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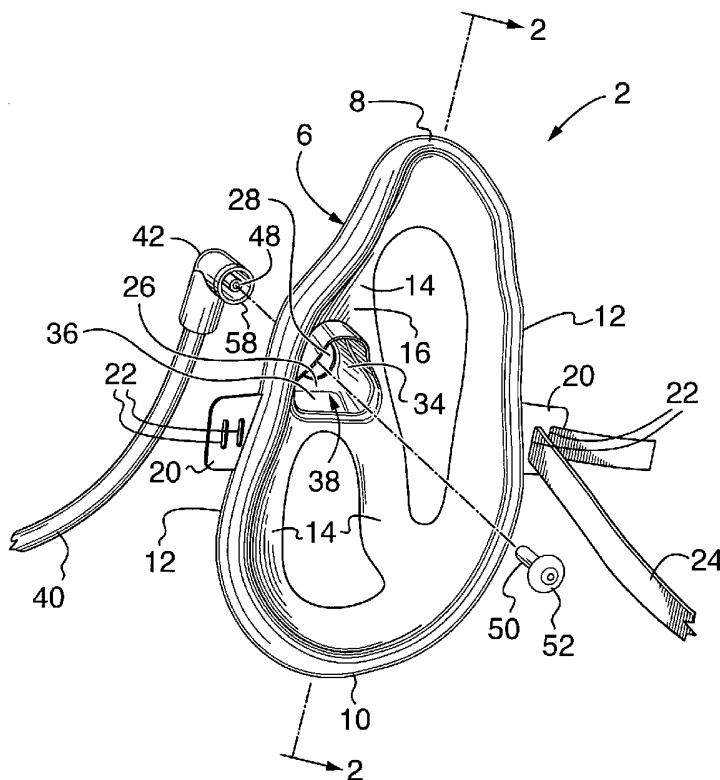
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(54) Title: PATIENT OXYGEN DELIVERY MASK



(57) Abstract: A mask for delivery of oxygen to a patient, the mask comprising a body having a peripheral portion, when in use to sit comfortably on a patient's face, a central portion, and bridge portions extending between the central portion and the peripheral portion and integral therewith, the central portion having an inner surface and an outer surface, the inner surface to be oriented towards the patient's face when the mask is in position and contoured so as to sit at a location spaced over the patient's nose and mouth, the inner surface of the central portion provided with a wall circumscribing a base, the wall and base being of generally concave configuration and circumscribing a centrally positioned oxygen delivery aperture extending through the central portion between the inner surface and the outer surface, the wall and base configured so as to act as an oxygen diffuser to direct the flow of oxygen generally towards the patient's nose and mouth when the mask is in use; a fastener to engage the mask to the patient so as to maintain contact between the patient's face and said peripheral portion when the mask is in use; the patient's head to hold the mask in position when in use; and means associated with the aperture and of the central portion releasably to receive and secure in position an oxygen delivery tube. In one embodiment,

exhaled carbon dioxide may also be collected for measurement. The mask according to the present invention provides comfort, reliability and efficiency in oxygen delivery to patients.



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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**PATIENT OXYGEN DELIVERY MASK****FIELD OF THE INVENTION**

The present invention relates to a novel mask for delivery of gases
5 such as oxygen or oxygen-enriched gas to a patient, and more particularly
to a mask which can be used to replace conventional oxygen masks and
nasal cannulae oxygen delivery systems.

BACKGROUND OF THE INVENTION

Conventional oxygen masks comprise tent like structures which are
10 strapped over the nose and mouth of the patient, often using an elastic
band or bands behind the patient's ears or head. Oxygen is fed from a
supply through a tube into the bottom portion of the mask at the front of
the patient. Many problems exist with such masks, including the fact that
many patients find them claustrophobic, the mask must be removed for the
15 patient to speak or eat, thereby discontinuing therapy, and the face mask
creates irregular and inefficient infusion of oxygen by the patient since
exhaled air from the patient is mixed with oxygen in the mask. Oxygen
masks can only be used for oxygen flows greater than 4 litres/minute
because exhaled gas accumulates in the mask, and, at lower flow rates
20 interferes with delivery of oxygen-enriched air to the patient.

Conventional nasal cannulae oxygen delivery systems employ an
oxygen delivery tube with tubular, open ended nasal prongs at the delivery
end of the tube for insertion into a patient's nasal passages. The oxygen
delivery tube and nasal cannulae are supported in position by a tube
25 wrapped about the patient's ears or head, making the system both difficult
to handle and uncomfortable since it applies downward pressure on the
patient's ears when the patient is in a seated position. As well, patients
often get nose bleeds from the dryness of the oxygen supplied through the
nasal cannulae. Patients also get sores on the ears, face and nose due to
30 the direct contact of the oxygen tubing with the skin. Nasal cannulae can
only deliver flows of 0.5 to 4 litres/minute.

- 2 -

Of background interest is Ketchedjian, U.S. Patent No. 6,247,470 issued June 19, 2001 which describes and illustrates an oxygen delivery apparatus comprising a headset to which is pivotally attached, for rotation in one plane, a flexible arm carrying tubular members for passing oxygen to a patient's mouth. The apparatus is also provided with a carbon dioxide monitoring system.

McCombs et al., U.S. Patent No. 6,065,473 issued May 23, 2000 describes a somewhat similar apparatus, for non-medical purposes, intended to dispense concentrated oxygen to users, the apparatus comprising an oxygen delivery nozzle attached by an arm extending from a flexible head band, to bathe the user's nose and mouth with oxygen, when in use. Laid-open German Application DE 43 07 754 A1, published April 7, 1994, teaches a system for controlled supply or removal of respiratory air from a user, which system incorporates a mask body held by a rigid air tube over the mouth and/or nose of the user, the air tube being pivotally adjustable in one plane, to enable proper positioning of the mask.

U.S. Patent No. 3,683,907 of Cotabish issued August 15, 1972 describes and illustrates a fresh air respirator, for use for example by miners, which comprises a cup, supported by pivotable arms in front of the face of the user, a stream of air being conducted to the cup to provide fresh air around the user's nose and mouth.

The applicant has developed a number of lightweight oxygen delivery systems for patients, as described for example in U.S. Patents Nos. 6,675,796 issued January 13, 2004, 6,595,207 issued July 22, 2003 and 6,450,166 issued September 17, 2002. Also, applicant's U.S. Design Patent Nos. D449,376 issued August 16, 2003 and D449,883 issued October 30, 2001 illustrate designs for such devices. All of these references feature oxygen diffuser devices, designed to create a turbulent oxygen flow, to be situated during use in front of the nose and mouth of a patient, and being held in that area by means of a mount such as a head band, to which is secured a rigid, but bendable oxygen delivery tube. The subject matter of each of these references is incorporated herein by reference.

Other references of general background interest include U.S. Patent No. 4,282,869 of Zidulka issued August 11, 1981, U.S. Patent No. 4,018,221

- 3 -

of Rennie issued April 19, 1977, U.S. Patent No. 5,687,715 of Landis et al. issued November 18, 1997, U.S. Patent No. 4,465,067 of Koch et al. issued August 4, 1984 and U.S. Patent No. 5,697,363 of Hart issued December 16, 1977, all of which describe and illustrate different types of head mounted
5 apparatus for delivering oxygen or other gases to a patient.

Most of these prior art devices intended for delivery of oxygen to a patient do not provide the ease of usage, both by health care workers and the patient, and reliability against unintended removal or dislodgement from position, as is required to permit widespread use by the health care
10 profession.

It is an object of the present invention to provide a more versatile, reliable and practical system for delivery of oxygen to patients.

SUMMARY OF INVENTION

In accordance with the present invention there is provided an
15 improved mask for delivery of oxygen to a patient. The mask comprises a body having a peripheral portion, when in use to sit comfortably on a patient's face, a central portion, and bridge portions extending between the central portion and the peripheral portion and integral therewith. The central portion has an inner surface and an outer surface. The inner surface
20 is oriented towards the patient's face, when the mask is in position, and is contoured so as to sit at a location spaced over the patient's nose and mouth. The inner surface of the central portion is provided with a wall circumscribing a base. The wall and base are of generally concave configuration and circumscribe a centrally positioned oxygen delivery
25 aperture which extends through the central portion between the inner surface and the outer surface. The wall and base are configured so as to act as an oxygen diffuser to direct the flow of oxygen generally towards the patient's nose and mouth when the mask is in use. The mask is provided with a fastener to engage the mask to the patient so as to maintain contact
30 between the patient's face and the peripheral portion of the mask when the mask is in use. Also, means are associated with the aperture and of the central portion releasably to secure in position an oxygen delivery tube.

- 4 -

In a further embodiment of the present invention, the mask additionally includes the oxygen delivery tube. It is releasably securable to the outer surface of the central portion of the mask so as to communicate with the oxygen delivery aperture. As well, a baffle is provided, the baffle
5 being constructed so as to be releasably seated over the oxygen delivery aperture on the inner surface of the central portion of the mask. The inner surface of the baffle is configured so as to assist, during use of the mask, in creating turbulence in an oxygen flow leaving the oxygen delivery aperture and assist in mixing oxygen with ambient air and thereby avoid a direct flow
10 of oxygen towards the patient's face.

In a yet a further embodiment of the present invention, the mask is further provided with an oxygen/carbon dioxide monitor tube releasably securable to the outer surface of the central portion of the mask, so as to communicate through the oxygen delivery aperture with an area above the
15 inner surface of the central portion during use of the mask for passage of air within the mask to an oxygen/dioxide monitor. The baffle is constructed so as to be releasably seated over the oxygen delivery aperture on the inner surface of the central portion of the mask. The baffle has a concave shaped wall and is configured and positioned so as to assist during use of the mask
20 in creating turbulence in an oxygen flow leaving the oxygen delivery aperture and assist in mixing oxygen with ambient air and thereby avoid a direct flow of oxygen toward a patient's face. A carbon dioxide intake is positioned within the concave shaped wall of the baffle, the carbon dioxide intake communicating with the carbon dioxide monitor tube.

25 The bridge portions of the mask, from a top of the mask to a bottom of the mask, may be configured in an inverted "Y" shape so that openings are provided towards the bottom and on both sides of the mask for unobstructed access to, and viewing of, a patient's mouth and other parts of the patient's face. The bridge portions comprise a web which is spaced
30 apart from the patient's face so as to maintain a sense of openness and lack of confinement for the patient. A suitable spacing is between 12 and 40 mm from the patient's face, for example when measured from the region between the patient's nose and upper lip.

- 5 -

In a further aspect, the oxygen delivery system comprises in general terms a mask for covering a portion of a patient's face, having a peripheral rim for contacting the patient's face to substantially surround only his mouth and nose region, a web-like mask body and an oxygen diffuser. A fastener
5 such as an elastic strap holds the mask firmly against the patient's face so as to cover the nose and mouth region. The body is semi-rigid or more preferably rigid and shaped so as to protrude outwardly away from the patient's face and to be spaced apart from the user's face when in use. The body may be provided in a range of sizes to fit different sized patients, such
10 as infants, normal adults and those with larger than average faces. The body supports a gas diffuser and a baffle or other gas obstructing member positioned in the path of the discharged gas so as to direct a turbulent stream of gas towards the user, such as oxygen or an oxygen-rich gas. The diffuser is positioned so as to direct the gas flow towards the user's mouth
15 and nose region at a non-oblique angle so as to generate a plume covering or substantially covering this portion of the patient's face. Thus, if the user is upright the gas flow is substantially horizontal. The mask body consists of a web having at least one and preferably a plurality of openings to permit access to the patient's face, for example an opening directly opposed to the
20 patient's mouth to permit the user to eat, drink and converse normally while wearing the mask, while also permitting medical personnel to easily administer medicines or a thermometer or the like to the patient's mouth. A plurality of relatively large openings also permits rapid discharge and dispersal of exhaled gases to the exterior of the mask. The location, size,
25 number and shape of the openings is dictated at least in part by the desired use of the mask, keeping in mind the comfort and convenience of the patient and the needs of medical staff for potential access to the patient's mouth or nose. For example, a mask for use by an infant may include openings which permit the feeding of the infant, wiping of his face and other
30 functions. Medical staff may also need to have quick access to the patient's nose or mouth, and it is convenient that this may be accomplished without removal of the mask. Another important aspect which dictates the opening size etc. relates to psychological factors, to minimize the anxiety felt by users. This can be particularly the case with infants, the elderly and those

- 6 -

suffering from diminished mental capacity. Thus, a variety of different configurations of the openings is contemplated. The mask has a size and shape so as to provide an oxygen-enriched zone surrounding the patient's nose and mouth.

5 The openings within the mask body can take on any convenient and suitable shape such as square, triangular or rectangular or other polygonal, or round or oval. The openings have a size range of between .25 square inches in area and 6.0 square inches and preferably between 0.25 and 3.0 square inches.

10 The size and shape of the mask is preferably optimized to minimize the surface area of the mask body and maximize the open area so as to enhance the user's comfort. This is accomplished by providing a relatively large area of the mask body being open, for example by providing a plurality of relatively large openings within the body. It has been found that the
15 mask body may comprise a relatively open structure if the gas diffuser mounted to the mask is positioned generally directly opposed to the patient's nose and mouth region and spaced apart from the patient's face so as to generate a plume of turbulent gas flow which covers the mouth and nose region of the patient. In order to generate this plume of turbulent
20 flow, the diffuser comprises a rear wall which fixedly receives a gas delivery tube, the mouth or nozzle of the tube discharging into the interior of the diffuser towards the patient's face. A gas flow disrupter is positioned between the nozzle and the patient's face. The rear portion is surrounded by a peripheral wall or flange extending towards the interior of the mask and
25 the patient's face so as to form a generally concave structure to assist in directing the gas flow towards the user. Preferably the rear portion is a wall which may be flat or curved, with the gas outlet entering the wall, for example at a generally central location. The peripheral wall may comprise a generally triangular shape as in the above-described embodiment, or any
30 other convenient shape. A turbulent flow pattern is generated by providing one or more obstructions associated with the diffuser in the path of the gas flow after it exits the gas delivery tube. For example, a baffle may be provided within the interior of the diffuser, which may comprise a mushroom-shaped structure which partly obstructs the path of the

- 7 -

discharged gas. Alternatively to a gas flow disrupter other means may provide a turbulent gas flow, for example the shaping of the nozzle itself.

The diffuser preferably conforms to dimensional and positional constraints described herein in order to provide a gas plume having

5 sufficient size to substantially cover the user's nose and mouth. It has been found that the diffuser should comprise a width of no more than 40 mm at its widest point and a maximum height of 80 mm and preferably smaller than this in both dimensions. The diffuser is positioned within the mask such that no part of the diffuser is outside of a region defined in relation to a

10 point between the base of the user's nose and his upper lip, the region comprising a space 40 mm above and below this point in the vertical plane when the mask is vertical and 20 mm on either side of this point in the side-to-side direction. Preferably the diffuser is centered in side to side and vertical dimensions relative to this point. Further, the diffuser is preferably

15 mounted so as to leave a gap between the diffuser and the user's face of between 12 and 40 mm as measured from the user's skin surface between upper lip and nose. Preferably, the diffuser is generally triangular in shape with base downwardly, and preferably the width and height are between 20-30 mm and 17-27 mm respectively.

20 It has been found that a mask which is provided with a diffuser that conforms to the dimensional requirements described above, and which is positioned within the mask as described, and wherein the mask body includes a plurality of openings or cut-outs, provides an optimal level of comfort, with minimal coverage of the face while still permitting an oxygen-

25 enriched zone fully covering the user's nose and mouth. Preferably the cut-outs within the mask body should comprise at least 30 per cent of the total surface area of the body when measured as if the body were considered to comprise a plane surface, and still more preferably the cut-outs comprise between 30 and 80 per cent of the total surface area. In a still more

30 preferred version the range is more narrowly defined as being between 50 and 80 per cent, and yet more narrowly as between 60 and 75 per cent.

An oxygen delivery mask permits an efficient delivery of oxygen-enriched gas in a comfortable fashion if it includes a combination of an oxygen diffuser opposed to the user's nose and mouth region and directed

- 8 -

so as to direct a turbulent gas plume directly at the nose and mouth, and at least one opening within the mask body. Without intending to be restricted to a theory, it is believed that the user's breath inspiration generates negative pressure within the mask interior thereby creating a mixing effect whereby exterior air is drawn through the mask openings into the turbulent gas plume via a venturi effect. Upon exhaling, a positive pressure is generated within the mask and the exhaled air is exhausted through the openings. This maintains within the mask interior an oxygen-rich and CO₂-poor environment, which enhances the user's comfort level and is more medically effective.

Positioning of the diffuser opposed to the nose and mouth region so as to direct a turbulent plume directly towards the user's nose and mouth also permits an oxygen-enriched gas plume which is of generally equal gas makeup over both the mouth and nose. Thus, user inhales the same gas mixture whether breathing through mouth or nose. The diffuser is suitable shaped and positioned so as to cover both the nose and mouth of a typical user.

The oxygen delivery mask of the present invention provides an easy to use, comfortable, reliable and efficient mask for delivery of oxygen to a patient. As well, since this mask construction does not provide complete enclosure over the patient's nose and mouth, there is less chance of claustrophobia.

In a further aspect, although the invention has been described for use for delivering oxygen or an oxygen-enriched gas it will be seen that with modifications it may be used for other medical applications such as delivery of anesthetic or other gases to a patient.

BRIEF DESCRIPTION OF THE DRAWINGS

These and other advantages of the invention will become apparent upon reading the following detailed description and upon referring to the drawings in which:

- 9 -

FIGURE 1 is an exploded perspective view from the rear of one embodiment of the oxygen delivery mask according to the present invention;

FIGURE 2 is an elevational section view of the mask of Figure 1, along
5 lines 2-2 of Figure 1;

FIGURE 3 is a perspective view from the front of the mask of Figure 1;

FIGURE 4 is an elevational section view of an alternative embodiment of oxygen delivery mask in accordance with the present invention, including
10 a carbon dioxide monitoring function; and

FIGURE 5 is a front elevation view of the mask of Figure 1.

FIGURES 6A, 6B and 6C are front elevational views of further alternative embodiments of the mask.

FIGURE 7 is a perspective view of a further embodiment of the mask;

15 FIGURE 8 is a further perspective view of the embodiment of Figure 7.

FIGURE 9 is a side elevational view of the embodiment of Figure 7.

While the invention will be described in conjunction with illustrated embodiments, it will be understood that it is not intended to limit the invention to such embodiments. On the contrary, it is intended to cover all
20 alternatives, modifications and equivalents as may be included within the spirit and scope of the invention as defined by the specification as a whole including the appended claims. It will be understood that dimensions and relative dimensions described and illustrated herein are intended to be by way of example only of specific embodiments and unless otherwise indicated
25 are not intended to limit the scope of the invention. References herein within both the description and claims to specific directions and positions such as "horizontal", "vertical", "forward" and the like are intended only to provide a convenient means of description and are intended to be in reference to the mask in an upright forward-facing position, as if it were
30 worn by a patient in a standing position. Naturally the mask may be used on a patient in any orientation.

- 10 -

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

In the following description similar features in the drawings have been given similar reference numerals.

Turning to Figures 1 and 2 there is illustrated an oxygen delivery mask 2 which represents an embodiment of the present invention. Mask 2 is made up of a body 4 having a peripheral portion 6 with a top 8 and a bottom 10. Sides 12 extend between top 8 and bottom 10. As can be seen in Figure 3, peripheral portion 6, when mask 2 is in use, rests on portions of a user's face both above the user's nose (top 8) and on the user's chin (bottom 10). Integrally formed with peripheral portion 6 are bridge portions 14 which integrally connect with a central portion 16. Bridge portions 14 and central portion 16 have an inverted "Y" shaped configuration (from top to bottom of the mask), when viewed from the front (Figure 5), providing unobstructed access to and viewing of the patient's mouth and other parts of the patient's face, so that for example, the patient may eat and drink without removing the mask. Of course other configurations of bridge portions may be provided as desired or appropriate, such as, for example, an "X" shape, a "+" shape or "T" shape. Peripheral portion 6, bridge portions 14 and central portion 16 are preferably made of a fairly soft, semi-rigid plastic material. The term "semi-rigid" refers to a material that is generally resilient but which provides sufficient rigidity to substantially maintain its shape when in normal use. The bridge portions 14 thus have sufficient rigidity to retain the diffuser 38, discussed below, in position. Suitable materials for the peripheral, bridge and central portions 6, 14 and 16 include plastics such as PVC, silicone, foam, polystyrene and any other thermoplastic elastomers. Tabs 20 extend outwardly from sides 12, and are provided with, for example, slots 22 in which may be adjustably secured ends of an elastic strap 24 for releasably securing the mask 2 in position on a user's face (Figure 3). As will be understood from Figures 1 and 3, peripheral portion 6, bridge portions 14 and central portion 16 are contoured so as to rise from base 26 of peripheral portion 16 in a curved contour so that central portion 16 sits spaced over the nose and mouth of the patient

- 11 -

when the mask 2 is in position. A circular aperture 28 extends through central portion 16 from outer surface 30 to inner surface 32.

Integrally formed on inner surface 32 of central portion 16, is a triangular wall 34, extending about a base 36 which circumscribes circular aperture 28. This wall 34 and base 36 together form a shape which is of generally concave configuration, with one of the apexes of the triangle formed by wall 34 being oriented towards top 8 of mask 2 and the other two apexes oriented towards bottom 10. Thus, a generally triangular shape is formed by the mask, albeit generally with rounded corners, which approximates the human nose and mouth region, so as to generate an oxygen enriched region within the mask which substantially covers the user's nose and mouth. This wall and base form a diffuser 38 which has a similar function to the diffuser construction described and illustrated in applicant's earlier patents and applications referred to previously herein. The diffuser 38 preferably has a maximum width of about 40 mm, with a more preferred maximum width of 30 mm and still more preferred maximum being 25 mm. A maximum height is preferred to be 80 mm, with increasing preferences being 27 mm and about 22 mm.

In the embodiment of mask illustrated in Figures 1 to 3, an oxygen delivery tube 40 is secured to a connection in the form of a rigid elbow 42, elbow 42 being rotatably secured by an appropriate, conventional securing means such as frictional engagement in aperture 28 or its snapping into an undercut 44 about aperture 28 on the outer surface 30, so that it can pivot about the circumference of aperture 28 (Figures 3 and 5). Elbow 42 provides a passageway 46 for delivery of oxygen, during operation of the device, into diffuser 38 on the inner surface 32 of central portion 16. The diffuser 38 and associated baffle are shaped so as to generate a plume of turbulent flow of oxygen-enriched gas which surrounds the user's nose and preferably also his mouth. The generation of this plume requires a suitable discharge velocity of gas through the nozzle. This may be accomplished by providing a gas discharge rate between 1- 15 liters/min. Formation of the plume is related to the physical design of the nozzle, diffuser and baffle. The blending of the oxygen and atmospheric oxygen is related to the gas velocity. Without wishing to be tied to any particular theory of operation,

- 12 -

the physical properties of the device required to generate a plume include the shape, size and positioning of the baffle which serves to deflect the gas flow back down, then bounce off sidewalls to create this plume. The direction of the gas flow exiting the diffuser is substantially directly towards the user's face so as to strike the user at an angle generally perpendicular to the user's face. The gas flow is thus non-oblique in relation to the user's face. The diffuser 38 is positioned so as to direct the flow most strongly at the user's nose and mouth region. It has been found that the configuration described herein efficiently provides an oxygen-enriched zone in the region surrounding the user's mouth and nose, while permitting relatively large cut-outs within the mask.

The baffle 50 is positioned within the path of the gas exiting the nozzle and has a shape and size which is suitable for interrupting the linear gas flow exiting the nozzle so as to generate turbulence. It will be seen that a variety of sizes and shapes will achieve this function. In the illustrated example, the baffle comprises an upstanding stem 48 within elbow 42 which provides a means for releasable attachment thereto of post 50 of mushroom shaped baffle 52. As can be seen in Figure 2, the inner end of baffle 52 has a curled back conical lip 54 on its head 56, the underside of this lip being in line with oxygen passing from aperture 26 at the inner end 58 of elbow 42. This curled back conical lip 54 is of a size and configuration, with respect to wall 34 of diffuser 38, such that turbulence is generated in the stream of oxygen passing from elbow inner end 58 and aperture 28, creating a plume of oxygen enriched air at the patient's nose and mouth when the mask is in position.

In the alternative embodiment of mask 2 illustrated in Figure 4, while mask body 4 and integral diffuser 38 are of a similar configuration to those of Figures 1, 2 and 3, in addition to an oxygen delivery tube 40 passing into elbow 42, elbow 42 is configured to have an oxygen/carbon dioxide monitor tube 60 secured to it, which tube communicates with a separate oxygen/carbon dioxide monitor passageway 62 extending within elbow 42 to its inner end 58. Oxygen/carbon dioxide monitor tube 60 and passageway 62 are separate and independent from oxygen delivery tube 40 and oxygen delivery passageway 46. Oxygen from delivery tube 40 is again delivered

- 13 -

through elbow 42 to aperture 28 and the inside of mask 2 and the wall 34 of diffuser 38 circumscribes this aperture 28 and directs the flow of oxygen generally outwardly from diffuser 38.

5 In this embodiment, baffle 64 has a hollow post 66, the hollow center communicating with an opening 68 on the inside of baffle 64, and with the oxygen/carbon dioxide monitor passageway 62 and tube 60.

Head 70 of the baffle 64 circumscribes the opening 68, the head being of a concave shape formed by wall 72. This head 70 fills a significant part of the interior of diffuser 38. Wall 72 extends outwardly beyond the
10 edges of wall 34, and generates the necessary oxygen turbulence to provide an effective plume of oxygen for delivery to the nose and mouth area of the patient when the mask 2 is in position. At the same time however, an effective oxygen/carbon dioxide monitoring of the patient's exhaled breath is permitted through the oxygen/carbon dioxide monitor opening 68 within
15 head 70.

In further embodiments shown in Figures 6A, 6B and 6C, a mask 2 comprises a body 4 having a peripheral rim 6. The rim 6 comprises a material such as PVC or silicone which provides sufficient rigidity to permit the mask to maintain its shape while permitting a degree of resiliency to
20 permit the mask to comfortably conform to a user's face. The rim is provided with an outer surface for contacting the patient's face of a material which is soft and pliable plastic with a Durometer range of 20-100 Shore A. The rim preferably has a roughly triangular shape to generally conform to a human nose and mouth region, namely a broad base and a narrow apex,
25 with rounded corners. The rim 6 may alternatively comprise a bendable material such as aluminum which retains its shape when flexed, which may be provided to the user in a shape which roughly follows the contours of a typical human face. The mask may be provided in a plurality of rim sizes to fit different classes of users, for example infants and small, medium and
30 large adults.

The body 4 comprises a web of shaped substantially rigid material. This may comprise a semi-rigid material such as that described above in connection with the first embodiment. However, if the body 4 is provided with a more lace-like structure defined by a large number of cut-outs,

- 14 -

described below, a more rigid material will be desired such as TPE-
"thermoplastic elastomer". In general, the body has sufficient rigidity to
maintain its generally cup-like configuration and to support the diffuser 38
which is fastened to and supported by the body, at a position spaced apart
5 from the user's face. Because the relative position of the diffuser 38 is
important to the functioning of this version, the body should have sufficient
rigidity to maintain the central position of the diffuser 38 during normal use
of the mask. The body 4 may possess a degree of resiliency to enhance
user comfort and daily functions and to permit the mask to better conform
10 to the user's face. The body comprises a plurality of cut-outs 60 which may
be of any shape and size suitable to serve several desired functions. These
functions include permitting the user to speak, eat and drink with a
minimum of obstruction, wipe or blow his nose, scratch or otherwise touch
his face, kiss and other normal activities. As well, it is contemplated that
15 others such as health professionals may rely on the cut-outs 60 to feed or
provide fluid to the user or for other functions. Thus, it is contemplated that
relatively large openings are provided in the region of the patient's nose and
mouth. However, other arrangements of openings are possible such a
larger number of smaller openings. The openings must not be so many or
20 large as limit the ability of the web to fixedly support the diffuser 38.

The diffuser 38 is mounted within the mask body 4 so as to be
centered vertically and horizontally (side-to-side) above a point which when
the mask is worn by a person, is about halfway between the base of the
nose and the upper lip.

25 Preferably, the cut-outs 60 comprise at least 50 per cent of the total
surface area of the body 6 (when measured with the surface area including
both open and closed areas) and still more preferably the cut-outs comprise
between 50 and 80 per cent of the total surface area. In a still more
preferred version the range is more narrowly defined as being between 60
30 and 75 per cent.

The diffuser 38 protrudes through and is supported by the body 4 at a
position generally opposed to the patient's nose and mouth region. The
diffuser and associated baffle are substantially as described in connection
with the embodiments. However, it has been found that if the diffuser body

- 15 -

conforms to particular size and positioning limits, it effectively generates a zone or region of oxygen-enriched gas in the region of the patient's nose and mouth, regardless of the number, size and shape of the cut-outs 60. For this purpose it is desirable to fabricate the body 4 from a relatively rigid

5 material such as polystyrene, thermoplastic elastomer or polycarbonate. The diffuser 38 receives a gas supply from supply line 40, connected via elbow 42 in the same manner as described above. A mushroom-shaped baffle within the interior of the diffuser 38 assists in the dispersal of gas. It will be further apparent to those skilled in the art that the diffuser 38 may

10 comprise a range of sizes and shapes. However, in order to generate the desired region of oxygen enriched gas the diffuser comprises a cup-like body opening towards the user. It should have a maximum size and positioning within the mask that permits the diffuser to fit entirely within a region of the mask defined by reference to an imaginary point on the user's face between

15 the base of the nose and upper lip, in the midline of the face, with the diffuser entirely fitting within the space defined by 20 mm on either side of this line horizontally and 40 mm above and below the this point vertically (when the user is upright). Preferably the diffuser is sufficiently small so as to permit some slippage of the mask while still remaining within this region,

20 for example as described above a generally triangular configuration about 25 mm wide at its base and about 22 mm high. The diffuser is also positioned within the mask such that the gas discharge nozzle is between 12 and 40 mm displaced forwardly of the user's face measured from the area between the upper lip and below the nose of a users face. The body 6 must

25 have sufficient forward protrusion so as to position the rear rim of the diffuser so that it does not contact the nose of the typical user. The spacing thus required will vary somewhat with different sizes of masks. For example, the diffuser may protrude rearwardly into the interior of the body 4 by about 2 mm, and the body thus has an overall depth of about 15 mm.

30 As well, the mask design of the present invention allows a patient to drink, eat, be suctioned and speak, without removal of the mask. Also, exhaled air does not collect in the area in front of the patient's nose and mouth and interfere with the mask's operation, as in the case of conventional oxygen masks, since exhaled air easily passes to the

- 16 -

surrounding environment through the spaces between the bridge portions and the peripheral portion of the mask.

A further embodiment of the mask is illustrated at Figures 7, 8, and 9, wherein the oxygen delivery mask 2 is provided with a removable packing support and retainer strap 100. The strap 100 is made from a semi-rigid material such as polypropylene or other plastic, and comprises a generally elongate body having opposing yoke-shaped ends defined by spaced apart arms 102. The arms 102 each have inside edges including inwardly-extending projections 104. The projections 104 are spaced apart to permit a portion of the mask to be inserted past the projections 104, into a partially enclosed space 105 within the yoke-shaped end regions which thereby retains the strap to the mask. The end regions 102 are intended to releasably receive spaced apart portions of the mask that conveniently fit within the interior space 105 and are retained therein by the projections 104. For example, the opposed ends may respectively receive the bridge portion 14 and the elbow 42 of the mask 2. It will be seen that the strap 100 must be bent into a substantially U-shaped form for this purpose. The strap 100 when flexed into this position is useful during shipping and storage of the mask to help maintain its proper shape and avoid compression or crushing. The strap 100 may be removed prior to conventional use of the mask. However, the strap 100 is also intended to serve a second function and for this purpose it is left in place on the mask, as will be described below.

The strap 100 includes a circular opening 106, located between the opposing ends 102. The opening 106 has a suitable dimension to snugly receive an end of a laryngeal mask 110. The laryngeal mask is a tube-shaped article intended to be inserted down the throat of a patient who has an obstruction in the throat, thereby assisting in the breathing for such patient. The laryngeal mask (shown in part in the figures) includes a first end to receive oxygen from the oxygen diffuser. The first end includes a rigid tubular insert 112, which is fitted within a flexible tube 114. The insert 112 includes an outwardly-protruding flange 116. The insert 112 fits snugly within the opening 106, up to the limit defined by flange 116, which serves as an effective stopper for the tube. The snug fit of the insert 112

- 17 -

provides a friction fit within the strap 100. As will be seen in Figure 9, the open end of insert 112 is spaced apart from and opens towards the diffuser 38, so as to receive gas from within the plume of gas generated by the diffuser 38, when in use.

5 The strap 100 thus serves two functions, the first being as a packing support member to prevent crushing or damage to the mask, for example during transportation, the second purpose being to retain a laryngeal mask, for use by patients in need of the same. Optionally, the strap 100 may serve either or both of these functions.

10 It will be seen that the strap 100 may be made of any suitable rigid or semi-rigid material, and polypropylene is described herein merely as an example of a suitable material. Similarly, the precise dimensions and shape of the strap 100 are subject to reasonably wide variation. For convenience, the strap is somewhat tapered so as to have a wider middle region and
15 narrower ends. This permits the strap to be easily fastened to the mask, at either end of the strap, while permitting the opening 106 within the middle region of the strap with sufficient material on either side of the opening to prevent tearing or cracking of the strap, in use.

 In tests which have been done and proven efficacy of the mask
20 designs according to the present invention, it has been determined that patients find the mask according to the present invention to be far more comfortable than conventional oxygen masks. Unlike conventional masks, users cannot feel oxygen being delivered to their nose and mouth area, and enjoy the compactness of the mask. Technically, lower flow rates of oxygen
25 to a patient through the mask of the present invention can be achieved, with as much or greater oxygen concentration in the air being delivered to the patient, as compared to conventional oxygen masks. In this manner, the mask according to the present invention provides both comfort and efficiency to patients which providing optimal blood oxygen saturation in a
30 cost effective manner. Flow rates ranging from .5 liters to 15 liters per minute have proven suitable providing a far greater range of possible flow rates than available through conventional oxygen delivery devices. The following examples describe tests performed with the mask.

- 18 -

Example 1

A study was conducted, comprising a randomized, cross-over, single-blind study of patients having the following inclusion criteria:

- 5 - chronic pulmonary disease requiring supplemental oxygen therapy;
- stable oxygen requirement (unchanging over a three hour period);
- 18-80 years of age.

10 Excluded were patients whose oxygen requirement was unstable (i.e. changing hourly) or who could not tolerate being deprived of supplemental oxygen for five minutes or less.

Protocol

Continuous monitoring of minute ventilation (almost non-traceable oxygen movement) by RespiTrace™, SaO₂ (% oxygen saturation of hemoglobin), HR, nasal/oral flow, PO₂ (mmHg of oxygen in arterial blood) and PCO₂ (% CO₂) at the lip, O₂ flow and TcPCO₂:

- 5-10 min washout (point when the waveform tracing begins to have a flatter peak) / 5 min baseline (room air);
- Mask #1 (supplemental oxygen; referred to herein as "OxyMask™" or "OM" and comprising an embodiment of the invention);
- 20 - 15-30 min at 4-5% above baseline SaO₂;
- 15-30 min at 8-9% above baseline SaO₂;
- 5-10 min washout / 5 min baseline (room air);
- Mask #2 (supplemental oxygen; referred to herein as "venturi" or "V", comprising a prior art mask);
- 25 - 15-50 min at 4-5% above baseline SaO₂;
- 15-30 min at 8-9% above baseline SaO₂.

Data analyzed using two-way analysis of variance (ANOVA) and paired t-test:

- 30 - p values <0.05 were considered statistically significant.

Patient Demographics

N=13 patients with chronic pulmonary disease using supplemental oxygen via nasal cannula.

4 male, 9 female.

5 age: 56 ± 16 (range: 28-79).

BMI: 35.0 ± 12.3 .

O2 requirement: 2.3 ± 1.3 Lpm (rest), 3.4 ± 1.6 Lpm (exercise).

Pulmonary Function Tests

| Spirometry | | |
|--|-------------------|---|
| | Measured | % Predicted (i.e. as compared to normal values) |
| FVC (forced vital capacity), l | 1.87 ± 0.66 | 57.38 ± 13.20 |
| FEV1 (forced expiratory volume in 1 second), l | 1.22 ± 0.56 | 51.54 ± 21.50 |
| FEV1/FVC | 65.06 ± 19.29 | - |
| V50 (flow at 50% expired FVC), l/sec | 1.38 ± 1.23 | 42.15 ± 34.71 |
| V25 (flow at the last 25% of expired FVC), l/sec | 0.40 ± 0.32 | 25.23 ± 18.66 |

- 20 -

| | | |
|------------------------------------|-------------|---------------|
| VC (vital capacity, not forced), l | 2.00 ± 0.80 | 60.20 ± 15.05 |
|------------------------------------|-------------|---------------|

| Arterial Blood Gases | |
|------------------------------------|--------------|
| pH | 7.38 ± 0.05 |
| H ion, nmol/l | 41.63 ± 5.01 |
| pCO ₂ , mmHg | 47.50 ± 6.65 |
| pO ₂ , mmHg | 50.88 ± 7.92 |
| Bicarbonate, mmol/l | 29.13 ± 4.70 |
| Measured O ₂ Saturation | 0.85 ± 0.06 |
| Base Excess, mmol/l | 2.26 ± 4.29 |

Results

| | Low Saturation | | High Saturation | | ANOVA | | |
|-----------------------------|----------------|------------|-----------------|------------|------------------|-------|--------------|
| | OxyMask | Venturi | OxyMask | Venturi | Saturation Level | Mask | Inter-action |
| SaO ₂ , % | 92.0 ± 3.6 | 91.7 ± 3.4 | 94.8 ± 3.2 | 94.9 ± 3.6 | - | NS | NS |
| Flow O ₂ , L/min | 0.9 ± 0.3 | 4.8 ± 1.5 | 2.1 ± 0.9 | 12.2 ± 3.9 | <0.01 | <0.01 | <0.01 |
| Ve, L/min | 9.1 ± 5.0 | 7.4 ± 4.2 | 10.6 ± 5.9 | 8.0 ± 4.4 | NS | <0.05 | NS |

- 21 -

| | | | | | | | |
|--|-----------------|-----------------|------------------|---------------------|-------|-------|-------|
| tPCO ₂ , mmHg | 51.9 ± 8.9 | 51.4 ± 7.6 | 51.3 ± 9.1 | 52.4 ± 8.0 | NS | NS | <0.05 |
| P _I O ₂ , mmHg | 229.7 ± 44.5 | 192.6 ± 11.9 | 459.5 ± 167.5 | 330.0 ± 126.6 | <0.01 | <0.01 | <0.05 |
| P _E O ₂ , mmHg | 164.4 ± 16.9 | 181.7 ± 12.2 | 209.2 ± 39.2 | 266.9 ± 52.4 | <0.05 | <0.01 | <0.05 |
| P _I CO ₂ , mmHg | 3.9 ± 1.5 | 1.6 ± 0.9 | 3.2 ± 0.6 | 1.4 ± 0.8 | <0.05 | <0.01 | NS |
| P _E CO ₂ , mmHg | 33.5 ± 8.9 | 11.3 ± 5.6 | 27.2 ± 8.9 | 11.6 ± 8.0 | <0.01 | <0.01 | <0.05 |
| HR, b/min | 77.9 ± 18.0 | 78.2 ± 17.9 | 78.9 ± 18.9 | 77.3 ± 17.7 | <0.05 | NS | NS |
| Nasal:Oral Flow | 1.20 ± 0.32 | 1.01 ± 0.26 | 1.11 ± 0.22 | 1.07 ± 0.26 | NS | NS | NS |

Summary

O₂ flow rate significantly lower with OM vs. V.

P_IO₂ significantly higher and P_EO₂ significantly lower with OM. vs. V

Ve significantly higher with OM vs. V while TcPCO₂ similar between
5 masks.

P_ICO₂ and P_ECO₂ significantly higher with OM vs. V.

Difference in O₂ flow remained significant when comparing patients
whose Ve increased by ≥10% with OM vs. V to those whose did not.

It is evident that many alternatives, modifications and variations will
10 be apparent to those skilled in the art in light of the foregoing description.

- 22 -

Accordingly, it is intended to embrace all such alternatives, modifications and variations as fall within the spirit and broad scope of the invention.

- 23 -

CLAIMS:

1. A mask for delivery of oxygen to a patient, the mask comprising:
a body having a peripheral portion, when in use to sit comfortably on a patient's face, a central portion, and bridge portions extending between the central portion and the peripheral portion and integral therewith, the central portion having an inner surface and an outer surface, the inner surface to be oriented towards the patient's face when the mask is in position and contoured so as to sit at a location spaced over the patient's nose and mouth, the inner surface of the central portion provided with a wall circumscribing a base, the wall and base being of generally concave configuration and circumscribing a centrally positioned oxygen delivery aperture extending through the central portion between the inner surface and the outer surface, the wall and base configured so as to act as an oxygen diffuser to direct the flow of oxygen generally towards the patient's nose and mouth when the mask is in use;
a fastener to engage the mask to the patient so as to maintain contact between the patient's face and said peripheral portion when the mask is in use; and
means associated with the aperture and of the central portion releasably to receive and secure in position an oxygen delivery tube.
2. A mask according to claim 1 in combination with an oxygen delivery tube releasably securable to the outer surface of the central portion of the mask so as to communicate with the oxygen delivery aperture.
3. A mask according to claim 2 further provided with a baffle, the baffle being constructed so as to be releasably seated over the oxygen delivery aperture on the inner surface of the central portion of the mask, the inner surface of the baffle being configured so as to assist during use of the mask in creating turbulence in an oxygen flow leaving the oxygen delivery aperture and assist in mixing oxygen with ambient air and thereby avoid a direct flow of oxygen towards the patient's face.

- 24 -

4. A mask according to claim 2 further provided with an oxygen/carbon dioxide monitor tube releasably securable to the outer surface of the central portion of the mask, so as to communicate through the oxygen delivery aperture with an area above the inner surface of the central portion, during use of the mask, for passage of air within the mask to an oxygen/carbon dioxide monitor, and a baffle being constructed so as to be releasably seated over the oxygen delivery aperture on the inner surface of the central portion of the mask, the baffle having a concave shaped wall and being configured and positioned so as to assist, during use of the mask, in creating turbulence in an oxygen flow leaving the oxygen delivery aperture and assist in mixing oxygen with ambient air and thereby avoid a direct flow of oxygen towards a patient's face, a carbon dioxide intake positioned within the concave shaped wall of the baffle, the carbon dioxide intake communicating with the carbon dioxide monitor tube.

5. A mask according to claim 1 wherein the bridge portions of the mask from a top of the mask to a bottom of the mask are configured in an inverted "Y" shape so that openings are provided towards the bottom and sides of the mask for unobstructed access to and viewing of a patient's mouth and other parts of the patient's face.

6. A mask according to claim 1 wherein the wall and base are of cup-shaped appearance, the walls ending in an edge of triangular peripheral contour so as to conform, when the mask is in position on a patient, to the shape of a patient's nose and mouth area.

7. A mask according to claim 3 wherein the baffle comprises a post to be releasably seated centrally within the oxygen delivery apparatus at an end of the post and having a curled back conical lip, at another end of the post, an underside of the lip being in line with a flow of oxygen when passing from the oxygen delivery aperture during use of the mask so as to generate turbulence in the oxygen flow and create, in conjunction with the diffuser, a plume of oxygen enriched air at the patient's nose and mouth.

- 25 -

8. A mask according to claim 4 wherein the baffle comprises a post at one end to be seated centrally within the oxygen outlet, the post having centrally through it a passageway for oxygen/carbon dioxide monitoring centrally through it, the passageway communicating with the oxygen/carbon dioxide intake of the baffle and the oxygen/carbon dioxide monitor tube.
9. A mask according to claim 3 wherein the wall and base are of cup-shaped appearance, the walls ending in an edge of triangular peripheral contour so as to conform, when the mask is in position on a patient, to the shape of a patient's nose and mouth area.
10. A mask according to claim 4 wherein the wall and base are of cup-shaped appearance, the walls ending in an edge of triangular peripheral contour so as to conform, when the mask is in position on a patient, to the shape of a patient's nose and mouth area.
11. A facial mask, for delivery of a gas to the nose and mouth region of a patient, comprising a substantially rigid mask body having a rim for contacting the patient's face substantially covering the nose and mouth of the patient, a fastener to engage the mask to a patient so as to maintain contact between the patient's face and said rim, a connector for connection to a supply of gas, a gas nozzle opening into the interior of said mask operatively connected to said gas connector, a diffuser at least partly surrounding said nozzle and fastened to and supported by said mask body at a position spaced apart from and not contacting the patient's face, and a plurality of openings within said body to permit access to the patient's face from the outside of said mask, said diffuser being positioned generally centrally within said mask and including a gas outlet nozzle, a gas flow disrupter between said nozzle and said patient's face so as to generate a turbulent plume of said gas substantially at a non-oblique angle towards the patient's nose and mouth and a peripheral wall surrounding said nozzle, said wall having a maximum top to bottom height when said mask is vertical of 40 mm above and below said nozzle and a maximum side to side width of

- 26 -

20 mm on either side of said nozzle, when said mask is positioned in an upright position.

12. A mask as defined in claim 11, wherein said body comprises material selected from the following: ABS, PVC, polypropylene, silicone polycarbonate.

13. A mask as defined in claim 11, wherein said diffuser is positioned within the mask so as to maintain a spacing of between 12 and 40 mm between an upper edge of said wall and said patient's face as face as measured from the skin surface between nose and mouth, when the mask is worn by the patient.

14. A mask as defined in claim 11, wherein said diffuser is positioned so as to be substantially centered in the side to side direction relative to said patient's face when the mask is worn by a patient.

15. A mask as defined in claim 1, wherein said openings comprise between 30 and 80 per cent of the total surface area of said body, said surface area comprising open and covered areas.

16. A mask as defined in claim 15, wherein said openings comprise between 60 and 75 per cent of the total surface area of said body.

17. A mask as defined in claim 1, wherein said diffuser is positioned so as to be centered over a point about halfway between the patient's nose and upper lip when the mask is worn by a patient.

18. A mask as defined in claim 1, further comprising a sensor for monitoring exhaled gases within the interior of said mask.

19. A mask as defined in claim 1, wherein at least one of said openings provides access from the exterior of the mask to the patient's mouth when the mask is worn by a patient.

- 27 -

20. A mask as defined in claim 1, wherein at least one of said openings provides access from the mask exterior to the patient's nostrils when the mask is worn by a patient.

21. A facial mask for delivery of a gas to the nose and mouth region of a patient, comprising a substantially rigid mask body having a rim for contacting the patient's face substantially covering the nose and mouth of the patient, a fastener to engage the mask to a patient so as to maintain contact between the user's face and said rim, a connection to a supply of gas, a gas nozzle opening into the interior of said mask operatively connected to said gas connection, a diffuser at least partly surrounding said nozzle and fastened to and supported by said mask body at a position spaced apart from and not contacting the patient's face, and a plurality of openings within said body to permit access to the patient's face from the outside of said mask, said diffuser being positioned generally centrally within said mask and including a gas outlet nozzle, means to generate a turbulent plume of said gas substantially at a non-oblique angle towards the user's nose and mouth and a peripheral flange surrounding said nozzle, said diffuser being positioned within said mask so as to fit entirely within a space defined in relation to a point on the patient's face about halfway between the base of the nose and uppermost edge of upper lip, said space defined in the vertical plane as 40 mm above and below said point, 20 mm on either side of said point and horizontally spaced apart from said point by between 12 and 40 mm when said patient is upright.

22. A mask as defined in claim 21, wherein said openings comprise between 30 and 80 per cent of the total surface area of said body, said surface area comprising open and covered areas.

23. A mask as defined in claim 22, wherein said openings comprise between 60 and 75 per cent of the total surface area of said body.

24. A mask as defined in claim 21, further comprising a sensor for monitoring exhaled gases within the interior of said mask.

- 28 -

25. A mask as defined in claim 21, wherein at least one of said openings provides access from the exterior of the mask to the patient's mouth when the mask is worn by a patient.

26. A mask as defined in claim 21, wherein at least one of said openings provides access from the mask exterior to the patient's nostrils when the mask is worn by a patient.

27. An oxygen delivery mask comprising a body for at least partly covering a patient's face, said body being generally concave, a fastener to engage the mask to a patient, an oxygen delivery opening to release a flow of oxygen within the concave interior of the mask towards the patient, and a removable mask support member comprising an at least partly rigid elongate member having opposed ends, each end including means for fastening to said mask whereby when fastened said member bows outwardly to partly fill said interior space of the mask to provide support during non-use, said member further comprising an opening shaped to snugly receive an end of a laryngeal mask in a position spaced apart from said oxygen delivery opening.

28. A mask as defined in claim 27, further comprising a laryngeal mask comprising an oxygen-receiving end for friction-fitting within said opening and a stop to retain said laryngeal mask in said spaced apart position.

AMENDED CLAIMS

Received by the International Bureau on 20 March 2006 (20.03.2006).

CLAIMS:

1. An oxygen delivery mask for wearing over the nose and mouth region of a patient, comprising:

a mask body for at least partially covering the patient's nose and mouth region, said mask body including a rim for contacting the patient's face surrounding the patient's nose and mouth;

a gas diffuser mounted to the body, said body having sufficient rigidity and shaped to space said diffuser away from the patient's face, said diffuser being connectable to a gas supply for generating a turbulent plume of gas towards the patient's nose and mouth region and comprising a concave structure opening towards the patient's face formed by a base and a wall; and

a fastener to maintain contact between the patient's face and said rim;

wherein said mask body has openings therein with suitable size, shape, and location to permit the patient to perform functions of eating or drinking and to permit others to deliver substances to the patient's mouth, without removing said mask.

2. A mask according to claim 1 in combination with an oxygen delivery tube releasably securable to the outer surface of the mask body so as to communicate with the diffuser.

3. A mask according to claim 2, further provided with a monitor tube for receiving carbon dioxide or oxygen from the interior of said mask body, and an intake positioned within the interior of said mask body in communication with said monitor tube.

4. A mask according to claim 1, wherein said mask body is configured in an inverted "Y" shape so that said openings are provided towards the bottom and sides of the mask for unobstructed access to and viewing of a patient's mouth and other parts of the patient's face.

5. A mask according to claim 1, wherein said diffuser comprises a concave wall seated over the gas supply, and a mushroom-shaped baffle positioned within the path of gas exiting the gas supply for interrupting the linear flow of said gas to generate turbulence, said baffle comprising a stem and a head, said head comprising a curled back conical lip to form a concave region facing said gas supply.

6. A mask as defined in claim 1, wherein said diffuser is positioned generally centrally within said mask and includes a gas outlet nozzle, means to generate a turbulent plume of said gas substantially at a non-oblique angle towards the user's nose and mouth and a peripheral wall surrounding said nozzle, said diffuser being positioned within said mask so as to fit entirely within a space defined in relation to a point on the patient's face about halfway between the base of the patient's nose and uppermost edge of the patient's upper lip, said space defined in the vertical plane as 40 mm above and below said point, 20 mm on either side of said point and horizontally spaced apart from said point by between 12 and 40 mm when said patient is upright.

7. A mask as defined in claim 6, wherein said diffuser is positioned within the mask so as to maintain a spacing of between 12 and 40 mm between an upper edge of said wall and said patient's face as measured from the skin surface between nose and mouth, when the mask is worn by the patient.

8. A mask as defined in claim 6, wherein said diffuser is positioned so as to be substantially centered in the side to side direction relative to said patient's face when the mask is worn by a patient.

9. A mask as defined in claim 1, wherein said openings comprise between 30 and 80 per cent of the total surface area of said body, said surface area comprising open and covered areas.

10. A mask as defined in claim 1, wherein said openings comprise between 60 and 75 per cent of the total surface area of said body.
11. A mask as defined in claim 1, wherein said diffuser is positioned so as to be centered over a point about halfway between the patient's nose and upper lip when the mask is worn by a patient.
12. A mask as defined in claim 1, further comprising a sensor for monitoring exhaled gases within the interior of said mask.
13. A mask as defined in claim 1, wherein at least one of said openings provides access from the mask exterior to the patient's nostrils when the mask is worn by a patient.
14. An oxygen delivery mask comprising a body for at least partly covering a patient's face, said body being generally concave and opening towards said patient's face, a fastener to engage the mask to a patient, an oxygen delivery opening to release a flow of oxygen within the concave interior of the mask towards the patient, and a removable mask support member comprising an at least partly rigid elongate member having opposed ends, each end including means for fastening to said mask whereby when fastened said member bows outwardly to partly fill said interior space of the mask to provide support during non-use, said member further comprising an opening shaped to snugly receive an end of a laryngeal mask in a position spaced apart from said oxygen delivery opening.
15. A mask as defined in claim 14, further comprising a laryngeal mask comprising an oxygen-receiving end for friction-fitting within said opening and a stop to retain said laryngeal mask in said spaced apart position.

1/11

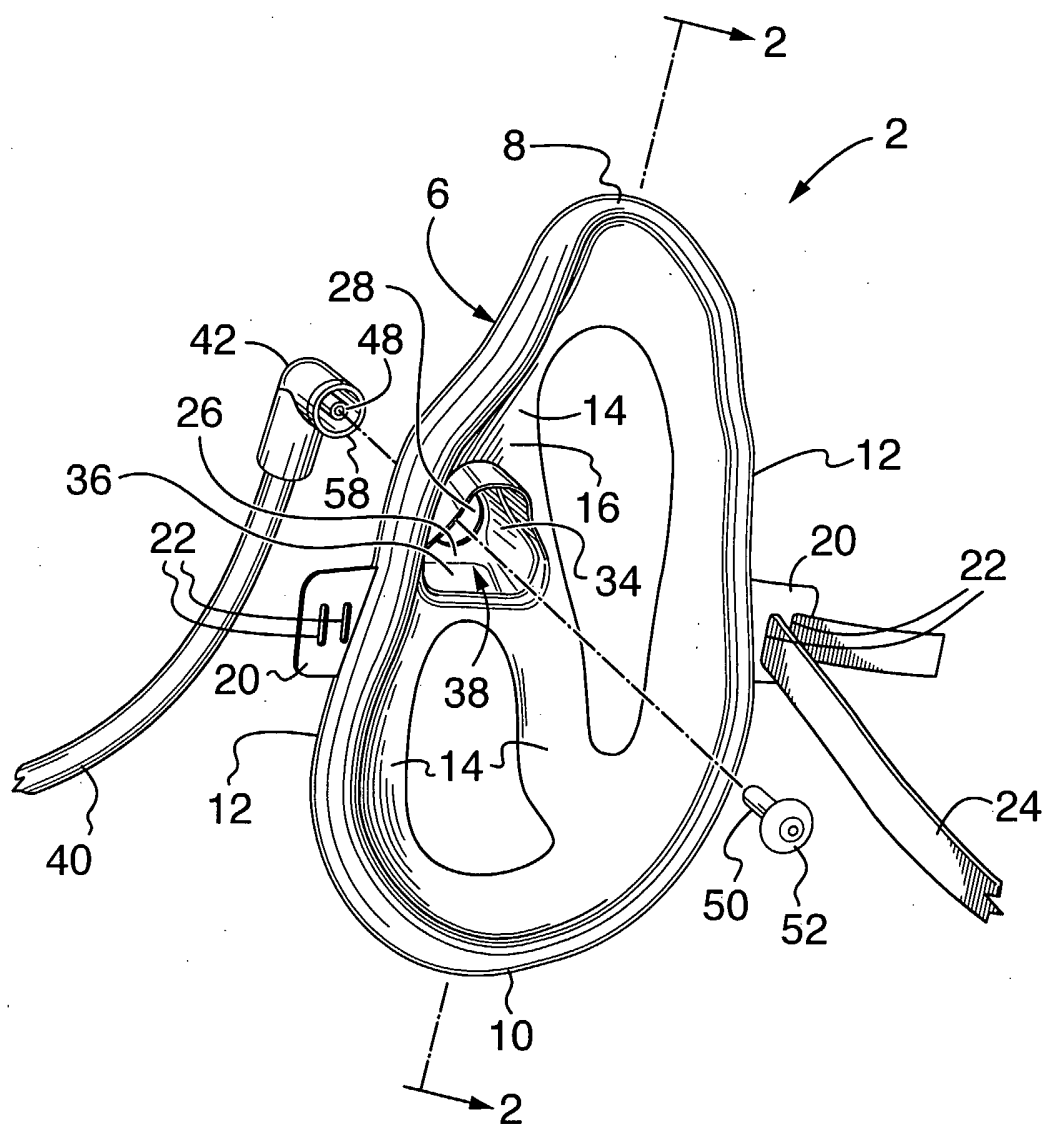


FIG. 1

2/11

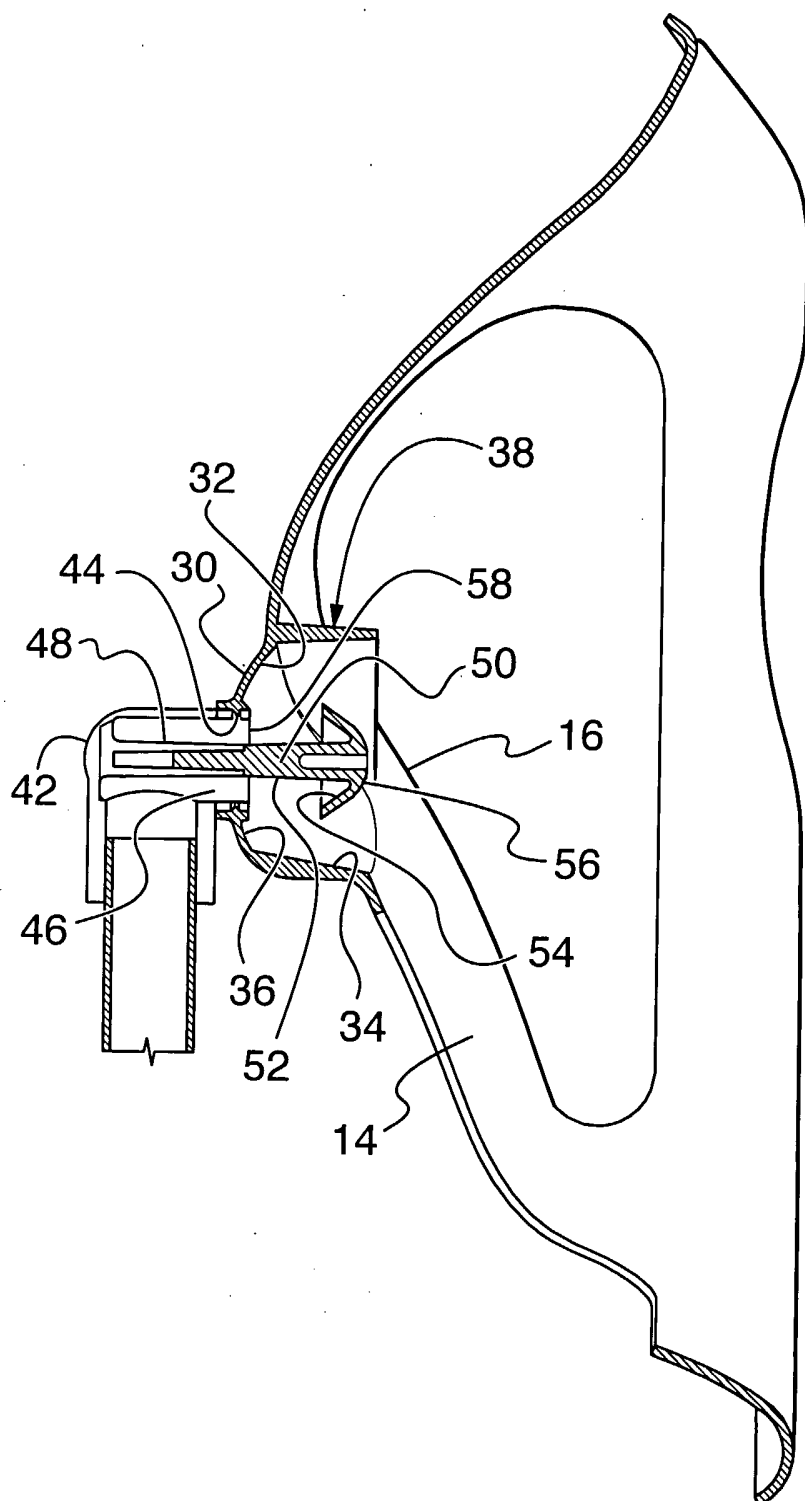


FIG. 2

3/11

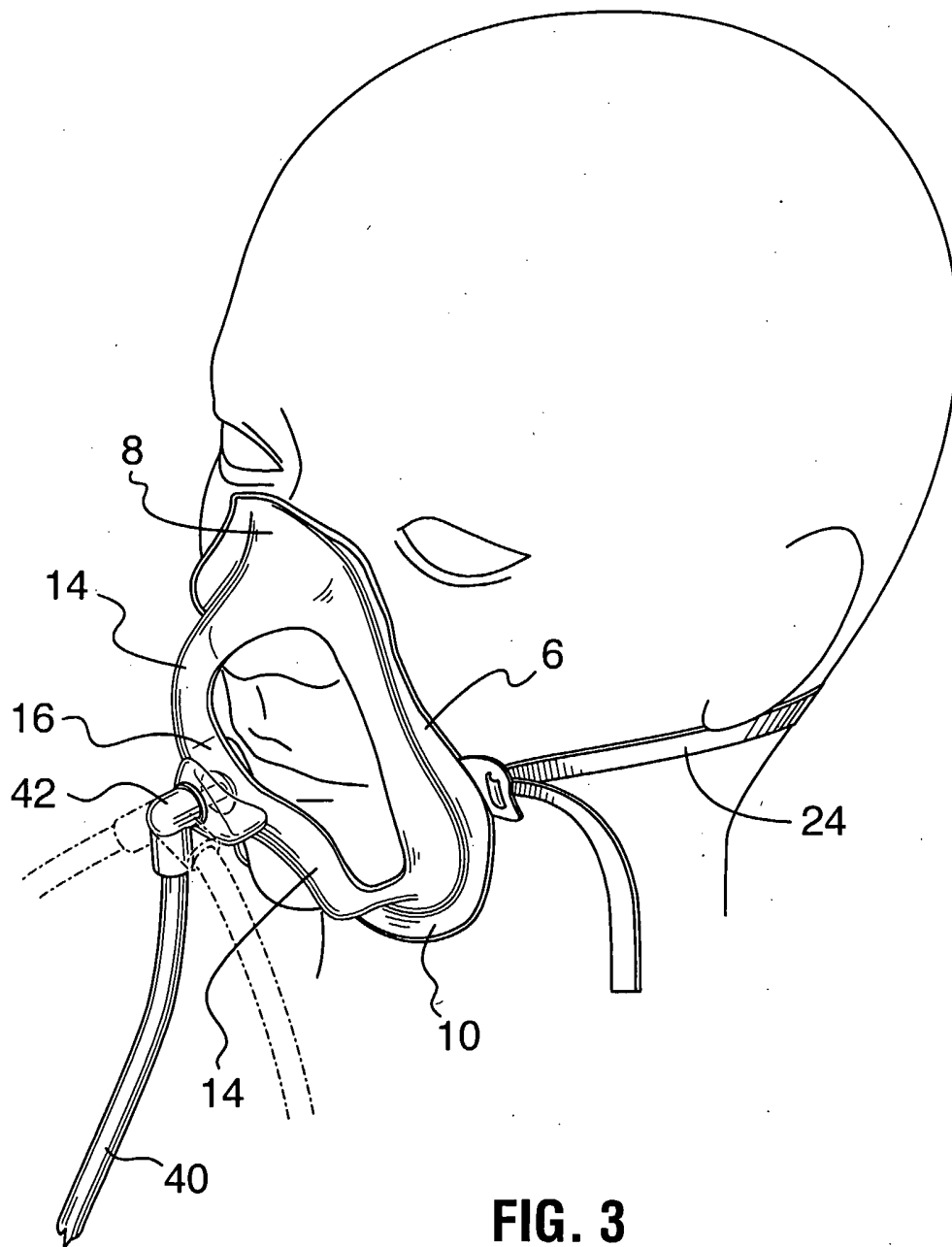


FIG. 3

4/11

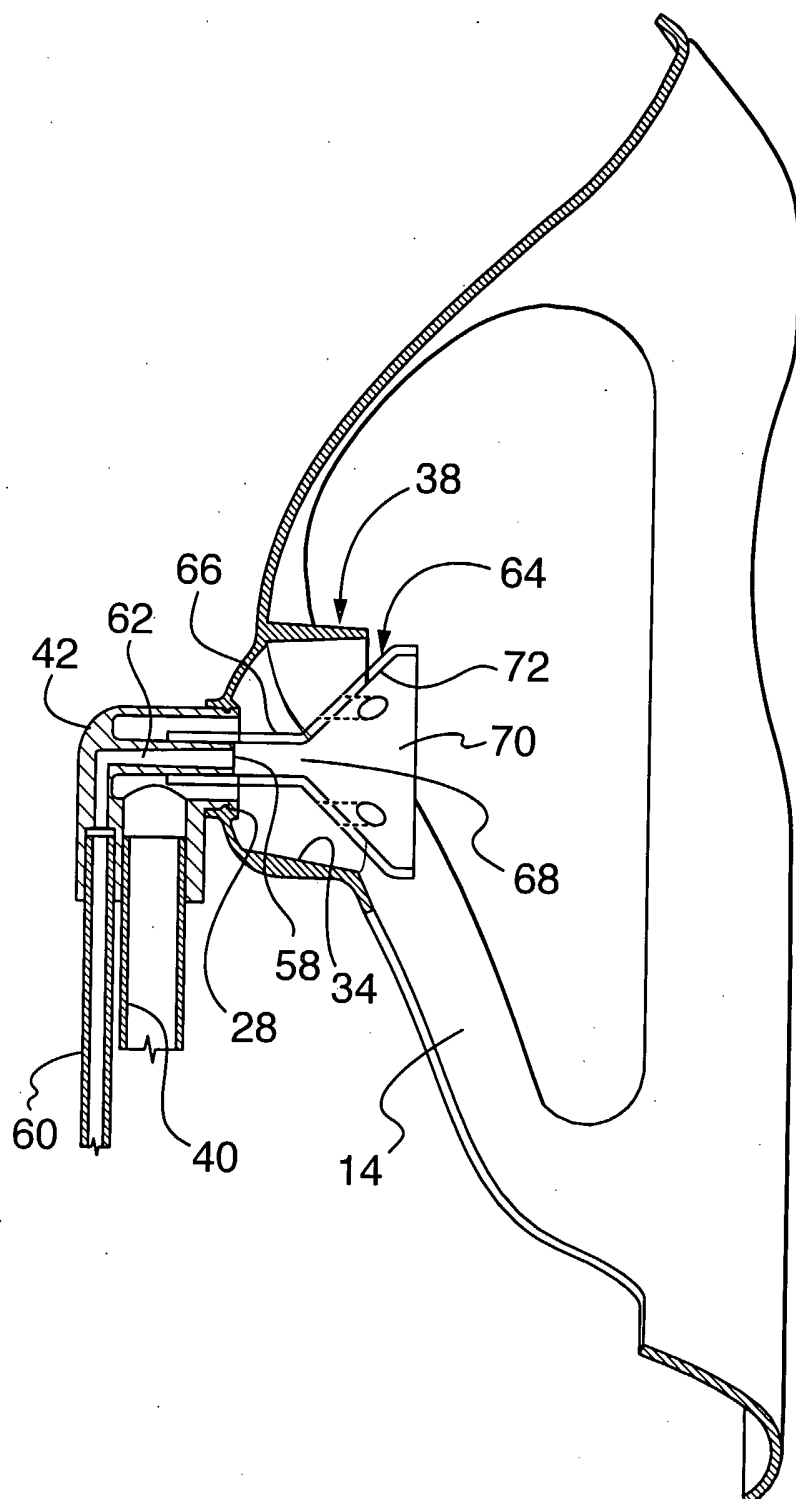
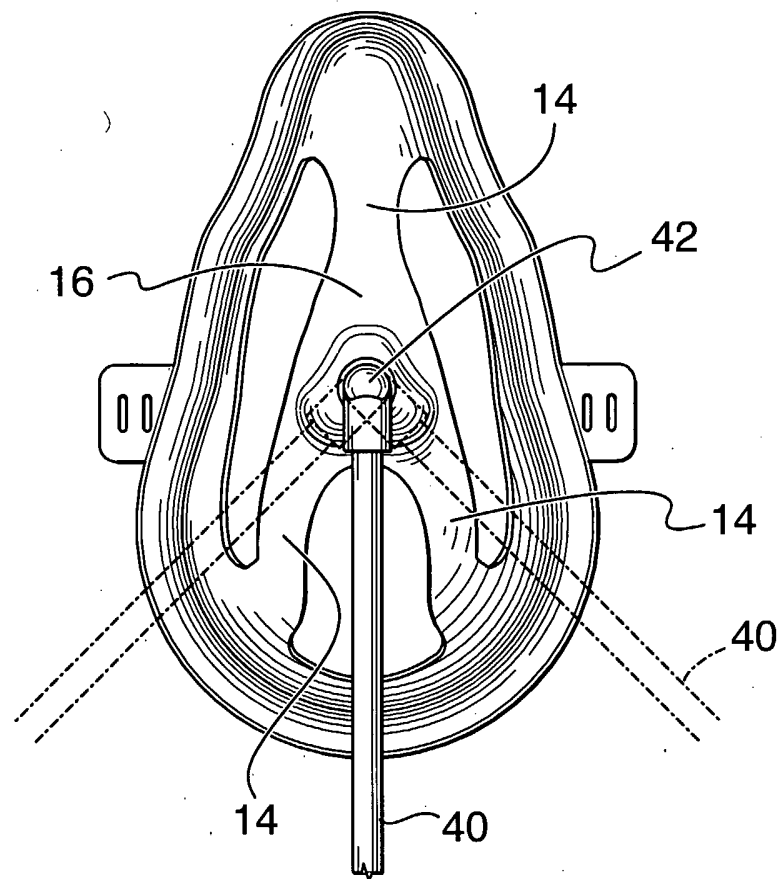
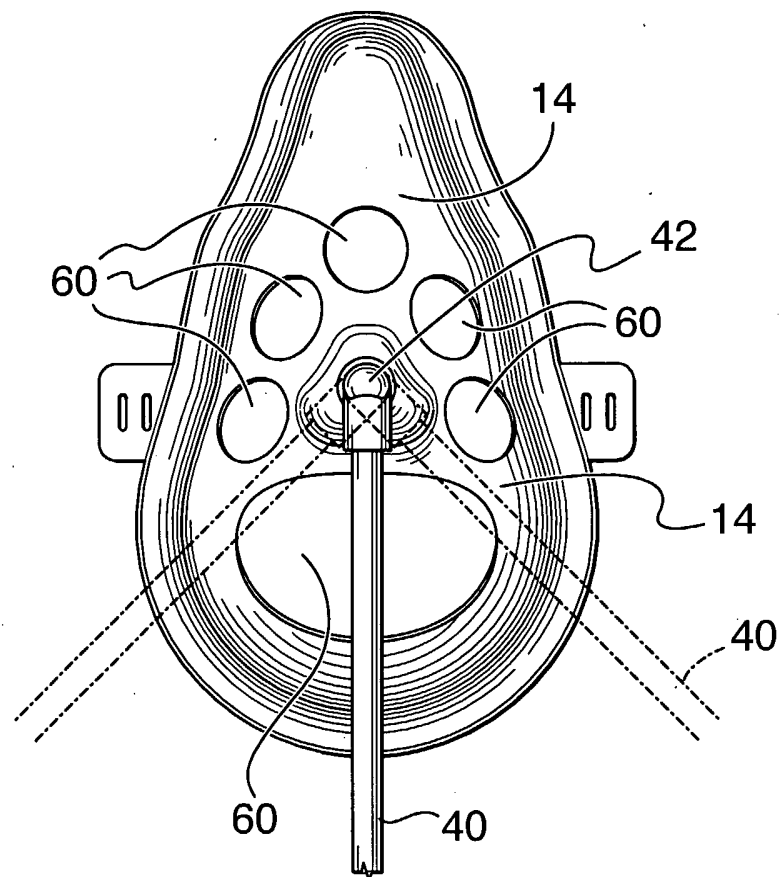


FIG. 4

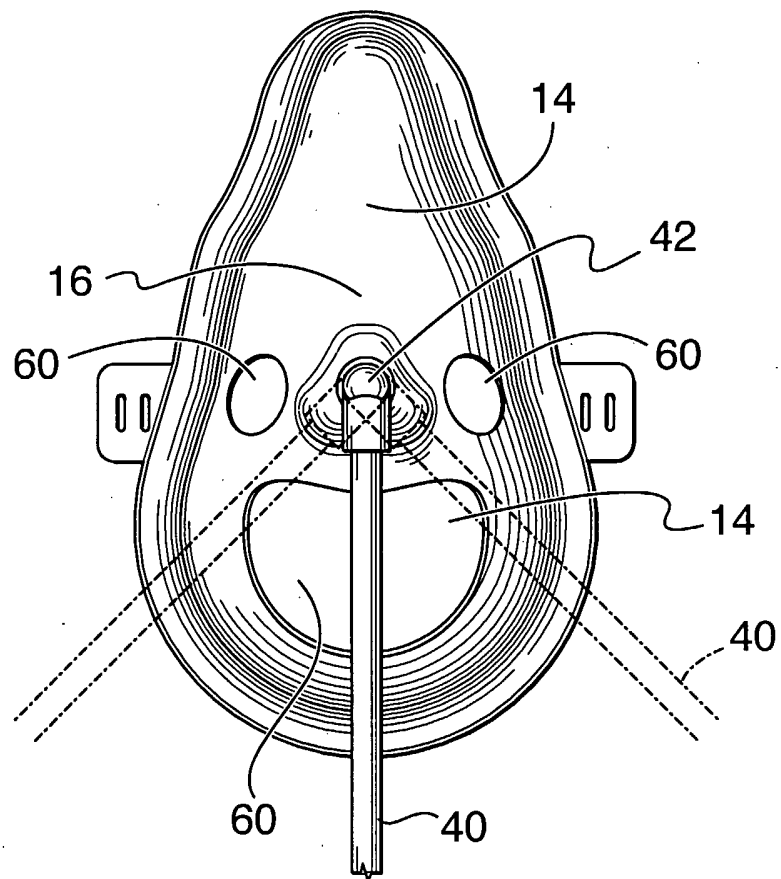
5/11

**FIG. 5**

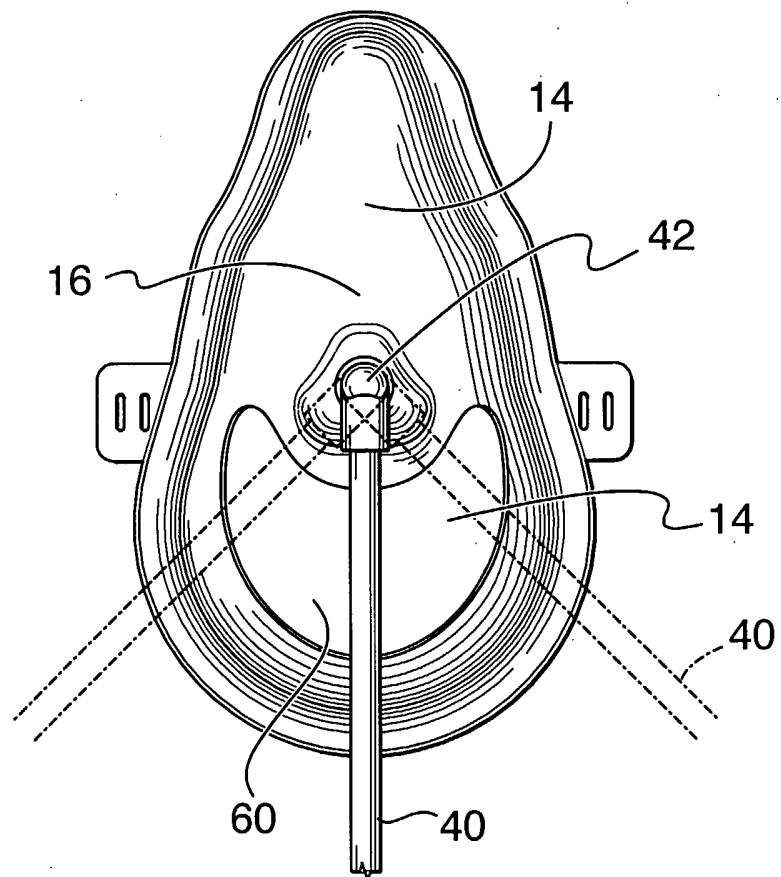
6/11

**FIG. 6A**

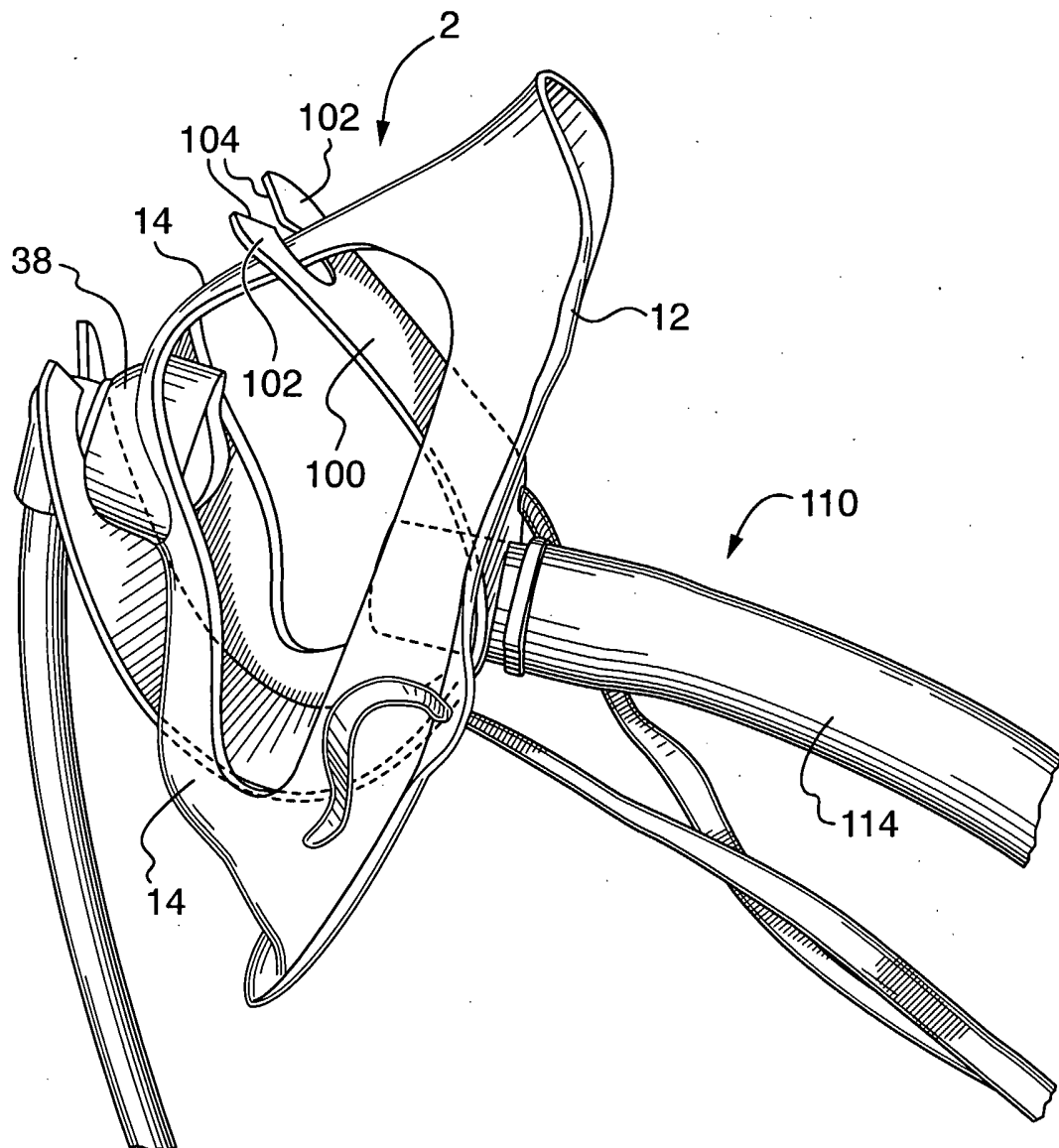
7/11

**FIG. 6B**

8/11

**FIG. 6C**

9/11

**FIG. 7**

10/11

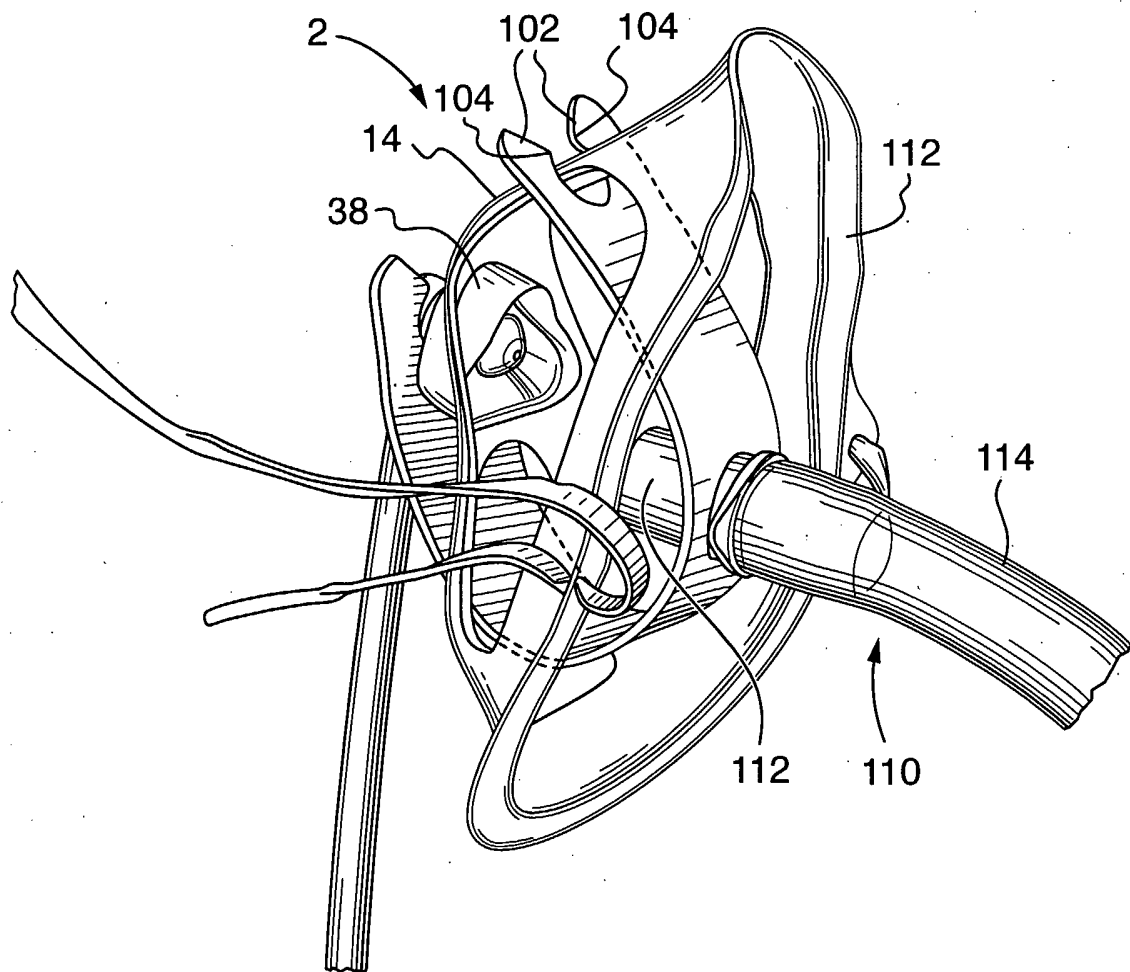
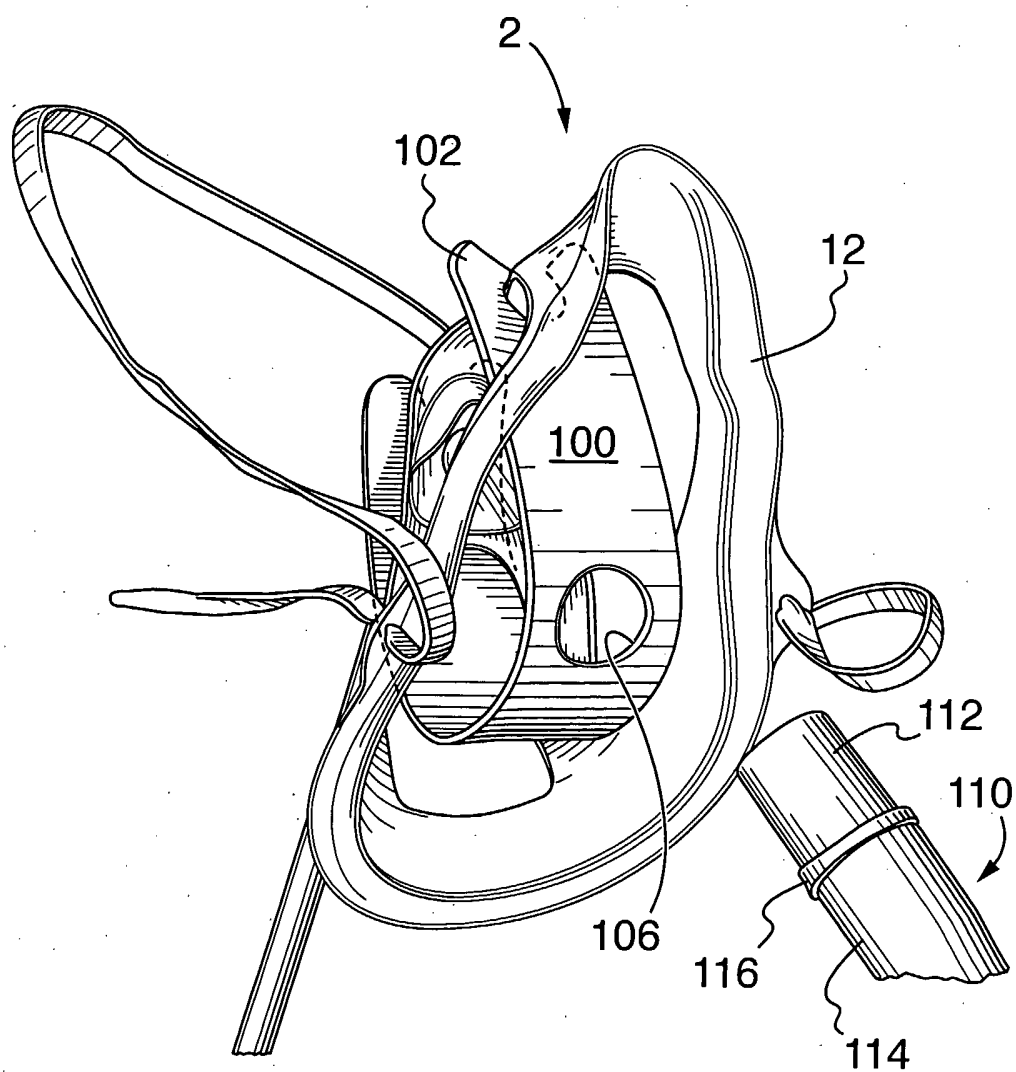


FIG. 8

11/11

**FIG. 9**

INTERNATIONAL SEARCH REPORT

International application No.
PCT/CA2005/001541

| A. CLASSIFICATION OF SUBJECT MATTER IPC: A61M 16/06 (2006.01) | | |
|---|---|--|
| 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): A61M 16/06, A61M 16/00 Canadian patent classification: 137/1.5; US classes: 201.19, 203.29, 206.21 and 206.28 | | |
| Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched | | |
| Electronic database(s) consulted during the international search (name of database(s) and, where practicable, search terms used) Canadian patent database, Delphion, USPTO, QPAT and Esp@cenet keywords: mask, diffuser, oxygen, openings, holes, access, dioxide, sensor | | |
| C. DOCUMENTS CONSIDERED TO BE RELEVANT | | |
| Category* | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
| Y | US 5 233 978 (Callaway) 10 Aug. 1993 (10-08-1993) Entire document | 1-4, 6-26 |
| Y | US 4 201 205 (Bartholomew) 6 May 1980 (06-05-1980) column 1, line 63 to column 3, line 11 and figures 1 to 4 | 1-4, 6-26 |
| Y | US 6 450 166 B1 (McDonald et al.) 17 Sep. 2002 (17-10-2002) column 3, line 62 to column 4, line 55 and figures 1 to 4 | 3, 7, 9, 11-14, 17-26 |
| Y | US 2004/0094160 A1 (McDonald et al.) 20 May 2004 (20-05-2004) paragraphs 38 to 42 and figures 1 to 6 | 4, 8, 10 |
| A | US 3 295 521 (Balch) 3 Jan. 1967 (03-01-1967) Entire document | 1-26 |
| <input type="checkbox"/> Further documents are listed in the continuation of Box C. <input checked="" type="checkbox"/> 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 | "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 |
| "E" | earlier application or patent but published on or after the international filing date | "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 |
| "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) | "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 |
| "O" | document referring to an oral disclosure, use, exhibition or other means | "&" document member of the same patent family |
| "P" | document published prior to the international filing date but later than the priority date claimed | |
| Date of the actual completion of the international search 30 December 2005 (30-12-2005) | | Date of mailing of the international search report 23 January 2006 (23-01-2006) |
| Name and mailing address of the ISA/CA Canadian Intellectual Property Office Place du Portage 1, C114 - 1st Floor, Box PCT 50 Victoria Street Gatineau, Quebec K1A 0C9 Facsimile No.: 001(819)953-2476 | | Authorized officer Eric Lafontaine (819) 956-9965 |

INTERNATIONAL SEARCH REPORT

International application No.
PCT/CA2005/001541

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of the first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons :

1. ☐ Claim Nos. :
because they relate to subject matter not required to be searched by this Authority, namely :

2. ☐ Claim Nos. :
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically :

3. ☐ Claim Nos. :
because they are dependant claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows :

Group A - Claims 1 to 26 are directed to a mask for delivery of gas to a patient comprising a plurality of openings permitting access to the patient's face and a gas nozzle with a diffuser and fastening means; and

Group B - Claims 27 and 28 are directed to a mask for delivery of gas to a patient comprising an oxygen delivery opening, a fastener and an elongate member with ends for fastening to the mask to provide support during non use and to provide an opening to receive an end of a laryngeal mask in a position spaced apart from the oxygen delivery opening.

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying additional fees, this Authority did not invite payment of additional fees.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claim Nos. :
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claim Nos. : 1 to 26

Remark on Protest ☐ The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.
☐ The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.
PCT/CA2005/001541

| Patent Document Cited in Search Report | Publication Date | Patent Family Member(s) | Publication Date |
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