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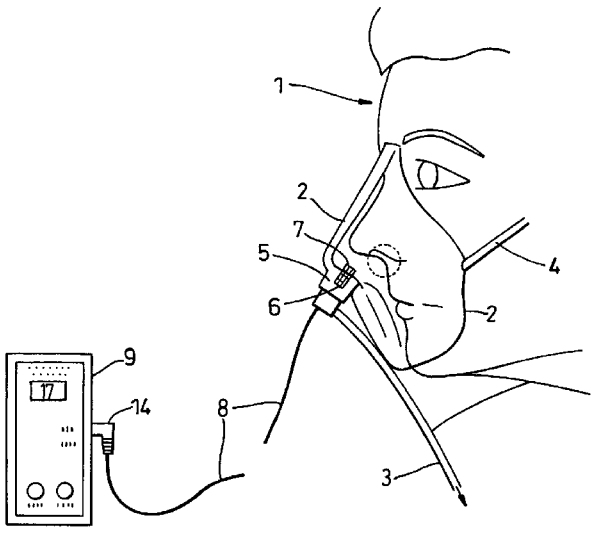
(54) Title: RESPIRATION MONITORING EQUIPMENT



(57) Abstract: Respiration monitoring equipment comprises, in one embodiment, a medical face mask (2), adapted to be located adjacent, or to cover a patient's nostrils and/or mouth; a transducer (7) adapted to be impinged by the patient's inspired and/or expired breaths, being sensitive to the presence and/or absence of a respiratory air flow, and being capable of emitting electrical signals in accordance with the presence and/or absence of a respiratory air flow; and a monitoring unit (9) electrically connected to the transducer (7) and responsive to the presence and/or absence of signals emitted by the transducer (7), and including means of triggering at least an alarm signal in the circumstances of non-detection of respiratory air flow within one or more predetermined parameters. Another embodiment provides for "cordless" monitoring, with a transmitter unit (11) at the face mask (2) and a receiver (12) at the monitoring unit (9).

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For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

Title of the Invention

Respiration monitoring equipment

Field of the Invention

This invention relates to respiration monitoring equipment particularly, but not exclusively for use with a subject, whether human or animal, potentially susceptible to sleep apnoea, extending also for use with patients in hospitals, ambulances and other paramedic situations. The equipment is equally suitable for veterinary use in the respiratory monitoring of animals.

Background of the Invention

Sleep apnoea is a major hazard in people of all ages and much innovative thinking has gone to develop various techniques for detecting the occurrence of apnoea. Many complex proposals have been made in efforts to provide suitable equipment. One example is described in EP 0484174, relies on battery power, on contacts and on other components that are subject to failure. Also, the possibility of using the piezoelectric and pyroelectric properties of PVDF (poly vinylidene flouride) films for developing transducers to sense the presence or absence of breathing, is described in US 5311875 which uses the PVDF film to sense the temperature differences between the inspired and expired breaths, and WO 97/05824. It is not known, however, whether any proposals have progressed beyond theory or experimentation.

Respiratory rate is one of the most important physiological parameters. It is a component of most medical and nursing records and is used in many clinical scoring

systems. Extremes of respiratory rate indicate the need for urgent intervention. Even today, in the case of non-intubated patients, the measurement of respiration rate is based on human observation alone, although this is known to be highly inaccurate.

Pyroelectric sensitive transducers incorporated in a face mask can be used to provide a quantitative measure of the respiratory rate in an easy and affordable manner. Furthermore, suitable circuitry for a respiratory monitor incorporating a PVDF sensor is described in our co-pending Patent Application GB 0201095.7.

As breathing could be wholly nasal, wholly oral, or part nasal and part oral, with changes between the modes, particularly during sleep, a problem for all workers in the field is optimum creation of a transducer to ensure that the patients' breath, if breathing, will always impinge upon the transducer, so obviating any false reading.

Object of the Invention

A basic object of the present invention is the provision of improved respiration monitoring equipment.

Summary of a First Aspect of the Invention

According to a first aspect of the invention, there is provided respiration monitoring equipment comprising:

(i) a medical face mask adapted to cover a patient's nostrils and mouth;

(ii) a transducer adapted to be impinged by the subject's inspired and/or expired breaths and sensitive to monitor the presence and/or absence of a respiratory air flow and capable of emitting electrical signals in accordance with the presence and/or absence of a respiratory air flow, and

(iii) a monitoring unit electrically connected to the transducer and responsive to the presence and/or absence of signals emitted by the transducer, and including means of triggering at least an alarm signal in the circumstances of non-detection of respiratory air flow within one or more predetermined parameters.

Preferred or Optional Features of the First Aspect

The transducer is of a pyroelectric and piezoelectric polymer.

The polymer is PVDF.

An encoded connector is provided between the face mask and the monitor.

The encoded connector, which assists the monitoring unit to differentiate between patient types, comprises a resistor, typically a 10K resistor being employed for an adult and a 2K resistor for a child. The encoded connector can also be used to differentiate between different locations (adult-forehead and child, cheek, skin); there could be a different algorithm for babies; finally it could also differentiate between different sensor types.

The transducer is carried by the face mask.

The face mask incorporates in an adaptor collar comprising a socket, in which socket the transducer is at least in part, housed.

The adaptor collar is of a synthetic material exhibiting elastomeric characteristics.

The transducer is a push fit into the socket of an elastomeric adaptor collar.

An electrical lead extends from the transducer to the monitoring unit.

The monitoring unit comprises means to provide a prescribed time period parameter.

The monitoring unit comprises means to provide minimum and maximum air flow

rate parameters.

In addition to triggering an alarm signal, the monitoring unit also comprises means to emit a "normal operation" signal, when respiratory air flow within the predetermined parameter(s) is sensed by the transducer.

Summary of a Second Aspect of the Invention

According to a second aspect of the invention, of independent significance, there is provided respiration monitoring equipment comprising:

(i) a device adapted to be located adjacent, and/or to cover, a patient's nose and/or mouth;

(ii) a transducer carried by the device in such a location as to be impinged by the patient's inspired and/or expired breaths, being sensitive to the presence and/or absence of a respiratory air flow, and being capable of emitting signals in accordance with detection of the presence and/or absence of a respiratory air flow;

(iii) a transmitter also carried by the device and activated by the presence or absence of signals from the transducer; and

(iv) a remotely located receiver and monitoring unit to receive signals from the transmitter and including means of triggering at least an alarm signal in the circumstance of non-detection of respiratory air flow within one or more predetermined parameters.

Advantages of the Second Aspect of the Invention

This aspect of the invention provides for "cordless" monitoring of a patient's

respiration, and consequently, ensures not only minimal interference with a patient, but also the absence of tubing, conduits, electrical leads etc avoids the presence of components that could become deranged, detached etc during sleep, or other mal-functions, resulting in the monitoring equipment falsely switching into an alarm mode.

Preferred or Optional Features of the Second Aspect

The device is an industry-standard, medical face mask.

The transducer is of a pyroelectric and piezoelectric polymer.

The polymer is PVDF.

The transducer is carried by the device.

The face mask incorporates an adaptor collar comprising a socket, in which socket the transducer is at least in part, housed.

The adaptor collar is of a synthetic plastics material exhibiting elastomeric characteristics.

The transducer is a push fit into the socket.

The transducer is housed in a length of tubing connected to, and extending from the device.

One parameter is a prescribed time period.

Other parameters are minimum and maximum air flow rates.

In addition to triggering an alarm signal, the monitoring unit also emits a "normal operation" signal, when respiratory air flow within the predetermined parameter(s) is sensed by the transducer.

Brief Description of the Drawings

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

Figure 1 is a diagrammatic side elevation of a first embodiment of equipment in accordance with the first aspect;

Figure 2 is a diagrammatic side elevation of equipment in accordance with the second aspect; and

Figure 3 is a diagrammatic side elevation of another embodiment of equipment.

Detailed Description of the Drawings

In all Figures, like components are accorded like reference numerals.

A head 1 of a subject is fitted with a transparent plastics, medical face mask 2 for the delivery of oxygen and/or allied therapy via a supply tube 3, the mask 2 extending over the subject's nose and mouth, and sealing against at least some portions of the subject's face under the influence of elastic loops 4 engaged over the subject's ears.

In the embodiment of Figure 1, the face mask 2 is provided with an adaptor collar 5 of elastomeric, synthetic plastics material. The adaptor collar 5 is provided with a socket 6, in which is located a transducer in the form of a PVDF sensor 7 responsive to the air flow of the patient's breathing rhythm, the sensor 7 being in close proximity to the patient's nostrils and mouth and the presence of the mask 2 ensuring that air flow resulting from expired breaths of the subject are directed towards, and impinge upon, the transducer/sensor 7. Any electrical output from the sensor 7 resulting from the sensing of the presence and/or absence of respiratory air flow within the predetermined

parameter(s) is fed by an electrical lead 8 to a monitoring unit 9 capable of switching to an alarm mode upon the sensor 7 emitting a signal indicative of lack of air flow within a predetermined parameter, such as a prescribed time period, or possibly lack of air flow above or below a predetermined flow rate threshold. The monitor 9 can be powered either by mains electricity or by battery and may emit a local audible and/or visual alarm, when triggered and/or, if required a local alarm eg at a nursing station. An encoded connector 14 is provided between the mask 2 and the monitoring unit 9. The encoded connector 14 comprises a changeable, or switchable resistor of eg 10K for an adult patient and 2K for a child patient. Selection of the resistor can also be used to differentiate between different locations (adult-forehead and child, cheek, skin); there could be a different algorithm for babies; finally it could also differentiate between different sensor types.

The embodiment of Figure 2 illustrates "cordless" or "wireless" respiration monitoring equipment, in which the sensor 7 is again embedded in adaptor collar 5, but the latter is provided with a transmitter unit 11 for any signals initiated by the sensor, and the monitoring unit 9 is provided with a receiver 12, the transmitter unit 11 being either battery or mains powered.

Whilst the use of a device in the form of a face mask 2 channels the patient's respiratory air flow to the sensor 7, it is a fact that a generally standard face mask 2 is unsuitable for some patients, in which case the arrangement illustrated in Figure 3 could provide a less intrusive solution, whereby the device comprises an electrical lead 8 secured eg by tapes 13 to the patient's face, extending from a sensor 7 located on the patient's upper lip to be impinged by air flow from the patient's nostrils and/or mouth, to the monitoring unit 9.

Output from the sensor 7 of Figure 2 could alternatively follow the teachings of Figure 2, ie by a "wireless" transmitter/receiver system.

CLAIMS

1. Respiration monitoring equipment comprising:

   (i) a medical face mask adapted to cover, a patient's nostrils and mouth;

   (ii) a transducer adapted to be impinged by the subject's inspired and/or expired breaths, being sensitive to the presence and/or absence of a respiratory air flow, and being capable of emitting electrical signals in accordance with the presence and/or absence of a respiratory air flow, and

   (iii) a monitoring unit electrically connected to the transducer and responsive to the presence or absence of signals emitted by the transducer, and including means of triggering at least an alarm signal in the circumstances of non-detection of respiratory air flow within one or more predetermined parameters.

2. Equipment as claimed in Claim 1, wherein the transducer is of a pyroelectric and piezoelectric polymer.

3. Equipment as claimed in Claim 2, wherein the polymer is PVDF.

4. Equipment as claimed in any preceding claim, comprising an encoded connector between the face mask and the monitoring unit.

5. Equipment as claimed in Claim 4, wherein the encoded connector comprises a resistor.

6. Equipment as claimed in any preceding claim, wherein the transducer is carried by the face mask.

7. Equipment as claimed in any preceding claim, wherein the face mask incorporates an adaptor collar comprising a socket, in which socket the transducer is at least in part, housed.

8. Equipment as claimed in Claim 7, wherein the adaptor collar is of a synthetic material exhibiting elastomeric characteristics.

9. Equipment as claimed in Claims 7 or 8, wherein the transducer is a push fit into the socket.

10. Equipment as claimed in any preceding claim, wherein an electrical lead extends from the transducer to the monitoring unit.

11. Equipment as claimed in any preceding claim, wherein the monitoring unit comprises means to provide a prescribed time period parameter.

12. Equipment as claimed in any preceding claim, wherein the monitoring unit comprises means to provide minimum and maximum air flow rate parameters.

13. Equipment as claimed in any preceding claim, wherein the monitoring unit also comprises means to emit a "normal operation" signal, when respiratory air flow

within the predetermined parameter(s) is sensed by the transducer.

14. Respiration monitoring equipment comprising:

(i) a device adapted to be located adjacent, and/or to cover, a patient's nostrils and/or mouth;

(ii) a transducer carried by the device in such a location as to be impinged by the patient's inspired and/or expired breaths to monitor the presence or absence of a respiratory air flow, and being capable of emitting signals in accordance with detection of the presence and/or absence of a respiratory air flow;

(iii) a transmitter also carried by the device and activated by the presence or absence of signals from the transducer, and

(iv) a remotely located receiver and monitoring unit to receive signals from the transmitter and including means of triggering at least an alarm signal in the circumstance of non-detection of respiratory air flow within one or more predetermined parameters.

15. Equipment as claimed in Claim 14, wherein the device is a medical face mask.

16. Equipment as claimed in Claims 14 or 15, wherein the transducer is of a pyroelectric and piezoelectric polymer.

17. Equipment as claimed in Claim 16, wherein the polymer is PVDF.

18. Equipment as claimed in any of Claims 14 to 17, wherein the transducer is carried by the device.

19. Equipment as claimed in Claim 15, and any claim appended thereto, wherein the face mask incorporates an adaptor collar comprising a socket, in which socket the transducer is at least in part, housed.

20. Equipment as claimed in Claim 19, wherein the adaptor collar is of a synthetic material exhibiting elastomeric characteristics.

21. Equipment as claimed in Claims 19 or 20, wherein the transducer is a push fit into the socket.

22. Equipment as claimed in any one of Claims 14 to 21, wherein the monitoring unit comprises means to provide a prescribed time period parameter.

23. Equipment as claimed in any one of Claims 14 to 22, wherein the monitoring unit comprises means to provide minimum and maximum air flow rate parameters.

24. Equipment as claimed in any one of Claims 14 to 23, wherein the monitoring unit also comprises means to emit a “normal operation” signal, when respiratory air flow within the predetermined parameter(s) is sensed by the transducer.

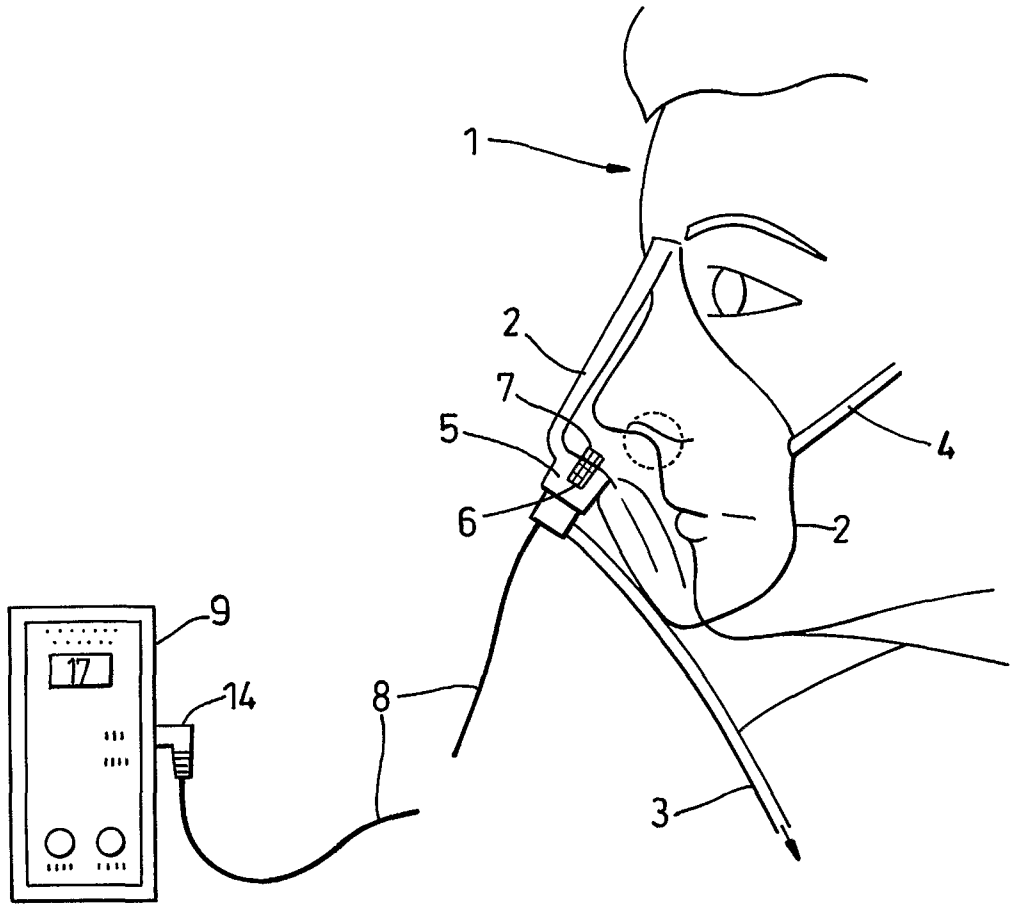

Fig. 1

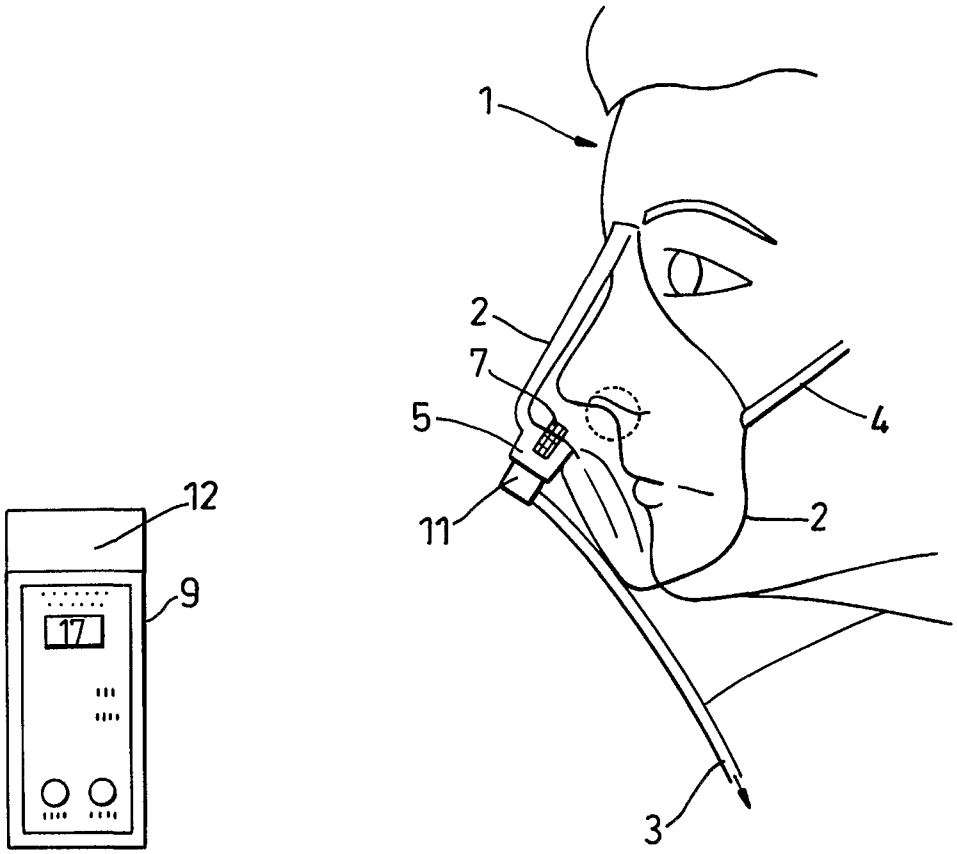

Fig. 2

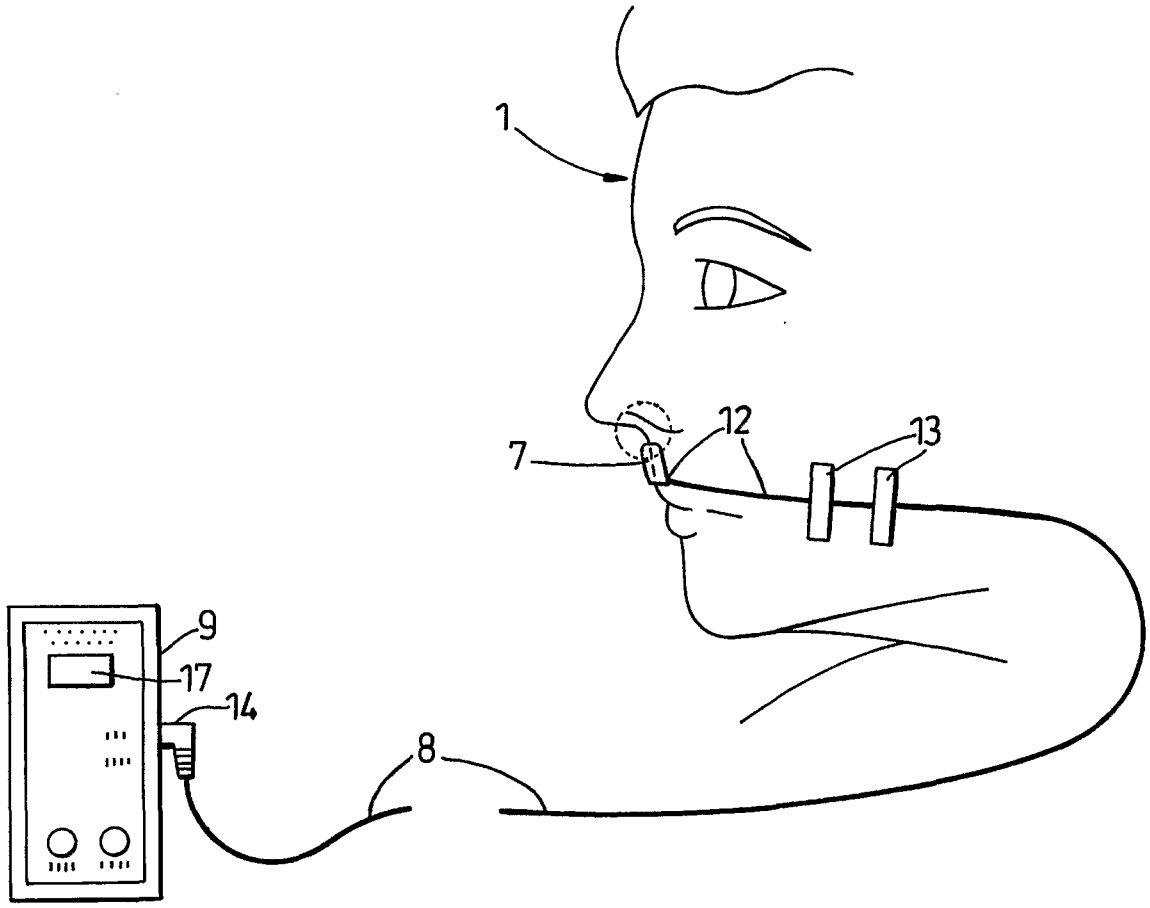

Fig. 3

INTERNATIONAL SEARCH REPORT

In al Application No
PCT/GB 02/02863

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61B5/087

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)
IPC 7 A61B A61M A61F

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, WPI Data, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 5 857 460 A (POPITZ MICHAEL D) 12 January 1999 (1999-01-12)	1-3,6, 10-18, 22-24
A	column 3, line 31 - line 45 column 3, line 66 -column 5, line 67 column 9, line 33 - line 65; tables 1-3	7-9, 19-21
Y	US 5 311 875 A (STASZ PETER) 17 May 1994 (1994-05-17) cited in the application abstract column 2, line 3 - line 38; tables 1-3	1-3,6, 10-18, 22-24

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[X] Further documents are listed in the continuation of box C.

[X] Patent family members are listed in annex.

° Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

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"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"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

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"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.

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
9 October 2002	16/10/2002

Name and mailing address of the ISA	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL – 2280 HV Rijswijk Tel. (+31–70) 340–2040, Tx. 31 651 epo nl, Fax: (+31–70) 340–3016	Weihs, J

INTERNATIONAL SEARCH REPORT

Inte[illegible]l Application No
PC[illegible]/GB 02/02863

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 01 43804 A (COMPUMEDICS SLEEP PTY LTD ;BURTON DAVID (AU)) 21 June 2001 (2001-06-21) abstract page 2, line 20 -page 3, line 23 page 5, line 24 -page 6, line 9 page 9, line 20 - line 24 page 10, line 10 - line 13; tables 1-4 -----	1,6,14, 18

INTERNATIONAL SEARCH REPORT

In ıal Application No
PCT/GB 02/02863

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5857460 A	12-01-1999	AU 2067497 A	01-10-1997
		WO 9733641 A1	18-09-1997
US 5311875 A	17-05-1994	NONE	
WO 0143804 A	21-06-2001	AU 1293201 A	25-06-2001
		EP 1237613 A1	11-09-2002
		WO 0143804 A1	21-06-2001

专利名称(译)	呼吸监测设备		
公开(公告)号	EP1404221A1	公开(公告)日	2004-04-07
申请号	EP2002732968	申请日	2002-06-19
[标]申请(专利权)人(译)	ÇLECT医疗		
申请(专利权)人(译)	C-LECT MEDICAL LTD.		
当前申请(专利权)人(译)	C-LECT MEDICAL LTD		
[标]发明人	DODDS DENNIS		
发明人	DODDS, DENNIS		
IPC分类号	A61B5/00 A61B5/087 A61B5/097		
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其他公开文献	EP1404221B1		
外部链接	Espacenet		

摘要(译)

在一个实施例中，呼吸监测设备包括医用面罩（2），其适于邻近或覆盖患者的鼻孔和/或口腔;换能器（7），适于受到患者的吸气和/或呼气呼吸的影响，对呼吸气流的存在和/或不存在敏感，并且能够根据是否存在发射电信号呼吸气流;监控单元（9）电连接到换能器（7）并响应换能器（7）发出的信号的存在和/或不存在，并包括在非环境中触发至少一个报警信号的装置检测一个或多个预定参数内的呼吸气流。另一个实施例提供“无线”监视，其中在面罩（2）处具有发射器单元（11）并且在监视单元（9）处具有接收器（12）。