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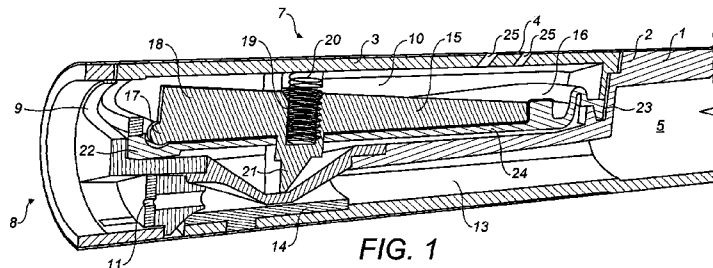
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(54) Title: AN INHALER



(57) Abstract: An inhaler comprising a reservoir (5) of an inhalable composition with an outlet (11) at one end (8) through which the inhalable composition is discharged. A non-metered breath-activated valve (7) is provided between the one end (8) and the reservoir (5), the breath-activated valve (7) comprising a flow path (13) extending from the reservoir (5) to the outlet end (8). At least a portion of the flow path is a deformable tube (14). A clamping member (21) pinches the deformable tube (14) closed when no suction force is applied to the device and releases the tube (14) to open the flow path when suction is applied at the outlet (11), to provide uninterrupted flow from the reservoir (5) to the outlet (11).



WO 2011/015825 A1

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AN INHALER

The present invention relates to an inhaler.

5 The invention has been specifically designed for a simulated cigarette device having a generally cigarette-shaped body. However, the invention relates to a development of the outlet valve for such a device which has broader applications in the field of inhalers, for example,
10 medicinal inhalers for oral drug delivery such as asthma inhalers.

 In the field of cigarette replacements, there have been a number of proposals to create a simulated cigarette. Such
15 a device has a number of advantages over traditional nicotine replacement therapies such as patches and gum in that they recreate the physical act of smoking which is psychologically important to a smoker, and also are able to deliver nicotine as a dose which more closely replicates the
20 pharmacokinetic effects of a cigarette that persistent smokers desire. Thus, a smoker is able to obtain the "hit" that is familiar from a cigarette, rather than having to deal with the slow release from a patch or gum which does not produce such a hit which leads to unpredictable dosings
25 and poor craving scores and cessation rates.

 A simulated cigarette has a reservoir of inhalable composition and an outlet valve which can be triggered, for example, by pushing a button or biting on the end of a
30 cigarette. However, a preferred mechanism for opening the valve is to provide a breath-activated valve as this ensures

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that the cigarette will only dispense when the user sucks on the device in a manner of a normal cigarette.

WO 02/45783 discloses a drug dispenser having a valve
5 arrangement suitable for dispensing a metered dose. This
takes the form of a flexible tube which is wrapped around a
valve element and has a pair of kinks. The portion of the
tube between the kinks is able to retain a metered dose of
the drug to be dispensed. When a user sucks on the
10 dispenser, the valve element moves opening the downstream
kink in order to dispense the metered dose. On exhalation,
the valve moves in the opposite direction closing the
downstream kink and opening the upstream kink to allow the
portion of the tube between the kinks to be re-charged with
15 the drug. This arrangement only ever allows a user to
inhale one metered dose no matter how long they suck on the
device.

US 4,393,884 discloses one such device which has a
20 large resilient "tongue" with a flow path passing through
it. This tongue is biased into a first position in which it
is out of alignment with the outlet of the cigarette and can
be sucked into a second position in which it aligns with the
outlet of the cigarette to provide a flow path from the
25 reservoir to the outlet. Such a device is difficult to seal
in the first position. It will also require considerable
force in order to suck the tongue to the open position
against the action of a retaining spring and the relatively
large mass of the tongue means that it will be difficult to
30 return to the closed position meaning that dispensing will
continue after the suction has been removed.

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US 6,889,687 discloses a further example of a simulated cigarette with a breath-activated valve. This discloses a number of examples. One of these has a pair of magnets, one of which is retained by a flexible membrane which allows the magnet to moved apart when suction is applied to a device. This opens up a flow path from the reservoir. However, the mechanism is reasonably complex, has a tortuous flow path which is likely to impede delivery of the composition from the reservoir, and is "binary" in nature in the sense that the valve is either open or closed. It does not provide the ability for the user to regulate the amount of flow by varying the degree of suction. A second example is a spring-biased plunger which is moved axially to open up a passageway in a central rod. Such an axially movable plunger is undesirable in practice as it has been found that the level of suction required to overcome the spring biasing force is too high to be usable in practice. Also, the flow path in the open position is out of the rod, into the plunger and back into the rod so that it is again somewhat tortuous. The third example relies on a magnetic interface where the force on which is overcome by a system of vanes which rotate about the axis of the device, thereby moving along a cam surface to pull the magnetic valve element from its seat. Again, this suffers from problems of complexity, lack of control and a tortuous flow path.

WO 2009/001082 which is our own earlier application discloses two different breath-activated devices. The first of these has a pair of vane systems rotatable about an axis perpendicular to the main axis of the device to align an orifice with the outlet orifice from the reservoir enabling dispensing. The second of these has a pair of hinged flaps

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which are sucked down against the action of return springs in order to open the flow path. While this solves some of the problems in that it provides a simple mechanism and an axial flow path, the force required as suction to trigger
5 the device is comparatively large and the user, as a result, can exert less control over the system to release a variable dose, small or large according to the inhalation intake.

The present invention is aimed at providing an improved
10 breath-operated valve for an inhaler and, particularly, a simulated cigarette.

According to the present invention, there is provided an inhaler comprising:

- 15 a reservoir of an inhalable composition;
an outlet at one end through which the inhalable composition is discharged; and
a non-metered breath-activated valve between the one end and the reservoir, the breath-activated valve comprising
20 a flow path extending from the reservoir to the outlet end, at least a portion of the flow path being a deformable tube, and a clamping member which pinches the deformable tube closed when no suction force is applied to the device and releases the tube to open the flow path when suction is
25 applied at the outlet, to provide uninterrupted flow from the reservoir to the outlet.

By clamping a deformable tube, the present invention has a number of advantages. The clamping mechanism is
30 simple as all that is required is a movable component to pinch the tube which need not be complex. Further, as the flow path is along an opened tube, there is no need for a

- 5 -

tortuous flow which may otherwise be required to flow around an unseated valve component such as those in US 6,889,687. The non-metered valve which allows uninterrupted flow allows a user to regulate the dose that they receive from the
5 inhaler, as it will continue to dispense for as long as the user sucks.

The pinched tube arrangement lends itself well to an unmetered dosage. Also, while the device may be provided
10 with the clamping member which effectively "snaps" open to provide a "binary" device which is either open or closed, the pinched deformable tube is well suited to a device in which the degree of opening of the tube is proportional to the strength of the suction. Such an outlet valve which is
15 breath-operable and of varied resistance according to the inhalation profile it is actuated by, allows the user to self-titrate the inhaled dose in a continuous manner. This has applications firstly for smoking cessation inhaler devices where users want to self-titrate their dose of
20 nicotine in a manner that fits their need, but also in other area such a pain management, diabetes, asthma and COPD where self-administered doses are preferable via the oral route.

The tube could be clamped at several locations but is
25 preferably clamped at a single location as that is all that is necessary to fulfil its function. The flow path is preferably substantially straight to ensure a smooth flow.

As mentioned above, the inhaler may be a drug delivery
30 inhaler for any inhalable pharmaceutical composition. However, the inhaler is preferably a simulated cigarette device having a generally cigarette-shaped body. In this

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case, the inhalable composition preferably includes nicotine or a nicotine derivative or salt thereof. On the other hand, it may be a simulated cigarette which replicates the physical act of smoking without requiring nicotine in the composition. Alternatively, the composition may include patient controlled analgesics, anti-inflammatory, bronchodilators, anti-spasmodics, retro-virals or opiates.

The clamping member may be formed of any suitable device which can release the deformable tube when suction is applied to the outlet. This could, for example, be a combination of an electronic flow sensor and a solenoid to move the clamping member. However, preferably, the clamping member is a mechanical member which is moved by a pressure differential caused when suction is applied at the outlet.

The mechanical member may take the form of the pair of vanes of the second example of WO 2009/001082 where it is pinched between the vanes in the closed position and is released as the vanes pivot towards the direction of the air through-flow.

However, preferably, the clamping member is an elongate vane which is pivotally mounted at one end and extends longitudinally within the inhaler, the vane being pivotable between a first position in which the tube is closed and a second position in which it releases the tube to provide the flow path. There may be a single vane which clamps the deformable tube against a fixed part of the inhaler, or there may be a pair of such vanes between which the tube is clamped.

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Preferably, the or each vane is biased into the closed position and the inhaler has a flow path creating a pressure differential across the vane which overcomes the biasing force.

5

The or each vane may be exposed to a through-flow of air such that, as a user sucks on the outlet, the through-flow of air impinges on an outer surface of the vane to move it inwardly. However, preferably, the or each vane is supported by a flexible membrane. This flexible membrane creates a chamber in which the vane is housed and which is open to the outlet, but is otherwise sealed, with the face of the membrane on the opposite side to the vane being open to atmospheric pressure. When a user sucks on the outlet, they create a negative pressure within the chamber to move the vane and the membrane flexes to allow this movement. This provides a very efficient mechanism for conveying the suction force to the vane.

20

Preferably, the deformable tube is a nozzle which is also provided integrally with an outlet orifice, which is the narrowest part of the flow path.

25

In this case, preferably the nozzle has an outwardly projecting annular flange at its upstream end which fits within a downstream end of an adjacent part of the flow path.

30

Examples of inhalers in accordance with the present invention will now be described with reference to the accompanying drawings, in which:

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Fig. 1 is a section through a perspective view of a first inhaler in a closed position;

Fig. 2 is a similar view in the open position;

Fig. 3 is an exploded perspective view of the inhaler
5 of Figs. 1 and 2;

Fig. 4 is a cross-section through a perspective view of a second inhaler in the closed position;

Fig. 5 is a similar view in the open position; and

Fig. 6 is a perspective view from the outlet end of the
10 second example shown with the outer housing removed to show the pinch mechanism; and

Fig. 7 is a cross-section of an alternative arrangement of deformable tubular element.

15 The present invention relates to an improvement of the outlet valve for a breath-activated cigarette and only this aspect of the invention will be specifically described here. For details of the construction of the remainder of the cigarette device and its refill mechanism, reference is made
20 to WO 2009/001078.

The first example of an inhaler in accordance with the present invention is shown in Figs. 1 to 3.

25 The device has a housing 1 made up of a main chassis 2 and a closure element 3 as shown in Fig. 1. This is held in place by label 4. Within the housing, there is a reservoir 5 containing the inhalable composition. This is preferably pressurised but could also work with a non-pressurised
30 reservoir in combination with a Venturi nozzle to generate an enhanced suction force on the reservoir, or a non-pressurised reservoir containing a substance that is prone

- 9 -

to evaporating at room temperature. It may be refillable as described in WO 2009/001082 through the filling valve 6, or the device may be a single use device, or may be arranged so that the reservoir 5 is a replaceable component.

5

The breath-activated valve 7 is positioned between an outlet end 8 and the reservoir 5. The breath-activated valve is arranged so that, when a user sucks on the outlet end 8, the breath-activated valve 7 opens to allow the
10 inhalable composition from the reservoir 5 to be inhaled.

The housing at the outlet end has two orifices. The first of these is the suction orifice 9 which communicates with a chamber 10 as will be described in greater detail
15 below and the second is an outlet orifice 11 from which the inhalable composition dispensed is also described in more detail below. As is apparent from Fig. 3, the outlet orifice 11 is provided on a separate component 12.

20 An outlet path 13 is defined between the reservoir 5 and outlet orifice 11.

A portion of the outlet path 13 is provided by deformable tubular element 14. This tubular element is
25 moved between the closed position shown in Fig. 1 and the open position shown in Fig. 2 by a mechanism which will now be described.

This mechanism comprises a pivotally mounted vane 15
30 and a membrane 16. The pivotally mounted vane has a pivot 17 at the end closest to the outlet end 8 and a central reinforcing rib 18 running along its length and tapering

- 10 -

away from the outlet end. At around the midpoint, the vane 15 is provided with a recess 19 for receiving a spring 20 which biases it into the closed position shown in Fig. 1. Below the recess 19 is a jaw 21 having a triangular cross-section which is configured to apply the force provided from the vane 15 to the deformable tube 14 over a narrow area. The vane 15 is supported by the diaphragm 16 which is sealed to the housing at its ends 22, 23. This seals off the chamber 10 other than to the suction orifice 9.

10

The underside 24 of the membrane 16 is open to atmospheric pressure as a leakage path exists through the housing 1 which is not shown in the drawings as it extends around the outlet path 1 and is therefore not shown in the plane of Figs. 1 and 2.

When a user sucks on the outlet end 8 with the device in the configuration shown in Fig. 1, the suction is communicated by the suction orifice 9 to the chamber 10 through orifices 25 thereby lowering the pressure in this chamber. This causes the vane 15 to be lifted against the action of the spring 20 to the position shown in Fig. 2 deforming the diaphragm into the configuration shown in Fig. 2 and lifting the jaw 21 to allow the deformable tube to open, thereby allowing the inhalable composition from the reservoir 5 along outlet path 13 through the deformable tube 14 and out through the outlet orifice 11. The degree of suction applied by the user will determine the extent to which the vane 15 moves and therefore the amount of composition that the user receives. As soon as a user stops sucking, atmospheric pressure will return to the chamber 10 via the suction orifice 9 and the spring 20 will return the

- 11 -

vane to the Fig. 1 position thereby pinching the tube 14 closed.

A second example of an inhaler is shown in Figs. 4 to 5 6. This is also provided with a deformable tube 14, a reservoir 5 and outlet end 8, but the mechanism is somewhat different. In the closed position of Fig. 4, the deformable tube is pinched between a pair of jaws 30 on a spring clip 31. This clip 31 is biased into the first position shown in 10 Fig. 4. The inhaler has an outlet orifice 32 which completes a flow path from the reservoir 5 via the deformable tube 14 through the outlet orifice 32, and a plurality of suction orifices 33 which provide suction to an internal chamber 34. In this example, there are a pair of 15 pivotally mounted vanes 35 extending longitudinally along opposite sides of the device. Each of the vanes is connected to an L-shaped bracket 36, one of which is shown in Fig. 6. This L-shaped bracket extends into the respective jaw 30 and effectively reaches around to the 20 opposite side of the deformable tube 14 for reasons described below.

The housing 1 is provided with a pair of inlet orifices 37 one for each vane. As a user sucks on the outlet end 8, 25 the suction force via suction orifices 34 draws air through inlet orifices 37 into chamber 34 thereby applying inward pressure to the vanes 35. As a result of inward pressure, the vanes pivot inwardly to the position shown in Fig. 5. The L-shaped brackets 36 mean that the downward movement of 30 the upper vane 35 moves the lower jaw 30 downwardly and the upward movement of the lower vane 35 moves the upper jaw 30

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downwardly, thereby opening the deformable tube 14 as shown in Fig. 5 to dispense the inhalable composition. When the suction force is removed, the pressure differential across the vanes 35 is removed and the resilience of the spring clip 31 returns the jaws 30 to the position shown in Fig. 4.

An alternative arrangement of a deformable tubular element will now be described with reference to Fig. 7.

10 The vane 15, membrane 16 and other components are broadly the same as those described with reference to the previous example. The main difference in this example is the configuration of the deformable tubular element 14' and these differences are all that will be described below.

15

Essentially, the orifice 11 which was previously in a separate component has now been integrated into the tubular element 14' as orifice 11'. This has some additional benefits. Firstly, by replacing the two components with a single component, the overall space required for the outlet path has been reduced. This allows other elements such as the vane 15 and membrane 16 to be increased in size. This, in turn, increases the sensitivity of the device as it is more efficient at converting small breath forces into a movement which opens the flow path. This is important for users who may have impaired lung function capacity. Secondly, by eliminating the requirement for a seal between the tubular element 14 and the plate with the orifice 11, the manufacturing of the device can be simplified. In addition, this eliminates the potential for leakage at this interface. Also, the manner in which the orifice plate 14' is sealed to the outlet path 13 at its upstream end has also

- 13 -

been modified. At its upstream end, the tubular element 14' is provided with an outwardly projecting annular flange 40. This fits within the downstream end of the outlet path 13. This provides a more reliable sealing arrangement than
5 bonding the tubular element 14 in place. As a result, the new nozzle design can contain a pressurised formulation without leakage for a much longer period, and thus increase the stability of the formulation within the device as well as retaining a higher capacity for a longer period.

10

The tubular element 14' can be different thicknesses at particular parts. For example, the hoop stresses will be greatest on the walls of the nozzle immediately downstream of the flange 40. However, for the mid-section of the
15 tubular element 14' where the jaw 21 is sealing the tubular member, the material can be of reduced thickness to allow an easier clamping action.

Preferably, the tubular element 14' has a shore
20 hardness of between 20A and 80A, most preferably 30A to 40A. At its thickest part, the wall can be 0.5mm thick and at its thinnest part can be 0.18mm thick. In order to deliver the optimum performance for the aerosol to reach the pulmonary system on inhalation, the outlet orifice 11 is preferably
25 between 0.1mm and 0.5mm wide, but preferably 0.2 to 0.3mm and most preferably 0.2mm wide. The inner channel in the tubular element 14' away from the outlet orifice 11 is preferably between 0.2mm and 0.6mm wide, preferably between 0.3mm and 0.5mm and most preferably 0.4mm.

30

CLAIMS:-

1. An inhaler comprising:
 - a reservoir of an inhalable composition;
 - 5 an outlet at one end through which the inhalable composition is discharged; and
 - a non-metered breath-activated valve between the one end and the reservoir, the breath-activated valve comprising a flow path extending from the reservoir to the outlet end,
 - 10 at least a portion of the flow path being a deformable tube, and a clamping member which pinches the deformable tube closed when no suction force is applied to the device and releases the tube to open the flow path when suction is applied at the outlet, to provide uninterrupted flow from
 - 15 the reservoir to the outlet.

2. An inhaler according to claim 1, wherein the inhaler is a simulated cigarette device having a generally cigarette-shaped body.
- 20 3. An inhaler according to claim 2, wherein the inhalable composition includes nicotine or a nicotine derivative or salt thereof.

- 25 4. An inhaler according to claim 1 or claim 2, wherein the clamping device is configured so that the degree of opening of the tube is proportional to the strength of the suction.

5. An inhaler according to any one of the preceding
- 30 claims, wherein the clamping member is a mechanical member which is moved by a pressure differential caused when suction is applied at the outlet.

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6. An inhaler according to claim 5, wherein the clamping member is an elongate vane which is pivotally mounted at one end and extends longitudinally within the inhaler, the vane being pivotable between a first position in which the tube is closed and a second position in which it releases the tube to provide the flow path.

7. An inhaler according to claim 6, wherein there is a single vane which clamps the deformable tube against a fixed part of the inhaler.

8. An inhaler according to claim 6 or claim, 7, wherein the or each vane is biased into the closed position and the inhaler has a flow path creating a pressure differential across the vane which overcomes the biasing force.

9. An inhaler according to any of claims 6 to 8, wherein, the or each vane is supported by a flexible membrane, wherein flexible membrane creates a chamber in which the vane is housed and which is open to the outlet, but is otherwise sealed, with the face of the membrane on the opposite side to the vane being open to atmospheric pressure.

10. An inhaler according to any one of the preceding claims, wherein the deformable tube is a nozzle which is also provided integrally with an outlet orifice, which is the narrowest part of the flow path.

11. An inhaler according to claim 10, wherein the nozzle has an outwardly projecting annular flange at its upstream

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end which fits within a downstream end of an adjacent part of the flow path.

12. An inhaler according to any one of the preceding
5 claims, wherein the path is substantially straight.

13. An inhaler according to any one of the preceding
claims, wherein the tube is clamped at a single location.

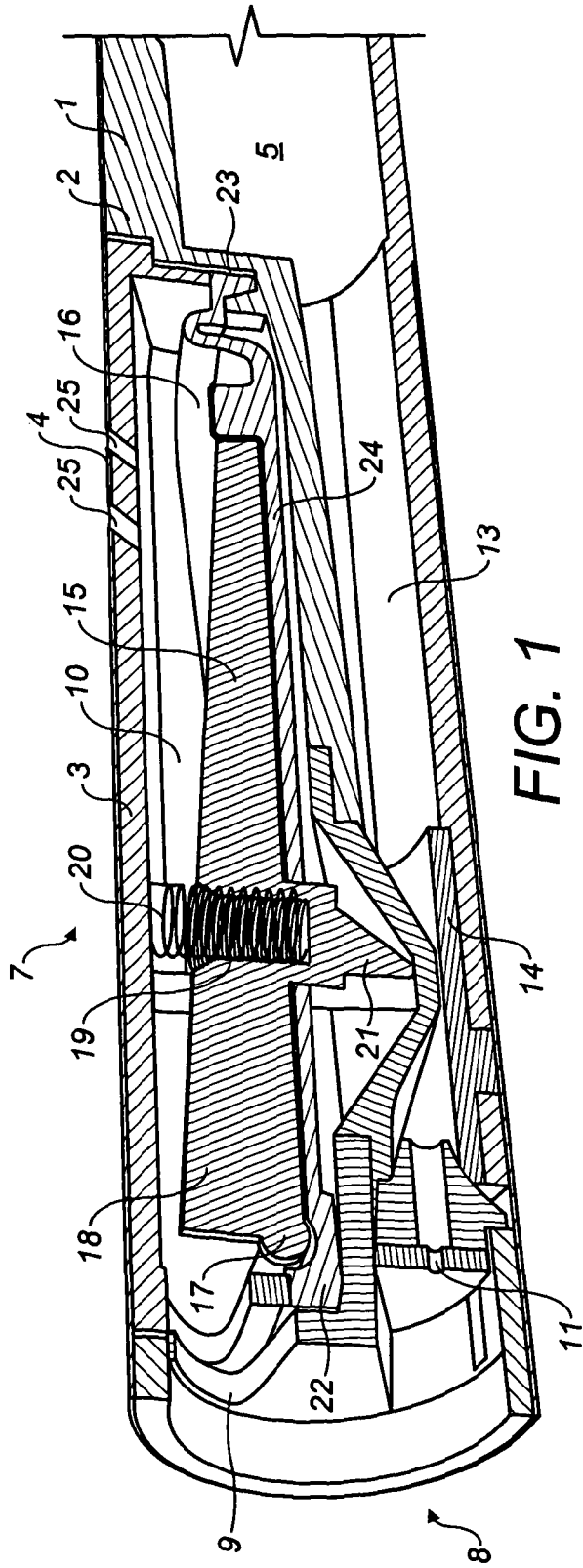


FIG. 1

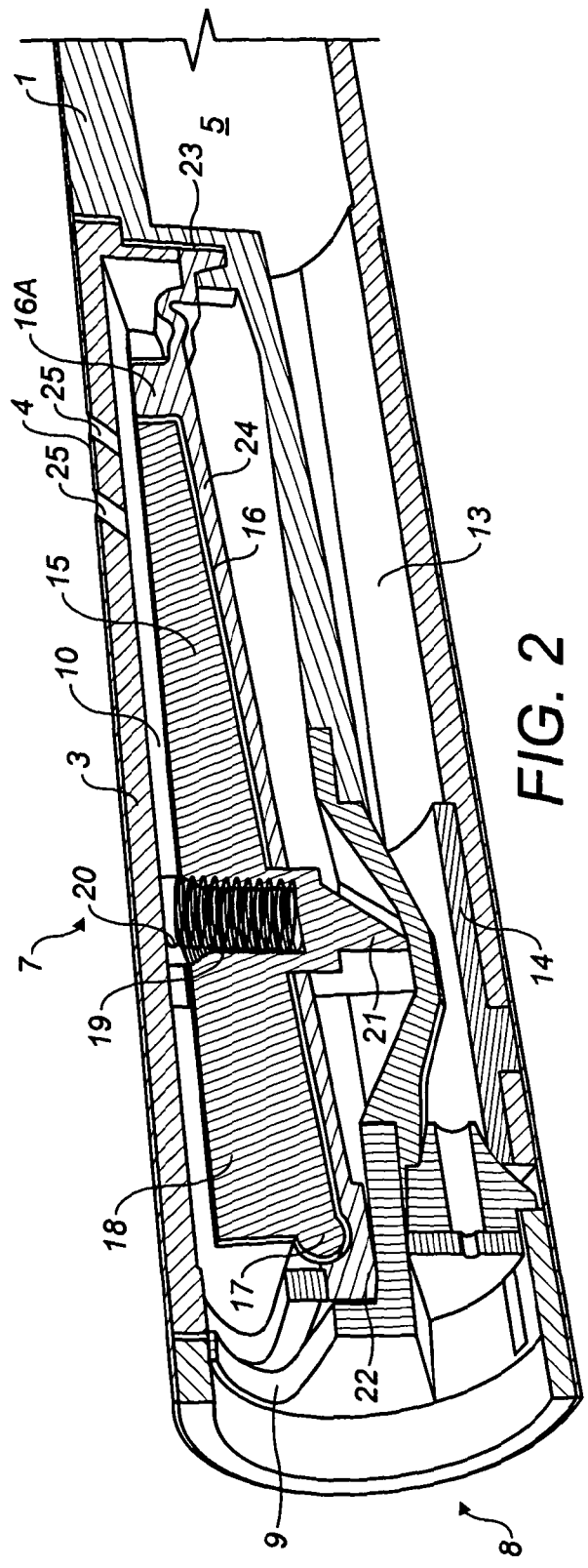


FIG. 2

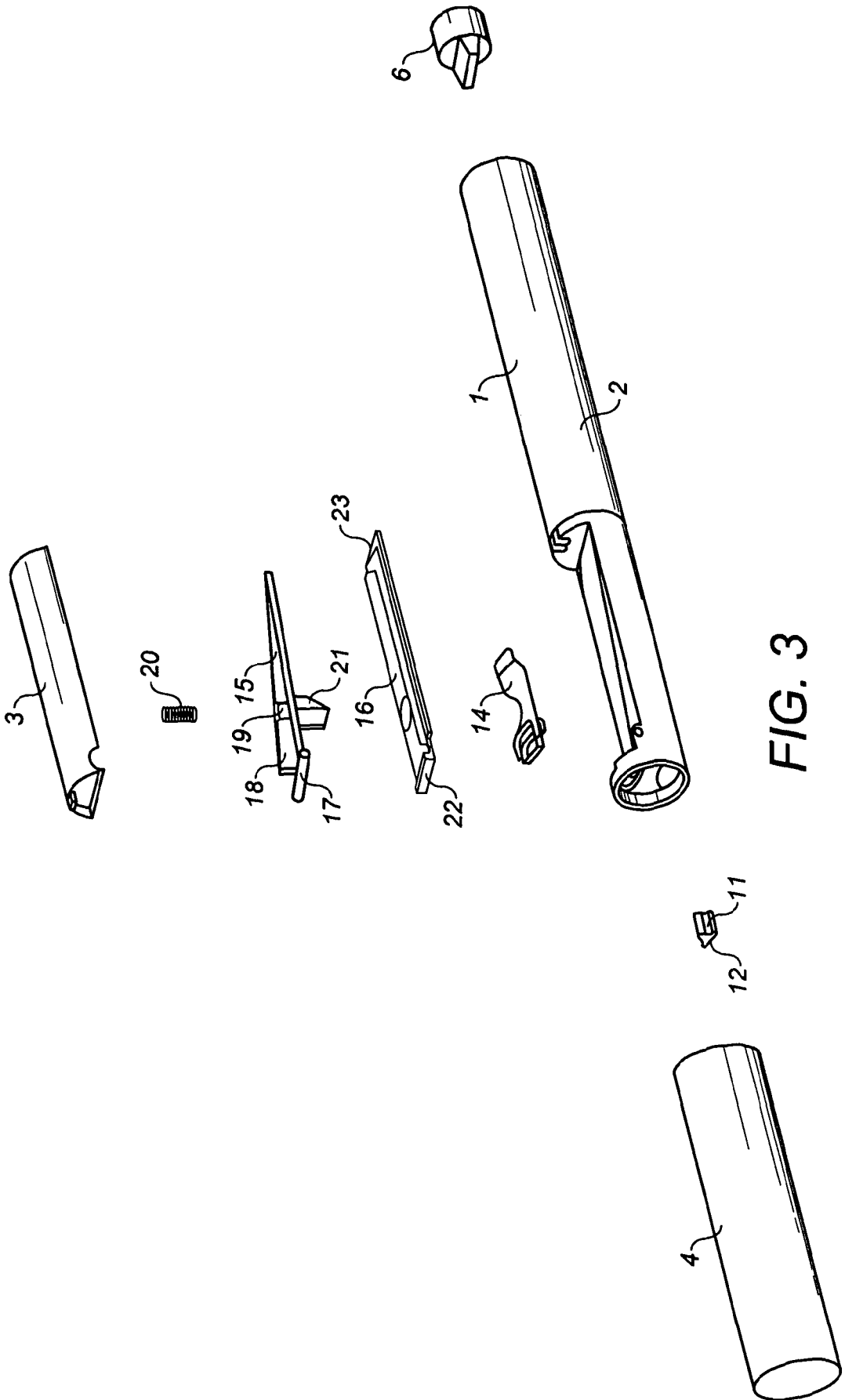
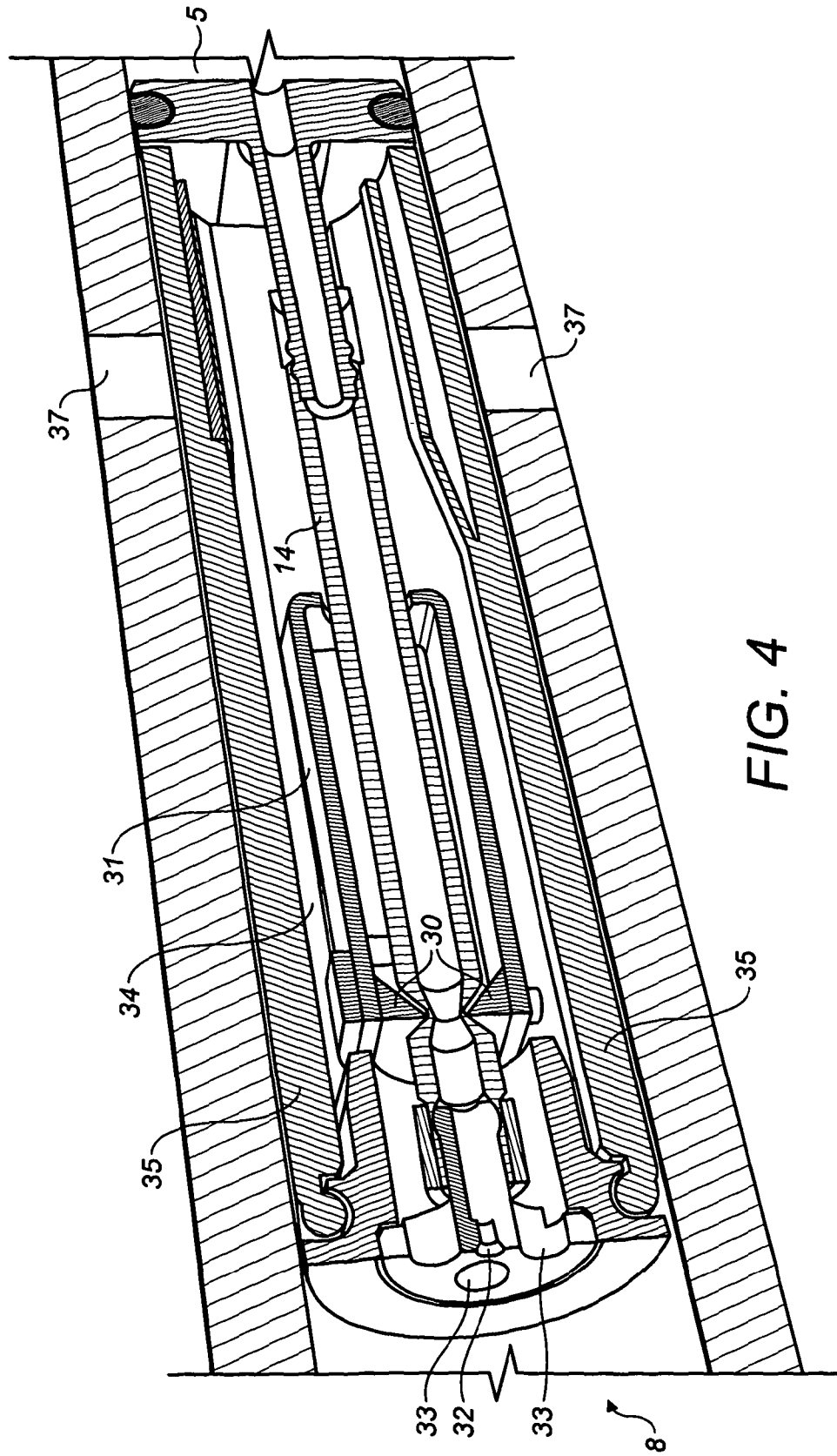


FIG. 3



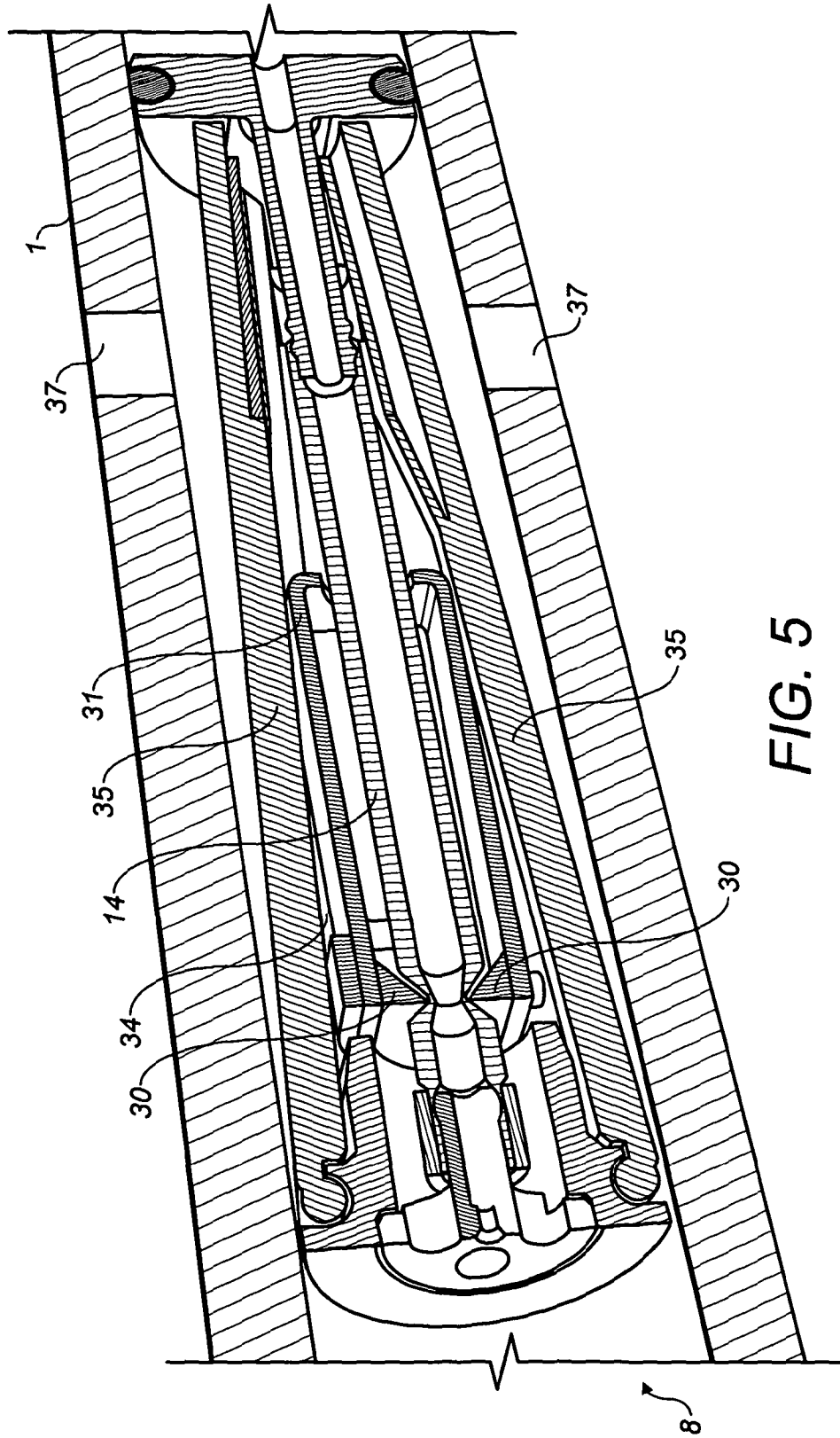


FIG. 5

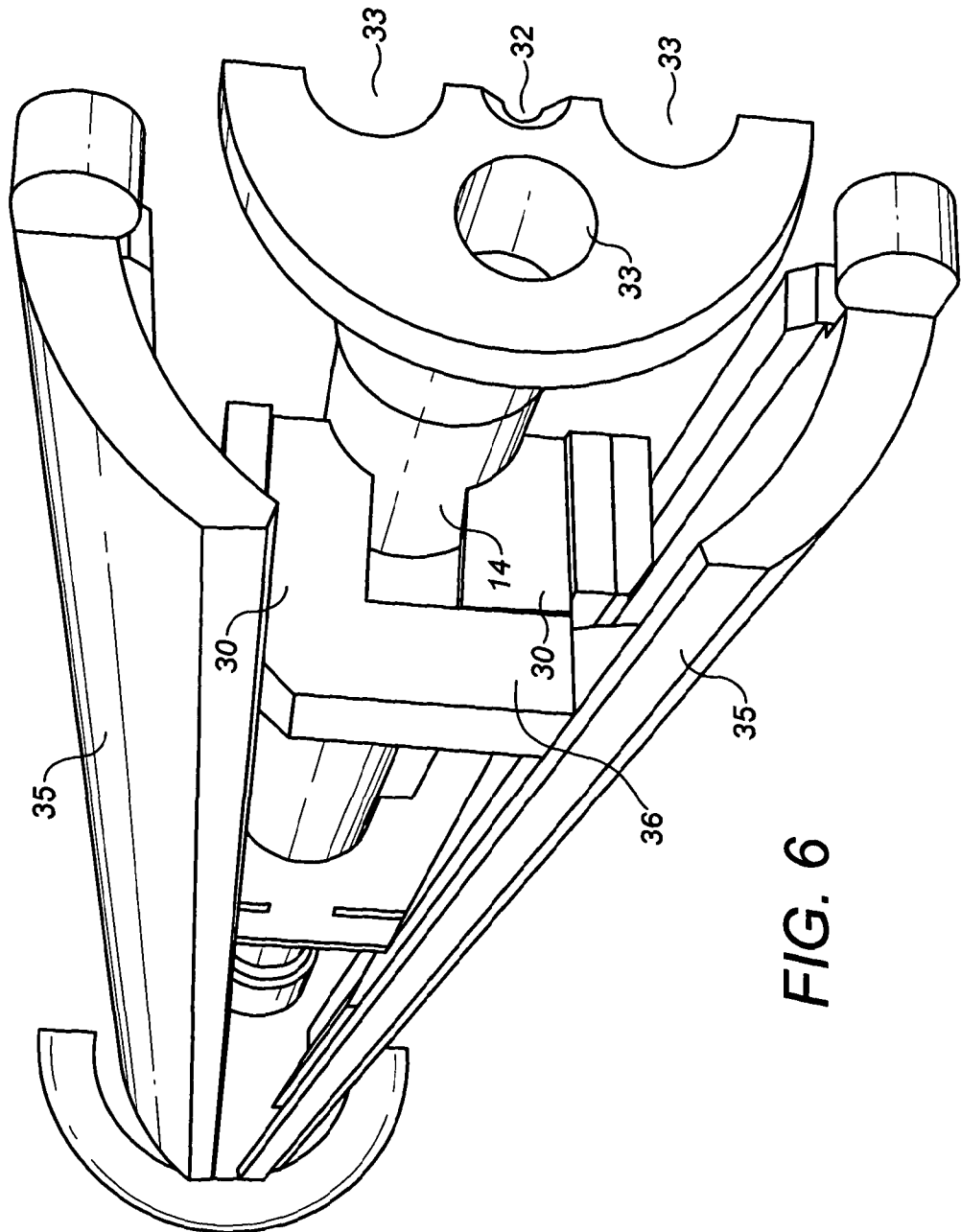


FIG. 6

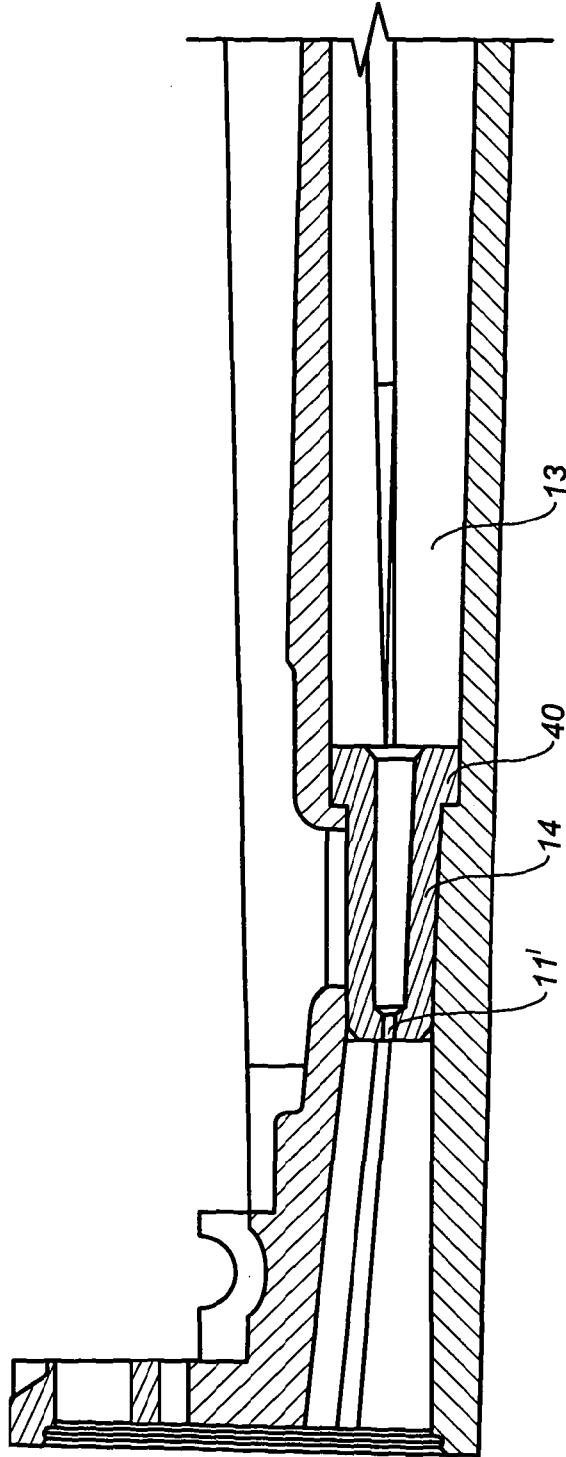


FIG. 7

INTERNATIONAL SEARCH REPORT

International application No
PCT/GB2010/001487

A. CLASSIFICATION OF SUBJECT MATTER

INV. A61M15/06 A24F47/00 A61M15/00 F16K7/06 F16K7/07
A61M16/20

ADD.

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

A61M A24F F16K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	FR 1 601 834 A (WERDING WINFRIED) 14 September 1970 (1970-09-14)	1, 2, 12, 13
Y	page 1, line 18 - page 3, line 18; figures 1-3	3-11
Y	----- US 2008/142008 A1 (POCOCK ANDREW [GB] ET AL) 19 June 2008 (2008-06-19) paragraph [0053] - paragraph [0057]; figures 1-4	4-11
Y	----- US 4 393 884 A (JACOBS ALLEN W [US]) 19 July 1983 (1983-07-19)	3
A	column 3, line 32 - column 4, line 19; figures 3,4	1, 2, 5, 6
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Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance
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Date of the actual completion of the international search

15 November 2010

Date of mailing of the international search report

23/11/2010

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INTERNATIONAL SEARCH REPORT

International application No

PCT/GB2010/001487

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 2009/001082 A1 (KIND CONSUMER LTD [GB]; HEARN ALEX [GB]) 31 December 2008 (2008-12-31) cited in the application page 7, line 21 - page 10, line 26; figures 1-5 -----	1-3
A	US 3 631 856 A (TAYLOR HAROLD V) 4 January 1972 (1972-01-04) column 1, line 63 - column 3, line 14; figures 1-8 -----	1,2,12, 13
A	US 6 629 528 B1 (WICKHAM PETER JOHN DEACON [AU] ET AL) 7 October 2003 (2003-10-07) column 3, line 64 - column 6, line 11; figures 1-6 -----	1,2,12, 13
A	US 2003/010946 A1 (FURUKAWA SHIGENOBU [JP] ET AL) 16 January 2003 (2003-01-16) paragraph [0065] - paragraph [0066]; figures 9-11 -----	1
A	US 6 581 590 B1 (GENOVA PERRY A [US] ET AL) 24 June 2003 (2003-06-24) column 4, line 23 - column 5, line 68; figures 1a-2 -----	1

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