).

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