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54) Title: A STAND-ALONE COUNTER FOR A MET		
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57) Abstract		43
A dose counter (40) for a metered dose inhaler comprising a medicament-containing vessel (30) and an		
actuator body (20) having a drug delivery outlet (25)	••	40 44
s adapted for attachment either to the medicament- containing vessel (30) or to the actuator (20). The		

> comprising a medicament-containing vessel (30) and an actuator body (20) having a drug delivery outlet (25) is adapted for attachment either to the medicamentcontaining vessel (30) or to the actuator (20). The dose counter (40) is provided with pressure sensing means (158, see Figs 2 and 3), counting means having a count display (43) and means (168, see Figs 2 and 3) to increment the count display (43) in direct response to the application of a predetermined pressure on the pressure sensing means. The invention does not rely on displacement triggering or the mechanical engagement between respective parts carried on the medicamentcontaining vessel and on the actuator housing.



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3M Company Exhibit 1014 3M Company, Merck & Co., Inc., and Merck Sharp & Dohme Corp., v. AptarFrance S.A.S. Page 2 of 20

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A STAND-ALONE COUNTER FOR A METERED DOSE INHALER

The present invention relates to a dose counter for a dispensing device of the type which is used for dispensing discrete amounts of a fluid or particulate material entrained in an air or other propellant stream. In particular, the 5 invention is concerned with dose counters for dispensing devices of the metered dose inhaler type which are well known in the art of medicine for treatment of, or alleviation of the effects of, respiratory complaints such as asthma.

Metered dose inhalers typically consist of a medicament-10 containing vessel and an actuator body having a drug delivery outlet.

The medicament-containing vessel may be a pressurised canister containing a mixture of active drug and propellant. Such canisters are usually formed from a deep drawn aluminium

15 cup portion having a crimped lid portion which carries a metering valve assembly. The metering valve assembly is provided with a protruding valve stem which, in use, is inserted as a tight push fit into a so-called "stem block" in the actuator body.

20 To actuate the "conventional" manually-operable inhaler, the user applies a compressive force to the closed end of the canister. The internal components of the metering valve assembly are spring loaded so that, typically, a compressive force of between 15 and 30 N is required to activate the

25 device. In response to this compressive force, the canister moves axially with respect to the valve stem by an amount varying between about 2.0 and 3.5 mm. This degree of axial movement is sufficient to actuate the metering valve and cause a metered quantity of the drug and propellant to be expelled

30 through the valve stem. This is then released into the mouthpiece via a nozzle in the stem block. A user inhaling through the drug delivery outlet of the device at this point will thus receive a dose of the drug.

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Metered dose inhalers as described above administer an accurate dose of medicament whenever required, which is particularly useful for users whose respiratory difficulties manifest themselves suddenly. Such has been the success of these devices that they are now used throughout the world, where they are exposed to a wide variety of climatic conditions.

A more recent development is the so-called "breathoperated actuator" which delivers a dose of drug through a 10 mouthpiece in response to inhalation by the user. This type of arrangement is particularly convenient in circumstances where the co-ordination between user inhalation and manual depression of the aerosol canister is imperfect. For example, children sometimes lack the necessary co-ordination to achieve 15 effective self-administration. At times of respiratory

- distress, adult users may also experience poor co-ordination. Unfortunately, one of the drawbacks of self-administration from an inhaler is that users often experience difficulty in determining when the charge in the medicament-containing vessel
- 20 has nearly run out. With aerosol canisters, part of the reason for this difficulty is that a surplus of propellant may remain in the canister even though the drug supply is nearly used up. Alternatively, the near-exhausted state may result in a surplus of drug in relation to propellant. Thus, the illusion is 25 created that the inhaler is still capable of providing useful doses of medicament simply because the canister contains liquid. This is potentially hazardous for the user since dosing becomes unreliable and because few people routinely

carry a back-up device.

30 Many users have several different inhalers for the treatment of a variety of conditions. Others keep inhalers at a number of different locations such as at school, home, work etc. In these circumstances it is particularly difficult for the user to keep track of the amount of usage extracted from 35 each individual inhaler apparatus.

Clearly there is a need for a counter mechanism which is capable of counting the number of doses delivered from or remaining in an inhaler. To this end, a number of counters have been proposed in recent times which aid the management of metered dosage. Such counters vary in complexity and sophistication, but they all have in common the feature that they detect relative movement between the medicament-containing vessel and the actuator body and increment in response to such movement.

10 One of the drawbacks of these known counters is that they rely on mechanical interaction between parts attached to the medicament-containing vessel and parts provided on the actuator body. This means that routine cleaning of the inhaler, which requires removal of the medicament-containing vessel from the actuator body, may result in damage to the counting mechanism.

- Also, the accumulated count may be corrupted by this operation. Another drawback is that such counters are difficult to manufacture with satisfactory tolerances. A factor which influences the effectiveness of known displacement-triggered
- 20 counters is the variation in length of typical aerosol canisters. This variation is attributable in part to the crimping operation used to connect the valve-carrying lid portion to the main cup portion. Although the technology involved is not especially demanding, it has been found that
- 25 the amount of travel effective to actuate the metering valve of a typical medicament-containing aerosol canister may fall in a tolerance band as small as 0.5 mm. Thus it is difficult to provide a generic counter which increments accurately in response to every actuation. This may be true even when the
- 30 counter, the aerosol canister and the inhaler housing have been specifically designed for use together. The problem is therefore likely to be worse in circumstances where different manufacturers' aerosol canisters, inhaler housings and displacement-triggered counters are used in combination.

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It is therefore an object of the present invention to provide a counter for a metered dose inhaler which does not rely on displacement triggering or the mechanical engagement between respective parts carried on the medicament-containing vessel and on the actuator housing.

It is a further object of the invention to provide a counter which allows removal of the medicament-containing vessel from the actuator body for routine cleaning without the risk of count corruption.

10 The invention is a dose counter for a metered dose inhaler comprising a medicament-containing vessel and an actuator body having a drug delivery outlet, characterised in that said dose counter is adapted for attachment to one of said medicamentcontaining vessel and said actuator body, the dose counter being provided with pressure (force) sensing means, counting

means having a count display and means to increment the count display in direct response to the application of a predetermined pressure upon the pressure sensing means.

One of the main advantages of a pressure-triggered counter compared to a displacement-triggered counter in the context of metered dose inhalers is that typical valve actuating <u>pressures</u> fall within a sensible range. This is in contrast to the valve actuating <u>travel</u> which, as mentioned above, may vary considerably from one inhaler to another. It is therefore

25 relatively simple to provide a pressure-triggered device having a predetermined pressure threshold which renders it universally applicable. This is much easier to achieve in engineering terms than a displacement-triggered counter which is required to increment in response to a relative displacement as low

30 as 0.5 mm.

counter may under-count.

It is preferred that the value of the predetermined pressure applied to the pressure sensing means which causes the count display to increment should be no greater than the compressive force required to actuate the metering value of the aerosol canister. If this criterion is not met, the dose

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Errors biased in this direction, as opposed to overcounting, could have serious consequences for the user. This is because the implication that effective doses of medicament remain when the drug supply is actually exhausted could be 5 potentially life-threatening.

Another important consideration is resistance to vibration. Often, the directions for use instruct the user to shake the dispenser prior to administration of the metered dose. The dose counter should therefore be able to withstand vigorous shaking without corruption of the recorded count. Similarly, the count should not be corrupted if the inhaler is accidentally dropped.

Preferably, the pressure sensing means is a solid state device, most preferably a piezo film sensor. In such an arrangement, the pressure sensing means generates an electrical output in proportion to the applied compressive force. The electrical output may then be fed into processing means such as a microprocessor in which the signal is analysed to determine whether the threshold of the predetermined minimum pressure has been reached. If so, an electrical impulse is sent to the counting means to effect incrementation of the count display.

Alternatively, the pressure sensing means may be a forcesensitive switch to which a predetermined force must be applied to overcome a biasing pressure that urges the switch contacts into an open (or closed) condition. When the switch contacts are closed (or opened), an electrical circuit is completed (or broken) which stimulus can be used to activate the counting means in order to increment the count display.

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In an especially preferred form of the invention, the count display is provided on a surface of the dose counter which is visible from the exterior of the inhaler. This means that the display is visible to the user both before and after the administration of a dose of the drug. Such an arrangement

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is most conveniently implemented by means of an electronic chip with associated display function.

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In one particularly preferred form of the invention, the dose counter is adapted to be attached to the closed end of an aerosol canister containing medicament and propellant under pressure. The dose counter in this arrangement is in the form of a button and actuation of the inhaler is effected by manually depressing the button. This, in turn, moves the aerosol canister axially and actuates the metering valve. The force exerted on the button also causes incrementation of the counter. Conveniently, the counter display is provided on the actuating surface of the button.

Preferably, the button is provided with a resilient skirt for attaching it to the closed end of the aerosol canister in a reliable fashion. In order to make the dose counter fit a variety of aerosol canisters of differing diameters, the skirt may be provided with auxiliary adaptor means. Preferably the auxiliary adaptor means is easily remeved and discourded if not

auxiliary adaptor means is easily removed and discarded if not required.

The button may incorporate a so-called "snap-dome" trigger which requires a consistent, minimum force to actuate. This 20 type of arrangement aids the user in applying an undeviating activation pressure which ensures that the pressure sensing means is positively actuated whenever the trigger is fired. When the user relaxes the compressive force, the dome snaps back to its original position. Many users prefer the positive 25 action which such a trigger imparts.

In another form of the invention, the dose counter is adapted for attachment to the actuator of the inhaler apparatus. This arrangement is preferred for breath-operated actuators, where manual depression of the counter may not be possible.

If desired, the pressure sensing means may be positioned at a remote location from other components of the counter, being connected thereto by trailing wires, for example. Such an arrangement allows for greater versatility in deployment of the counter and applicability to more varieties of inhaler.

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Preferably, the dose counter is provided with a resetting mechanism operable to reset the count display when the counter is transferred from an exhausted inhaler apparatus. This may be implemented, for example, by means of a reset button which is adapted to make contact with a resetting circuit provided in the counter device. Once this contact is made, the count display is returned to its starting value. In order to avoid accidental resetting, it is preferable for the reset button to be concealed or recessed. For example, the reset button could

10 be in the form of a recessed pin which is operable by means of an everyday object of narrow section, such as the tip of a ball-point pen.

The invention will now be described by way of example only with reference to the drawings, in which:

- 15 Figure 1 shows a schematic cross-sectional view of a metered dose inhaler apparatus fitted with a dose counter in accordance with the invention.
 - Figure 2 is an exploded view showing the components of a first embodiment of the dose counter according to the invention, and
 - Figure 3 is a side sectional view of a second embodiment of dose counter in accordance with the invention.

Referring now to Figure 1, reference numeral 10 denotes a metered dose inhaler comprising an actuator body 20 having 25 a drug delivery outlet 25, and an aerosol canister 30. The aerosol canister 30 is formed from a deep drawn aluminium cup section 31 to which a lid portion 32 is attached by crimping. The lid portion 32 carries a metering valve assembly having a protruding valve stem 33, the end of which is received as a

30 tight push fit in a stem block 21 of the actuator body 20. Stem block 21 has a nozzle 22 communicating with the drug delivery outlet 25 so that, upon actuation of the metering valve assembly, a charge of the drug is emitted through the nozzle 22 into the drug delivery outlet 25. A user inhaling 35 through the drug delivery outlet 25 at this time will receive a metered dose of the drug.

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The closed end 35 of aerosol canister 30 is provided with a dose counter 40 which comprises a body portion 41 and a depending skirt 42. The skirt 42 is formed of a resilient material which grips the outside of the aerosol canister 30 to maintain the dose counter 40 in position. Body portion 41 is

- in the form of a button which includes pressure sensing means and which has, on its top surface 44, a count display 43. Body portion 41 includes a processing chip and associated electronics (not shown) which are configured to determine
- 10 whether a predetermined force has been applied to the pressure sensing means. When this force threshold is reached, the chip sends an electrical impulse to the count display 43 in order to effect incrementation thereof.

Turning now to Figure 2, the dose counter 40 of Figure 1 15 will be described in more detail with reference to the exploded view of its components.

A skirt element 142 is integrally moulded with a seat portion 152 on which a first resilient member 154 is placed. An intermediate housing 156 is adapted for placement on first

- 20 resilient member 154, the upper surface of housing 156 being configured to receive a piezo film element 158 in a well 157. Overlying the piezo film element 158 is a second resilient member and a conducting element 166 which conveys electrical impulses from the piezo film element 158 to integrated circuit
- 25 168. Integrated circuit 168 is powered by a battery 170 which activates a liquid crystal counter display 174 through the integrated circuit 168 via connector 172. The assembly is completed by an outer cover 176 having a window 178 through which the counter display 174 is visible. The respective housing parts 152, 156 and 176 may be held together by bolts

or other suitable fastening means.

In use, the fully-assembled device is installed on an aerosol canister as shown in Figure 1, with substantially no dead space between the closed end of the canister and the underside of the seat portion 152. Then, application of

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pressure to outer cover 176 in the direction of arrow A causes deflection of the piezo film element 158 inside the assembly at the same time as the canister is caused to translate within its holder. Deflection of the piezo film element 158 generates an electrical impulse which is transmitted to the integrated circuit 168.

A piezo film such as the element 158 is a force-sensitive device, so the degree of deflection will determine the strength of the generated electrical impulse. If the condition is satisfied that the electrical impulse exceeds a minimum

threshold value, the count display 174 is caused to increment. Ideally, the threshold value is set at a level which ensures that the dose counter does not under-count. In other words, the activation threshold of the counter should be equal to or less than the activation threshold of the metering valve assembly of the aerosol canister to which it is attached in use.

When the applied pressure is removed, the piezo film element 158 relaxes to a neutral position, aided by first and 20 second resilient elements 154, 164.

Referring now to Figure 3, this view shows an embodiment of the present invention adapted for mounting in the side wall of an inhaler actuator, for example a breath-operated actuator. In this Figure, the features of the device which are in common with the Figure 2 embodiment are denoted by the same reference

numerals.

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In this arrangement, base element 256 corresponds to intermediate housing 156 of the Figure 2 embodiment and includes a well 257 in which piezo film element 158 is received. Base element 256 is provided with an arm 255 having a tip portion 253 directed towards the piezo film element 158. In its relaxed condition, the arm 255 overlies the piezo film element 158 with a small gap between the tip portion 253 and the plane occupied by the piezo film element 158. This condition is shown in ghost outline in the Figure.

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On the other side of the piezo film element 158 from the arm 255, a resilient element 264 is provided. This provides a restoring force for the piezo film element 158 when it is deflected by the arm 255, as will be described in more detail

5 below. Also on this side of the piezo film element 158 is a conducting element 166 which conveys electrical impulses to integrated circuit 168.

As previously described in relation to Figure 2, integrated circuit 168 is powered by battery 170 which 10 activates a liquid crystal counter display 174 through the integrated circuit 168 *via* connector 172. An outer cover 176 is also provided, said cover having a window 178 through which

Operation of this device is similar to that for the Figure 2 embodiment, except that pressure is applied in the direction of arrow B instead. This causes the tip portion 253 of arm 255 to engage the piezo film element 158, resulting in deflection thereof and consequent generation of an electrical impulse. If the strength of the impulse exceeds a minimum threshold value, the count display 174 is caused to increment.

counter display 174 is visible.

Arm 255 may be arranged to flex in response to engagement by a nib element (not shown) provided on a so-called "force holding unit" of a breath-actuated inhaler. This is the unit which moves in response to user inhalation to effect actuation of the metering valve assembly previously described.

Although the invention has been particularly described above in relation to certain specific embodiments, it will be understood by persons skilled in the art that its application is not limited to inhalers which use pressurised aerosol

30 dispensers. For example, the invention may be applied to dry powder inhalers provided that their mechanism of operation is such that pressure sensing is a legitimate parameter to equate with dose counting. Other variations and modifications are possible and will be apparent to skilled persons without 35 departing from the scope of the claims which follow.

CLAIMS

1. A dose counter for a metered dose inhaler comprising a medicament-containing vessel and an actuator body having a drug delivery outlet, characterised in that said dose counter is adapted for attachment to one of said medicament-containing vessel and said actuator body, the dose counter being provided with pressure sensing means, counting means having a count display and means to increment the count display in direct response to the application of a predetermined pressure upon the pressure sensing means.

2. A dose counter as claimed in claim 1 in which the pressure sensing means is a solid state device such as a piezo film sensor.

3. A dose counter as claimed in claim 1 or claim 2 wherein the pressure sensing means is a force-sensitive switch the contacts of which are held open by a biasing force corresponding to said predetermined pressure.

4. A dose counter as claimed in any preceding claim in which a microprocessor is used to determine whether the threshold of the minimum predetermined pressure has been reached.

5. A dose counter as claimed in any preceding claim wherein the count display is provided on a surface of the dose counter which is visible from the exterior of the inhaler.

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6. A dose counter as claimed in any preceding claim adapted for attachment to the closed end of a medicament-containing vessel in the form of a pressurised aerosol canister, wherein the dose counter is configured as button means.

7. A dose counter as claimed in claim 6 wherein said button means has a resilient skirt depending therefrom for attaching the button means to said canister.

8. A dose counter as claimed in any one of claims 1 to 5 adapted for attachment to the actuator of a breath-operated inhaler.

9. A dose counter as claimed in claim 8, wherein the dose counter is configured as a module receivable in a wall of the breath-operated inhaler.

10. A dose counter as claimed in any preceding claim having resetting means operable to reset the count when the counter is transferred to a new inhaler apparatus.

11. A dose counter as claimed in claim 10 wherein the resetting means is recessed to prevent accidental actuation thereof during normal use.



FIGURE 1

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FIGURE 2

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FIGURE 3

INTERNATIONAL SEARCH REPORT

Interna l Application No PCT/GB 95/00725

A. CLASSIFICATION OF SUBJECT MATTER IPC 6 A61M15/00

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

B. FIELDS SEARCHED

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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)

C. DOCUM	IENTS CONSIDERED TO BE RELEVANT		
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X	WO-A-92 17231 (INNOMED INC) 15 Oc 1992	tober	1,3,5,6, 10,11
Y	see abstract; figures 1,4,5 see page 4, line 9 - page 6, line	2 4	2,4,8
Y	US-A-5 034 648 (GASTGEB) 23 July see abstract; figures 1,2 see column 3, line 4 - line 37	1991	2,4
Y	EP-A-O 461 281 (ATOCHEM NORTH AME December 1991 see abstract; figures 1,6,7	RICA) 18	8
A	see column 5, line 12 - line 28 see column 7, line 40 - line 56		2,4,6
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Date of the	actual completion of the international search	Date of mailing of the international se	earch report
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C.(Continua Category °	ation) DOCUMENTS CONSIDERED TO BE RELEVANT Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US-A-5 020 527 (DESSERTINE PAULINE L) 4 June 1991 see abstract; figures see column 3, line 40 - line 52	5,6,9
Ą	see abstract; figures	8

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INTERNATIONAL SEARCH REPORT information on patent family members			Interna al Application No PCT/GB 95/00725	
Publication date			Publication date	
15-10-92	NONE			
23-07-91	NONE			
18-12-91	NONE			
04-06-91			01-06-95 25-09-91	
09-12-93			30-12-93 08-12-93	
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