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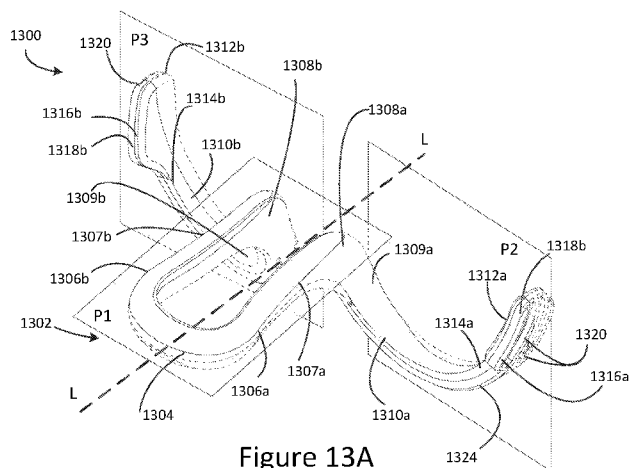
(54) **Title:** NASAL DILATOR DEVICES

Figure 13A

(57) **Abstract:** A nasal dilator device comprises a substantially U-shaped body including a central portion arranged to span a septum of a nose when worn by a user and first and second leg members extending from the central portion in a first plane, a first cantilever rib member extending outward from the U-shaped body in a second plane and a second cantilever rib member extending outward from the U-shaped body in a third plane, wherein the first and second cantilever rib members extend away from each other. The nasal dilator device further comprises a first intermediate section connecting an end of the first leg member to a proximal end of the first cantilever rib member, wherein the first intermediate section extends from the first plane to the second plane; and a second intermediate section connecting an end of the second leg member to a proximal end of the second cantilever rib member, wherein the second intermediate section extends from the first plane to the third plane and at least one projection protruding from and extending along at least a portion of a length of each of the first and second cantilever rib members.



Nasal dilator devices

Technical Field

[0001] Described embodiments generally relate to nasal dilator devices for facilitating respiration. Some embodiments relate to nasal dilator devices to be fitted to the nose to facilitate or improve respiration during sleeping and/or sporting activities and/or for general day-to-day wear. Some embodiments relate to nasal dilator devices including filtration mechanisms to filter airflow during respiration and other embodiments relate to nasal dilator devices including agent delivery mechanisms for delivery of fragrances and/medicaments to the nose during respiration.

Background

[0002] Nasal dilator devices are worn by users to dilate their nasal cavities when sleeping and/or partaking in sporting activities to thereby facilitate respiration. However, many nasal dilator devices are uncomfortable to wear and/or become easily dislodged from a user's nose during such activities.

[0003] In particular, activities that are percussive, such as running, cause nasal dilator devices to move from their position as intended by the user requiring readjustment. In the field of professional sports, such adjustments are a physical and mental distraction that can negate any other intended benefit of such devices.

[0004] It is desired to address or ameliorate one or more shortcomings of prior nasal dilator devices, or to at least provide a useful alternative thereto.

[0005] Throughout this specification the word "comprise", or variations such as "comprises" or "comprising", will be understood to imply the inclusion of a stated element, integer or step, or group of elements, integers or steps, but not the exclusion of any other element, integer or step, or group of elements, integers or steps.

[0006] Any discussion of documents, acts, materials, devices, articles or the like which has been included in the present specification is not to be taken as an admission that any or all of these matters form part of the prior art base or were common general knowledge in the field relevant to the present disclosure as it existed before the priority date of each claim of this application.

Summary

[0007] Some embodiments relate to a nasal dilator device comprising a substantially U-shaped body including a central portion arranged to span a septum of a nose when worn by a user and first and second leg members extending from the central portion in a first plane, a first cantilever rib member extending outward from the U-shaped body in a second plane, a second cantilever rib member extending outward from the U-shaped body in a third plane, wherein the first and second cantilever rib members extend away from each other, a first intermediate section connecting an end of the first leg member to a proximal end of the first cantilever rib member, wherein the first intermediate section extends from the first plane to the second plane and a second intermediate section connecting an end of the second leg member to a proximal end of the second cantilever rib member, wherein the second intermediate section extends from the first plane to the third plane, and at least one projection protruding from and extending along at least a portion of a length of each of the first and second cantilever rib members.

[0008] In some embodiments, the at least one projection may comprise first and second projections, each forming a flange disposed at respective elongate edges of a major surface of the nasal dilator device. The at least one projection may be arranged, in use, to engage with an inner surface of a nose.

[0009] In some embodiments, the second and third planes may be orthogonal or perpendicular to the first plane. The first intermediate section and the second intermediate section may be right angled sections. The first and second intermediate sections may be arranged, in use, to engage with the septum and extend from the

septum behind the columella and alar fibrofatty tissue of the nose, allowing the first and second cantilever rib members, in use, to extend along respective nasal orifices to an inner wall of the nostrils.

[0010] The first and second cantilever rib members may be arcuate cantilever rib members, each having a curvature along its length. The first and second intermediate sections may be arcuate intermediate sections, each having a curvature along its length. The second and third planes may be converging planes such that the first and second cantilever rib members are angled and/or extend substantially toward the central portion of the U-shaped body. The second and third planes may be diverging planes such that the first and second cantilever rib members are angled and/or extend substantially toward the central portion of the U-shaped body. The second and third planes may be the same plane.

[0011] In some embodiments, the first and second cantilever rib members may exhibit an elongate arched profile which approximates at least a portion of one of a circle, ellipse or parabola. The first and second leg members may be inclined towards each other such that a relatively greater distance is provided between the first and second leg members towards the central portion accommodate a columella of a nose when donned by the user. The first and second intermediate sections may be inclined away from each other to assist in urging the respective first and second cantilever rib members against inner walls of respective nostrils when worn by the user.

[0012] In some embodiments, the first and second cantilever rib members may comprise respective first and second nostril engaging elements for engaging with an inner wall of a respective nostril. The first and second nostril engaging elements may extend arcuately from the respective first and second distal ends of the respective first and second rib members. For example, the first and second nostril engaging elements may extend from the first and second planes respectively and in a direction away from the central portion of the U-shaped body. The first and second nostril engaging elements may be disposed at distal ends of the first and second cantilever rib members, respectively.

[0013] In some embodiments, a series of protrusions may be disposed on the first and second nostril engaging elements. For example, the first and second nostril engaging elements may be substantially elongate and the series of protrusions may extend along a length of the first and second nostril engaging elements. The series of protrusions may comprise a plurality of elongated v-shaped protrusions.

[0014] In some embodiments, enlarged pads may be disposed on the first and second nostril engaging elements to engage with inner walls of the nostrils. For example, the enlarged pads may comprise the series of protrusions and/or the series of protrusions may be formed from an overmould material. In some embodiments, the at least one projection may be integrated with and extend from a corresponding protrusion of the series of protrusions. The at least one projection may be formed from an overmould material.

[0015] In some embodiments, the nasal dilator may comprise a first and second releasable attachment mechanism for releasably attaching the first and second cantilever rib members, for example, distal ends of the first and second cantilever rib members, to the U-shaped body and/or the first and second intermediate section respectively. For example, the releasable attachment mechanisms may be arranged to releasably attach first and second nostril engaging elements disposed at distal ends of the first and second cantilever rib members to the first and second leg members, respectively. The releasable attachment mechanisms may be arranged to releasably attach the first and second nostril engaging elements disposed at distal ends of the first and second cantilever rib members to the first and second intermediate sections, respectively.

[0016] The releasable attachment mechanisms may each comprise an arm and a socket arranged to receive and engage the arm. A stopper or hook may be disposed at an end of the arm to hinder the arm from withdrawing from the socket. A notch may be disposed on an inner surface of the socket and may be configured to engage with the stopper or hook on the arm to hinder the arm from withdrawing from the socket.

[0017] In some embodiments, the arms may be disposed on an inner surface of first and second nostril engaging elements of the first and second cantilever rib members, respectively, and the sockets may be disposed on the first and second leg members. In some embodiments, the arms may be disposed on an inner surface of first and second nostril engaging elements of the first and second cantilever rib members, respectively, and the sockets may be disposed on the first and second intermediate sections. In some embodiments, the sockets may be disposed on an inner surface of first and second nostril engaging elements of the first and second cantilever rib members, respectively, and the arms may be disposed on the first and second leg members. In some embodiments, the sockets may be disposed on an inner surface of first and second nostril engaging elements of the first and second cantilever rib members, respectively, and the arms may be disposed on the first and second intermediate sections. In some embodiments, the nasal dilator device may comprise first and second arm support members projecting from respective first and second intermediate sections, the sockets may be disposed on an inner surface of first and second nostril engaging elements of the first and second cantilever rib members, respectively, and the arms may be disposed on the first and second arm support members.

[0018] The sockets may comprise substantially elongate housings which taper along their length and provide a relatively broad opening for receiving the arms. Each of the sockets may be provided with apertures in side walls of the housing.

[0019] Some embodiments relate to a nasal dilator device comprising a substantially U-shaped body including a central portion arranged to span a septum of a nose when worn by a user and first and second leg members extending from the central portion in a first plane, a first cantilever rib member extending outward from the U-shaped body in a second plane, a second cantilever rib member extending outward from the U-shaped body in a third plane, wherein the first and second cantilever rib members extend away from each other, a first intermediate section connecting an end of the first leg member to a proximal end of the first cantilever rib member, wherein the first intermediate section extends from the first plane to the second plane, a second intermediate section connecting an end of the second leg member to a proximal end of

the second cantilever rib member, wherein the second intermediate section extends from the first plane to the third plane, a first releasable attachment mechanism for releasably attaching a distal end of the first cantilever rib member to the first intermediate section and a second releasable attachment mechanism for releasably attaching a distal end of the second cantilever rib member to the second intermediate section.

[0020] The second and third planes may be orthogonal or perpendicular to the first plane. The first intermediate section and the second intermediate section are right angled sections. The first and second cantilever rib members may comprise respective first and second nostril engaging elements for engaging with an inner wall of a respective nostril.

[0021] In some embodiments, the releasable attachment mechanisms may be arranged to releasably attach first and second nostril engaging elements disposed at the distal ends of the first and second cantilever rib members to the first and second leg members, respectively. The releasable attachment mechanisms may each comprise an arm and a socket arranged to receive and engage the arm. A stopper or hook may be disposed at an end of the arm to hinder the arm from withdrawing from the socket. A notch may be disposed on an inner surface of the socket and may be configured to engage with the stopper or hook on the arm to hinder the arm from withdrawing from the socket.

[0022] In some embodiments, the arms may be disposed on an inner surface of first and second nostril engaging elements of the first and second cantilever rib members, respectively, and the sockets may be disposed on the first and second intermediate sections. In some embodiments, the sockets may be disposed on an inner surface of first and second nostril engaging elements of the first and second cantilever rib members, respectively, and the arms may be disposed on the first and second intermediate sections. In some embodiments, the nasal dilator device may further comprise first and second arm support members projecting from respective first and second intermediate sections and the arms may be disposed on the first and second arm support members.

[0023] In some embodiments, the sockets may comprise substantially elongate housings which taper along their length and provide a relatively broad opening for receiving the arms. Each of the sockets may be provided with apertures in side walls of the housing.

[0024] Some embodiments relate to a nasal dilator device comprising a substantially U-shaped body including a central portion arranged to span a septum of a nose when worn by a user and first and second leg members extending from the central portion in a first plane, a first cantilever rib member extending outward from the U-shaped body in a second plane, a second cantilever rib member extending outward from the U-shaped body in a third plane, wherein the first and second cantilever rib members extend away from each other, a first intermediate section connecting an end of the first leg member to a proximal end of the first cantilever rib member and a second intermediate section connecting an end of the second leg member to a proximal end of the second cantilever rib member.

[0025] In some embodiments, the first intermediate section may extend from the first plane to the second plane and the second intermediate section may extend from the first plane to the third plane. The second and third planes may be orthogonal to the first plane.

[0026] For example, the first intermediate section and the second intermediate section may be right angled sections. In some embodiments, the first and second intermediate sections are arranged, in use, to engage with the septum and extend from the septum behind the columella and alar fibrofatty tissue of the nose allowing the first and second cantilever rib members, in use, to extend along respective nasal orifices to an inner wall of the nostrils.

[0027] In some embodiments, the first intermediate section may extend between the first plane and the second plane and the second intermediate section may extend between the first plane and the third plane. For example, the first and second intermediate sections may be arranged, in use, to extend along a length of the septum

and the first and second cantilever rib members are each arranged, in use, to extend from a floor of a respective nasal orifice to an inner wall of the nostrils.

[0028] Some embodiments relate to a nasal dilator device comprising a substantially U-shaped body including a central portion arranged to span a septum of a nose when worn by a user and first and second leg members extending from the central portion in a first plane, a first cantilever rib member extending outward from an end of the first leg member of the U-shaped body in a second plane, a second cantilever rib member extending outward from an end of the second leg member of the U-shaped body in a third plane, wherein the first and second cantilever rib members extend away from each other and at least one projection protruding from and extending along at least a portion of a length of each of the first and second cantilever rib members. For example, the second and third planes may be orthogonal to the first plane.

[0029] Some embodiments relate to a nasal dilator device comprising a substantially U-shaped body including a central portion arranged to span a septum of a nose when worn by a user and first and second leg members extending from the central portion, first and second cantilever rib members extending outward from the U-shaped body and away from one another, a first intermediate section connecting an end of the first leg member to a proximal end of the first cantilever rib member and a second intermediate section connecting an end of the second leg member to a proximal end of the second cantilever rib member, wherein the first and second leg members are arranged, in use, to extend inward of respective nasal orifices along the septum, the first and second intermediate sections are arranged, in use, to engage with the septum and extend from the septum behind the columella and alar fibrofatty tissue of the nose allowing the first and second cantilever rib members, in use, to extend along the respective nasal orifices to an inner wall of the nostrils.

[0030] Some embodiments relate to a nasal dilator device comprising a substantially U-shaped body including a central portion arranged to span a septum of a nose when worn by a user and first and second leg members extending from the central portion in a first plane a first cantilever rib member extending outward from the U-shaped body in

a second plane a second cantilever rib member extending outward from the U-shaped body in a third plane wherein the first and second cantilever rib members extend away from each other a first intermediate section connecting an end of the first leg member to a proximal end of the first cantilever rib member, wherein the first intermediate section extends between the first plane and second plane and a second intermediate section connecting an end of the second leg member to a proximal end of the second cantilever rib member, wherein the second intermediate section extends between the first plane and the third plane.

[0031] In some embodiments, the first and second cantilever rib members may be arcuate cantilever rib members, each having a curvature along its length. In some embodiments, the first and second intermediate sections are arcuate intermediate sections, each having a curvature along its length.

[0032] The first and second intermediate sections may be arranged, in use, to extend along a length of the septum and the first and second cantilever rib members may be each arranged, in use, to extend from a floor of a respective nasal orifice to an inner wall of the nostrils.

[0033] In some embodiments, the first and second intermediate portions may extend obtusely from the ends of the first and second leg members. The second and third planes may be converging planes. In some embodiments, the first and second cantilever rib members may exhibit an elongate arched profile which approximates at least a portion of one of a circle, ellipse or parabola.

[0034] In some embodiments, the first and second leg members may be inclined towards each other such that a relatively greater distance is provided between the first and second leg members towards the central portion to accommodate a columella of a nose when donned by the user. In some embodiments, the first and second intermediate sections may be inclined away from each other to assist in urging the respective first and second cantilever rib members against inner walls of respective nostrils when worn by the user.

[0035] The first and second cantilever rib members may comprise respective first and second nostril engaging elements for engaging with an inner wall of a respective nostril. The first and second nostril engaging elements may be disposed at distal ends of the first and second cantilever rib members, respectively. A series of protrusions is disposed on the first and second nostril engaging elements. The first and second nostril engaging elements may be substantially elongate and the series of protrusions may extend along a length of the first and second nostril engaging elements. For example, the series of protrusions comprises a plurality of an elongated v-shaped protrusions. Enlarged pads may be disposed on the first and second nostril engaging elements to engage with inner walls of the nostrils. The enlarged pads may comprise the series of protrusions. The series of protrusions are formed from an overmould material.

[0036] In some embodiments, the nasal dilator device may comprise at least one projection protruding from and extending along at least a portion of a length of the first and second cantilever rib members. The nasal dilator device may comprise first and second projections, each forming a flange disposed at respective elongate edges of a major surface of the nasal dilator device and extending along at least a portion of a length of the first and second cantilever rib members. The nasal dilator device may comprise at least one projection protruding from and extending along at least a portion of a length of the first and second cantilever rib members, wherein the at least one projection is integrated with and extends from a corresponding protrusion of the series of protrusions. The at least one projection may be formed from an overmould material.

[0037] In some embodiments, the nasal dilator device may further comprise a first and second releasable attachment mechanism for releasably attaching the first and second cantilever rib members, respectively, to the U-shaped body. For example, the releasable attachment mechanisms may be arranged to releasably attach the first and second nostril engaging elements to the first and second leg members, respectively. In some embodiments, the releasable attachment mechanisms may be arranged to releasably attach the first and second nostril engaging elements to the first and second intermediate sections, respectively.

[0038] The releasable attachment mechanisms may each comprise an arm and a socket arranged to receive and engage the arm. A stopper or hook may be disposed at an end of the arm to hinder the arm from withdrawing from the socket. A notch may be disposed on an inner surface of the socket and is configured to engage with the stopper or hook on the arm to hinder the arm from withdrawing from the socket.

[0039] In one embodiment, the arms may be disposed on an inner surface of first and second nostril engaging elements of the first and second cantilever rib members, respectively, and the sockets may be disposed on the first and second leg members. In another embodiment, the arms may be disposed on an inner surface of first and second nostril engaging elements of the first and second cantilever rib members, respectively, and the sockets may be disposed on the first and second intermediate sections. In another embodiment, the sockets may be disposed on an inner surface of first and second nostril engaging elements of the first and second cantilever rib members, respectively, and the arms may be disposed on the first and second leg members. In another embodiment, the sockets may be disposed on an inner surface of first and second nostril engaging elements of the first and second cantilever rib members, respectively, and the arms may be disposed on the first and second intermediate sections. In another embodiment, the nasal dilator device may comprise first and second arm support members projecting from respective first and second intermediate sections and the sockets may be disposed on an inner surface of first and second nostril engaging elements of the first and second cantilever rib members, respectively, and the arms may be disposed on the first and second arm support members.

[0040] In some embodiments, the sockets may comprise substantially elongate housings which taper along their length and provide a relatively broad opening for receiving the arms. The socket may be provided with apertures in side walls of the housing.

[0041] In some embodiments, a capsule may be provided within the socket and may be arranged to be activated by the arm when the arm is received by the socket. The capsule may include at least one of a medicament or compound. The arm may

comprise a coating disposed thereon arranged to release a scent in response to abrasion of the coating. An aperture may be disposed in each of the first and second nostril engaging elements. The aperture may be arranged to receive at least one of a compound, a medicament, and a capsule comprising a medicament or compound emanating a scent.

[0042] Some embodiments relate to a nasal dilator device a nasal dilator device comprising a substantially U-shaped body including a central portion arranged to span a septum of a nose when worn by a user and first and second leg members extending from the central portion in a first plane, a first closed loop structure extending outward from a longitudinal axis of the U-shaped body in a second plane and defining a first aperture; a second closed loop structure extending outward from a longitudinal axis of the U-shaped body in a third plane and defining a second aperture, wherein the first and second closed loop structures extend away from each other, a first intermediate section connecting an end of the first leg member to a proximal end of the first loop structure, wherein the first intermediate section extends between the first plane and second plane, and a second intermediate section connecting an end of the second leg member to a proximal end of the second loop structure, wherein the second intermediate section extends between the first plane and the third plane.

[0043] In some embodiments, the first loop structure may comprise a first flange portion and the second loop structure may comprise a second flange portion, wherein the first and second flange portions are arranged to form a seal with the walls nasal passage in use. For example, the first and second flange portions may project from an outer surface of the first and second loop structures, respectively, and extend along at least a portion of a circumference of the first and second loop structures, respectively.

[0044] In some embodiments, the first loop structure may comprise two or more flange portions and the second loop structure may comprise two or more flange portion, wherein the flange portions are arranged to restrict dislodgement of the nasal dilator from a nose in use.

[0045] In some embodiments, the first and second loop structures may each comprise a filter spanning the first and second apertures defined by the first and second loop structures. The filters may be arranged to snap fit into the first and second loop structures. The filters may be welded to the first and second loop structures.

[0046] In some embodiments, the first and second intermediate sections are arcuate intermediate sections, each having a curvature along its length. The first and second intermediate portions may extend obtusely from the ends of the first and second leg members.

[0047] The first and second intermediate sections may be arranged, in use, to extend along a length of the septum and the first and second loop structures may be each arranged, in use, to extend from a floor of a respective nasal orifice along an inner wall of the nostrils such that the first and second apertures are aligned with a nasal passage of the nose.

[0048] In some embodiments, the second and third planes may be converging planes.

[0049] The first and second leg members may be inclined towards each other such that a relatively greater distance is provided between the first and second leg members towards the central portion to accommodate a columella of a nose when donned by the user. The first and second intermediate sections may be inclined away from each other to assist in urging the respective first and second loop structures against inner walls of respective nostrils when worn by the user.

[0050] In some embodiments, the nasal dilator device may further comprise a film disposed on a surface of the nasal dilator and a removable seal provided on the film to mitigate release of a compound from the film.

[0051] In some embodiments, the nasal dilator device may further comprise an overmould disposed on at least one of the central portion, the leg members, the intermediate sections and the arcuate cantilever rib members. The overmould may be

infused with a compound, a medicament, a fragrance or an aroma. The nasal dilator device may be composed of a substrate material infused with a medicament, a fragrance or an aromatic agent.

[0052] In some embodiments, the central portion comprises a tab extending in a direction substantially opposite to the first and second leg members to assist with insertion, removal and/or placement of the nasal dilator device. The tab may be removeable from the nasal dilator device.

[0053] Some embodiments relate to a nasal dilator device comprising a substantially U-shaped body including: a central portion arranged to span a septum of a nose when worn by a user, and first and second leg members extending from the central portion, first and second cantilever rib members extending outward from a longitudinal axis of the U-shaped body and away from one another, a first intermediate section connecting an end of the first leg member to a proximal end of the first cantilever rib member, and a second intermediate section connecting an end of the second leg member to a proximal end of the second cantilever rib member, wherein the first and second leg members are arranged, in use, to extend inward of respective nasal orifices along the septum, the first and second intermediate sections are arranged, in use, to extend along a length of the septum behind the columella and alar fibrofatty tissue of the nose and the first and second cantilever rib members are each arranged, in use, to extend from a floor of the respective nasal orifices to an inner wall of the nostrils.

Brief Description of Drawings

[0054] Embodiments are described in further detail below, by way of example, with reference to the accompanying drawings, in which:

[0055] Figure 1A is front perspective view of a nasal dilator device according to some embodiments;

[0056] Figure 1B is a further front perspective view of the nasal dilator device of Figure 1A;

[0057] Figure 1C is a front view of the nasal dilator device of Figure 1A;

[0058] Figure 1D is a rear perspective view of the nasal dilator device of Figure 1A;

[0059] Figure 1E is a partial side view of the nasal dilator device of Figure 1A;

[0060] Figure 2 is a front perspective view of a nasal dilator device including fin-like structures disposed thereon, according to some embodiments;

[0061] Figure 3A is a perspective view of a user donning the nasal dilator device of Figures 1A to 1E;

[0062] Figure 3B is a side view of the user of Figure 3A;

[0063] Figure 4A is a rear perspective view of a nasal dilator device according to some embodiments;

[0064] Figure 4B is a front perspective view of the nasal dilator device of Figure 4A;

[0065] Figure 4C is a front view of the nasal dilator device of Figure 4A in a closed configuration;

[0066] Figure 4D is a front view of the nasal dilator device of Figure 4A in a partially closed configuration;

[0067] Figure 5 is a front perspective view of a user donning the nasal dilator device of Figures 4A to 4D;

[0068] Figure 6A is a front view of a nasal dilator device in a partially closed configuration, wherein the nasal dilator device includes a capsule, according to some embodiments;

[0069] Figure 6B is a front view of a nasal dilator device of Figure 6A in a closed configuration;

[0070] Figure 7A is a front view of a nasal dilator device including a film according to some embodiments;

[0071] Figure 7B is a front perspective view of the nasal dilator device of Figure 7A;

[0072] Figure 8A is a front perspective view of a nasal dilator device according to some embodiments;

[0073] Figure 8B is a further front perspective view of the nasal dilator device of Figure 8A;

[0074] Figure 8C is a top view of the nasal dilator device of Figure 8A;

[0075] Figure 9 is a rear perspective view of a nasal dilator device with an attachable filter, according to some embodiments;

[0076] Figure 10 is a rear perspective view of a nasal dilator device including a filter, according to some embodiments;

[0077] Figure 11 is front perspective view of a nasal dilator device, according to some embodiments;

[0078] Figure 12A is front perspective view of a nasal dilator device, according to some embodiments;

[0079] Figure 12B is a front view of the nasal dilator device of Figure 12A in a substantially closed configuration;

[0080] Figure 12C is a rear perspective view of the nasal dilator device of Figure 12A;

[0081] Figure 13A is front perspective view of a nasal dilator device, according to some embodiments;

[0082] Figure 13B is back view of the nasal dilator device of Figure 13A;

[0083] Figure 14A is front perspective view of a nasal dilator device, according to some embodiments;

[0084] Figure 14B is a top view of the nasal dilator device of Figure 14A;

[0085] Figure 14C is a front view of the nasal dilator device of Figure 14A;

[0086] Figure 14D is a side view of the nasal dilator device of Figure 14A;

[0087] Figure 14E is a bottom view of the nasal dilator device of Figure 14A;

[0088] Figure 14F is a front perspective view of the nasal dilator device of Figure 14A in a closed state; and

[0089] Figure 15 is a front perspective view of a nasal dilator device, according to some embodiments.

Description of Embodiments

[0090] Described embodiments generally relate to nasal dilator devices for facilitating respiration. Some embodiments relate to nasal dilator devices to be fitted to the nose to facilitate or improve respiration during sleeping and/or sporting activities and/or for

general day-to-day wear. Some embodiments relate to nasal dilator devices including filtration mechanisms to filter airflow during respiration and other embodiments relate to nasal dilator devices including agent delivery mechanisms for delivery of fragrances and/medicaments to the nose during respiration.

[0091] Referring to Figure 1A to 1E, there is illustrated a nasal dilator device, generally indicated at 100 and substantially symmetrical about a longitudinal axis L, according to some embodiments. The nasal dilator device 100 comprises a generally U-shaped body 102 having a central portion 104 and first and second leg members, 106a and 106b, respectively, extending from the central portion 104 in a first plane P1.

[0092] The nasal dilator device comprises a first intermediate section 108a extending from an end 107a of the first leg member 106a and a second intermediate section 108b extending from an end 107b of the second leg member 106b. In some embodiments, and as depicted in Figures 1A to 1E, the first and second intermediate portions 108a, 108b, may be curved or arcuate along their length. In other embodiments, the first and second intermediate portions 108a, 108b may be substantially straight along their length or may comprise a plurality of angled or arcuate portions. The first and second intermediate portions 108a, 108b may extend obtusely from the first plane P1. For example, the first and second intermediate portions 108a, 108b may extend obtusely from the first and second ends 107a, 107b, for example, substantially at an angle of between approximately 95° and 130° to the longitudinal axis. For example, the first and intermediate sections 108a, 108b may deviate by approximately 100° from the longitudinal axis.

[0093] Referring again to Figures 1A to 1E, the nasal dilator device 100 comprises a first rib member 110a projecting from the first intermediate section 108a in a second plane P2 and a second rib member 110b projecting from the second intermediate section 108b in a third plane P3. In some embodiments, the first and second rib members 110a, 110b may project substantially outward or laterally of the longitudinal axis of the U-shaped body 102. For example, the first and second rib members 110a, 110b may be cantilever rib members that extend from the first and second intermediate

sections 108a, 108b, respectively outwardly from the longitudinal axis and away from one another in a substantially cantilever manner. In some embodiments, the first and second rib members 110a, 110b may be arcuate rib members 110a, 110b or arcuate cantilever rib members 110a, 110b.

[0094] In some embodiments, the first and second rib members 110a, 110b may exhibit an elongate arched or bow-like profile which may approximate at least a portion of a circle, ellipse or parabola. For example, the first and second rib members 110a, 110b may extend arcuately along the second and third planes, P2 and P3, respectively in a direction substantially toward the first plane P1.

[0095] The first and second rib members 110a, 110b may be flexible and resiliently biased away from the first and second intermediate sections 108a, 108b, respectively, to allow the first and second rib members 110a, 110b to be compressed for insertion into the nose of a user and to reform once placed inside the nose to thereby dilate the nostrils as discussed in more detail below with reference to Figures 3A and 3B. As best depicted in Figures 1A and 1B, the first intermediate section 108a may extend or transition between the first plane P1 and the second plane P2 to interconnect the end 107a of the first leg member 106a to a proximal end 109a of the first rib member 110a and the second intermediate section 108b may extend or transition between the first plane and the third plane to interconnect the end 107b of the second leg member 106b to a proximal end 109b of the second rib member 110b.

[0096] In some embodiments, the configuration of the first and second intermediate sections 108a, 108b may be associated with an orientation or location of the first and second rib members 110a, 110b with respect to the U-shaped body 104. For example, the configuration of the first and second intermediate sections 108a, 108b may dictate or define an angle between the first plane P1 and the second plane P2 and between the first plane P1 and the third plane P3, respectively. The second and third planes, P2 and P3, may each form an acute angle, a right angle, or substantially right angle or an obtuse angle with the first plane P1. For example, the second and third planes P2 and P3, may be converging planes and may each form an obtuse angle of approximately 95°

to 130° with the first plane P1 such that the first and second intermediate sections 108a, 108b take the form of obtuse arcuate sections. In some embodiments, the first, second and third planes, P1, P2, P3 may be different from each other and in some embodiments, the second and third planes, P2, P3 may be the same plane and may be different to the first plane P1.

[0097] The first and second intermediate sections 108a, 108b may be inclined away from or diverge from one another to assist in urging the respective first and second rib members 110a, 110b against inner walls of respective nostrils when worn by the user.

[0098] As depicted in Figures 1A to 1E, the first and second rib members 110a, 110b, of the nasal dilator device 100 may comprise respective first and second nostril engaging elements, 112a and 112b, disposed at distal ends 114a, 114b, of the first and second arcuate rib members 110a, 110b, respectively, for engaging with inner walls of respective nostrils when worn by a user. In some embodiments, the first and second nostril engaging elements, 112a, 112b may comprise relatively large surface areas 116a, 116b with respect to the first and second arcuate rib members 110a, 110b.

[0099] In some embodiments, the first and second nostril engaging elements 112a, 112b may have pads 118a, 118b, disposed thereon, to engage with the inner walls of the nostrils. For example, the pads 118a, 118b may be disposed on the relatively large major surface areas 116a, 116b of the nostril engaging elements, 112a and 112b and may be enlarged with respect to the first and second arcuate rib members 110a, 110b, and/or the nostril engaging elements, 112a and 112b.

[0100] Referring now to Figure 2, there is illustrated a nasal dilator device, generally indicated at 200, according to some embodiments. The nasal dilator device 200 may comprise similar components and elements to those of nasal dilator device 100 depicted in Figures 1A to 1E and accordingly those similar components and elements are denoted like numerals.

[0101] In some embodiments, as depicted in Figure 2, the pads 118a, 118b of the nasal dilator device 200 may be composed of a relatively soft overmould material, for example a polymer material such as thermoplastic elastomer (TPE) and/or may be provided with a series of protrusions, fins or fin-like structures 220 to provide a comfortable and/or grippable surface for engaging with the inner walls of the nostrils. In some embodiments, such an overmould material may be provided on at least a portion of the rib members 110a, 110, and/or on at least a portion of the intermediate sections 108a, 108b.

[0102] The nasal dilator device 100, 200 may be configured to be orientated in a manner such that the first and second nostril engaging elements 112a, 112b may be positioned at a junction of the greater alar cartilage and lateral nasal cartilage, providing improved support for dilation of the nasal passage 308, as discussed in more detail with reference to Figures 3A and 3B below.

[0103] Figure 3A is a perspective view of a user, generally indicated at 300, wearing or donning the nasal dilator device 100 of Figure 1A and Figure 3B is a cross sectional view taken along a midline A-A of the nose of the user of Figure 3A.

[0104] As depicted in Figures 3A and 3B, the nasal dilator device 100 is configured to be orientated such that the central portion 104 spans a septum 302, and in particular, a columella 310 (the terminal section or fleshy external end of the septum) of a nose 304 and is positioned toward a tip 306 of the nose 304 and the first and second leg members 106a, 106b extend inward, along a nasal passage 308. For example, the first and second leg members 106a, 106b, may extend inward at an angle of approximately 30 to 40 degrees to a midline A-A of the nose 304. The first and second intermediate sections 108a, 108b may extend along a length of the septum 302 behind the columella 310 and the fibrofatty tissue 305 or bulbous region around the base of the nostrils 314 and the first and second rib members 110a, 110b, each may extend from a floor 312 of the nasal passage 308 behind the columella 310 and the fibrofatty tissue 305 or bulbous region around the base of the nostrils 314 to an inner wall (not shown) of the nostrils 314. In this way, the nasal dilator device 100 may be securely retained within the nose

304 with little or no pinching of or pressure being exerted on the septum 302.

Furthermore, the ergonomic shape of the intermediate portions 108a, 108b allows the nasal dilator device to sit within the nose in a manner that may accommodate various shapes and sizes of noses, including those having hanging columellas 310.

[0105] In some embodiments, the first and second rib members 110a, 110b of the nasal dilator device 100 are composed of a flexible material and are generally squeezed or compressed by a user into a compressed state to allow insertion into the nasal passages 308 of the nose 304. The first and second rib members 110a, 110b may be biased to reform or revert to a natural uncompressed state and once inserted into the nasal passage 308, the first and second rib members 110a, 110b may each exert an outward force on the inner wall (not shown) of the nostril 314 and on the floor 312 of the nose 304, to thereby dilate the nasal passage 308. Thus, as opposed to exerting pressure on the septum 302 to dilate the nasal passage 308, the intermediate portions 108a, 108b, of nasal dilator device 100 are effective to cause the first and second rib members 110a, 110b to use the floor 312 of the nose 304 as a support structure for dilation of the nostrils 314. By using the floor 312 of the nose 304 as a support structure or anchor from which the first and second rib members 110a, 110b may launch or push off from, any pinching or exertion of force on the septum may be mitigated or avoided and a more comfortable and natural or anatomical fit may be achieved.

[0106] The nasal dilator device 100 is configured to cooperate with internal contours of the nose 304 and sit securely and comfortably in the nose, whilst mitigating obstruction of air flow through the nasal passage 308. For example, the rib members 110a, 110b, may be curved or arcuate along their length to correspond with the internal contours of the nose 304 and provide a more comfortable fit. In some embodiments, the first and second leg members 106a, 106b may be inclined toward each other or converge such that a relatively greater distance is provided between the first and second leg members 106a, 106b towards the central portion 104 in order to accommodate the columella 310 and to assist in holding the nasal dilator device 100 in place when worn.

[0107] Referring now to Figures 4A to 4D, there is illustrated a nasal dilator device, generally indicated at 400, according to some embodiments. The nasal dilator device 400 may comprise similar components and elements to those of nasal dilator device 100 depicted in Figures 1A to 1E and accordingly those similar components and elements are denoted like numerals.

[0108] In addition to those similar components and elements of nasal dilator device 100, nasal dilator device 400 may comprise a first and second releasable attachment mechanism 402a and 402b, respectively. The first and second releasable attachment mechanism 402a, 402b may comprise mating or interlocking components and may be employed to releasably attach the first and second rib members, 110a and 110b, respectively, to the U-shaped body 102, to thereby define first and second adjustable looped structures, 411a, and 411b, respectively.

[0109] In some embodiments, the first and second releasable attachment mechanisms 402a, 402b may comprise respective arms 404a, 404b, such as pins, extending from respective reverse or inner surfaces 406a, 406b of the first and second nostril engaging elements 112a, 112b. The first and second releasable attachment mechanisms 402a, 402b may comprise respective sockets 408a, 408b for receiving and/or engaging the respective arms 404a, 404b. The first and second releasable attachment mechanisms 402a, 402b may be configured to allow a user to selectively adjust a degree of dilation or expansion and contraction of the first and second rib members 110a and 110b with respect to the U-shaped body 102.

[0110] For example, and as best illustrated in Figures 4C and 4D, the arms 404a, 404b may include at least one of or a series of serrations, detents or protrusions 410 arranged to engage with at least one of or a series of grooves or ridges 412 provided on or within the sockets 408a, 408b. For example, the grooves or ridges 412 may extend downwardly from an upper jaw portion 414 of the sockets 408a, 408b and/or may extend upwardly from a lower jaw portion 416.

[0111] Application of sufficient force by a user to the first and second releasable attachment mechanisms 402a, 402b may be effective to move the arms 404a, 404b with respect to the sockets 408a, 408b and overcome a restrictive force between the detents 410 and the grooves 412 to allow the detents 410 and/or the grooves 412 to deform and the degree or level of dilation to be adjusted. The engagement of the detents 410 with the grooves 412 may provide a sufficient restrictive force to hold the arms 404a, 404b fixed when provided in the nose 304, as depicted in Figure 5.

[0112] The arms 404a, 404b may comprise stoppers 414 at their ends to prevent or hinder the arms 404a, 404b from disengaging from or withdrawing from the respective sockets 408a, 408b. For example, application of a relatively large pulling force may be sufficient to cause the arms 404a, 404b to withdraw from the sockets 408a, 408b. In some embodiments, the stoppers 414 may be arrow shaped.

[0113] In some embodiments, the sockets 408a, 408b may be disposed on the first and second intermediate sections 108a, 108b and extend therefrom towards the respective arms 404a, 404b. The releasable attachment mechanisms 402a, 402b may be arranged to releasably attach or lock the first and second nostril engaging elements 112a, 112b to the first and second intermediate sections 108a, 108b.

[0114] In other embodiments, the sockets 408a, 408b may be disposed on the first and second leg members 106a, 106b and extend therefrom towards the respective arms 404a, 404b. The releasable attachment mechanisms 402a, 402b may be arranged to releasably attach or lock the first and second nostril engaging elements 112a, 112b to the first and second leg members 106a, 106b.

[0115] In other embodiments, the sockets 408a, 408b may be disposed on the first and second rib members 110a, 110b and extend therefrom towards the respective arms 404a, 404b. The attachment mechanisms 402a, 402b may be arranged to releasably attach or lock the first and second nostril engaging elements 112a, 112b to the first and second rib members 110a, 110b.

[0116] In other embodiments, the first and second releasable attachment mechanisms 402a, 402b may comprise respective sockets 408a, 408b, extending from the respective reverse or inner surfaces 406a, 406b, of the first and second nostril engaging elements 112a, 112b and respective arms 404a, 404b extending from the first and second intermediate sections 108a, 108b, the first and second leg members 106a, 106b, or the first and second rib members 110a, 110b.

[0117] As illustrated in Figure 4C, the arms 404a, 404b may be fully or substantially fully inserted into the respective sockets 408a, 408b to enable the nasal dilator device 400 to adopt or assume a fully closed or substantially fully closed state, to thereby tighten or contract the looped structures 411a, 411b.

[0118] As illustrated in Figure 4D, the arms 404a, 404b may be partially inserted into the sockets 408a, 408b to enable the nasal dilator device 400 to adopt or assume a partially closed state, to provide for looser or less tight looped structures 411a, 411b and accommodate variations in nasal passage sizes.

[0119] Referring to Figures 6A and 6B, there is depicted a nasal dilator device 600 according to some embodiments. The nasal dilator device 600 may comprise similar components and elements to those of nasal dilator device 400 depicted in Figures 4A to 4D and accordingly those similar components and elements are denoted like numerals.

[0120] The nasal dilator device 600 comprises at least one capsule 602 disposed within respective sockets 408a, 408b. The capsule 602 may include an agent such as a medicament and/or a fragrance or aromatic agent. As depicted in Figure 6B, the arms 404a, 404b are configured to activate, pierce or burst the capsules 602 to release the agent, medicament and/or fragrance or aromatic agent when inserted into the sockets 408a, 408b. In this way, the medicament and/or fragrance or aromatic agent is released only when the capsule 602 is activated, pierced or burst, thereby increasing a longevity or “shelf-life” and/or protecting the integrity of the medicament and/or aromatic agent. For example, the agent may be an aromatic scent such as an essential oil blend or synthetic fragrance blend to provide an olfactory and/or physiological response such as

decongesting the nasal passages 318, promoting relaxation, promoting sleepiness, suppressing appetite or a medicament such as a drug to reduce pain such as a migraine.

[0121] Referring to Figures 7A and 7B, there is depicted the nasal dilator device 700 according to some embodiments. The nasal dilator device 700 may comprise similar components and elements to those of nasal dilator device 400 depicted in Figures 4A to 4D and accordingly those similar components and elements are denoted like numerals. The nasal dilator device 700 comprises at least one coating or film 702 arranged to release a fragrance, aroma or medicament. In some embodiments, the film 702 is arranged to release a fragrance, aroma or medicament in response to abrasion, such as scratching, scraping. The film 702 may be provided with an outer cover, seal or strip 704 to protect the film 702 from unintended abrasion, as depicted in Figures 7A and 7B at two separate stages of removal from the nasal dilator device 400.

[0122] In other embodiments, the coating or film 700 may be arranged to release a fragrance, aroma or medicament in response to the removal or peeling off of the outer cover, strip or seal 704. In some embodiments, a fragrance, aroma or medicament may be provided or retained between two strips or films 702 forming a blister.

[0123] The coating or film 702 may be comprise a polymer or a fibre. The coating or film 702 may be in the form of a “scratch and sniff” technology or peel off technology.

[0124] In some embodiments, as depicted in Figures 7A and 7B, the coating or film 702 may be disposed on a surface of at least one of the attachment mechanisms 402a, 402b, such as on an inner surface of the looped structures 411a, 411b. In other embodiments, the coating or film 702 may be disposed on the central portion 102, the first and second leg members 106a, 106b, the first and second intermediate sections 108a, 108b, the rib members 110a, 110b, and/or the first and second nostril engaging elements 112a, 112b.

[0125] In some embodiments, an aperture (not shown) is disposed in each of the first and second nostril engaging elements and is arranged to receive an agent, a compound,

a medicament, a capsule, and/or a housing or compact arranged to receive an agent, medicament and/or a fragrance or aromatic agent. The agent may be absorbed by the inner walls of the nostrils transdermally and/or may be absorbed by mucosa in the nostrils 314.

[0126] Referring to Figures 8A and 8B, there is depicted a nasal dilator device, generally indicated at 800 and substantially symmetrical about a longitudinal axis L, according to some embodiments. The nasal dilator device 800 may comprise a generally U-shaped body 802 having a central portion 804 and first and second leg members, 806a and 806b, respectively, extending from the central portion 804 in a first plane P1.

[0127] The nasal dilator device 800 comprises a first intermediate section 808a extending from an end 807a of the first leg member 806a and a second intermediate section 808b extending from an end 807b of the second leg member 806b. In some embodiments, and as depicted in Figures 8A and 8B, the first and second intermediate portions 808a, 808b, may be curved or arcuate along their length. In other embodiments, the first and second intermediate portions 808a, 808b may be substantially straight along their length or may comprise a plurality of angled or arcuate portions. For example, the first and second intermediate portions 808a, 808b may extend obtusely from the first and second ends 807a, 807b, for example, substantially at an angle of between approximately 95° and 130° to the longitudinal axis.

[0128] As depicted in Figures 8A to 8C, the nasal dilator device 800 comprises a first loop structure 811a projecting from the first intermediate section 808a in a second plane P2 and a second loop structure 811b projecting from the second intermediate section 808b in a third plane P3. In some embodiments, the first and second loop structures 811a, 811b may project substantially outward or lateral of the longitudinal axis of the generally U-shaped body 802 and away from one another. In some embodiments, the loop structure 811a, 811b may exhibit an elongate arched or curved profile which may substantially take the form of a circle, ellipse or parabola.

[0129] In some embodiments, the first intermediate section 808a may extend or transition between the first plane P1 and the second plane P2 to interconnect the end 807a of the first leg member 806a to a proximal end 809a of the first loop structure 811b and the second intermediate section 808b may extend or transition between the first plane P1 and the third plane P3 to interconnect the end 807b of the second leg member 806b to a proximal end 809b of the second loop structure 811b.

[0130] In some embodiments, the configuration of the first and second intermediate sections 808a, 808b may be associated with an orientation or location of the first and second loop structures 811a, 811b with respect to the U-shaped body 804. For example, the configuration of the first and second intermediate sections 808a, 808b may dictate or define an angle between the first and second planes, P1 and P2 and between the first and third planes, P1 and P3, respectively. The second and third planes, P2 and P3, may each form an acute angle, a right angle, or substantially right angle or an obtuse angle with the first plane P1. For example, the second and third planes P2 and P3, may be converging planes or intersecting planes and may each form an obtuse angle of approximately 95° to 130° with the first plane P1 such that the first and second intermediate sections 108a 108b take the form of obtuse arcuate sections. In some embodiments, the first, second and third planes, P1, P2, P3 may be different from each other and in some embodiments, the second and third planes, P2, P3 may be the same plane and may be different to the first plane P1.

[0131] In some embodiments, the first and second leg members 806a, 806b may be inclined toward each other or converge such that a relatively greater distance is provided between the first and second leg members 806a, 806b towards the central portion 804 in order to accommodate the columella 310 and to assist in holding the nasal dilator device 800 in place when worn.

[0132] In some embodiments, the first and second intermediate sections 808a, 808b may be inclined away from or diverge from one another to assist in urging the respective first and second loop structures 811a, 811b against inner walls of the nose when worn by the user. In some embodiments, the first and second loop structures 811a

and 811b may comprise first and second flanged portions, 812a and 812b, respectively. For example, first and second flanged portions 812a and 812b may project from an outer surface of the loop structures 811a and 811b, respectively, and extend along at least a portion of a circumference of the loop structures 811a and 811b. In some embodiments, the first and second loop structures 811a and 811b may each comprise two or more flanged portions (not shown). For example, the more than two or more flanged portions (not shown) may project from an outer surface of the loop structures 811a and 811b and extend along at least a portion of a circumference of the loop structures 811a and 811b. The flanged portions 812a and 812b may provide additional compliance to the loop structures 811a, 811b and may provide for or create a drag effect which may improve retention of the nasal dilator device 1300 in a desired position despite movement of the user.

[0133] In some embodiments, the first and second flanged portions 812a and 812b may be comprise an overmould material, for example, flexible TPE, to thereby provide an improve sealing of the loop structures 811a and 811b to the nasal orifices.

[0134] Referring to Figure 9, there is depicted the nasal dilator device 900 according to some embodiments. The nasal dilator device 900 may comprise similar components and elements to those of nasal dilator device 800 depicted in Figures 8A to 8C and accordingly those similar components and elements are denoted like numerals.

[0135] The first and second loop structures 811a, 811b, of the nasal dilator device 900 may be each arranged to receive a filter 902a, and 902b, respectively. The filters 902a, 902b, may be arranged or configured to span apertures defined by the first and second loop structures 811a, 811b.

[0136] The filters 902a, 902b may be composed of a fine woven mesh or an open celled porous material, such as a foam or compressed fibre. The filters 902a, 902b may be employed to filter out airborne particles such as bacteria, dust, pollens, and/or other allergens.

[0137] In some embodiments, as depicted in Figure 9, the filters 902a, 902b, may be replaceable and may be arranged to be removeably connected to the first and second loop structures 811a, 811b respectively. For example, the filters 902a, 902b, may be configured to “snap-fit” into the first and second loop structures 811a, 811b respectively.

[0138] Referring to Figure 10, there is depicted the nasal dilator device 1000 according to some embodiments. The nasal dilator device 1000 may comprise similar components and elements to those of nasal dilator device 800 depicted in Figures 8A to 8C and accordingly those similar components and elements are denoted like numerals

[0139] The filters 1002a, 1002b of the nasal dilator device 1000 may be fixed to the first and second loop structures 811a, 811b respectively. For example, the filters 1002a, 1002b may be integrally formed with the first and second loop structures 811a, 811b or may be welded or ultrasonically welded to the first and second loop structures 811a, 811b.

[0140] In some embodiments, the nasal dilator device 100, 200, 400, 600, 700, 800, 900, 1000 may comprise an overmould disposed on at least one of the central portion, the leg members, the sections and the rib members. The overmould may be infused with a medicament and/or fragrance.

[0141] In some embodiments, the nasal dilator devices 100, 200, 400, 600, 700, 800, 900, 1000 may comprise a tab (not shown) extending outward from the central portion in a direction substantially opposite to the first and second leg members to assist with insertion, removal and/or placement of the nasal dilator device 100, 200, 400, 600, 700, 800, 900, 1000. The tab (not shown) may be removeable from the nasal dilator device, for example, by tearing the tab along a perforated line connecting the tab to the central portion 104, 804.

[0142] The U-shaped body 102, 802, the intermediate sections 108a, 108b, 808a, 808b, the rib members 110a, 110b, and the loop structure 811a, 811b may be composed

of a polymer material such as thermoplastic elastomer (TPE) and/or thermoplastic polypropylene (PP). In some embodiments, the U-shaped body 102 and/or the intermediate sections 108a, 108b may be configured to be more rigid than the rib members 110a, 110b. For example, the U-shaped body 102 and/or the intermediate sections 108a, 108b and/or the rib members 110a, 110b may be composed of different materials or materials having differing hardness or stiffness. In some embodiments, the relative flexibility of the rib members 110a, 110b with respect to the U-shaped body 102 and/or the intermediate sections 108a, 108b may be derived from the length and/or thickness of the rib members 110a, 110b.

[0143] In some embodiments, an overall width of the nasal dilator device 100, 200, 400, 600, 700, 800, 900, 1000 may be in a range of approximately 20mm to 35mm when fully closed and approximately 25mm to 40mm when fully open, a length of the central portion 102, 802 may be in a range of approximately 5mm to 10mm, a length of the leg members 106a, 106b, 806a, 806b may be within a range of approximately 5mm to 12mm, and a length of the intermediate sections 108a, 108b, 808a, 808b may be in a range of approximately 7mm to 15mm and the rib members 110a, 110b, 810a, 810b may be in a range of approximately 15mm to 30mm. For example, in one embodiment, the overall width of the nasal dilator device 100, 200, 400, 600, 700, 800, 900, 1000 may be 25mm when fully closed, 27.4mm when open and the length of the leg members 106a, 106b, 806a, 806b may be 14.6mm. In another embodiment, the overall width of the nasal dilator device 100, 200, 400, 600, 700, 800, 900, 1000 may be 27.2mm when fully closed, 29.3mm when open and the length of the leg members 106a, 106b, 806a, 806b may be 17.5mm. In another embodiment, the overall width of the nasal dilator device 100, 200, 400, 600, 700, 800, 900, 1000 may be 29mm when fully closed, 31.6mm when open and the length of the leg members 106a, 106b, 806a, 806b may be 20.4mm.

[0144] Referring now to Figure 11, there is illustrated a nasal dilator device, generally indicated at 1100, according to some embodiments. The nasal dilator device 1100 may comprise similar components and elements to those of nasal dilator device 100 depicted

in Figures 1A to 1E and accordingly those similar components and elements are denoted like numerals.

[0145] In some embodiments, as depicted in Figure 11, the relatively large surface areas 116a, 116b of the nostril engaging members 112a, 112b are provided with a series of protrusions 1102. The nostril engaging members 112a, 112b may be substantially elongate, for example, extending from the distal ends 114a, 114b of the rib members 110a, 110b, and the protrusions 1102 may extend along a length of the surface areas 116a, 116b of the nostril engaging members 112a, 112b. In some embodiments, the surface areas 116a, 116b of the nostril engaging members 112a, 112b may be substantially oval, rectangular, triangular or truncated triangular in shape. In some embodiments, the series of protrusions 1102 may form a u-shaped pattern, a v-shaped pattern, or elongated v or ✓ (tick) shaped pattern, such as a chevron design, comprising a plurality of aligned u-shaped, v-shaped or elongated v or ✓ (tick) shaped protrusions 1102.

[0146] As depicted in Figure 11, the rib members 110a, 110b of the nasal dilator device 1100 may comprise one or more projections 1104, flanges or ridges protruding from a major surface 1106 of the nasal dilator device 1100 and extending along at least a portion of a length of the rib members 110a, 110b. For example, the major surface 1106 of the nasal dilator device 1100 may be the surface of the nasal dilator device 1100 configured, in use, to engage with or face nasal passage walls of a user's nose. In some embodiments, as depicted in Figure 11, the nasal dilator device 1100 may comprise first and second projections 1104, each forming a flange disposed at respective elongate edges 1106a, 1106b of the major surface 1106 of the nasal dilator device 1100.

[0147] The one or more projections 1104 extend from the distal ends 114a, 114b of the rib members 110a, 110b toward the intermediate sections 108a, 108b. For example, as depicted in Figure 11, the one or more projections 1104 may integrate with and extend from corresponding protrusions 1102 of the series of protrusions 1102 provided on the surface areas 116a, 116b of the nostril engaging members 112a, 112b. In some

embodiments, the one or more projections 1104 may be disposed on and project from the major surface 1106 of the intermediate sections 108a, 108b and/or the leg members 106a, 106b and/or the central portion 104 of the nasal dilator device 1100.

[0148] In some embodiments, the nasal dilator device 1100 may comprise pads 118a, 118b disposed on the enlarged surface areas 116a, 116b of the nostril engaging members 114a, 114b. The pads 118a, 118b may be composed of a relatively soft overmould material, for example, a polymer material such as thermoplastic elastomer (TPE). In some embodiments, the pads 118a, 118b may be configured to cooperate with and conform to the surface area 116a, 116b and the series of protrusions 1102 such that the overmoulded series of protrusions 1102 project or protrude from the overmoulded surface area 116a, 116b. In other embodiments, the surface areas 116a, 116b of the nostril engaging members 114a, 114b may be substantially smooth or flat and the pads 118a, 118b may comprise or form the series of protrusions 1102. For example, the pads 118a, 118b may be substantially elongate, for example, extending from the distal ends 114a, 114b of the rib members 110a, 110b, and the protrusions 1102 may extend along a length of the pads 118a, 118b.

[0149] In some embodiments, an overmould material may be provided on at least a portion of the rib members 110a, 110b, on at least a portion of the intermediate sections 108a, 108b, and/or on at least a portion of the u-shaped body 102. For example, the overmould may be configured to cooperate with and conform to the major surface 1106 of the nasal dilator device 1100 and the one or more projections 1104 such that the overmoulded projections 1104 project or protrude from the overmoulded major surface 1106. In other embodiments, the major surface 1106 of the nasal dilator device 1100 may be substantially smooth or flat and an overmould disposed thereon may comprise or form the one or more projections or ridges 1104.

[0150] The series of protrusions 1102 and/or the one or more projections 1104 may provide a comfortable and/or grippable surface for engaging with the inner walls of the nose in use. The series of protrusions 1102 and/or the one or more projections 1104 may provide for or create a drag effect which may improve retention of the nasal dilator

device 1100 in a desired position in a user's nose, in use, despite movement or motion of the user.

[0151] Referring now to Figures 12A, 12B and 12C, there is illustrated a nasal dilator device, generally indicated at 1200, according to some embodiments. The nasal dilator device 1200 may comprise similar components and elements to those of nasal dilator device 1100 depicted in Figure 11 and accordingly those similar components and elements are denoted like numerals.

[0152] In addition to those similar components and elements of nasal dilator device 1100, nasal dilator device 1200 may comprise a first and second releasable attachment mechanism 1202a and 1202b, respectively. The first and second releasable attachment mechanism 1202a, 1202b may comprise mating or interlocking components and may be employed to releasably attach the first and second rib members, 110a and 110b, respectively, to the U-shaped body 102, to thereby define first and second adjustable looped structures, 1204a, and 1204b, respectively. The first and second releasable attachment mechanisms 1202a, 1202b may allow a user to selectively adjust a degree of dilation or expansion and contraction of the first and second rib members 110a and 110b with respect to the U-shaped body 102.

[0153] The first and second releasable attachment mechanisms 1202a, 1202b may comprise respective arms 1206a, 1206b, such as pins, and respective sockets 1208a, 1208b for receiving and/or engaging the respective arms 1206a, 1206b.

[0154] As illustrated in Figures 12A, 12B and 12C, the arms 1206a, 1206b may be disposed on and extend from respective first and second intermediate sections 108a, 108b toward respective sockets 1208a, 1208b provided on opposite or inner surfaces 1210a, 1210b of the first and second nostril engaging elements 112a, 112b. In some embodiments, as shown in Figures 12A and 12C, the first and second arms 1206a, 1206b extend from respective first and second arm supports 1207a, 1207b projecting from respective first and second intermediate sections 108a, 108b. The releasable attachment mechanisms 1202a, 1202b may therefore be arranged to releasably attach or

lock the first and second nostril engaging elements 112a, 112b to the first and second intermediate sections 108a, 108b.

[0155] In other embodiments, the first and second arms 1206a, 1206b may be disposed on and extend from respective first and second rib members 110a, 110b toward respective first and second sockets 1208a, 1208b to allow the releasable attachment mechanisms 1202a, 1202b to releasably attach or lock the first and second nostril engaging elements 112a, 112b to the first and second rib members 110a, 110b or the first and second arms 1206a, 1206b may be disposed on and extend from respective first and second leg members 106a, 106b toward respective first and second sockets 1208a, 1208b to allow the releasable attachment mechanisms 1202a, 1202b to releasably attach or lock the first and second nostril engaging elements 112a, 112b to the first and second leg members 106a, 106b.

[0156] In other embodiments, the first and second releasable attachment mechanisms 1202a, 1202b may comprise first and second arms 1206a, 1206b extending from the respective opposite or inner surfaces 1210a, 1210b of the first and second nostril engaging elements 112a, 112b and first and second sockets 1208a, 1208b extending from the first and second intermediate sections 108a, 108b, the first and second rib members 110a, 110b, or the first and second leg members 106a, 106b.

[0157] Referring again to Figures 12A, 12B and 12C, the first and second arms 1206a, 1206b may include at least one or a series of serrations, detents or protrusions 1212 arranged to engage with at least one or a series of grooves or ridges 1214 provided on or within the sockets 1208a, 1208b. For example, the grooves or ridges 1214 may extend downwardly from an upper jaw portion 1216 of the sockets 1208a, 1208b and/or may extend upwardly from a lower jaw portion 1218.

[0158] Application of sufficient force by a user to the first and second releasable attachment mechanisms 1202a, 1202b may be effective to move the arms 1206a, 1206b with respect to the sockets 1208a, 1208b and overcome a restrictive force between the detents 1212 and the grooves 1214 to allow the detents 1212 and/or the grooves 1214 to

deform and the degree or level of dilation to be adjusted. The engagement of the detents 1212 with the grooves 1214 may provide a sufficient restrictive force to hold the arms 1206a, 1206b fixed when provided in a user's nose.

[0159] As depicted in Figure 12, the arms 1206a, 1206b may each comprise a stopper or catch 1220 provided at their free ends to prevent or hinder the arms 1206a, 1206b from disengaging from or withdrawing from the respective sockets 1208a, 1208b. For example, in some embodiments, the catch 1220 comprises a hook portion 1222 which is configured to cooperate with a corresponding notch 1224 disposed in an upper or lower jaw portion 1216, 1218 of the socket. Application of a relatively large pulling force may be sufficient to cause the catch 1220 to disengage with the notch 1224 and to cause the arms 1206a, 1206b to withdraw from the sockets 1208a, 1208b.

[0160] In some embodiments, as best shown in Figures 12A and 12C, the sockets 1208a, 1208b may take the form of substantially elongate housings 1226 provided with apertures 1228 or gaps in side walls of the housing 1226. The housings 1226 may be substantially wedge shaped and taper along their length towards opposite or inner surfaces 1210a, 1210b, of the first and second nostril engaging elements 112a, 112b to provide a relatively broad opening to receive the arms 1206a, 1206b. The elongate or wedge shaped housing 1226 may provide for an improved engagement between the arms 1206a, 1206b and sockets 1208a, 1208b and more robust releasable attachment mechanisms 1202a, 1202b. For example, the elongate or wedge shaped housing 1226 may allow a face 1230 of the socket housing to engage with and lie substantially flush with the arm supports 1207a, 1207b, the intermediate sections 108a, 108b, rib members 110a, 110b, or leg members 106a, 106b, when the arms 1206a, 1206b are fully engaged within the sockets 1208a, 1208b, when the nasal dilator device 1200 assumes a closed state. For example, in use, such a configuration may mitigate contact between the sockets 1208a, 1208b and the septum and relative movement or twisting of the sockets as a result of contact with the septum. The elongate or wedge shaped housing 1226 may facilitate improved or easier adjustment of the first and second releasable attachment mechanisms 1202a, 1202b.

[0161] The arms 1206a, 1206b may be fully or substantially fully inserted into the respective sockets 1208a, 1208b to enable the nasal dilator device 1200 to adopt or assume a fully closed or substantially fully closed state, to thereby tighten or contract the looped structures 1211a, 1211b. The arms 1206a, 1206b may be partially inserted into the sockets 1208a, 1208b to enable the nasal dilator device 1200 to adopt or assume a partially closed state, to provide for looser or less tight looped structures 1204a, 1204b and accommodate variations in nasal passage sizes.

[0162] As depicted in figures 12A, 12B, and 12C, the nasal dilator device 1200 may include a series of protrusions 1102 disposed on the nostril engaging members 112a, 112b, and/or one or more projections 1104 protruding from the major surface 1106 of the nasal dilator device 1200, as discussed with reference to Figure 11.

[0163] Similar to the embodiment of nasal dilator device 600 depicted in Figure 6, in some embodiments, the nasal dilator device 1200 may comprise at least one capsule (not shown), which may include an agent such as a medicament and/or a fragrance or aromatic agent, disposed within respective sockets 1208a, 1208b. The arms 1206a, 1206b may be configured to activate, pierce or burst the capsule (not shown) to release the agent, medicament and/or fragrance or aromatic agent when they are inserted into the sockets 1208a, 1208b. In this way, the medicament and/or fragrance or aromatic agent is released only when the capsule (not shown) is activated, pierced or burst, thereby increasing a longevity or “shelf-life” and/or protecting the integrity of the medicament and/or aromatic agent.

[0164] Similar to the embodiment of nasal dilator device 700 depicted in Figure 7, in some embodiments, the nasal dilator device 1200 may comprise at least one coating or film (not shown) arranged to release a fragrance, aroma or medicament. For example, the film (not shown) may be disposed on a surface of at least one of the attachment mechanisms 1202a, 1202b, such as on an inner surface of the looped structures 1204a, 1204b, on the central portion 102, on the first and second leg members 106a, 106b, on the first and second intermediate sections 108a, 108b, on the rib members 110a, 110b, and/or on the first and second nostril engaging elements 112a, 112b.

[0165] In some embodiments, the film (not shown) is arranged to release a fragrance, aroma or medicament in response to abrasion, such as scratching, scraping and may be provided with an outer cover, seal or strip to protect the film (not shown) from unintended abrasion. In other embodiments, the coating or film (not shown) may be arranged to release a fragrance, aroma or medicament in response to the removal or peeling off of an outer cover, strip or seal. In some embodiments, a fragrance, aroma or medicament may be provided or retained between two strips or films (not shown) forming a blister. For example, the coating or film (not shown) may comprise a polymer or a fibre and/or may be in the form of a “scratch and sniff” technology or peel off technology.

[0166] Referring to Figure 13A and 13B, there is illustrated a nasal dilator device, generally indicated at 1300. The nasal dilator device 1300 is substantially symmetrical about a longitudinal axis L, according to some embodiments, and comprises a generally U-shaped body 1302 having a central portion 1304 and first and second leg members, 1306a and 1306b, respectively, extending from the central portion 1304 in a first plane P1.

[0167] The nasal dilator device comprises a first intermediate section 1308a extending from an end 1307a of the first leg member 1306a and a second intermediate section 1308b extending from an end 1307b of the second leg member 1306b. In some embodiments, and as depicted in Figures 13A and 13B, the first and second intermediate portions 1308a, 1308b, may be angled, curved or arcuate along their length and/or may comprise a plurality of angled or arcuate portions. In some embodiments, the first and second intermediate portions 108a, 108b may comprise L-shaped or right angled sections. For example, the first and second intermediate portions 108a, 108b may extend from the first and second ends 107a, 107b in a substantially perpendicular direction to the longitudinal axis L to form substantially right angled intermediate portions.

[0168] The nasal dilator device 1300 comprises a first rib member 1310a projecting from the first intermediate section in a second plane P2 and a second rib member 1310b

projecting from the second intermediate section 1308b in a third plane P3. For example, and as best depicted in Figure 13A, the first intermediate section 1308a may connect the end 1307a of the first leg member 1306a to a proximal end 1309a of the first rib member 1310a and the second intermediate section 1308b may connect the end 1307b of the second leg member 1306b to a proximal end 1309b of the second rib member 110b. In some embodiments, the first intermediate section 1308a extends from the first plane P1 to the second plane P2 and the second intermediate section 1308b extends from the first plane P1 to the third plane P3.

[0169] In some embodiments, the first and second rib members 1310a, 1310b may project substantially outward or laterally of the longitudinal axis of the U-shaped body 1302. For example, the first and second rib members 1310a, 1310b may be cantilever rib members that extend from the first and second intermediate sections 1308a, 1308b, respectively outwardly from the longitudinal axis and away from one another in a substantially cantilever manner. In some embodiments, the first and second rib members 1310a, 1310b may be arcuate rib members 1310a, 1310b or arcuate cantilever rib members 1310a, 1310b.

[0170] In some embodiments, the first and second rib members 1310a, 1310b may exhibit an elongate arched or bow-like profile which may approximate at least a portion of a circle, ellipse or parabola. For example, the first and second rib members 1310a, 1310b may extend arcuately along the second and third planes, P2 and P3, respectively in a direction substantially toward the first plane P1.

[0171] The first and second rib members 1310a, 1310b may be flexible and resiliently biased away from the first and second intermediate sections 1308a, 1308b respectively, to allow the first and second rib members 1310a, 1310b to be compressed for insertion into the nose of a user and to reform once placed inside the nose to thereby dilate the nostrils.

[0172] In some embodiments, the configuration of the first and second intermediate sections 1308a, 1308b may be associated with an orientation, positioning or location of

the first and second rib members 1310a, 1310b with respect to the U-shaped body 1302. For example, the configuration of the first and second intermediate sections 1308a, 1308b may dictate or define a relationship or an angle between the first plane P1 and the second plane P2 and between the first plane P1 and the third plane P3, respectively. The second and third planes, P2 and P3, may each form an acute angle, a right angle, or substantially right angle or an obtuse angle with the first plane P1.

[0173] For example, in some embodiments, the first and second intermediate sections 1308a 1308b may take the form of substantially right angled sections such that the second and third planes P2, P3 each form a right angle with the first plane P1. In other embodiments, the first and second intermediate sections 1308a 1308b may take the form of obtuse angled or arcuate sections such that the second and third planes P2 and P3, each form an obtuse angle with the first plane P1. In other embodiments, the first and second intermediate sections 1308a 1308b may take the form of acute angled or arcuate sections such that the second and third planes P2 and P3, each form an acute angle with the first plane P1.

[0174] In some embodiments, the second and third planes P2 and P3, may be converging planes such that the first and second cantilever rib members are angled and/or extend substantially toward the central portion of the U-shaped body. In other embodiments, the second and third planes are diverging planes such that the first and second cantilever rib members are angled and/or extend substantially toward the central portion of the U-shaped body.

[0175] In some embodiments, the first, second and third planes, P1, P2, P3 may be different from each other and in some embodiments, the second and third planes, P2, P3 may be the same plane and may be different to the first plane P1. In some embodiments, the first and second planes P1, P2 may be orthogonal to the first plane P1.

[0176] For example, in some embodiments, the nasal dilator device 1300 is configured to be orientated such that, in use, the central portion 1304 spans a septum

302, and in particular, a columella (the terminal section or fleshy external end of the septum) of a nose and is positioned toward a tip of the nose and the first and second leg members 1306a, 1306b extend inward of respective nasal orifices, along a nasal passage and/or the septum of the nose. The first and second leg members 1306a, 1306b, may extend inward at an angle of approximately 30 to 40 degrees to a midline of the nose. The first and second intermediate sections 1308a, 1308b may engage with the septum and extend from the septum behind the columella and alar fibrofatty tissue or bulbous region around the base of the nostrils of the nose allowing the first and second rib members, in use, to extend along the respective nasal orifices to an inner wall of the nostrils behind the columella and the fibrofatty tissue or bulbous region around the base of the nostrils to an inner wall of the nostrils. In this way, the nasal dilator device 1300 may be securely retained within the nose with little or no pinching of or pressure being exerted on the septum.

[0177] The first and second intermediate sections 1308a, 1308b may be inclined away from or diverge from one another to assist in urging the respective first and second rib members 1310a, 1310b against inner walls of respective nostrils when worn by the user.

[0178] As depicted in Figures 13A and 13B, the first and second rib members 1310a, 1310b, of the nasal dilator device 1300 may comprise respective first and second nostril engaging elements, 1312a and 1312b, disposed at distal ends 1314a, 1314b, of the first and second arcuate rib members 1310a, 1310b, respectively, for engaging with inner walls of respective nostrils when worn by a user. In some embodiments, the first and second nostril engaging elements, 1312a, 1312b may comprise relatively large surface areas 1316a, 1316b with respect to the first and second arcuate rib members 1310a, 1310b.

[0179] In some embodiments, the first and second nostril engaging elements, 1312a, 1312b, may extend at an angle to or arcuately from the respective first and second distal ends 1314a, 1314b, of the respective first and second rib members 1310a, 1310b. For example, the first and second nostril engaging elements, 1312a, 1312b, may extend

from the first and second planes, P2, P3, respectively and may extend in a direction away from the central portion 104 of the U-shaped body 1302.

[0180] In some embodiments, the first and second nostril engaging elements 1312a, 1312b may have pads 1318a, 1318b, disposed thereon, to engage with the inner walls of the nostrils of a user, in use. For example, the pads 1318a, 1318b may be disposed on the relatively large major surface areas 1316a, 1316b of the nostril engaging elements, 1312a and 1312b and may be enlarged with respect to the first and second arcuate rib members 1310a, 1310b, and/or the nostril engaging elements, 1312a, 1312b. The pads 1318a, 1318b of the nasal dilator device 1300 may be composed of a relatively soft overmould material, for example a polymer material such as thermoplastic elastomer (TPE) to provide a comfortable and/or grippable surface for engaging with the inner walls of the nostrils in use.

[0181] In some embodiments, as depicted in Figures 13A and 13B, the relatively large surface areas 1316a, 1316b of the nostril engaging members 1312a, 1312b are provided with a series of protrusions 1320. The nostril engaging members 1312a, 1312b may be substantially elongate, for example, extending from the distal ends 1314a, 1314b of the rib members 1310a, 1310b, and the protrusions 1320 may extend along a length of the surface areas 1316a, 1316b of the nostril engaging members 1312a, 1312b. In some embodiments, the surface areas 1316a, 1316b of the nostril engaging members 1312a, 1312b may be substantially oval, rectangular, triangular or truncated triangular in shape. In some embodiments, the series of protrusions 1320 may form a u-shaped pattern, a v-shaped pattern, or elongated v or ✓ (tick) shaped pattern, such as a chevron design, comprising a plurality of aligned u-shaped, v-shaped or elongated v or ✓ (tick) shaped protrusions 1320.

[0182] As depicted in Figures 13A and 13B, the rib members 1310a, 1310b of the nasal dilator device 1300 may comprise one or more projections 1304, flanges or ridges protruding from a major surface 1324 of the nasal dilator device 1300 and extending along at least a portion of a length of the rib members 1310a, 1310b. For example, the major surface 1324 of the nasal dilator device 1300 may be the surface of the nasal

dilator device 1300 configured, in use, to engage with or face nasal passage walls of a user's nose. In some embodiments, as depicted in Figure 13A and 13B, the nasal dilator device 1300 may comprise first and second projections 1322, each forming a flange disposed at respective elongate edges of the major surface 1324 of the nasal dilator device 1300.

[0183] The one or more projections 1322 may extend from the distal ends 1314a, 1314b of the rib members 1310a, 1310b toward the intermediate sections 1308a, 1308b. For example, as depicted in Figure 13A, the one or more projections 1322 may integrate with and extend from corresponding protrusions 1320 of the series of protrusions 1320 provided on the surface areas 1316a, 1316b of the nostril engaging members 1312a, 1312b. In some embodiments, the one or more projections 1322 may be disposed on and project from the major surface 1324 of the intermediate sections 1308a, 1308b and/or the leg members 1306a, 1306b and/or the central portion 1304 of the nasal dilator device 1300.

[0184] In some embodiments, the pads 1318a, 1318b may be configured to cooperate with and conform to the surface area 1316a, 1316b and the series of protrusions 1320 such that the overmoulded series of protrusions 1320 project or protrude from the overmoulded surface area 1316a, 1316b. In other embodiments, the surface areas 1316a, 1316b of the nostril engaging members 1314a, 1314b may be substantially smooth or flat and the pads 1318a, 1318b may comprise or form the series of protrusions 1320. For example, the pads 1318a, 1318b may be substantially elongate, for example, extending from the distal ends 1314a, 1314b of the rib members 1310a, 1310b, and the protrusions 1320 may extend along a length of the pads 1318a, 1318b.

[0185] In some embodiments, an overmould material may be provided on at least a portion of the rib members 1310a, 1310, on at least a portion of the intermediate sections 1308a, 1308b, and/or on at least a portion of the u-shaped body 1302. For example, the overmould may be configured to cooperate with and conform to the major surface 1322 of the nasal dilator device 1300 and the one or more projections 1322 such that the overmoulded projections 1322 project or protrude from the overmoulded

major surface 1324. In other embodiments, the major surface 13024 of the nasal dilator device 1300 may be substantially smooth or flat and an overmould disposed thereon may comprise or form the one or more projections or ridges 1322.

[0186] The series of protrusions 1320 and/or the one or more projections 1322 may provide a comfortable and/or grippable surface for engaging with the inner walls of the nose in use. For example, activities that are percussive, such as running, are often associated with the production of skin perspiration which may reduce the mechanical and frictional attachment of other nasal devices to the nasal passage of the user causing the devices to move from a position intended by the user and therefore requiring readjustment. However, the series of protrusions 1320 and/or the one or more projections 1322 may provide for or create a drag effect which may improve retention of the nasal dilator device 1300 in a desired position in a user's nose, in use, despite motion or skin perspiration of the user.

[0187] Referring now to Figures 14A, 14B, 14C, 14D, 14E and 14E, there is illustrated a nasal dilator device, generally indicated at 1400, according to some embodiments. The nasal dilator device 1400 may comprise similar components and elements to those of nasal dilator device 1300 depicted in Figures 13A and 13B and accordingly those similar components and elements are denoted like numerals.

[0188] In addition to those similar components and elements of nasal dilator device 1300, nasal dilator device 1400 may comprise a first and second releasable attachment mechanism 1402a and 1402b, respectively. The first and second releasable attachment mechanism 1402a, 1402b may comprise mating or interlocking components and may be employed to releasably attach the first and second rib members, 1310a and 1310b, respectively, to the U-shaped body 1302, to thereby define first and second adjustable looped structures, 1404a, and 1404b, respectively. The first and second releasable attachment mechanisms 1402a, 1402b may allow a user to selectively adjust a degree of dilation or expansion and contraction of the first and second rib members 1310a and 1310b with respect to the U-shaped body 1302.

[0189] The first and second releasable attachment mechanisms 1402a, 1402b may comprise respective arms 1406a, 1406b, such as pins, and respective sockets 1408a, 1408b for receiving and/or engaging the respective arms 1406a, 1406b.

[0190] As illustrated in Figure 14A to 14F, the arms 1406a, 1406b may be disposed on and extend from respective first and second intermediate sections 1308a, 1308b toward respective sockets 1408a, 1408b provided on opposite or inner surfaces 1410a, 1410b of the first and second nostril engaging elements 1312a, 1312b. The releasable attachment mechanisms 1402a, 1402b may therefore be arranged to releasably attach or lock the first and second nostril engaging elements 1312a, 1312b to the first and second intermediate sections 1308a, 1308b.

[0191] In other embodiments, the first and second arms 1406a, 1406b may be disposed on and extend from respective first and second rib members 1310a, 1310b toward respective first and second sockets 1408a, 1408b to allow the releasable attachment mechanisms 1402a, 1402b to releasably attach or lock the first and second nostril engaging elements 1312a, 1312b to the first and second rib members 1310a, 1310b or the first and second arms 1406a, 1406b may be disposed on and extend from respective first and second leg members 1306a, 1306b toward respective first and second sockets 1408a, 1408b to allow the releasable attachment mechanisms 1402a, 1402b to releasably attach or lock the first and second nostril engaging elements 1312a, 1312b to the first and second leg members 1306a, 1306b.

[0192] In other embodiments, as depicted in Figure 15, nasal dilator device 1500 having first and second releasable attachment mechanisms 1402a, 1402b may comprise first and second arms 1406a, 1406b extending from the respective opposite or inner surfaces 1410a, 1410b of the first and second nostril engaging elements 1312a, 1312b and first and second sockets 1408a, 1408b extending from the first and second intermediate sections 1308a, 1308b. In yet other embodiments, the first and second releasable attachment mechanisms 1402a, 1402b may comprise first and second arms 1406a, 1406b extending from the respective opposite or inner surfaces 1410a, 1410b of the first and second nostril engaging elements 1312a, 1312b and first and second

sockets 1408a, 1408b extending from the first and second rib members 1310a, 1310b or the first and second leg members 1306a, 1306b.

[0193] Referring again to Figures 14A to 14E, the first and second arms 1406a, 1406b may include at least one or a series of serrations, detents or protrusions 1412 arranged to engage with at least one or a series of grooves or ridges 1414 provided on or within the sockets 1408a, 1408b. For example, the grooves or ridges 1414 may extend downwardly from an upper jaw portion 1416 of the sockets 1408a, 1408b and/or may extend upwardly from a lower jaw portion 1418.

[0194] Application of sufficient force by a user to the first and second releasable attachment mechanisms 1402a, 1402b may be effective to move the arms 1406a, 1406b with respect to the sockets 1408a, 1408b and overcome a restrictive force between the detents 1412 and the grooves 1414 to allow the detents 1412 and/or the grooves 1414 to deform and the degree or level of dilation to be adjusted. The engagement of the detents 1412 with the grooves 1414 may provide a sufficient restrictive force to hold the arms 1406a, 1406b fixed when provided in a user's nose.

[0195] As depicted in Figures 14A to 14E, the arms 1406a, 1406b may each comprise a stopper or catch 1420 provided at their free ends to prevent or hinder the arms 1406a, 1406b from disengaging from or withdrawing from the respective sockets 1408a, 1408b. For example, in some embodiments, the catch 1420 comprises a hook portion 1422 which is configured to cooperate with a corresponding notch 1424 disposed in an upper or lower jaw portion 1416, 1418 of the socket 1408a, 1408b. Application of a relatively large pulling force may be sufficient to cause the catch 1420 to disengage with the notch 1424 and to cause the arms 1406a, 1406b to withdraw from the sockets 1408a, 1408b.

[0196] In some embodiments, as best shown in Figures 14A and 14B, the sockets 1408a, 1408b may take the form of substantially elongate housings 1426 provided with apertures 1428 or gaps in side walls of the housing 1426. The housings 1426 may be substantially wedge shaped and taper along their length towards opposite or inner

surfaces 1410a, 1410b, of the first and second nostril engaging elements 1312a, 1312b to provide a relatively broad opening to receive the arms 1406a, 1406b. The elongate or wedge shaped housing 1426 may provide for an improved engagement between the arms 1406a, 1406b and sockets 1408a, 1408b and more robust releasable attachment mechanisms 1402a, 1402b. For example, the elongate or wedge shaped housing 1426 may allow a face 1430 of the socket housing 1426 to engage with and lie substantially flush to the intermediate sections 1308a, 1308b, rib members 1310a, 1310b, or leg members 1306a, 1306b, when the arms 1306a, 1306b are fully engaged within the sockets 1408a, 1408b, when the nasal dilator device 1400 assumes a closed configuration or state.

[0197] The arms 1406a, 1406b may be fully or substantially fully inserted into the respective sockets 1408a, 1408b to enable the nasal dilator device 1400 to adopt or assume a fully closed or substantially fully closed state, to thereby tighten or contract the looped structures 1411a, 1411b. The arms 1406a, 1406b may be partially inserted into the sockets 1408a, 1408b to enable the nasal dilator device 1400 to adopt or assume a partially closed state, to provide for looser or less tight looped structures 1404a, 1404b and accommodate variations in nasal passage sizes.

[0198] The particular configuration of the releasable attachment mechanisms 1402a, 1402b allows for self-adjustment of the looped structures, 1404a, and 1404b of the nasal dilator device 1400 whilst the device is in place of the nose of the user. This may be of particular benefit to a user partaking in percussive activities. For example, percussive activities, such as running or cycling, may cause a reduction in mechanical and frictional attachment of the nasal dilator device to a nasal passage of a user, for example, due to motion and/or skin perspiration, causing the device to move from a position intended by the user and therefore requiring readjustment.

[0199] As depicted in Figures 14A to 14F, the nasal dilator device 1400 may include a series of protrusions 1320 disposed on the nostril engaging members 1312a, 1312b, and/or one or more projections 1322 protruding from the major surface 1324 of the nasal dilator device 1400, as discussed with reference to Figure 13.

[0200] Similar to the embodiment of nasal dilator device 600 depicted in Figure 6, in some embodiments, the nasal dilator device 1400 may comprise at least one capsule (not shown), which may include an agent such as a medicament and/or a fragrance or aromatic agent, disposed within respective sockets 1408a, 1408b. The arms 1406a, 1406b may be configured to activate, pierce or burst the capsule (not shown) to release the agent, medicament and/or fragrance or aromatic agent when they are inserted into the sockets 1408a, 1408b. In this way, the medicament and/or fragrance or aromatic agent is released only when the capsule (not shown) is activated, pierced or burst, thereby increasing a longevity or “shelf-life” and/or protecting the integrity of the medicament and/or aromatic agent.

[0201] Similar to the embodiment of nasal dilator device 700 depicted in Figure 7, in some embodiments, the nasal dilator device 1400 may comprise at least one coating or film (not shown) arranged to release a fragrance, aroma or medicament. For example, the film (not shown) may be disposed on a surface of at least one of the attachment mechanisms 1402a, 1402b, such as on an inner surface of the looped structures 1404a, 1404b, on the central portion 1302, on the first and second leg members 1306a, 1306b, on the first and second intermediate sections 1308a, 1308b, on the rib members 1310a, 1310b, and/or on the first and second nostril engaging elements 1312a, 1312b.

[0202] In some embodiments, the film (not shown) is arranged to release a fragrance, aroma or medicament in response to abrasion, such as scratching, scraping and may be provided with an outer cover, seal or strip to protect the film (not shown) from unintended abrasion. In other embodiments, the coating or film (not shown) may be arranged to release a fragrance, aroma or medicament in response to the removal or peeling off of an outer cover, strip or seal. In some embodiments, a fragrance, aroma or medicament may be provided or retained between two strips or films (not shown) forming a blister. For example, the coating or film (not shown) may comprise a polymer or a fibre and/or may be in the form of a “scratch and sniff” technology or peel off technology.

[0203] It will be appreciated by persons skilled in the art that numerous variations and/or modifications may be made to the above-described embodiments, without departing from the broad general scope of the present disclosure. The present embodiments are, therefore, to be considered in all respects as illustrative and not restrictive.

Claims

1. A nasal dilator device comprising:
a substantially U-shaped body including:
a central portion arranged to span a septum of a nose when worn by a user; and
first and second leg members extending from the central portion in a first plane;
a first cantilever rib member extending outward from the U-shaped body in a second plane;
a second cantilever rib member extending outward from the U-shaped body in a third plane;
wherein the first and second cantilever rib members extend away from each other;
a first intermediate section connecting an end of the first leg member to a proximal end of the first cantilever rib member, wherein the first intermediate section extends from the first plane to the second plane; and
a second intermediate section connecting an end of the second leg member to a proximal end of the second cantilever rib member, wherein the second intermediate section extends from the first plane to the third plane; and
at least one projection protruding from and extending along at least a portion of a length of each of the first and second cantilever rib members.
2. The nasal dilator device of claim 1, wherein the at least one projection comprises first and second projections, each forming a flange disposed at respective elongate edges of a major surface of the nasal dilator device.
3. The nasal dilator device of claim 1 or claim 2, wherein the at least one projection is arranged, in use, to engage with an inner surface of a nose.
4. The nasal dilator device of any preceding claim, wherein the second and third planes are orthogonal to the first plane.

5. The nasal dilator device of any preceding claim, wherein the first intermediate section and the second intermediate section are right angled sections.

6. The nasal dilator device of any preceding claim, wherein the first and second intermediate sections are arranged, in use, to engage with the septum and extend from the septum behind the columella and alar fibrofatty tissue of the nose, allowing the first and second cantilever rib members, in use, to extend along respective nasal orifices to an inner wall of the nostrils.

7. The nasal dilator device of any preceding claim, wherein the first and second cantilever rib members are arcuate cantilever rib members, each having a curvature along its length.

8. The nasal dilator device of any preceding claim, wherein the first and second intermediate sections are arcuate intermediate sections, each having a curvature along its length.

9. The nasal dilator device of any preceding claim, wherein the second and third planes are converging planes such that the first and second cantilever rib members are angled and/or extend substantially toward the central portion of the U-shaped body.

10. The nasal dilator device of any one of claims 1 to 9, wherein the second and third planes are diverging planes such that the first and second cantilever rib members are angled and/or extend substantially toward the central portion of the U-shaped body.

11. The nasal dilator device of any one of claims 1 to 9, wherein the second and third planes are the same plane.

12. The nasal dilator device of any preceding claim, wherein the first and second cantilever rib members exhibit an elongate arched profile which approximates at least a portion of one of a circle, ellipse or parabola.

13. The nasal dilator device of any preceding claim, wherein the first and second leg members are inclined towards each other such that a relatively greater distance is provided between the first and second leg members towards the central portion accommodate a columella of a nose when donned by the user.

14. The nasal dilator device of any preceding claim, wherein the first and second intermediate sections are inclined away from each other to assist in urging the respective first and second cantilever rib members against inner walls of respective nostrils when worn by the user.

15. The nasal dilator device of any preceding claim, wherein the first and second cantilever rib members comprise respective first and second nostril engaging elements for engaging with an inner wall of a respective nostril.

16. The nasal dilator device of claim 15, wherein the first and second nostril engaging elements extend arcuately from the respective first and second distal ends of the respective first and second rib members.

17. The nasal dilator device of claim 16, wherein the first and second nostril engaging elements extend from the first and second planes respectively and in a direction away from the central portion of the U-shaped body.

18. The nasal dilator device of any preceding claim, wherein the first and second nostril engaging elements are disposed at distal ends of the first and second cantilever rib members, respectively.

19. The nasal dilator device of any one of claims 15 to 18, further comprising a series of protrusions disposed on the first and second nostril engaging elements.

20. The nasal dilator device of claim 19, wherein the first and second nostril engaging elements are substantially elongate and the series of protrusions extends along a length of the first and second nostril engaging elements.

21. The nasal dilator device of claim 19 or 20, wherein the series of protrusions comprises a plurality of elongated v-shaped protrusions.

22. The nasal dilator device of any one of claims 15 to 21, wherein enlarged pads are disposed on the first and second nostril engaging elements to engage with inner walls of the nostrils.

23. The nasal dilator device of claim 22, wherein the enlarged pads comprise the series of protrusions.

24. The nasal dilator device of any one of claims 19 to 23, wherein the series of protrusions are formed from an overmould material.

25. The nasal dilator device of any one of claims 19 to 24, wherein the at least one projection is integrated with and extends from a corresponding protrusion of the series of protrusions.

26. The nasal dilator device of any preceding claim, wherein the at least one projection is formed from an overmould material.

27. The nasal dilator device of any preceding claim, further comprising a first and second releasable attachment mechanism for releasably attaching the first and second cantilever rib members, respectively, to the U-shaped body.

28. The nasal dilator device of claim 27, wherein the releasable attachment mechanisms are arranged to releasably attach first and second nostril engaging

elements disposed at distal ends of the first and second cantilever rib members to the first and second leg members, respectively.

29. The nasal dilator device of claim 27, wherein the releasable attachment mechanisms are arranged to releasably attach the first and second nostril engaging elements disposed at distal ends of the first and second cantilever rib members to the first and second intermediate sections, respectively.

30. The nasal dilator device of any one of claims 27 to 29, wherein the releasable attachment mechanisms each comprise an arm and a socket arranged to receive and engage the arm.

31. The nasal dilator device of claim 30, wherein a stopper or hook is disposed at an end of the arm to hinder the arm from withdrawing from the socket.

32. The nasal dilator device of claim 31, wherein a notch is disposed on an inner surface of the socket and is configured to engage with the stopper or hook on the arm to hinder the arm from withdrawing from the socket.

33. The nasal dilator device of any one of claims 30 to 32, wherein the arms are disposed on an inner surface of first and second nostril engaging elements of the first and second cantilever rib members, respectively, and the sockets are disposed on the first and second leg members.

34. The nasal dilator device of any one of claims 30 to 32, wherein the arms are disposed on an inner surface of first and second nostril engaging elements of the first and second cantilever rib members, respectively, and the sockets are disposed on the first and second intermediate sections.

35. The nasal dilator device of any one of claims 30 to 32, wherein the sockets are disposed on an inner surface of first and second nostril engaging elements of the first

and second cantilever rib members, respectively, and the arms are disposed on the first and second leg members.

36. The nasal dilator device of any one of claims 30 to 32, wherein the sockets are disposed on an inner surface of first and second nostril engaging elements of the first and second cantilever rib members, respectively, and the arms are disposed on the first and second intermediate sections.

37. The nasal dilator device of any one of claims 30 to 32, further comprising first and second arm support members projecting from respective first and second intermediate sections, and wherein the sockets are disposed on an inner surface of first and second nostril engaging elements of the first and second cantilever rib members, respectively, and the arms are disposed on the first and second arm support members.

38. The nasal dilator device of any one of claims 30 to 37, wherein the sockets comprise substantially elongate housings which taper along their length and provide a relatively broad opening for receiving the arms.

39. The nasal dilator device of claim 38, wherein each of the sockets is provided with apertures in side walls of the housing.

40. The nasal dilator device of any one of claims 30 to 39, wherein a capsule is provided within the socket and is arranged to be activated by the arm when the arm is received by the socket.

41. The nasal dilator device of claim 40, wherein the capsule includes at least one of a medicament or compound.

42. The nasal dilator device of any one of claims 30 to 41, wherein the arm comprises a coating disposed thereon arranged to release a scent in response to abrasion of the coating.

43. The nasal dilator device of any one of claims 15 to 42, wherein an aperture is disposed in each of the first and second nostril engaging elements.
44. The nasal dilator device of claims 43, wherein the aperture is arranged to receive at least one of a compound, a medicament, and a capsule comprising a medicament or compound emanating a scent.
45. The nasal dilator device of any preceding claim further comprising a film disposed on a surface of the nasal dilator and a removable seal provided on the film to mitigate release of a compound from the film.
46. The nasal dilator device of any preceding claim further comprising an overmould disposed on at least one of the central portion, the leg members, the intermediate sections and the arcuate cantilever rib members.
47. The nasal dilator device of claim 46, wherein in the overmould is infused with a compound.
48. The nasal dilator device of claim 46 or 47, wherein in the overmould is infused with a medicament, a fragrance or an aroma.
49. The nasal dilator device of any preceding claim, wherein in the nasal dilator device is composed of a substrate material infused with a medicament, a fragrance or an aromatic agent.
50. The nasal dilator device of any preceding claim, wherein the central portion comprises a tab extending in a direction substantially opposite to the first and second leg members to assist with insertion, removal and/or placement of the nasal dilator device.

51. The nasal dilator device of claim 51, wherein the tab is removeable from the nasal dilator device.

52. A nasal dilator device comprising:
a substantially U-shaped body including:
a central portion arranged to span a septum of a nose when worn by a user; and
first and second leg members extending from the central portion in a first plane;
a first cantilever rib member extending outward from the U-shaped body in a second plane;
a second cantilever rib member extending outward from the U-shaped body in a third plane;
wherein the first and second cantilever rib members extend away from each other;
a first intermediate section connecting an end of the first leg member to a proximal end of the first cantilever rib member, wherein the first intermediate section extends from the first plane to the second plane;
a second intermediate section connecting an end of the second leg member to a proximal end of the second cantilever rib member, wherein the second intermediate section extends from the first plane to the third plane;
a first releasable attachment mechanism for releasably attaching a distal end of the first cantilever rib member to the first intermediate section; and
a second releasable attachment mechanism for releasably attaching a distal end of the second cantilever rib member to the second intermediate section.

53. The nasal dilator device of claim 52, wherein the second and third planes are orthogonal to the first plane.

54. The nasal dilator device of claim 52 or 53, wherein the first intermediate section and the second intermediate section are right angled sections.

55. The nasal dilator device of any one of claims 52 to 54, wherein the first and second cantilever rib members comprise respective first and second nostril engaging elements for engaging with an inner wall of a respective nostril.

56. The nasal dilator device of claim 55, wherein the releasable attachment mechanisms are arranged to releasably attach first and second nostril engaging elements disposed at the distal ends of the first and second cantilever rib members to the first and second leg members, respectively.

57. The nasal dilator device of any one of claims 55 or 56, wherein the releasable attachment mechanisms each comprise an arm and a socket arranged to receive and engage the arm.

58. The nasal dilator device of claim 57, wherein a stopper or hook is disposed at an end of the arm to hinder the arm from withdrawing from the socket.

59. The nasal dilator device of claim 58, wherein a notch is disposed on an inner surface of the socket and is configured to engage with the stopper or hook on the arm to hinder the arm from withdrawing from the socket.

60. The nasal dilator device of any one of claims 57 to 59, wherein the arms are disposed on an inner surface of first and second nostril engaging elements of the first and second cantilever rib members, respectively, and the sockets are disposed on the first and second intermediate sections.

61. The nasal dilator device of any one of claims 57 to 59, wherein the sockets are disposed on an inner surface of first and second nostril engaging elements of the first and second cantilever rib members, respectively, and the arms are disposed on the first and second intermediate sections.

62. The nasal dilator device of any one of claim 61, further comprising first and second arm support members projecting from respective first and second intermediate sections, and wherein the arms are disposed on the first and second arm support members.

63. The nasal dilator device of any one of claims 57 to 62, wherein the sockets comprise substantially elongate housings which taper along their length and provide a relatively broad opening for receiving the arms.

64. The nasal dilator device of claim 63, wherein each of the sockets is provided with apertures in side walls of the housing.

65. A nasal dilator device comprising:
a substantially U-shaped body including:
a central portion arranged to span a septum of a nose when worn by a user; and
first and second leg members extending from the central portion in a first plane;
a first cantilever rib member extending outward from the U-shaped body in a second plane;
a second cantilever rib member extending outward from the U-shaped body in a third plane;
wherein the first and second cantilever rib members extend away from each other;
a first intermediate section connecting an end of the first leg member to a proximal end of the first cantilever rib member; and
a second intermediate section connecting an end of the second leg member to a proximal end of the second cantilever rib member.

66. The nasal dilator device of claim 65, wherein the first intermediate section extends from the first plane to the second plane and wherein the second intermediate section extends from the first plane to the third plane.
67. The nasal dilator device of claim 65 or claim 66, wherein the second and third planes are orthogonal to the first plane.
68. The nasal dilator device of any one of claims 65 to 67, wherein the first intermediate section and the second intermediate section are right angled sections.
69. The nasal dilator device of any one of claims 65 to 68, wherein the first and second intermediate sections are arranged, in use, to engage with the septum and extend from the septum behind the columella and alar fibrofatty tissue of the nose allowing the first and second cantilever rib members, in use, to extend along respective nasal orifices to an inner wall of the nostrils.
70. The nasal dilator device of claim 65, wherein the first intermediate section extends between the first plane and the second plane and the second intermediate section extends between the first plane and the third plane.
71. The nasal dilator device of claim 70, wherein the first and second intermediate sections are arranged, in use, to extend along a length of the septum and the first and second cantilever rib members are each arranged, in use, to extend from a floor of a respective nasal orifice to an inner wall of the nostrils.
72. A nasal dilator device comprising:
a substantially U-shaped body including:
a central portion arranged to span a septum of a nose when worn by a user; and
first and second leg members extending from the central portion in a first plane;

a first cantilever rib member extending outward from an end of the first leg member of the U-shaped body in a second plane;

a second cantilever rib member extending outward from an end of the second leg member of the U-shaped body in a third plane;

wherein the first and second cantilever rib members extend away from each other; and

at least one projection protruding from and extending along at least a portion of a length of each of the first and second cantilever rib members.

73. The nasal dilator device of claim 72, wherein the second and third planes are orthogonal to the first plane.

74. A nasal dilator device comprising:

a substantially U-shaped body including:

a central portion arranged to span a septum of a nose when worn by a user; and

first and second leg members extending from the central portion;

first and second cantilever rib members extending outward from the U-shaped body and away from one another;

a first intermediate section connecting an end of the first leg member to a proximal end of the first cantilever rib member; and

a second intermediate section connecting an end of the second leg member to a proximal end of the second cantilever rib member;

wherein the first and second leg members are arranged, in use, to extend inward of respective nasal orifices along the septum, the first and second intermediate sections are arranged, in use, to engage with the septum and extend from the septum behind the columella and alar fibrofatty tissue of the nose allowing the first and second cantilever rib members, in use, to extend along the respective nasal orifices to an inner wall of the nostrils.

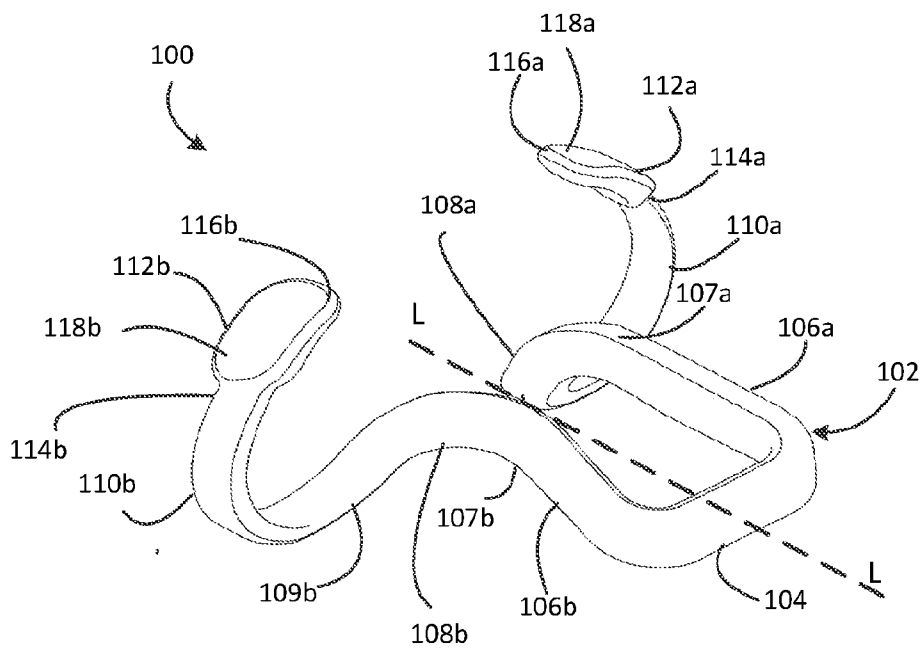


Figure 1A

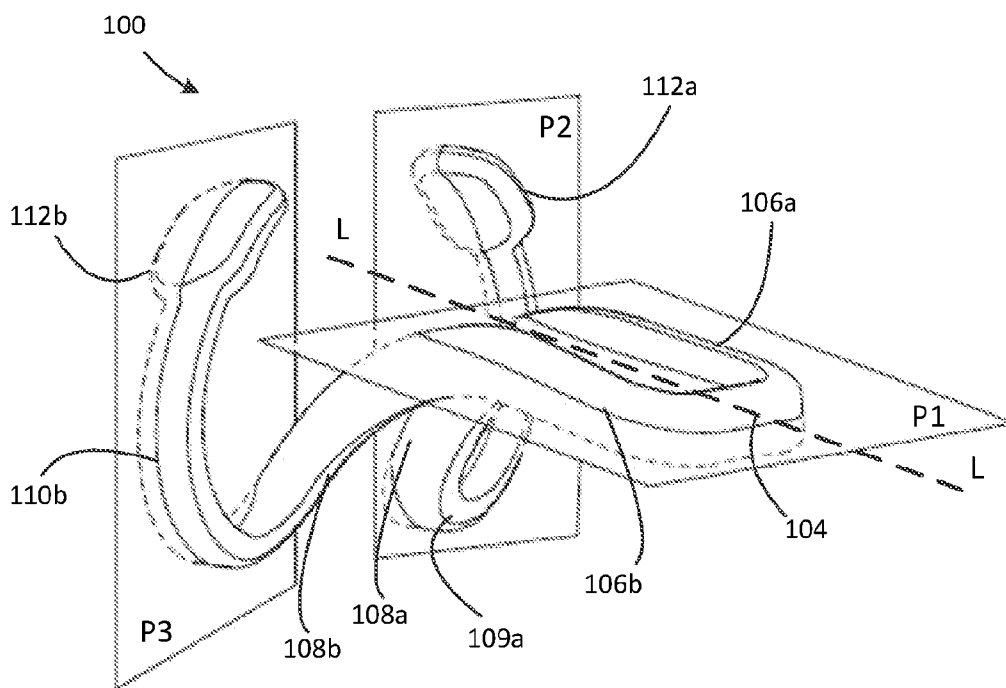


Figure 1B

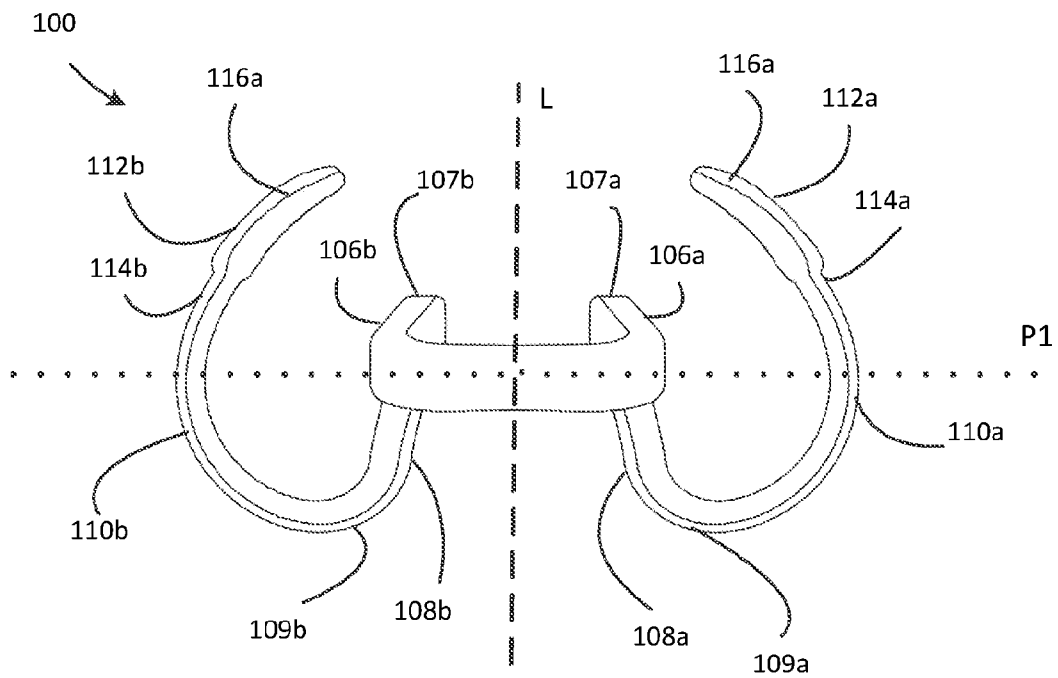


Figure 1C

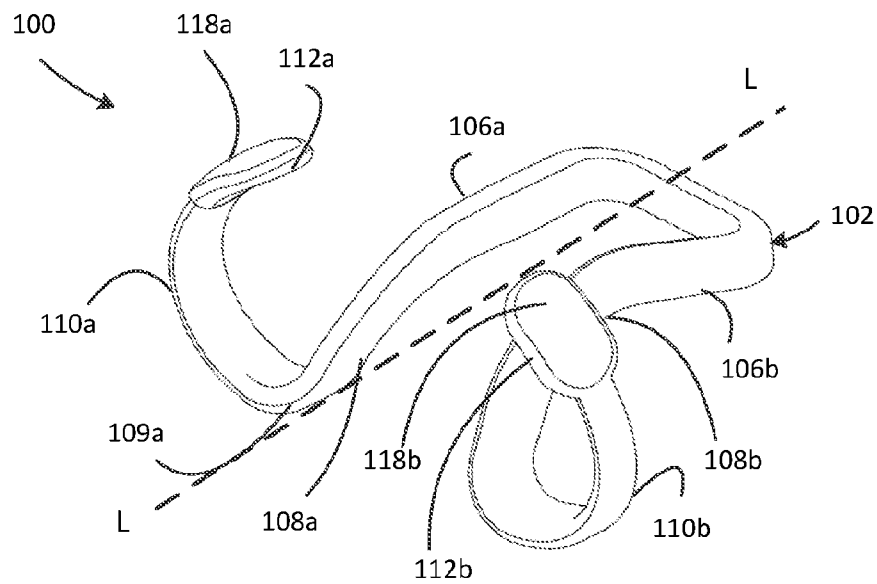


Figure 1D

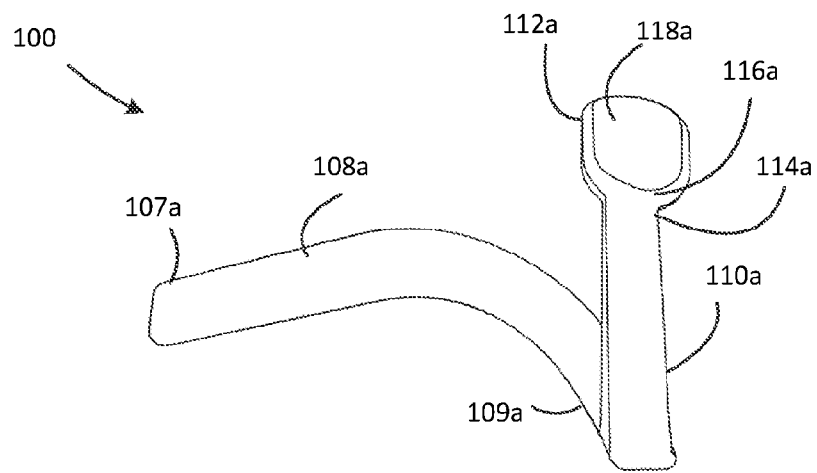


Figure 1E

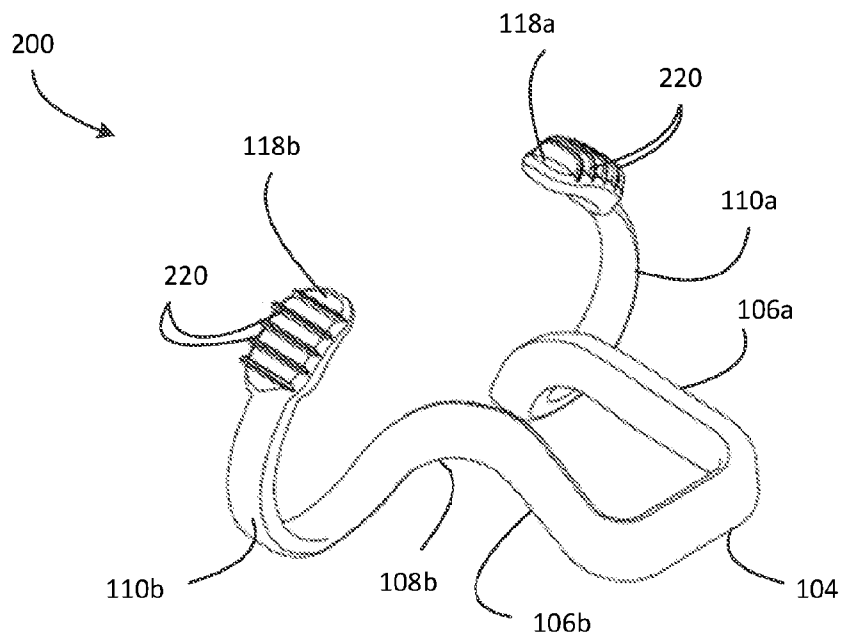


Figure 2

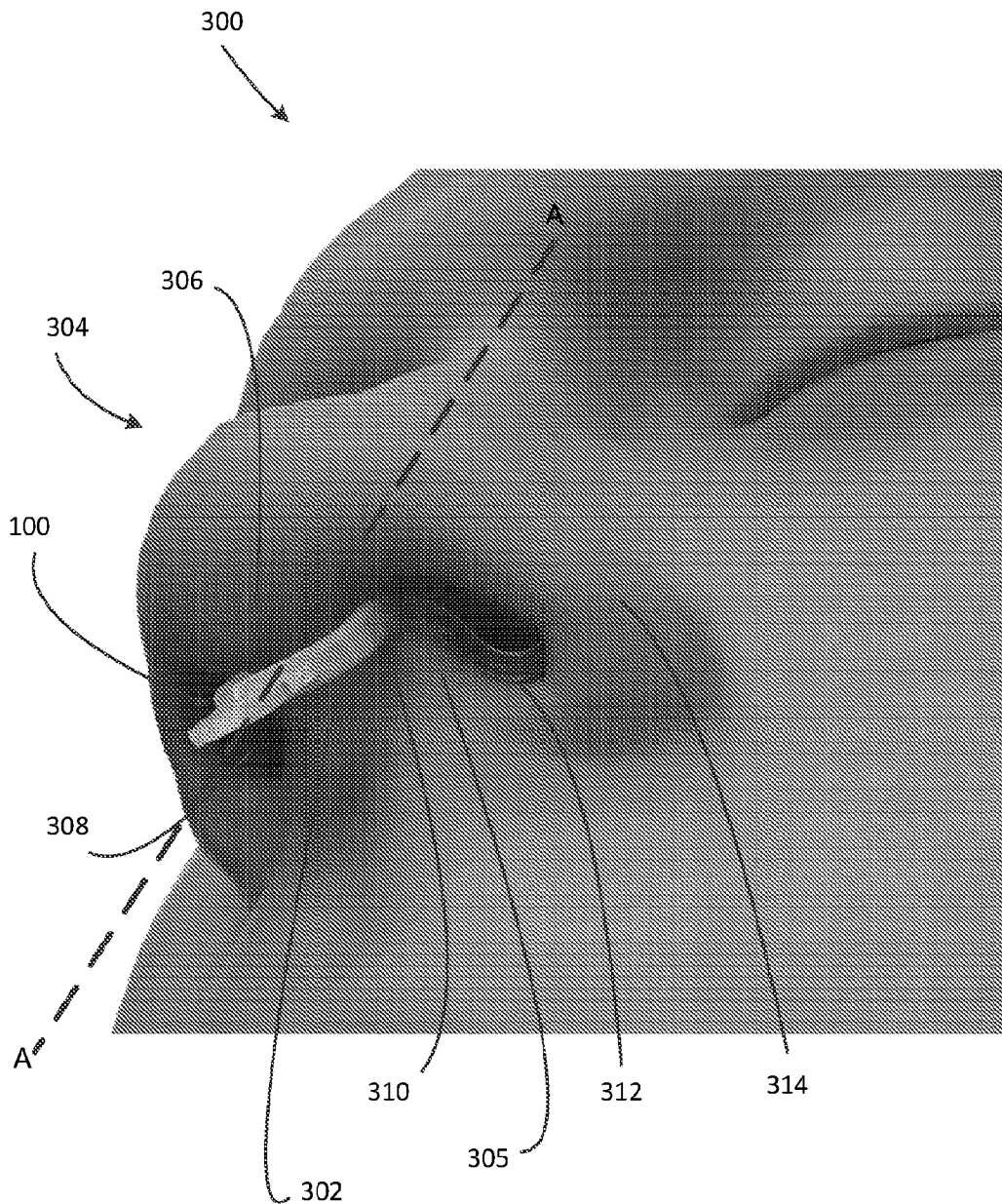


Figure 3A

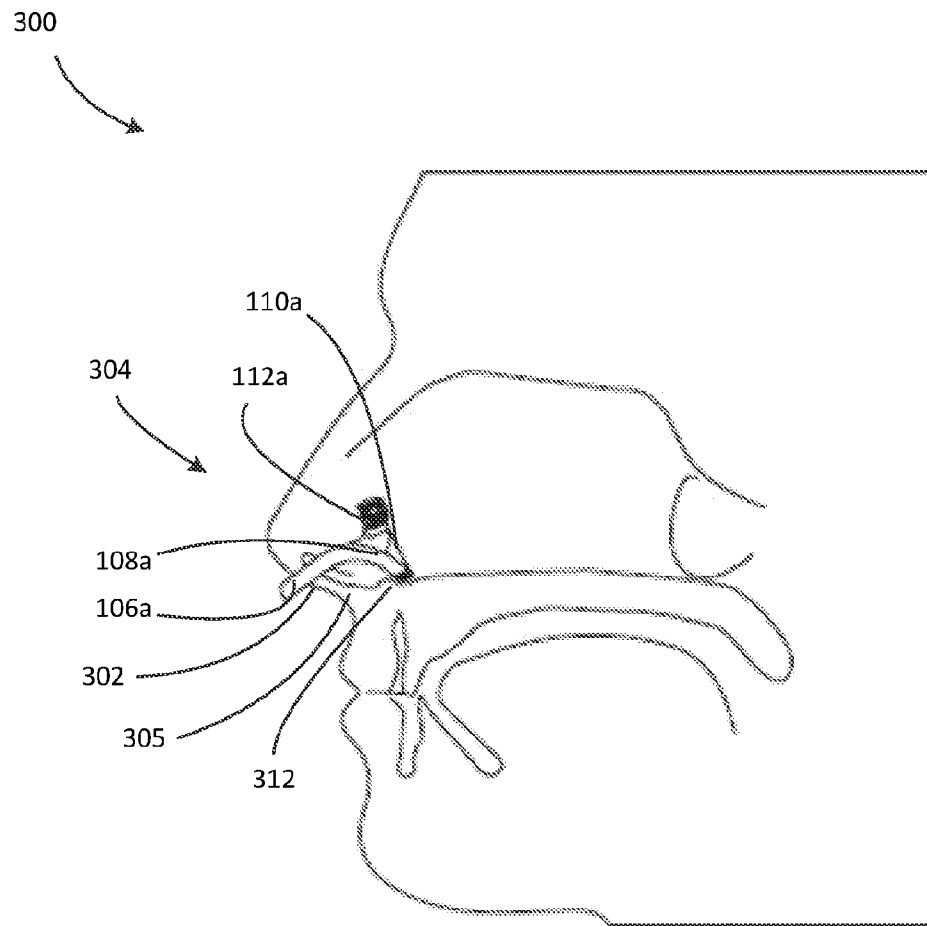


Figure 3B

7/22

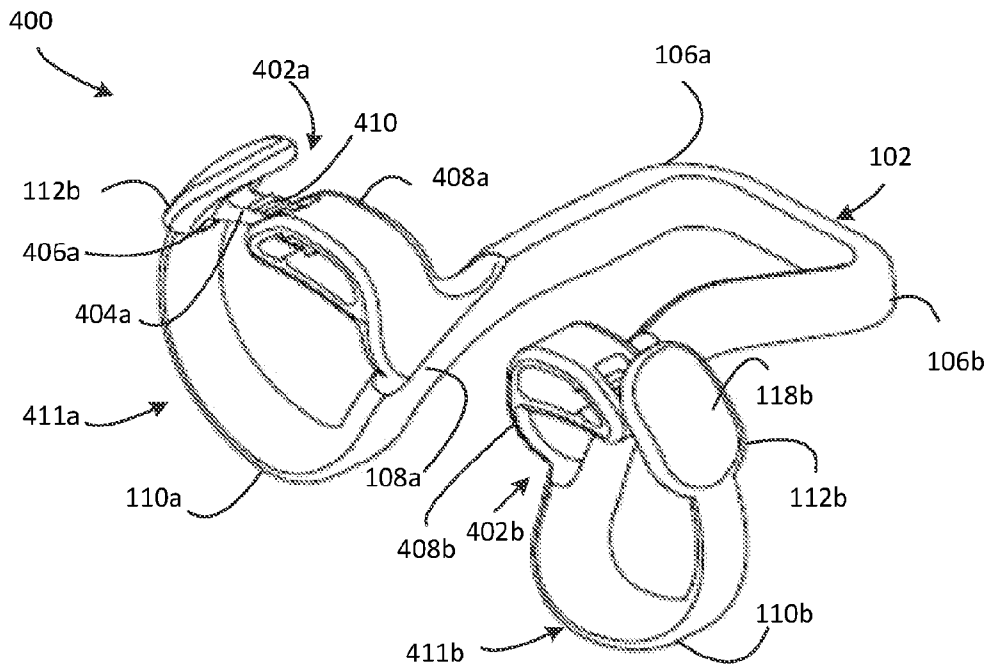


Figure 4A

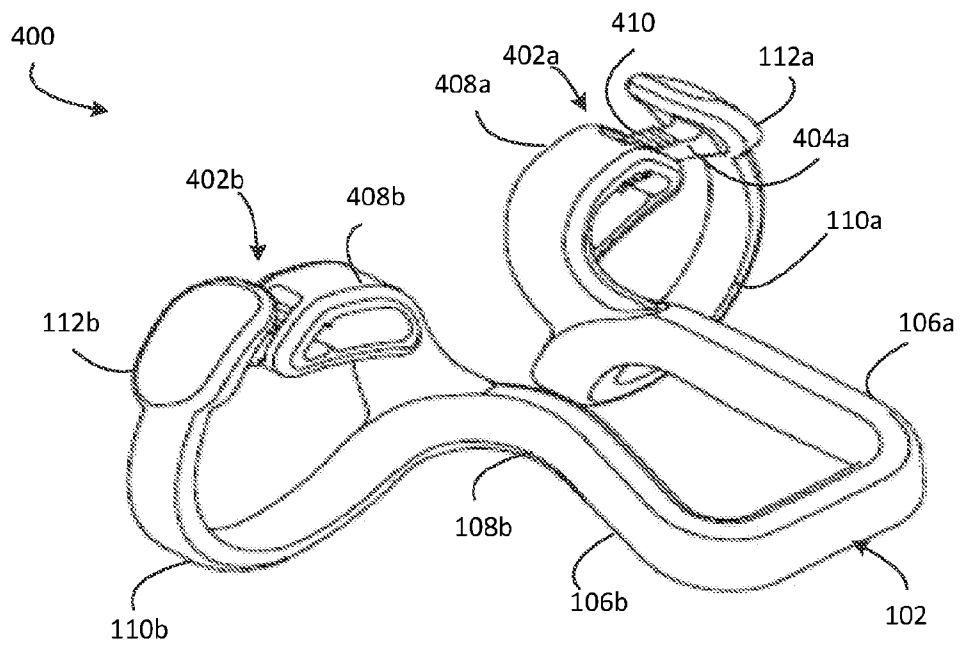


Figure 4B

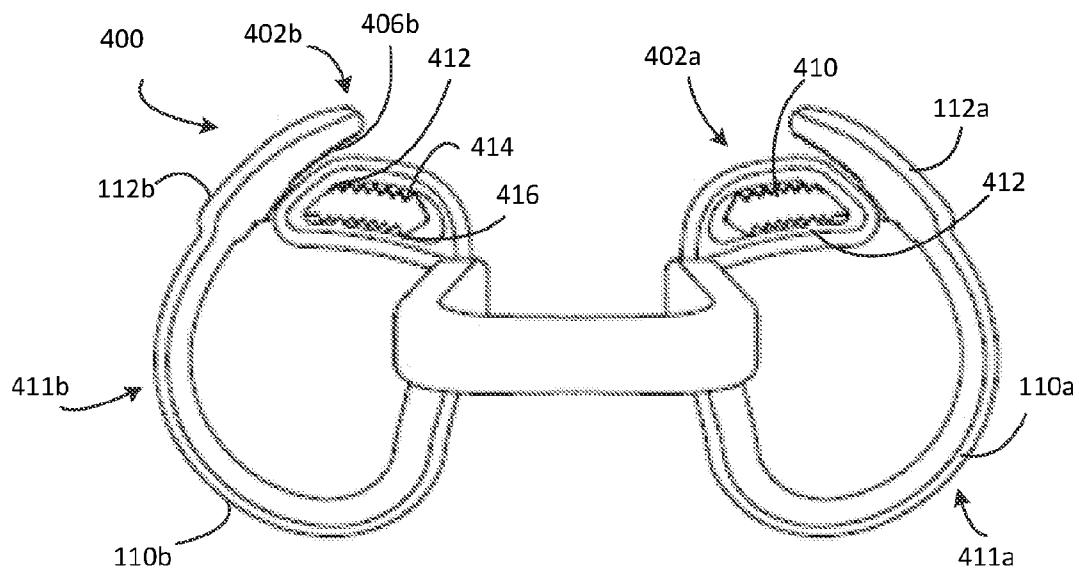


Figure 4C

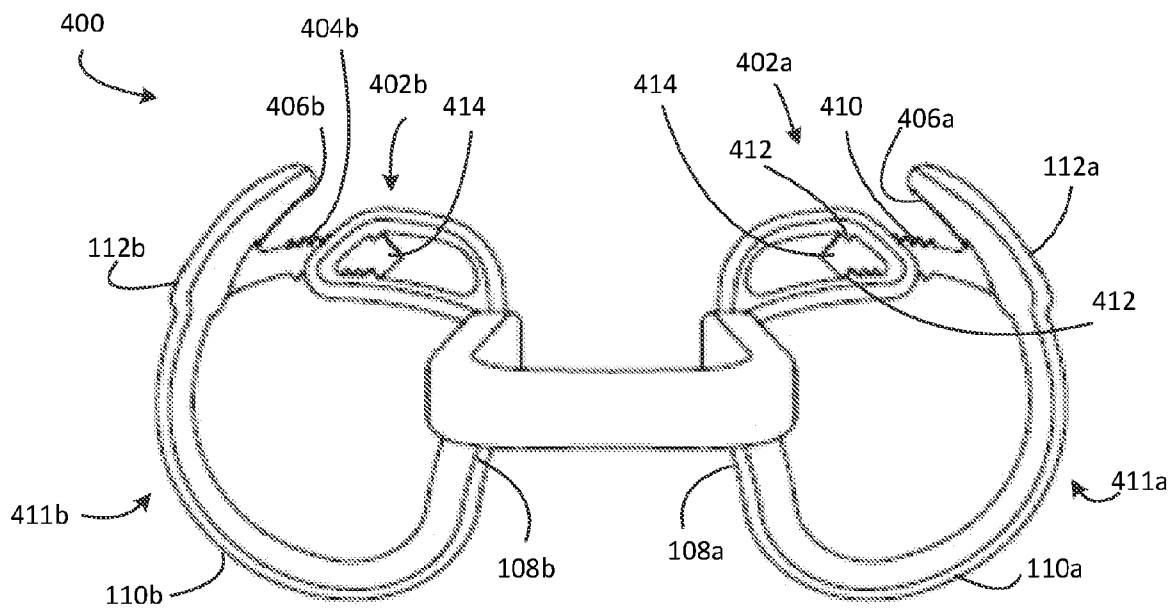


Figure 4D

9/22

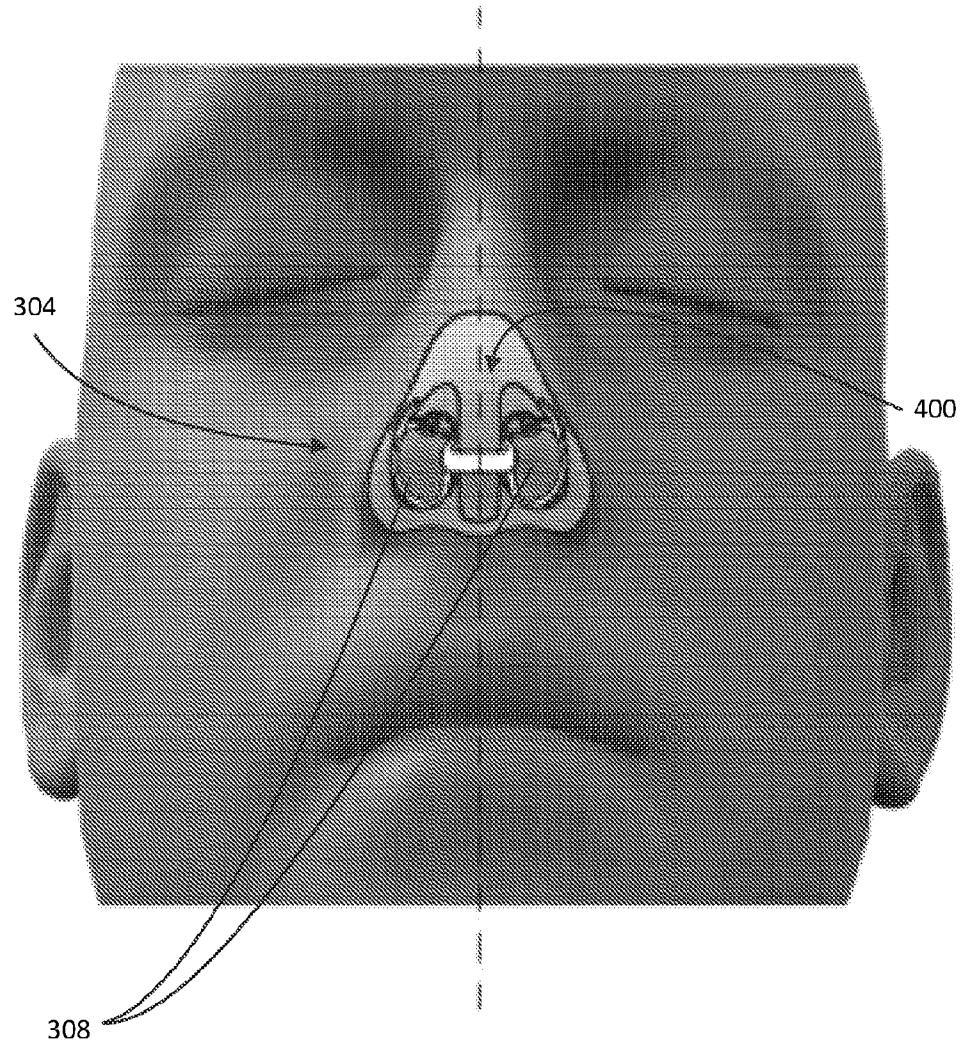


Figure 5

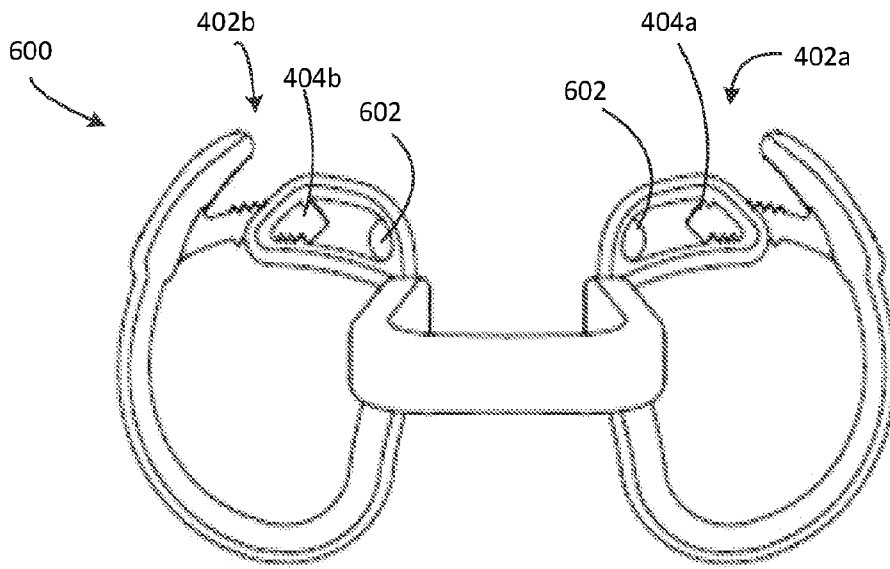


Figure 6A

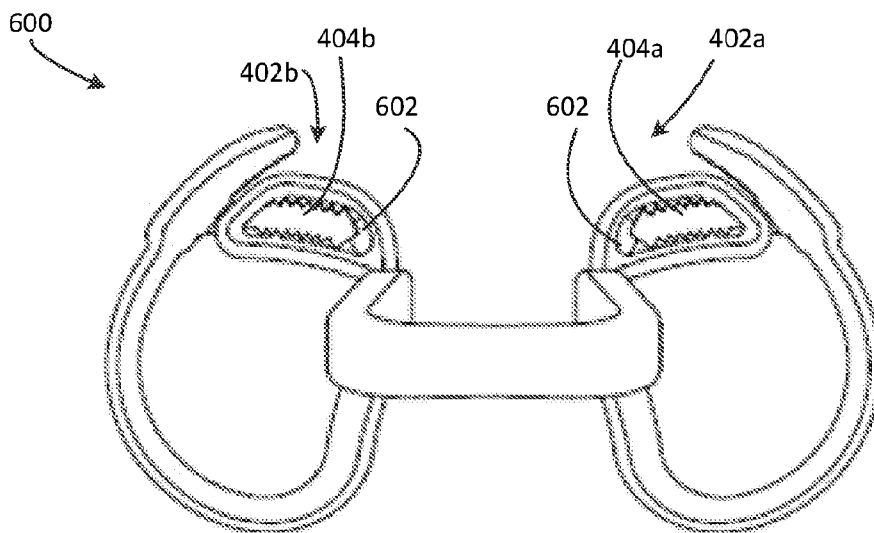


Figure 6B

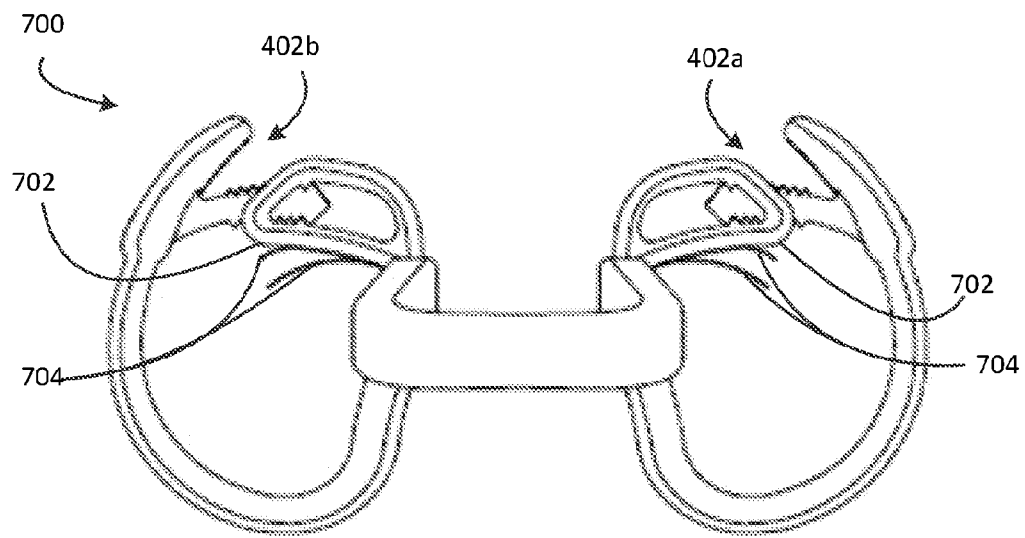


Figure 7A

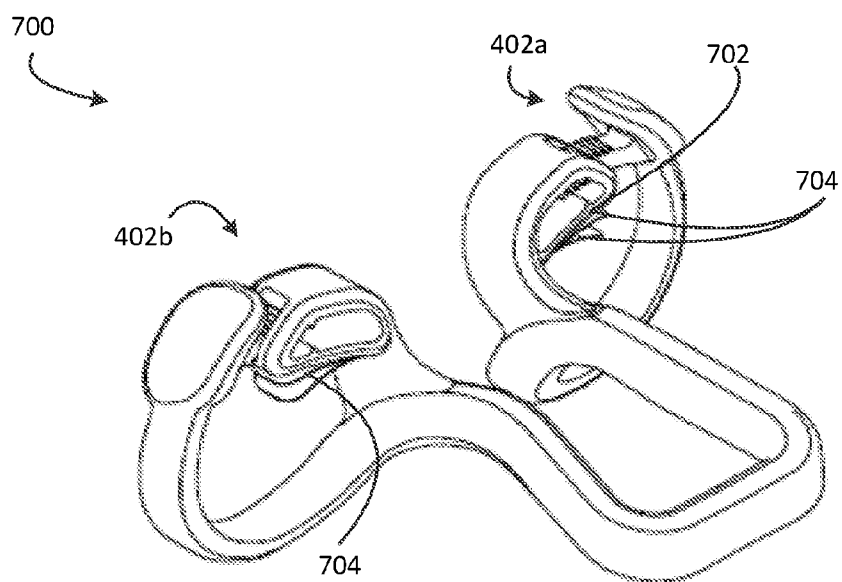


Figure 7B

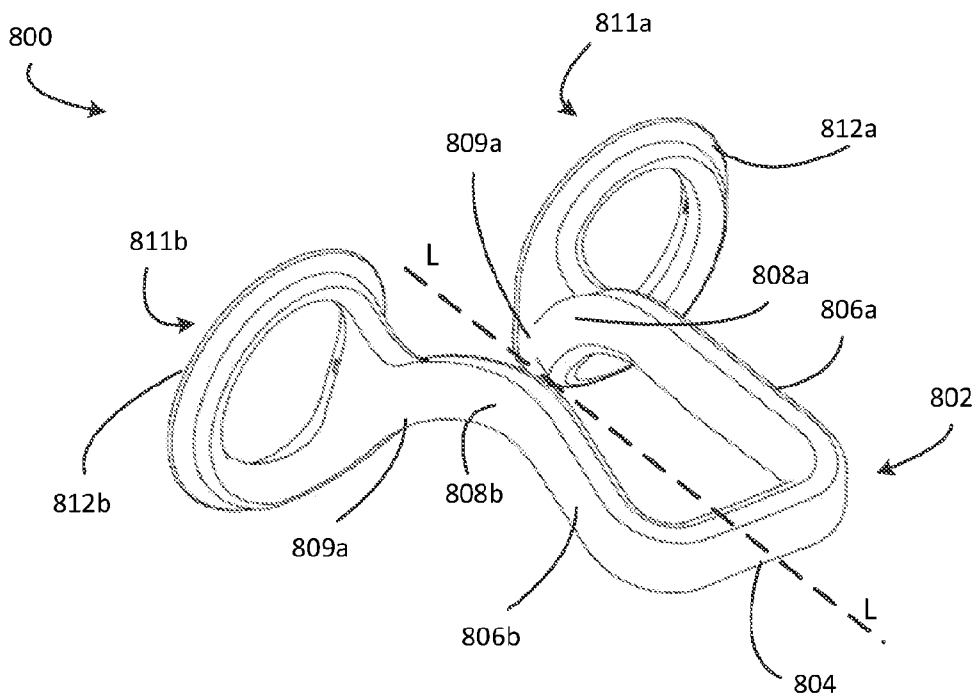


Figure 8A

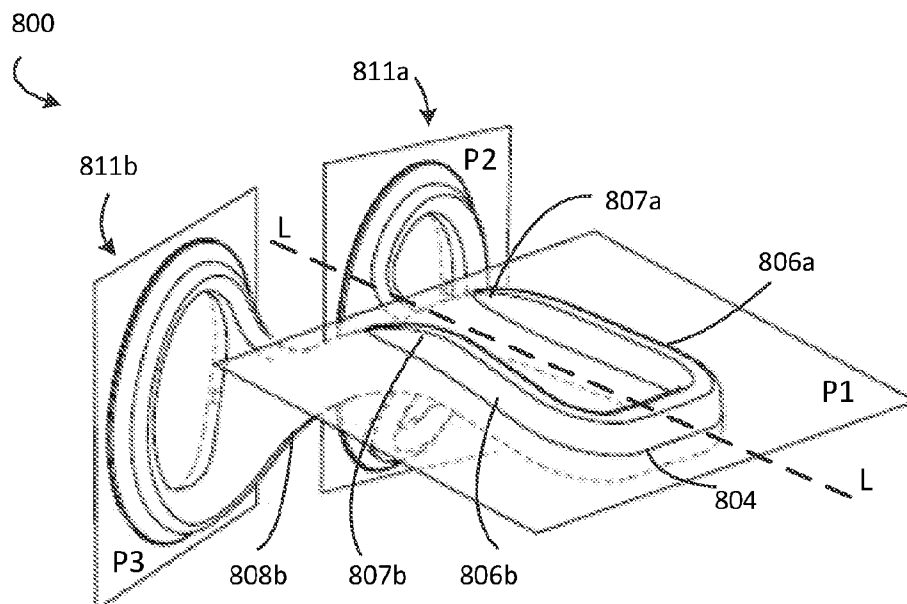


Figure 8B

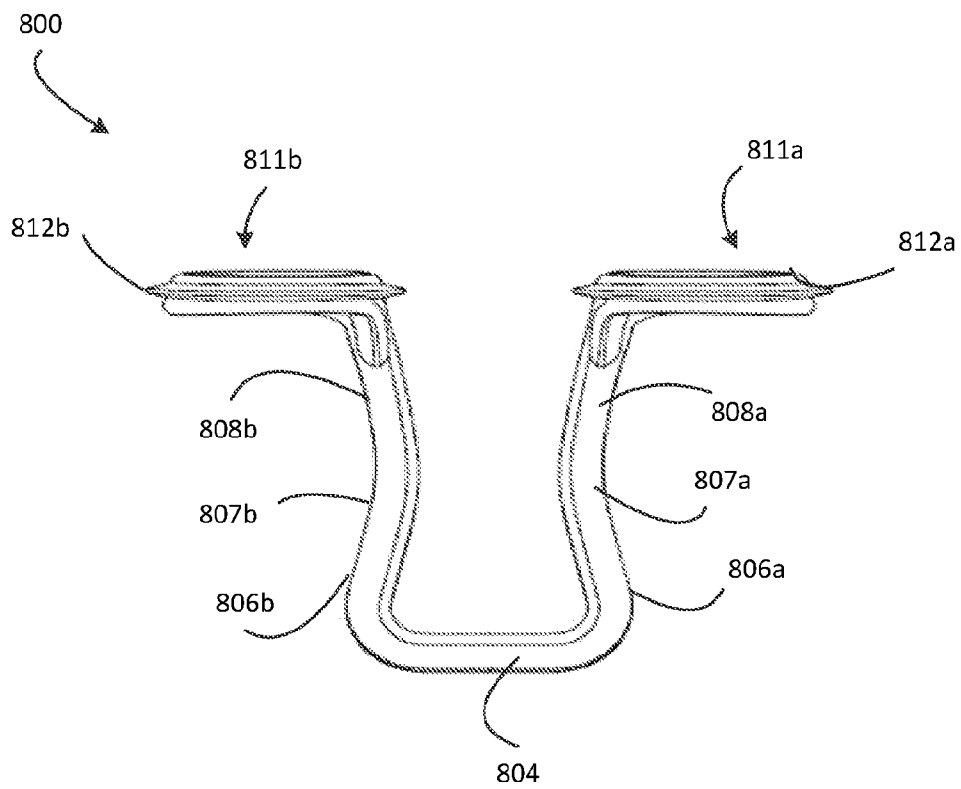


Figure 8C

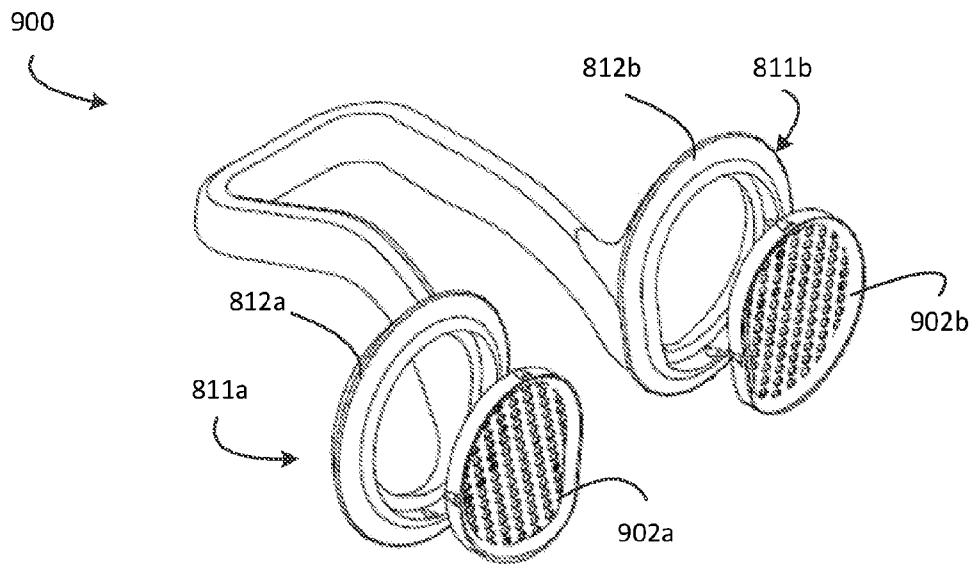


Figure 9

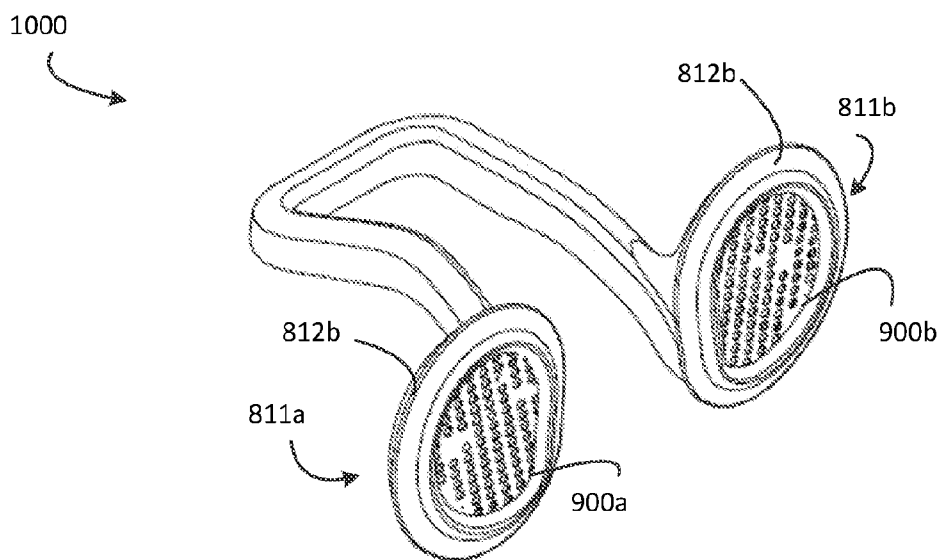


Figure 10

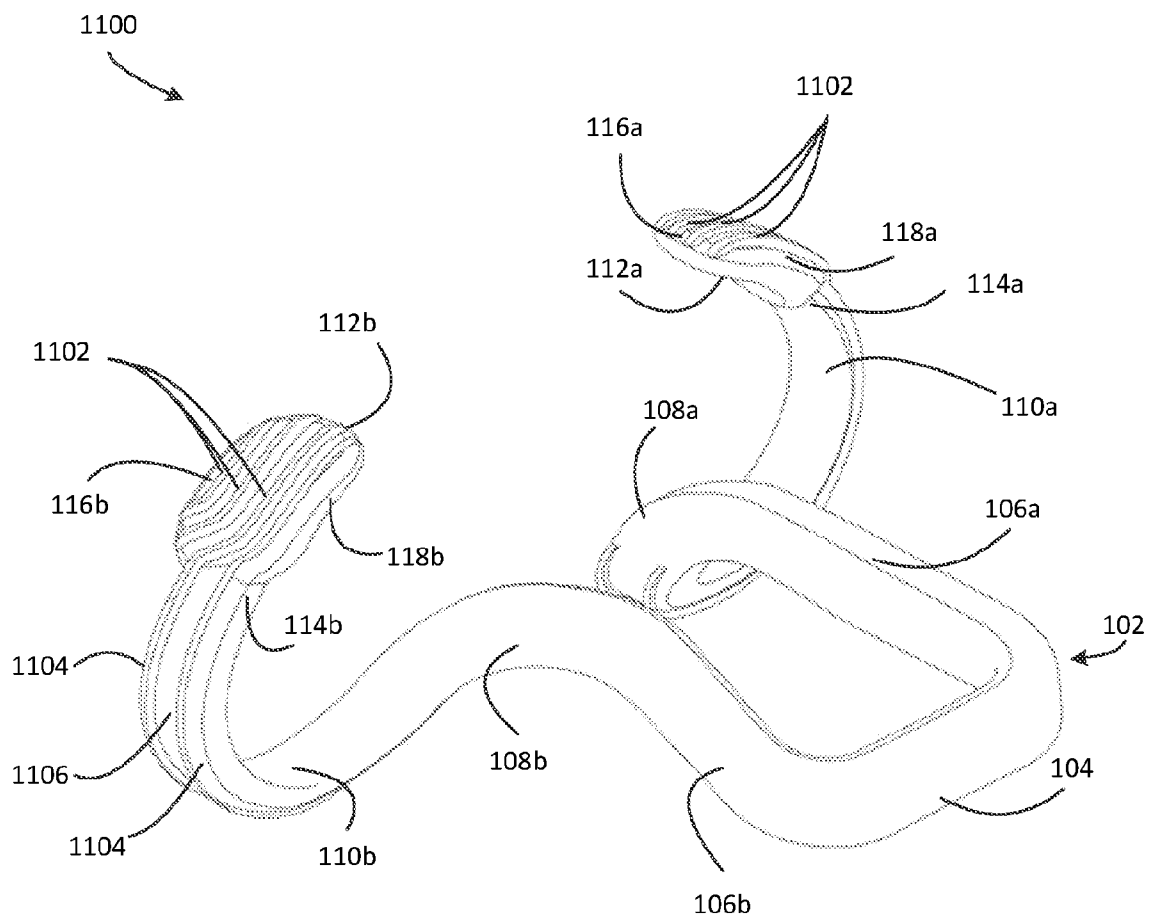


Figure 11

16/22

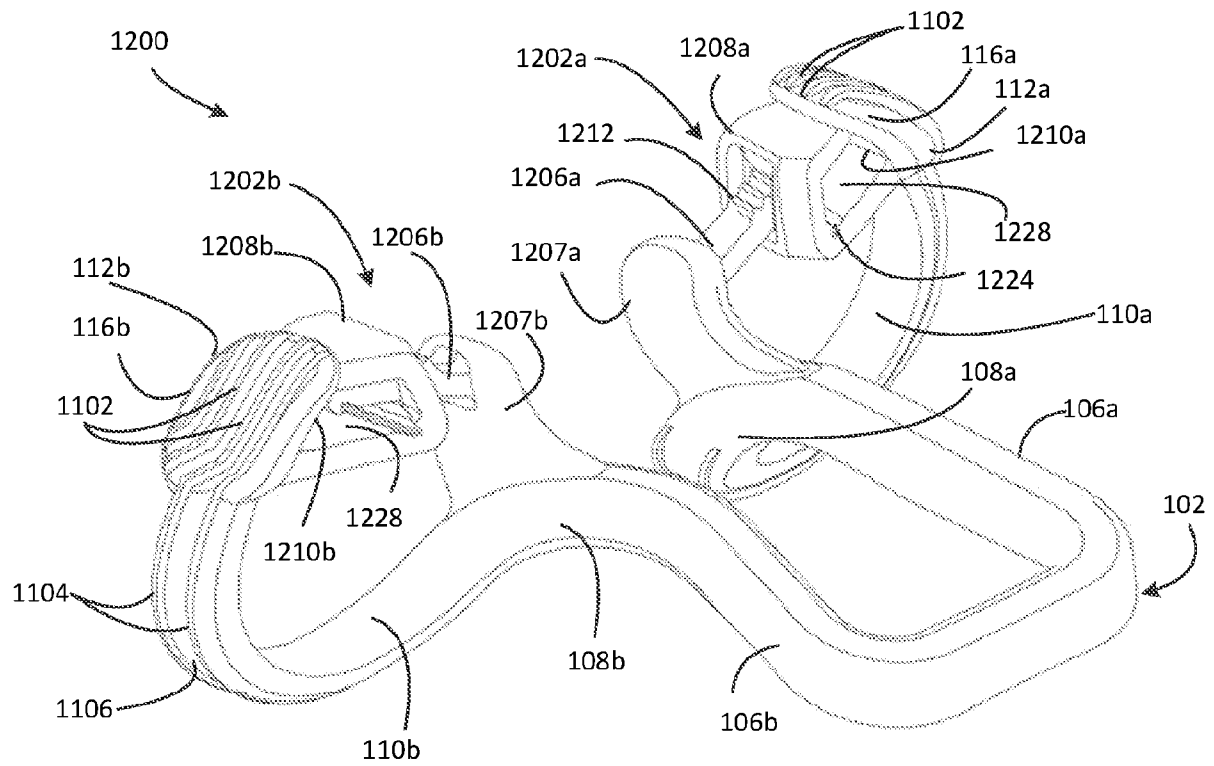


Figure 12A

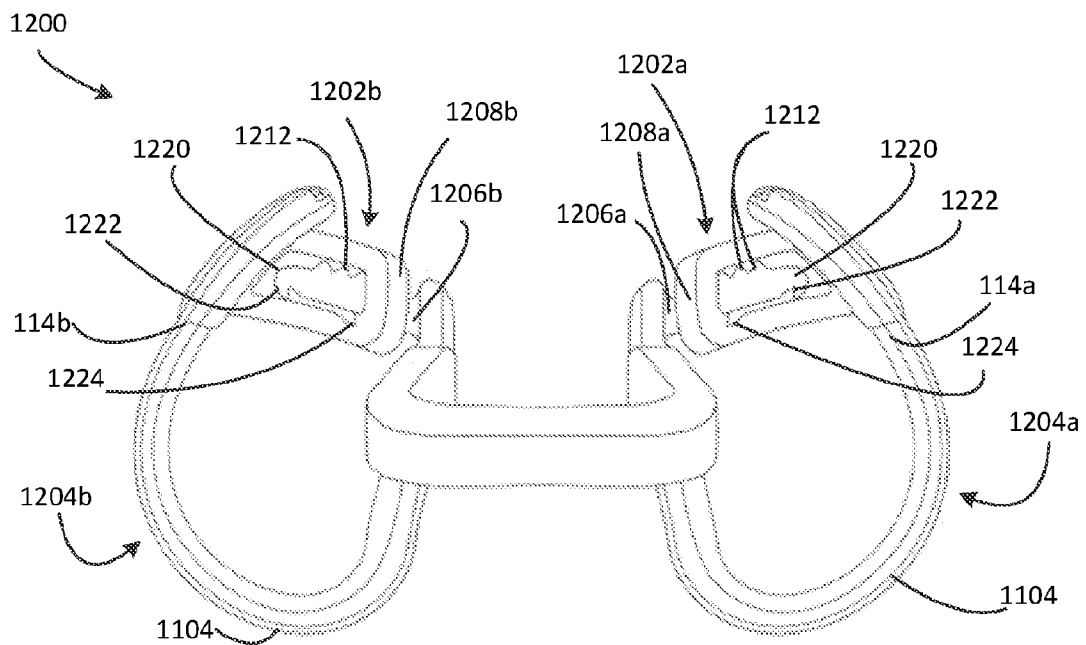


Figure 12B

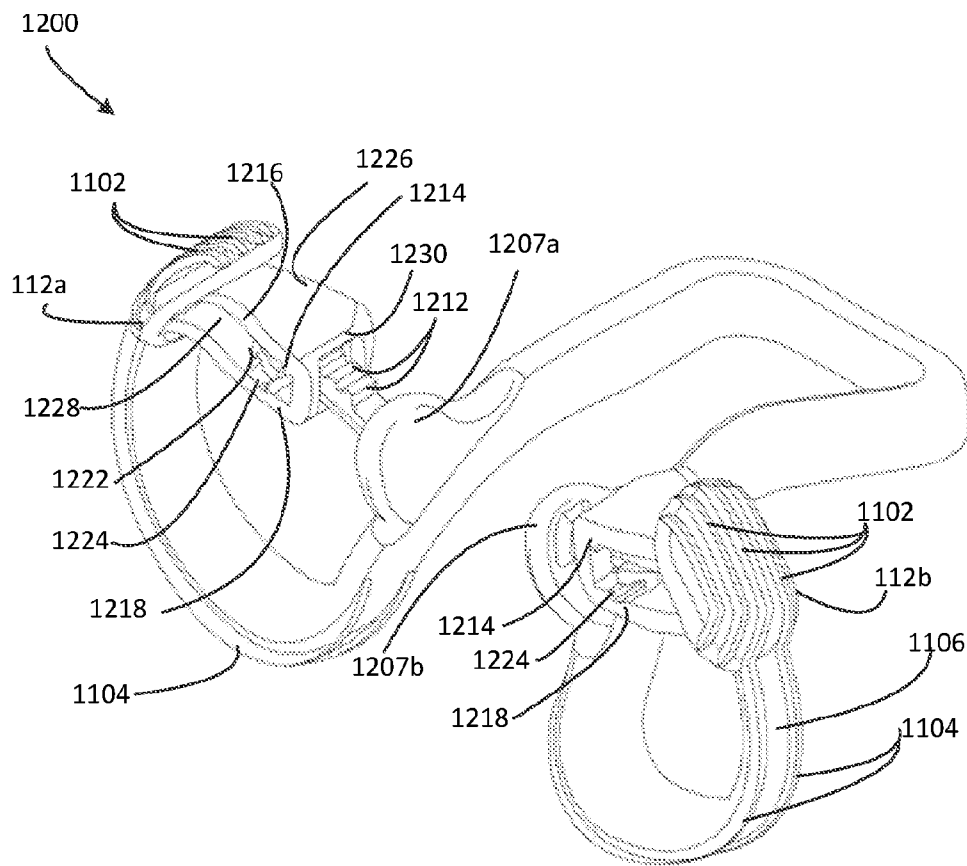
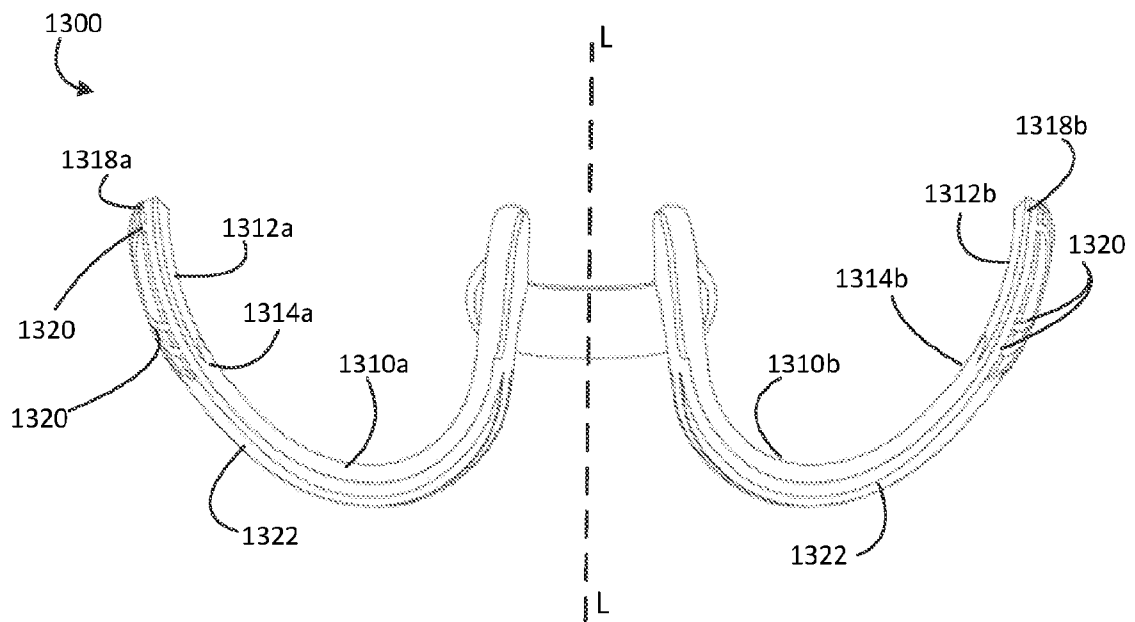
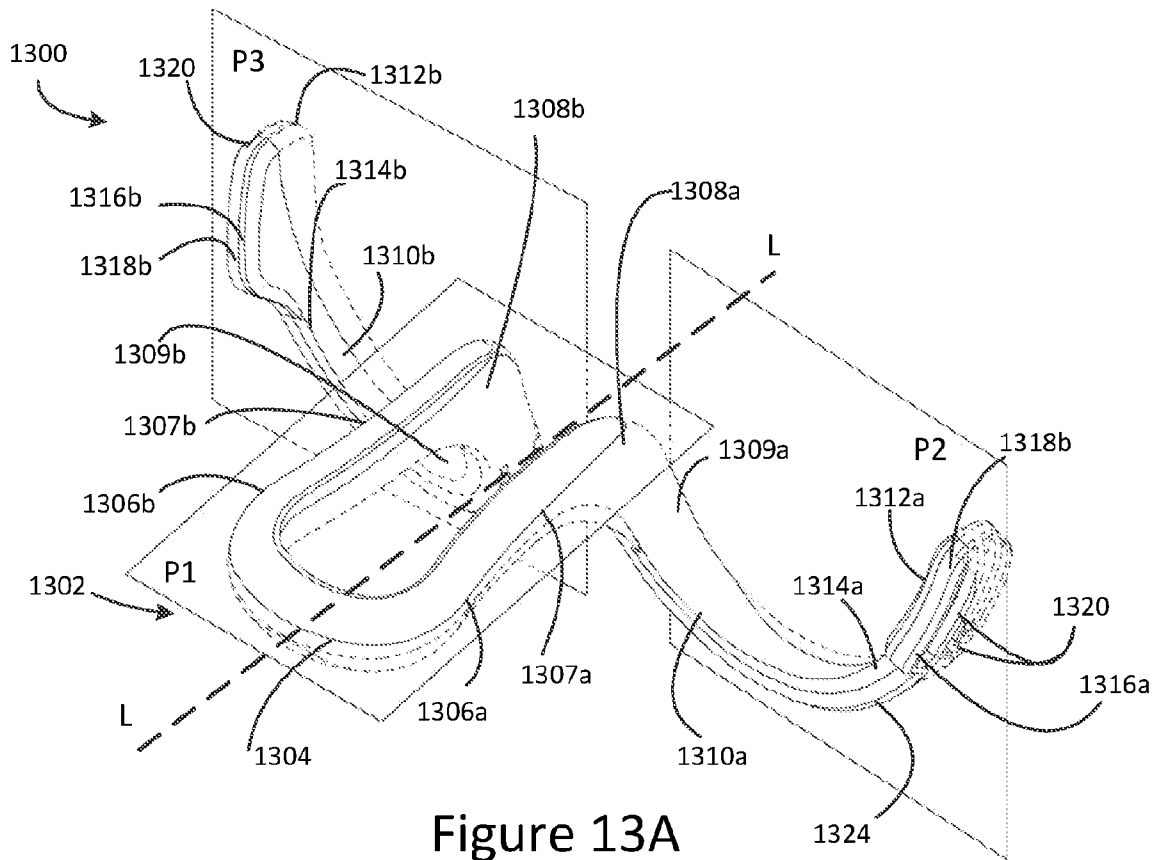


Figure 12C



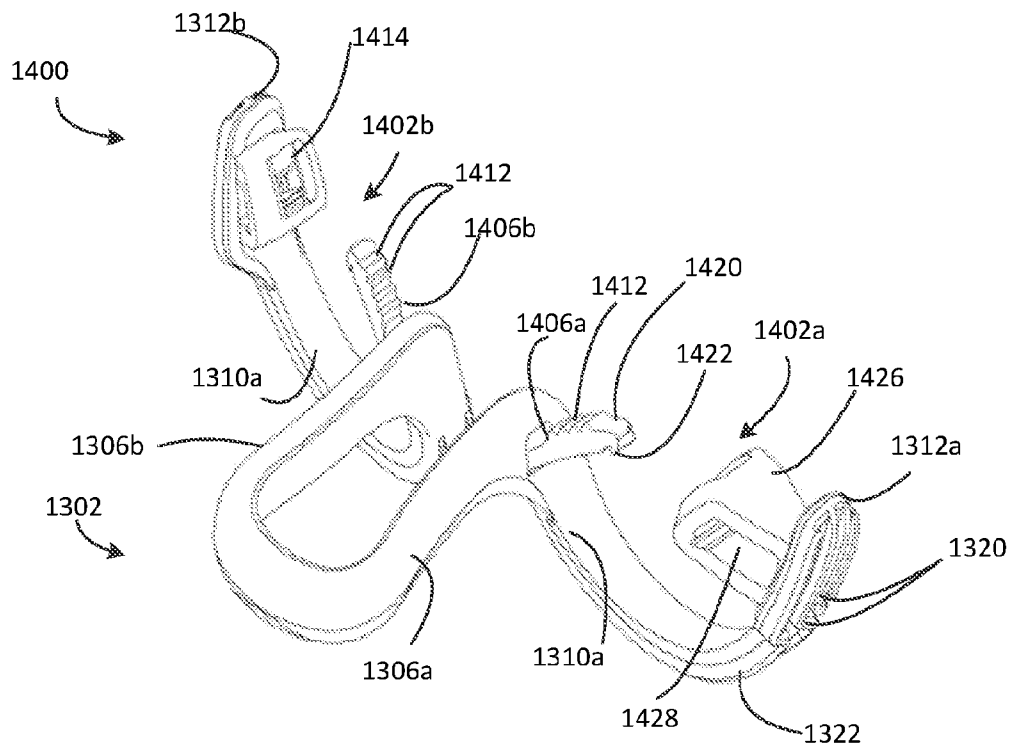


Figure 14A

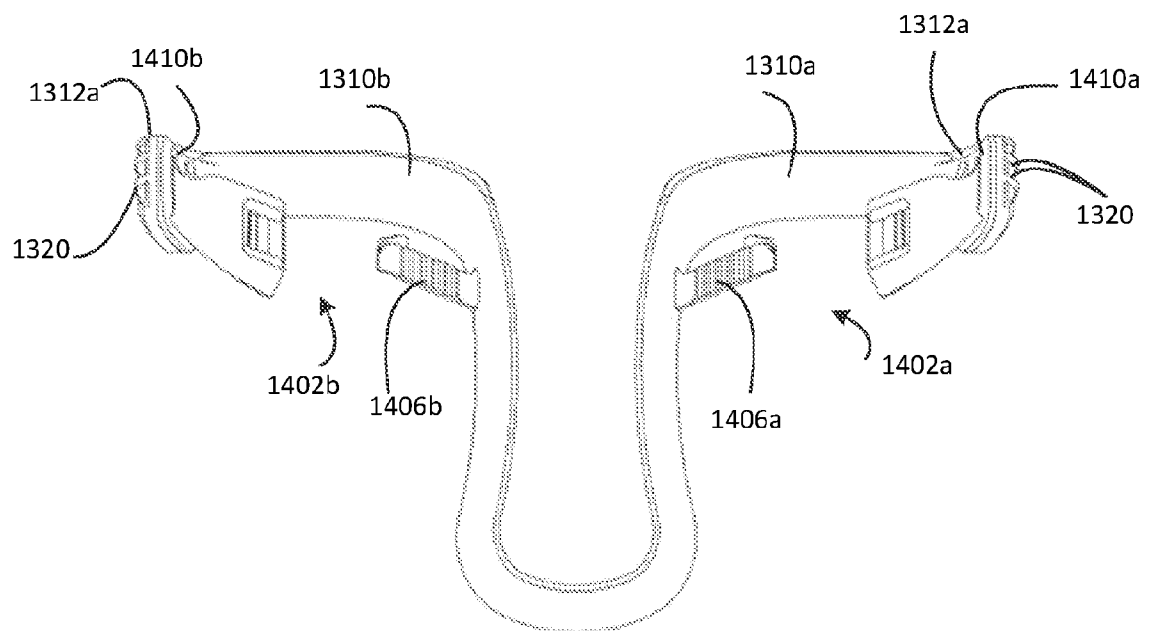


Figure 14B

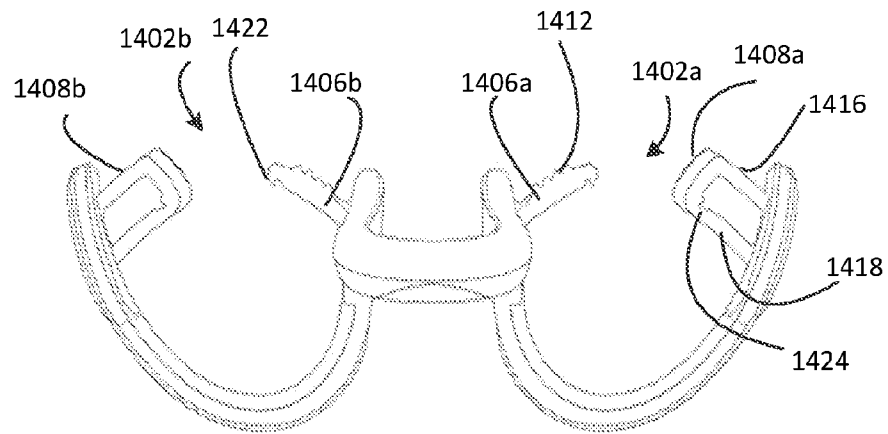


Figure 14C

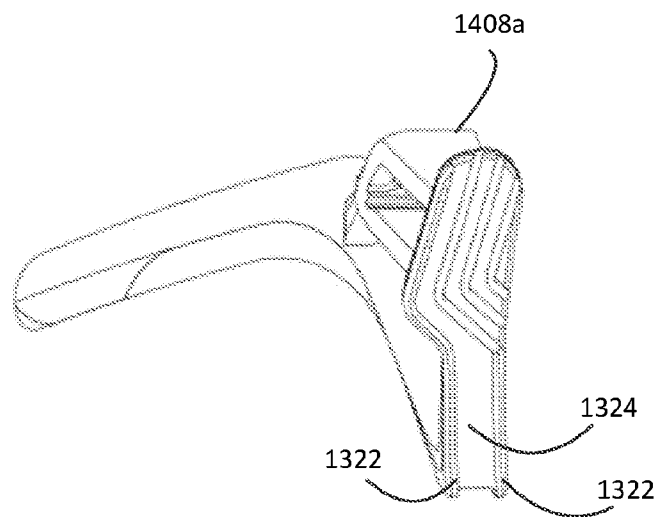


Figure 14D

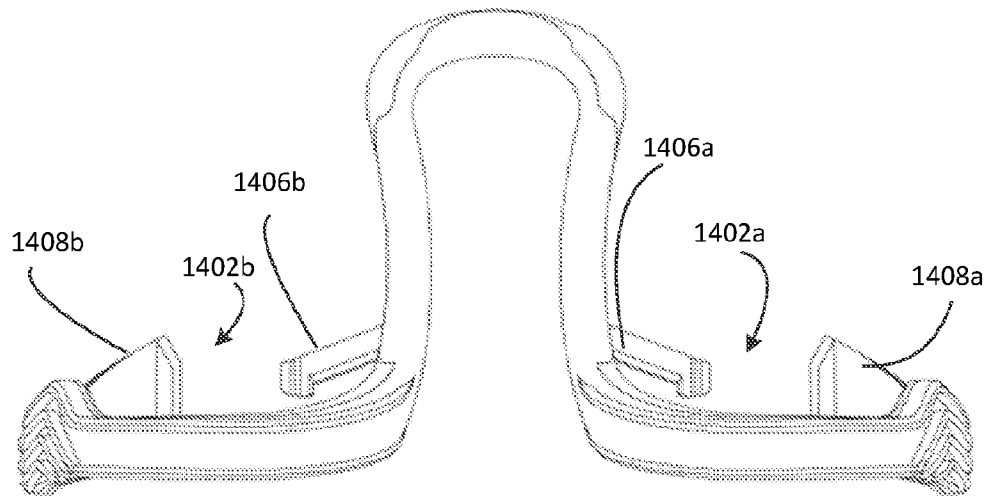


Figure 14E

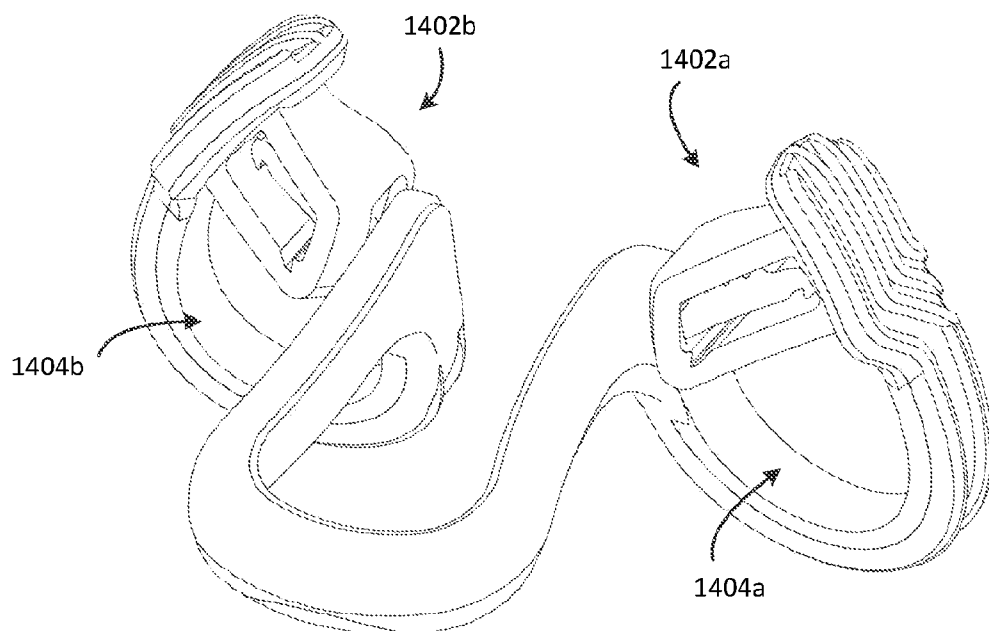


Figure 14F

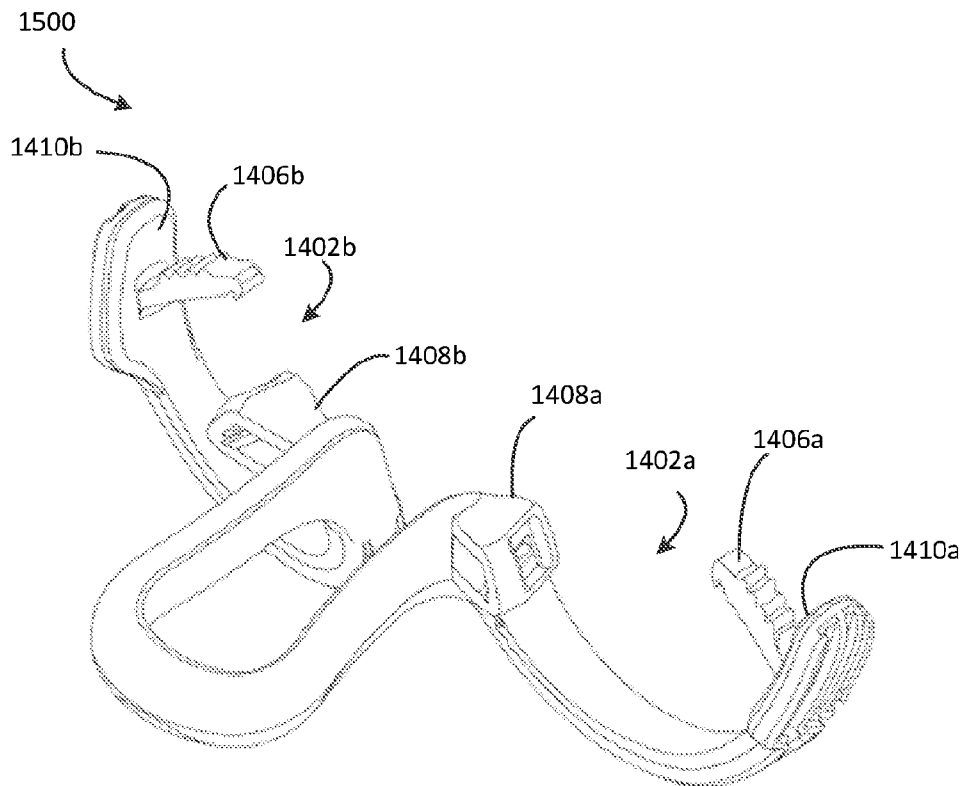


Figure 15

INTERNATIONAL SEARCH REPORT

International application No.
PCT/AU2015/050032

A. CLASSIFICATION OF SUBJECT MATTER

A61F 5/08 (2006.01) A61F 5/56 (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)

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 practicable, search terms used)

WPI, EPODOC: A61F5/08, A61M29/00, A61F5/56 and Keywords (Nostril, Bridge, Expand, Cantilever and like terms).

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
	Documents are listed in the continuation of Box C	

☒ Further documents are listed in the continuation of Box C

☒ See patent family annex

* "A"	Special categories of cited documents: 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
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"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
17 April 2015

Date of mailing of the international search report
17 April 2015

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INTERNATIONAL SEARCH REPORT		International application No.
C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		PCT/AU2015/050032
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 2004/026391 A1 (ASAP BREATHE ASSIST PTY LTD) 01 April 2004 Abstract, Figure 27, 28A, Page 3 Line 13-Page 12 Line 16	1-74
A	AU 2013205674 A1 (CONSEGNA GROUP LIMITED) 13 February 2014 Abstract, Figure 1A-1D, Page 2 Line 19-Page 11 Line 27	1-74
A	US 2010/0042134 A1 (WIEN) 18 February 2010 Whole Document	1-74
A	US 5931852 A (BRENNAN) 03 August 1999 Whole Document	1-74
P,A	US 8834512 B1 (BROWN et al) 16 September 2014 Whole Document	1-74

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INTERNATIONAL SEARCH REPORT		International application No.	
Information on patent family members		PCT/AU2015/050032	
This Annex lists known patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.			
Patent Document/s Cited in Search Report		Patent Family Member/s	
Publication Number	Publication Date	Publication Number	Publication Date
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Form PCT/ISA/210 (Family Annex)(July 2009)			

INTERNATIONAL SEARCH REPORT Information on patent family members		International application No. PCT/AU2015/050032	
This Annex lists known patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.			
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Publication Number	Publication Date	Publication Number	Publication Date
US 8834512 B1	16 September 2014	EP 1105047 A1 WO 0078223 A1	13 Jun 2001 28 Dec 2000
End of Annex			
Due to data integration issues this family listing may not include 10 digit Australian applications filed since May 2001. Form PCT/ISA/210 (Family Annex)(July 2009)			