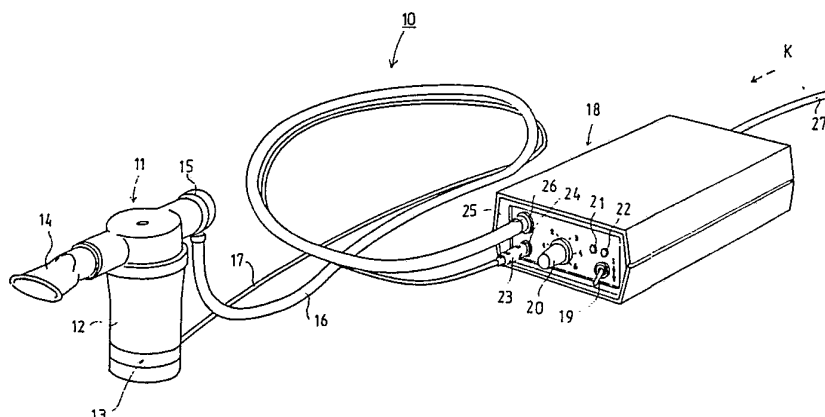




## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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**(54) Title:** ULTRASONIC ATOMIZER**(57) Abstract**

The invention concerns an ultrasonic atomizer (10), which is intended for an inhalation treatment apparatus for persons suffering from respiratory diseases. The ultrasonic atomizer (10) comprises an atomizer device (11) provided with a drug container (12) as well as an ultrasonic oscillator (13) connected to the container (12), preferably an oscillating crystal. The atomizer device (11) is provided with a duct passing into the patient, advantageously a mouth piece and an air-inlet duct (15). The atomizer device (11) comprises a pressure detector (36) directly connected to it or a duct (16) that is connected to the atomizer device (11) and passes to the pressure detector (36) and transmits the pressure of the breathing air. The pressure detector (36) detects the changes in pressure resulting from inhalation or exhalation of the patient, whereby, starting from the beginning of the inhalation stage, atomizing can be switched on by the apparatus. The ultrasonic atomizer (10) comprises a regulating device (18) connected to the atomizer (11), by means of which said regulator the operation of the ultrasonic oscillator (13) is regulated. The regulating device (18) and the atomizing device (11) are interconnected by means of an electric connection (17), whereby, through said connection (17), an electric oscillation can be supplied to the crystal present in the ultrasonic oscillator. Said electric oscillation can be converted to mechanical oscillation of the crystal. The apparatus comprises means for switching-on of said ultrasonic oscillator (13) at least when the inhalation stage of the patient begins. Also, a timing device (44) is provided, by means of which the duration of operation of the ultrasonic oscillator (13) can be regulated as desired.

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## 1 Ultrasonic atomizer

5 The invention concerns an ultrasonic atomizer, which is intended for an inhalation treatment apparatus for persons suffering from respiratory diseases, said ultrasonic atomizer comprising an atomizer device provided with a drug container as well as an ultrasonic oscillator connected to the container, preferably an oscillating crystal, and said atomizer device being provided with a  
10 duct passing into the patient, advantageously a mouth piece and an air-inlet duct, and said atomizer device comprising a pressure detector directly connected to the atomizer device or a duct that is connected to the atomizer device and passes to the pressure  
15 detector and transmits the pressure of the breathing air, whereat the pressure detector detects the changes in pressure resulting from inhalation or exhalation of the patient, whereby, starting from the beginning of the inhalation stage, atomizing can be switched on by the apparatus, and said ultrasonic atomizer comprising a regulating device connected to the atomizer device, by  
20 means of which said regulating device the operation of the ultrasonic oscillator is regulated.

Water or drug mist that is used in respiration treatment can also  
25 be produced by means of ultrasonic oscillations. By means of intensive oscillation, a field of waves is produced on the surface of liquid, in which the velocity of the liquid particles in the waves becomes so high that it surpasses the effects of gravity and of surface tension forces, and small particles are detached from  
30 the liquid surface into the air. The drop size is determined by the properties of the liquid and by the ultrasonic frequency used in the ultrasonic oscillator. Most commonly, the ultrasonic frequency in treatment atomizers is of an order of 1 to 2 MHz. In such a case the drop size of water mist becomes 2 to 5  $\mu\text{m}$ . The  
35 atomizing capacity of the ultrasonic atomizer depends on the size of the oscillating crystal and on the magnitude of the electric power supplied to the crystal. For example, with a crystal of a

1 diameter of about 15 mm and with a power of about 12 W, the water  
atomizing capacity that is obtained is about 250 ml/h.

5 It can be considered that a drawback of the prior-art ultrasonic  
atomizers is the short service life of the oscillator crystal.  
Nor has a precise adjustment of the atomizing been possible in the  
prior-art ultrasonic atomizers. It would be essential that the  
atomizing time could be adjusted so that it is advantageous in  
10 view of the treatment of the patient. Thereat, such a solution of  
equipment would be advantageous in which the optimal atomizing  
time can be adjusted individually for each patient. In view of the  
service life of the apparatus, it would also be advantageous to be  
able to form an ultrasonic atomizer in which it is possible to  
increase the service life of the ultrasonic crystal considerably,  
15 yet, without deterioration of the treatment result.

The objects of the invention have been achieved by means of a  
solution of apparatus which is mainly characterized in that the  
regulating device and the atomizing device are interconnected by  
20 means of an electric connection, whereby, through said connection,  
an electric oscillation can be supplied to the crystal present in  
the ultrasonic oscillator (13), whereby said electric oscillation  
can be converted to mechanical oscillation of the crystal, and  
that means are provided for switching on said ultrasonic oscillator  
25 at least when the inhalation stage of the patient begins and that  
a timing device is provided by means of which the duration of  
operation of the ultrasonic oscillator can be regulated.

An ultrasonic atomizer in accordance with the invention includes a  
30 control between the pressure detector and the timing device, whereby  
the pressure detector detects any change in pressure resulting  
from the inhalation of the patient, and further, the signal thereat  
arriving from the pressure detector is fitted to start the counting  
of the timing device from the beginning of the inhalation stage.  
35 By means of the regulating device, the maximum time that the ultra-  
sonic oscillator oscillates can be set in advance into the timing  
device. The ultrasonic atomizer in accordance with the invention

1 also includes a control logic which detects if the liquid in the  
container is about to end, in which case the control logic detects  
the quantity of liquid present in the container in a change in the  
current or voltage of the oscillating crystal. When current or  
5 voltage values higher or lower than certain predetermined values  
are reached, the control logic switches the crystal off oscillation.  
The control logic also includes means by which it is ensured that  
the crystal cannot be switched on oscillation until after the main  
current circuit has first been switched off once. Thereby it is  
10 ensured that liquid is added to the liquid vessel.

The ultrasonic atomizer in accordance with the invention also  
includes an infrared detector, which detects whether the duct  
passing to the pressure detector and transmitting negative pressure  
15 is connected to the regulator device. When the connector of the  
duct is not connected to the regulator device, the infrared detector  
detects this, and then the apparatus atomizes constantly.

The apparatus in accordance with the invention also includes con-  
20 nector means by which the ultrasonic oscillator can be connected  
detachably to a liquid container. In this way, the same ultrasonic  
oscillator can be used in connection with several different atom-  
izers or liquid containers.

25 In the following, the invention will be described with reference  
to some preferred embodiments of the invention illustrated in the  
figures in the accompanying drawing, the invention being, however,  
not supposed to be confined to said embodiments alone.

30 Figure 1A is an axonometric view of an ultrasonic atomizer in  
accordance with the invention.

Figure 1B shows the device of Fig. 1A with the duct of negative  
pressure disconnected from the regulator device.

35 Figure 1C shows the device of Fig. 1A as viewed from the rear and  
in the direction of the arrow K in Fig. 1A.

1     Figure 2A is a side view of the atomizer with the flow ducts inside  
the apparatus shown with broken lines.

5     Figure 2B shows the apparatus of Fig. 2A with the ultrasonic  
oscillator disconnected.

Figure 3 shows an ultrasonic atomizer in accordance with the  
invention as an operational block diagram.

10    Fig. 1A is an axonometric view of an ultrasonic atomizer 10 in  
accordance with the invention. The ultrasonic atomizer 10 comprises  
an atomizer device 11 and a liquid container 12 provided therein,  
an ultrasonic oscillator 13 being fitted at the bottom of the  
container.

15    The ultrasonic oscillator 13 comprises an oscillating crystal, the  
oscillation energy being transferred from the oscillating crystal  
to the surface of the liquid, preferably a drug, contained in the  
liquid vessel, whereby a wave field is formed on the surface of  
20    the liquid. The velocity of the liquid particles in the waves  
becomes so high that it surpasses the effects of gravity and surface  
tension forces, and small particles are detached from the liquid  
surface into the air. The drops are carried from the liquid vessel  
further into the air space passing into the patient.

25    The atomizer device 11 further comprises a connecting duct 14  
passing to the patient, as well as an air-inlet duct 15 connected  
to the atomizer. The connecting duct 16 connected to the pressure  
detector is fitted so that it is advantageously connected to the  
30    inlet duct 15. The connecting conduit passing the oscillating  
power to the crystal is denoted with the reference numeral 17, and  
said conduit is favourably an electric conduit, preferably a coaxial  
cable.

35    The ultrasonic atomizer 10 in accordance with the invention includes  
a regulator device 18, which contains all essential regulating  
circuits and regulating functions related to the apparatus in a

1 centralized form.

The regulator device 18 includes a main current switch 19 and a power regulator 20 for the ultrasonic oscillator. Further, located  
5 on the front panel of the regulator device, there is a first signal light 21, which is the signal light of the main current circuit and, when it is on, indicates that the current is on, and when the signal light is off, the current has been switched off. At the side of the first signal light, in the front panel, there is a second  
10 signal light 22, which indicates that the liquid is being exhausted in the liquid vessel 12.

As is shown in Fig. 1B, the conduit 17 is provided with a connector 23, by means of which the conduit 17 can be connected to the  
15 counter-connector 24, provided on the regulator device 18, detachably.

Correspondingly, the connecting duct 16 of the pressure detector 36 is provided with a connector 25, by means of which the duct 16  
20 can be connected to the counter-connector 26, provided on the regulator device 18, detachably. The reference numeral 27 denotes the voltage cable through which the operating power is supplied into the apparatus. It is also self-evident that it is possible to connect an accumulator or any other source of power to the regulator  
25 device 18.

Fig. 1C is a rear view of the regulator device 18, seen in the direction of the arrow K in Fig. 1A. In the way shown in Fig. 1C, the voltage cable 27 includes a connector 28, which can be connected  
30 further to the counter-connector 27 provided on the regulator device 18. As was already explained above, an accumulator or any other source of power can be connected to the counter-connector 29 directly.

35 In the illustration shown in Fig. 1C the adjusting knob 30 of the atomizing-time timing device 44, placed on the rear panel of the device, is shown. The adjusting knob 30 comprises an indicator

1 arrow 31 or equivalent and an adjusting slot 32, by means of which  
the adjusting knob can be turned and the atomizing time be adjusted  
to the desired level. The rear panel is provided with a time scale  
33, from which the desired atomizing time can be read as indicated  
5 by the indicator arrow 31.

Fig. 2A is a side view of the atomizing device 11 to be connected  
to the regulator device 18, the air ducts inside the apparatus  
being shown by broken lines. When the patient breathes through the  
10 connecting duct, i.e. mouth piece 14, passing into the patient,  
air is absorbed from the outer air in the way shown by the arrow  
 $L_1$  through the duct 15 and the drug is carried from the liquid  
vessel 12 along with the air flow  $L_1$  in the way shown by the arrow  
 $L_2$  into the connecting duct 14 that passes into the patient, and  
15 further the air + atomized substance are carried into the patient  
as is shown by the arrow  $L_3$ . As the patient inhales through the  
connecting duct 14, a flow is produced in the duct 15, and further  
a negative pressure is formed at the connection point 16a, to  
which the connection hose 16 passing to the pressure detector is  
20 connected. The negative pressure (P) is transmitted through the  
connection hose 16 further to the pressure detector in the regulator  
device 18.

The bottom portion of the container part 12a of the liquid con-  
25 tainer 12 is constituted by the top portion 13a of the ultrasonic  
oscillator 13. Thus, said bottom portion 13a, which transmits the  
oscillation of the crystal, acts as the bottom portion of the  
liquid container 12, favourably the drug container. In this em-  
bodiment of the invention, a liquid container 12 is spoken of,  
30 which means a container which may contain water or drug to be  
atomized or any other treatment agent to be atomized.

Along the cable 17, a variable electric voltage is passed to the  
crystal in the ultrasonic oscillator 13.

35

As is shown in Fig. 2B, the ultrasonic oscillator 13 can be detached  
from the liquid container 12. Thus, the same ultrasonic oscillator



1 can be used for different atomizers 11 and for different liquid containers 12, e.g., for administration of different medicines.

As is shown in Fig. 2B, the apparatus includes connecting means  
5 34,35, by which the ultrasonic oscillator 13 can be connected to the liquid vessel 12 detachably. As is shown in Fig. 2B, the side faces of the drug container 12 are provided with attachments 35, which, as is shown in the figure, consist of two cavities 35a,35b, into which the attaching members 34 that grasp the ultrasonic  
10 oscillator 13 are coupled detachably. As is shown in Fig. 2B the attaching members 34 consist of screws 34a,34b, which said screws attach the ultrasonic oscillator 13 to the liquid container 12 when they are engaged into the screw cavities 35a and 35b.

15 Fig. 3 shows a functional block diagram of the ultrasonic atomizer. According to the invention, a quartz crystal or any other, corresponding oscillating crystal is made to oscillate depending on the inhalation stage of the patient. When the patient breathes through the connecting duct 14, i.e. mouth piece, passing into the patient,  
20 an air flow passes through the duct 15 opening into the open air, and the pressure is lowered in said duct 15. Said negative pressure (P) is transmitted through the connection 16a along the hose 16 to the pressure detector 36 provided in the regulator device 18.

25 Within the scope of the invention, such an embodiment of the invention is also possible in which the pressure detector 36 is placed directly in the atomizing device 11, e.g., in its air-inlet duct 15. In such a case, the conduit passing from the pressure detector to the regulator device 18 may be an electric one.

30

According to the invention, the apparatus further includes a timing device 44, in which the desired duration of atomizing can be preset by means of the atomizing-time regulator 30. When inhalation begins, the pressure in the duct 15 is lowered and the negative pressure  
35 concerned is detected by the pressure detector 36. The pressure detector 36 is connected to the timing device 44 electrically, and when the inhalation begins and the pressure detector 36 detects

1 said start of inhalation, the information is transmitted to the  
timing device 44, which, by means of the control logic and by the  
intermediate of the power oscillator 51 keeps the quartz crystal  
oscillating for the time preset in the timing device 44 by means  
5 of the regulator 30. When the time preset by means of the atomizing-  
time regulator 30 is exceeded, the control logic switches off the  
power oscillator and the ultrasonic oscillator 13 stops oscillating.  
Thereat, transfer of treatment agent into the air passing into the  
patient is also prevented.

10

The source of power may be either an accumulator or a mains supply  
connection unit, which is connected to the voltage connector placed  
in the rear panel of the device.

15 The apparatus has dual action. By means of the apparatus it is  
possible to perform atomization regulated by means of breathing or  
constant atomization.

In the atomization controlled by breathing, the little negative  
20 pressure produced at the beginning of the inhalation stage of the  
patient is transferred along the hose 16 to the regulator device  
18 and to the pressure detector 36 provided therein, said detector  
producing voltage data on the basis of the pressure. Said voltage  
is amplified in an amplifier 39, and that starts the timing device  
25 44. At the same time, the control logic 46 switches on the power  
oscillator 51 to oscillate at an ultrasonic frequency. This electric  
oscillation is passed along a coaxial cable 17 to the crystal,  
preferably a quartz crystal, which is placed in the ultrasonic  
oscillator denoted with the reference numeral 13. Said quartz  
30 crystal converts the electric oscillation to mechanical oscillation.  
The mechanical oscillation is transferred into the liquid placed  
in the liquid container 12 in its space 12a, which said liquid is  
atomized the more efficiently the higher the power regulator 20  
has been set.

35

With the voltage data received at the beginning of the exhalation  
stage, the timing circuit is reset to zero and the control logic

1 switches the power oscillator off operation and the atomizing is discontinued.

5 If the inhalation stage lasts longer than the time pre-set in the timing device, in such a case the timing device switches off the atomizing upon the preset time. The infrared detector 47 detects when the hose connector 25 is in its place inserted in the corresponding connector 26 of the regulator device 18. Thereat the apparatus operates automatically controlled by breathing.

10 In the case of constant atomization, the hose connector 25 of the pressure detector is not inserted in the corresponding connector 26 of the regulator device 18. In such a case, the apparatus atomizes constantly independently from breathing. In the case of constant atomizing, the power regulation operates in a similar way as  
15 in the case of atomization controlled by breathing.

The current monitoring electronics 62 operate with both modes of operation. When the liquid is exhausted in the liquid container  
20 12, the input current is lowered and thereby the control logic 46 switches off the atomizing. The indicator light 22 for exhausting of liquid indicates for the operator that the liquid has been exhausted. The apparatus does not restart atomizing until the current switch 19 has been turned to the off position for a moment.  
25 The current indicator light 21 indicates whether the current is on or off in the apparatus.

In the following, based on Fig. 3, the connections between the block diagrams will be described. Negative pressure arrives in the  
30 pressure detector 36 along the duct 37. The pressure detector transmits the voltage along the signal path 38 to the signal amplifier 39, which transmits the amplified voltage further along the signal path 40 to the trigger and reset device. From the trigger and reset device 41 there are outputs for triggering along the signal  
35 path 43 and for reset along the signal path 42. Said signal paths are connected to the timing device 44 as inputs. From the atomizing-time regulator 30, the preset atomizing time is passed along

1 the signal path 30a to the timing device.

From the timing device there is a signal path 45 further to the control logic 36. Further inputs to the control logic are the  
5 signal path 49 from the infrared detector 47, and the infrared detector 47 receives the signal detecting the presence of the connector 25 along the signal path 48 as an input. From the current monitoring electronics 62 the control logic 46 receives an input along the signal path 63. The output 50 of the control logic is  
10 connected to the power oscillator 51, to which the power regulation is also connected along the path 52. The current from the connector 29 comes through a fuse 55 along the current conduit 54 to the current switch 19 and further, from the current switch 19 the current is passed along the current conduit 56 to the current  
15 distribution point 57, and from there further to the current monitoring electronics 62 along the current conduit 58 and to the power oscillator 51 along the current conduit 59. From the power oscillator 51 there is an output 53 through the connectors 23,24 to the coaxial cable 17 passing to the crystal.

20

From the control logic 46 there is a connection to the signal light 22 indicating exhaustion of the liquid along the signal path 61, and correspondingly there is a signal path 60 to the current signal light 21 from the branch 56 connected to the current switch  
25 19.

30

35

1 WHAT IS CLAIMED IS:

1. Ultrasonic atomizer (10), which is intended for an inhalation  
treatment apparatus for persons suffering from respiratory diseases,  
5 said ultrasonic atomizer (10) comprising an atomizer device (11)  
provided with a drug container (12) as well as an ultrasonic  
oscillator (13) connected to the container (12), preferably an  
oscillating crystal, and said atomizer device (11) being provided  
10 with a duct passing into the patient, advantageously a mouth piece  
and an air-inlet duct (15), and said atomizer device (11) comprising  
a pressure detector (36) directly connected to the atomizer device  
or a duct (16) that is connected to the atomizer device (11) and  
passes to the pressure detector (36) and transmits the pressure of  
15 the breathing air, whereat the pressure detector (36) detects the  
changes in pressure resulting from inhalation or exhalation of the  
patient, whereby, starting from the beginning of the inhalation  
stage, atomizing can be switched on by the apparatus, and said  
ultrasonic atomizer (10) comprising a regulating device (18)  
connected to the atomizer device (11), by means of which said regu-  
20 lating device the operation of the ultrasonic oscillator (13) is  
regulated, c h a r a c t e r i z e d in that the regulating device  
(18) and the atomizing device (11) are interconnected by means of  
an electric connection (17), whereby, through said connection (17),  
an electric oscillation can be supplied to the crystal present in  
25 the ultrasonic oscillator (13), whereby said electric oscillation  
can be converted to mechanical oscillation of the crystal, and that  
means (36,39,41,46,51) are provided for switching on said ultrasonic  
oscillator (13) at least when the inhalation stage of the patient  
begins and that a timing device (44) is provided by means of which  
30 the duration of operation of the ultrasonic oscillator (13) can be  
regulated.

2. Ultrasonic atomizer as claimed in claim 1, c h a r a c t e r -  
i z e d in that the ultrasonic atomizer (10) comprises a control  
35 (39,43) between the pressure detector (36) and the timing device  
(44), whereby the signal arriving from the pressure detector (36)  
is fitted to start the counting of the timing device (44) from the

1 beginning of the inhalation stage, and that a regulator device  
(30) is provided for the timing device (44), whereby the maximum  
time of oscillation of the ultrasonic oscillator (13), preferably  
a crystal, can be preset in the timing device (44).

5

3. Ultrasonic atomizer as claimed in claim 1 or 2, c h a r a c -  
t e r i z e d in that the ultrasonic atomizer (10) comprises a  
control logic (46), which detects whether there is liquid in the  
liquid container (12), in which connection the control logic (46)  
10 detects a change in the quantity of liquid present in the container  
(12) on the basis of a change in the current or voltage of the  
oscillating crystal, and when a value above or below a certain  
current or voltage value is reached, the control logic (46) switches  
the crystal off oscillation, and that the control logic (46)  
15 comprises means by which it is ensured that the crystal cannot be  
switched on oscillation until after the main current circuit has  
first been switched off, whereby it is ensured that liquid is  
added to the liquid vessel (12).

20 4. Ultrasonic atomizer as claimed in any of the preceding claims,  
c h a r a c t e r i z e d in that the ultrasonic atomizer (10)  
comprises an infrared detector (47), which detects whether the  
duct (16) that passes to the pressure detector (36) and transmits  
the negative pressure (P) has been connected to the regulator  
25 device(18), and if the connector (25) of the duct (16) has not been  
connected to the regulator device (18), the infrared detector (47)  
detects this, and the apparatus (10) atomizes constantly.

5. Ultrasonic atomizer as claimed in any of the preceding claims,  
30 c h a r a c t e r i z e d in that connecting means (34,35) are  
provided, by means of which the ultrasonic atomizer (13) can be  
connected to the liquid container (12) detachably.

6. Ultrasonic atomizer as claimed in any of the preceding claims,  
35 c h a r a c t e r i z e d in that there are signalling means,  
preferably a signal light (22), which signals when the liquid level  
in the liquid container (12) has been lowered below a certain

13

1 minimum level or when the liquid has been exhausted from the liquid vessel (12).

5

10

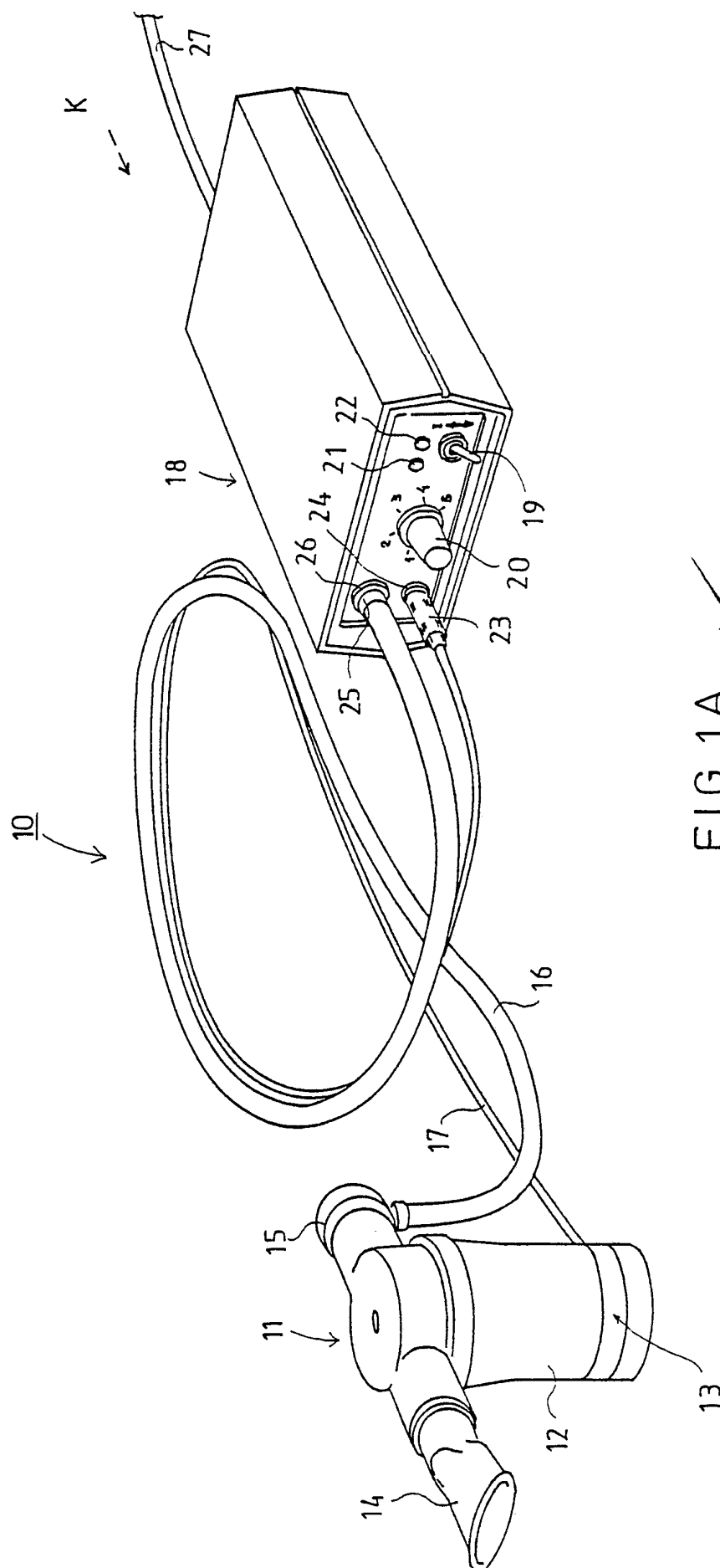
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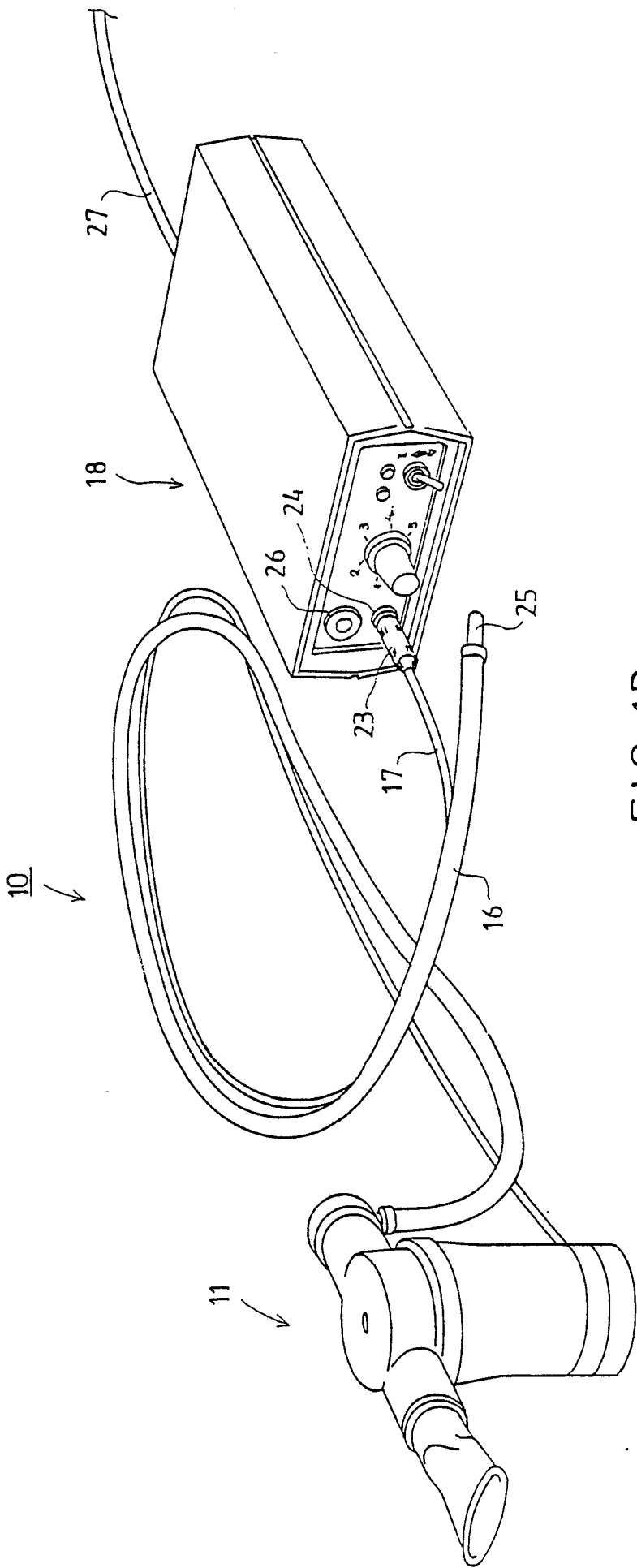


FIG. 1B

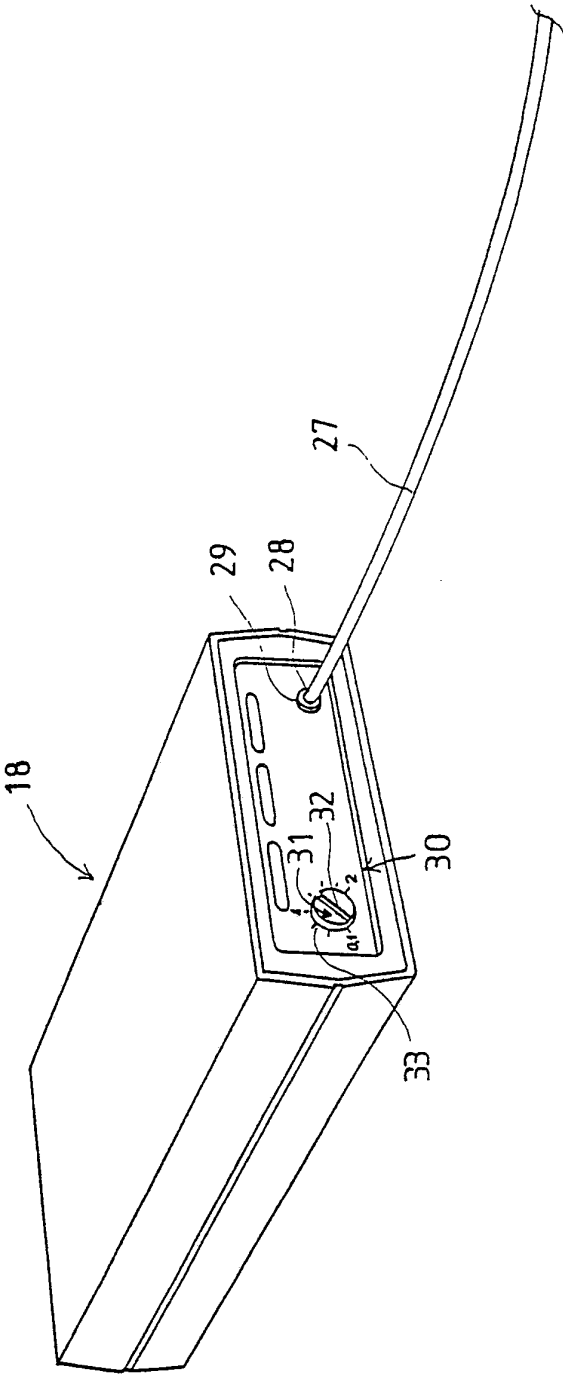


FIG. 1C

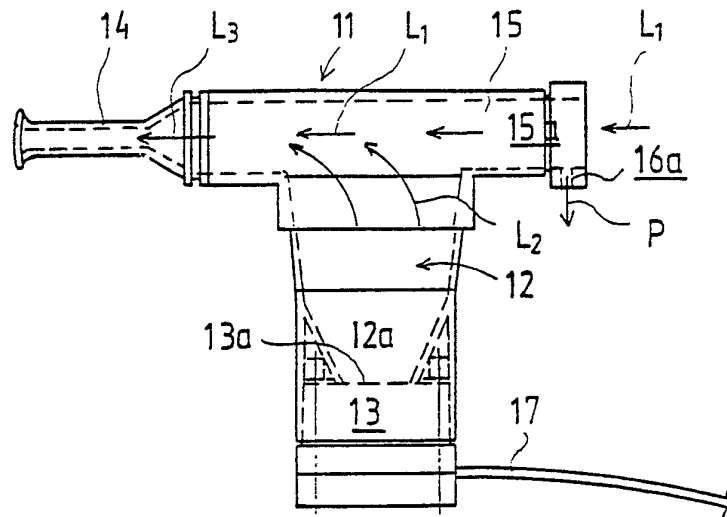


FIG. 2A

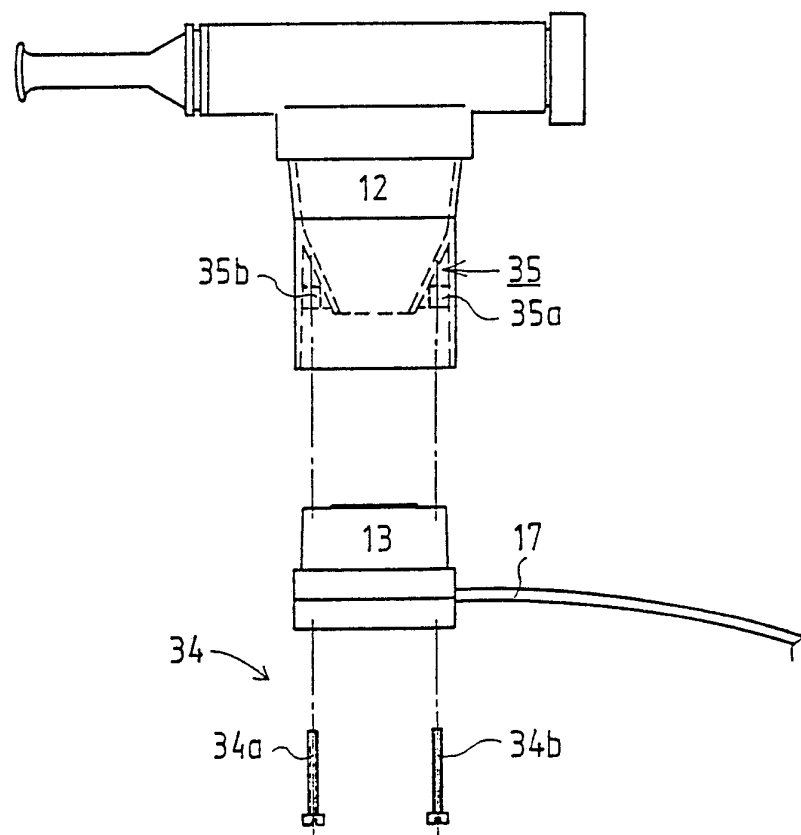


FIG. 2B

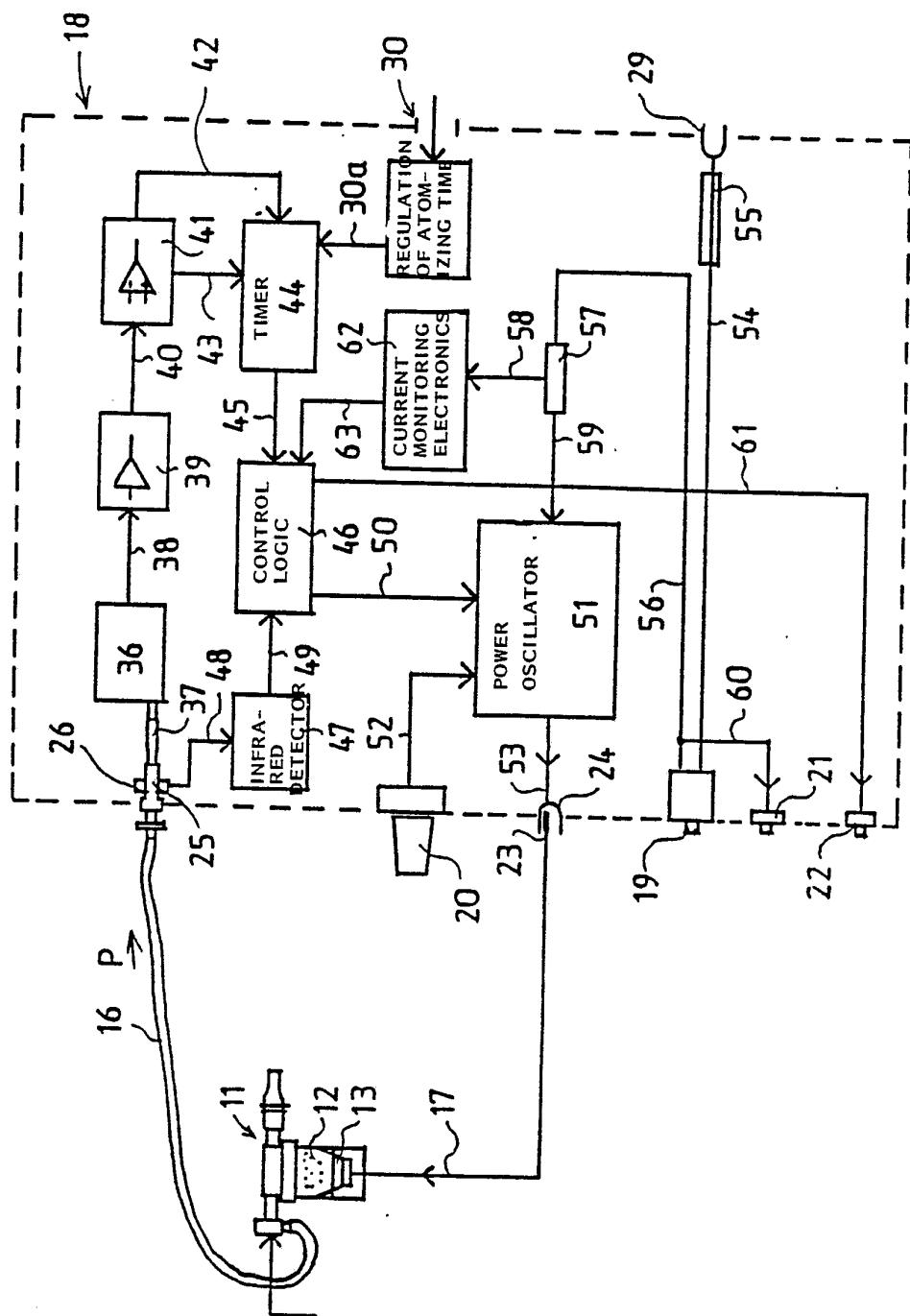



FIG. 3

# INTERNATIONAL SEARCH REPORT

International Application No PCT/FI88/00176

<b>I. CLASSIFICATION OF SUBJECT MATTER</b> (if several classification symbols apply, indicate all) *		
According to International Patent Classification (IPC) or to both National Classification and IPC 4		
A 61 M 15/00		
<b>II. FIELDS SEARCHED</b>		
Minimum Documentation Searched 7		
Classification System 1	Classification Symbols	
IPC 4	A 61 M 11/00-/08, 13/00, 15/00-/08; B 05 B 17/00-/06	
US C1	128:185, 186, 194, 200.11-16; 261:78, 81	
Documentation Searched other than Minimum Documentation to the extent that such Documents are included in the Fields Searched *		
SE, NO, DK, FI classes as above		
<b>III. DOCUMENTS CONSIDERED TO BE RELEVANT *</b>		
Category *	Citation of Document, 11 with indication, where appropriate, of the relevant passages 12	Relevant to Claim No. 13
A	DE, A1, 2 831 970 (STERN-ELEKTRONIK GMBH) 7 February 1980	1-2
Y	US, A, 3 561 444 (RAYMOND MARCEL GUT BOUCHER) 9 February 1971	1, 5
Y	US, A, 3 812 854 (ALAN S. MICHAELS ET AL) 28 May 1974	1, 3
A	US, A, 3 861 386 (RALEIGH J. HARRIS ET AL) 21 January 1975	1, 3, 5
A	US, A, 4 113 809 (RAYMOND L. ABAIR ET AL) 12 September 1978	1, 3
Y	US, A, 4 396 015 (ROBERT J. JOHNSON) 2 August 1983	1
<div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p>* Special categories of cited documents: 10</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="width: 45%;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"Δ" document member of the same patent family</p> </div> </div>		
<b>IV. CERTIFICATION</b>		
Date of the Actual Completion of the International Search	Date of Mailing of this International Search Report	
1989-04-10	1989 -04 1 2	
International Searching Authority	Signature of Authorized Officer	
Swedish Patent Office	 Leif Karnsäter	