

| Justification: | Function: | Electronic signature justification: | Signed: Date (yyyy-mm-dd) - Time (hh:mm): |
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| Issued: | QA | Abdallah Almashharawi - ABDALM | 2021-12-28 - 10:46 |
| Reviewed: | QA | Elin Algotson - ELIALG | 2021-12-28 - 11:28 |
| Approved: | DD | Jon Berg - JONBER | 2021-12-29 - 07:00 |
| Released: | QA | Abdallah Almashharawi - ABDALM | 2022-04-04 - 09:27 |

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Provox® HME Cap



Product description:

Provox HME Cap is a dome-shaped titanium ring for rehabilitation after total laryngectomy. It allows use of the Provox FreeHands HME Cassettes without the Provox FreeHands HME Speech Valve. The front opening of the cap can be occluded manually to speak.

Product Information

| | | | |
|--------------------------------------|---|-----------------|----|
| Document ID: | PF019-01-TechInfo | Edition: | 06 |
| Manufacturer: | Atos Medical AB Kraftgatan 8 SE-242 35 Hörby, Sweden | | |
| Classification: (EU) 2017/745 | Class I (1.1 Rule) | | |
| Intended Use: | <p>Provox HME Cap is a single patient use, dome-shaped titanium ring, that allows use of Provox FreeHands HME cassette (REF 8220, 8221) without Provox FreeHands FlexiVoice.</p> <p>Provox HME Cap is only intended for use when using Provox FreeHands FlexiVoice is not recommended, i.e. when sleeping.</p> <p>Provox HME Cap cannot be used with any other type of HME cassette.</p> <p>The front opening of the cap can be occluded manually to speak.</p> <p>Provox HME Cap can be cleaned and reused.</p> | | |
| Use specifications: | <p>Intended medical indication: Product for rehabilitation after total laryngectomy.</p> <p>Intended patient population: Male and female of any age. Cognitive ability: by a clinician judged as sufficient. Manual dexterity: by a clinician judged as sufficient. Not intended for patients with mechanical ventilation. Not intended for patients with a low tidal volume.</p> <p>Intended usage: Single patient use, prescription.</p> <p>Intended part of the body/type of tissue applied to or interacted with: The product is placed in front of the tracheostoma. The tissue contact is indirect via inhaled air.</p> <p>Intended user profile: The product is supposed to be handled by the patient but is also handled by physicians, trained nurses, SLPs.</p> <p>Intended conditions of use: Environment: Home use (normal daily environment without any hygienic or environmental restrictions regarding temperature, moisture etc.). Outpatient clinic use. Hospital use.</p> | | |
| Contraindications: | There are no known contraindications. | | |
| CE Mark: | Yes. Devices are CE-marked. | | |
| GMDN code: | 58705 (Tracheostoma protective filter) | | |
| Sterilization: | Non-Sterile | | |
| Raw material: | Titanium | | |
| Latex information: | Not manufactured with natural rubber latex. | | |
| Biological origin: | The device is not manufactured with materials derived from human or animal source. | | |
| Handling and storage: | Store the product dry and away from sunlight at room temperature. Excursions permitted between 2°C - 42°C. | | |

Product Information

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|-------------------------------------|--|
| Waste handling and disposal: | Waste handling and disposal should be carried out in agreement with medical practice and applicable national laws and legislations. Used product may be a potential biohazard. |
| Hazardous components: | None |
| Expiration date: | 5 years after manufacturing. |
| Packaging: | HME Cap is separately packed in a mini grip plastic bag made of low-density polyethylene together with instructions for use in a cardboard box. |

Product Information

Devices under Basic UDI-DI: 7331791-HME-A-000-0002-F2

| REF | Name | UDI-DI |
|------|----------------|----------------|
| 7730 | Provox HME Cap | 07331791003011 |

Atos Medical AB compatible products:

| Range | BASIC UDI-DI |
|-------------------------------|---------------------------|
| Provox FreeHands HME Cassette | 7331791-HME-0-000-0003-XJ |

| Justification: | Function: | Electronic signature justification: | Signed: Date (yyyy-mm-dd) - Time (hh:mm): |
|----------------|-----------|-------------------------------------|---|
| Issued: | QA | Peter Sundsten - X-PETSUN | 2020-04-22 - 10:12 |
| Reviewed: | DD | Jon Berg - JONBER | 2020-04-22 - 15:00 |
| Approved: | DD | Fredrik Calais - FRECAL | 2020-04-23 - 09:11 |
| Released: | QA | Peter Sundsten - X-PETSUN | 2020-05-13 - 14:27 |

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Released

Product Information

Provox® FreeHands HME



Product description:

Provox FreeHands HMEs are designed to allow automatic tracheostoma closure (instead of digital stoma occlusion) for voice prosthesis users (e.g. Provox, Voice Prosthesis), while providing heat-and-moisture-exchange (HME, via the Provox FreeHands Cassette) for pulmonary rehabilitation. The device should be adjusted for the end user by a clinician trained in voice and pulmonary rehabilitation, e.g. a speech pathologist, before it can be used without supervision – *this does not apply for Provox FreeHands FlexiVoice.*

Product Information

| | | | |
|--|--|-----------------|----|
| Document ID: | PF022-02-TechInfo | Edition: | 07 |
| Manufacturer: | Atos Medical AB Kraftgatan 8, P.O. Box 183, SE-242 22 Hörby, Sweden | | |
| Classification: (EU) 2017/745 | Class I, Rule 1 | | |
| Intended Use: | Provox FreeHands HME Cassette/Moist/Flow is intended for single use for spontaneously breathing laryngectomized patients, utilizing a voice prosthesis and must be used in combination with a Provox FreeHands speaking valve, a Provox cap or Digitop O2. Provox FreeHands combines pulmonary rehabilitation with its Heat and Moisture Exchanging functionality with voice rehabilitation using an Automatic Speaking Valve or Manual Occlusion. The HME conditions inhaled air by retaining heat and moisture from the exhaled air. The device also partially restores lost breathing resistance. | | |
| CE Mark: | Yes, the devices are CE marked. | | |
| GMDN code: | 58705 (Tracheostoma protective filter) | | |
| Sterilization: | Non-sterile | | |
| Raw material: | Cassette: Styrene-ethylene-butadiene-styrene (SEBS) and Polyurethane (PUR). Foam: Polyurethane (PUR) with calcium chloride (CaCl ₂) | | |
| Latex information: | Not manufactured with natural rubber latex | | |
| Biological origin: | The device is not manufactured with materials derived from human or animal source. | | |
| Handling and storage: | Store the product dry and away from sunlight at room temperature. Excursions permitted between 2°C - 42°C. | | |
| Waste handling and disposal: | Waste handling and disposal should be carried out in agreement with medical practice and applicable national laws and legislations. Used product may be a potential biohazard. | | |
| Hazardous components: | None | | |
| Expiration date: | 3 years after manufacturing. | | |
| Packaging: | HMEs are packed in plastic bags made of polyethylene that are placed in a cardboard box. The following configurations are present: | | |

| REF | #HMEs per plastic bag | #Plastic bags in cardboard box | #HMEs in product |
|---------------------|-----------------------|--------------------------------|------------------|
| 8220, 8221 | 10 | 3 | 30 |
| 8220-18, 8221-18 | 10 | 3 | 30 |
| 8222-18, 8223-18 | 10 | 2 | 20 |

Product Information

Devices under Basic UDI-DI: 7331791-HME-0-000-0003-XJ

| REF | Name | UDI-DI |
|---------|-------------------------------------|----------------|
| 8220 | Provox FreeHands HME Moist (30 pcs) | 07331791008368 |
| 8221 | Provox FreeHands HME Flow (30 pcs) | 07331791008375 |
| 8220-18 | Provox FreeHands HME Moist (30 pcs) | 07331791012372 |
| 8221-18 | Provox FreeHands HME Flow (30 pcs) | 07331791013553 |
| 8222-18 | Provox FreeHands HME Moist (20 pcs) | 07331791013560 |
| 8223-18 | Provox FreeHands HME Flow (20 pcs) | 07331791013577 |

Atos Medical AB compatible products:

| Range | BASIC UDI-DI |
|-----------------------------|---------------------------|
| Provox StabiliBase | 7331791-ADH-0-000-0000-CQ |
| Provox XtraBase | 7331791-ADH-0-000-0000-CQ |
| Provox StabiliBase OptiDerm | 7331791-ADH-0-000-0000-CQ |
| Provox Flexiderm | 7331791-ADH-0-000-0000-CQ |
| Provox Optiderm | 7331791-ADH-0-000-0000-CQ |
| Provox LaryTube | 7331791-LTU-0-000-0002-3E |
| Provox LaryButton | 7331791-LTU-0-000-0000-38 |
| Provox HME Cassette Adaptor | 7331791-HME-A-000-0003-F5 |
| Provox HME Cap | 7331791-HME-A-000-0002-F2 |
| Provox FreeHands FlexiVoice | 7331791-HME-0-000-0003-XJ |

| Justification: | Function: | Electronic signature justification: | Signed: Date (yyyy-mm-dd) - Time (hh:mm): |
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| Issued: | QA | Carolina Johansson - SEHRBJNC | 2022-04-22 - 08:34 |
| Reviewed: | QA | Karolina Nilsson - KARNIL | 2022-04-22 - 09:16 |
| Approved: | DD | Mikael Melefors - SEHRBSGM | 2022-04-25 - 10:55 |
| Released: | QA | Carolina Johansson - SEHRBJNC | 2022-05-19 - 15:04 |

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Product Information

Provox® FreeHands FlexiVoice

Provox® Life FreeHands FlexiVoice

Provox® FreeHands FlexiVoice Arch



Figure 1- Provox FreeHands FlexiVoice/Provox Life FreeHands FlexiVoice



Figure 2- Provox FreeHands FlexiVoice Arch

Product description:

The Provox FreeHands FlexiVoice/Provox Life FreeHands FlexiVoice consists of two parts assembled together, a speaking valve for single patient use and a disposable HME cassette. The speaking valve is made of plastic and the membrane is made of silicone. The HME cassette is also made of plastic and a salt treated polyurethane foam.

The speaking valve has two modes; Automatic Speaking Mode and Locked Mode. Rotating the top of the speaking valve moves the device into the automatic speaking or the locked position. Speaking can be done both by using the automatic speaking valve and by manual occlusion of the opening in the front. Manual occlusion is possible in both modes.

The Provox FreeHands FlexiVoice Arch to be used together with the Provox FreeHands FlexiVoice to keep fabric away from the opening of the device.

Product Information

| | | | |
|--------------------------------------|---|-----------------|----|
| Document ID: | PF058-01-TechInfo | Edition: | 06 |
| Manufacturer: | Atos Medical AB Kraftgatan 8 SE-242 35 Hörby,Sweden | | |
| Classification: (EU) 2017/745 | Class I, Rule 1 | | |
| Intended Use: | The Provox FreeHands FlexiVoice/ Provox Life FreeHands FlexiVoice combines pulmonary rehabilitation using a Heat and Moisture Exchanger with voice rehabilitation using an Automatic Speaking Valve or Manual Occlusion, in laryngectomized patients using a voice prosthesis. | | |
| Use specifications: | <p>Intended medical indication: Total laryngectomy with a voice prosthesis in situ.</p> <p>Intended patient population: Male and female of any age. Cognitive ability, by a clinician judged as sufficient. Manual dexterity, by a clinician judged as sufficient. Not intended for patients with mechanical ventilation.</p> <p>Intended usage: Single patient use, prescription.</p> <p>Intended part of the body/type of tissue applied to or interacted with: The device will contact intact skin and as external communicating device the contact mode with tissue is indirect via air.</p> <p>Intended user profile: The product is supposed to be handled by the patient but is also handled by physicians, trained nurses, SLPs, clinicians and caregivers.</p> <p>Intended conditions of use: The HME cassette /moist/flow/Life FreeHands HME is for single use. The Speaking valve, Arch and HME Removal Aid are for single patient use. Should only be used when awake. Only intended for laryngectomized patients using a voice prosthesis. Only intended for laryngectomized patients who can tolerate using an HME. Environment including hygienic requirements: Primarily: Home use. Secondary: Hospital use. Frequency of use: Daily continuous use, not during sleep. Location: Ambient temperature -20°C to +45°C. Mobility: Only intended for use by patients with sufficient manual dexterity to handle rotation of the speaking valve, assembly/disassembly of the device and insertion/removal of the device.</p> | | |
| Contraindications: | The product shall not be used by patients with a decreased level of consciousness, patients with reduced mobility of the arms and/or hands, or patients who are unable to remove the device themselves. | | |
| CE Mark: | Yes. Devices are CE-marked. | | |
| GMDN code: | 36071 | | |
| Sterilization: | Non-sterile | | |
| Raw material: | Speaking valve: PP, Silicone and POM. Arch: MABS | | |

Product Information

| | |
|-------------------------------------|---|
| Latex information: | Not manufactured with natural rubber latex |
| Biological origin: | The device is not manufactured with materials derived from human or animal source. |
| Handling and storage: | Store the product dry and away from sunlight at room temperature. Excursions permitted between 2°C - 42°C. |
| Waste handling and disposal: | Waste handling and disposal should be carried out in agreement with medical practice and applicable national laws and legislations. Used product may be a potential biohazard. |
| Hazardous components: | None |
| Expiration date: | 3 years after manufacturing. |
| Packaging: | <p>Ref 7757: Provox FreeHands FlexiVoice, 1 pc/model (light, medium, strong) is packed in a plastic jar of polypropylene. Provox FreeHands HME Cassette, 30pcs (10pcs/bag) are packed in a plastic bag of polyethylene. Removal Aid, 1pc is packed in plastic bag of polyethylene. FlexiVoice Arch, 2pcs are packed in plastic bag of polyethylene. The products and instructions for use are packed in a cardboard box.</p> <p>Ref 7760: Provox FreeHands FlexiVoice (light, medium, strong) 1 pc/model is packed in a plastic jar of polypropylene. Removal Aid, 1pc is packed in plastic bag of polyethylene. FlexiVoice Arch, 2pcs are packed in plastic bag of polyethylene. The products and instructions for use are packed in a cardboard box.</p> <p>Ref 8161: Provox FreeHands FlexiVoice Light, 1pc is packed in a plastic jar of polypropylene. Removal Aid, 1pc is packed in plastic bag of polyethylene. FlexiVoice Arch, 1pc are packed in plastic bag of polyethylene. The products and instructions for use are packed in a cardboard box.</p> <p>Ref 8162: Provox FreeHands FlexiVoice Medium, 1pc is packed in a plastic jar of polypropylene. Removal Aid, 1pc is packed in plastic bag of polyethylene. FlexiVoice Arch, 1pc are packed in plastic bag of polyethylene. The products and instructions for use are packed in a cardboard box.</p> <p>Ref 8163: Provox FreeHands FlexiVoice Strong, 1pc is packed in a plastic jar of polypropylene. Removal Aid, 1pc is packed in plastic bag of polyethylene. FlexiVoice Arch, 1pc are packed in plastic bag of polyethylene. The products and instructions for use are packed in a cardboard box.</p> |

Product Information

Ref 8165:

FlexiVoice Arch, 5pcs (1pc/bag) are packed in plastic bag of polyethylene.
The products and a informative insert are packed in a cardboard box.

Ref 8166:

Provox FreeHands FlexiVoice XtraStrong, 1pc is packed in a plastic jar of polypropylene.

Removal Aid, 1pc is packed in plastic bag of polyethylene.

FlexiVoice Arch, 1pc are packed in plastic bag of polyethylene.

The products and instructions for use are packed in a cardboard box.

Ref 8210:

Provox Life FreeHands HME, 20pcs (10pcs/blister) are packed in blister of PET.

Provox FreeHands FlexiVoice (light, medium, strong) 1 pc/model is packed in a plastic jar of polypropylene.

FlexiVoice Arch, 2pcs are packed in plastic bag of polyethylene.

The products and instructions for use are packed in a cardboard box.

Product Information

Devices under Basic UDI-DI: 7331791-HME-0-000-0007-XW

| REF | Name | UDI-DI |
|---------|--|----------------|
| 7757 | Provox FreeHands FlexiVoice Set Plus | 07331791008276 |
| 7760 | Provox FreeHands FlexiVoice Set | 07331791008283 |
| 8161 | Provox FreeHands FlexiVoice Light | 07331791008290 |
| 8162 | Provox FreeHands FlexiVoice Medium | 07331791008306 |
| 8163 | Provox FreeHands FlexiVoice Strong | 07331791008313 |
| 8165 | Provox FreeHands FlexiVoice Arch (5 pcs) | 07331791010408 |
| 8166 | Provox FreeHands FlexiVoice XtraStrong | 07331791010668 |
| 7757-18 | Provox FreeHands FlexiVoice Set Plus | 07331791013508 |
| 7760-18 | Provox FreeHands FlexiVoice Set | 07331791013515 |
| 8161-18 | Provox FreeHands FlexiVoice Light | 07331791013461 |
| 8162-18 | Provox FreeHands FlexiVoice Medium | 07331791012365 |
| 8163-18 | Provox FreeHands FlexiVoice Strong | 07331791013485 |
| 8166-18 | Provox FreeHands FlexiVoice XtraStrong | 07331791013492 |
| 8210 | Provox Life FreeHands FV Set Plus | 07331791015175 |

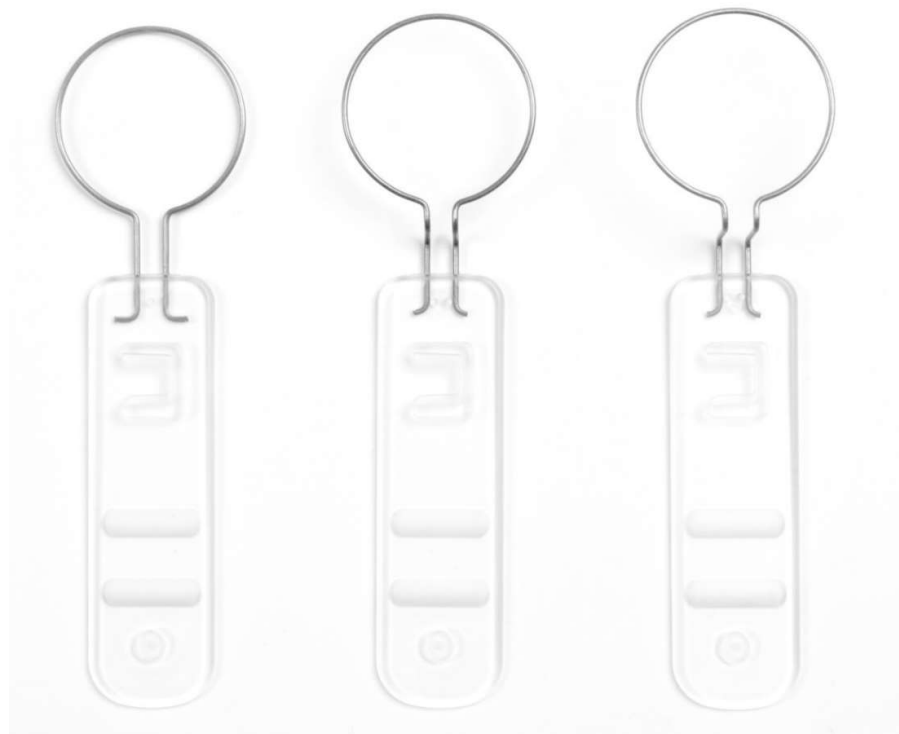

Atos Medical AB compatible products:

| Range | BASIC UDI-DI |
|---------------------------|---------------------------|
| Provox FreeHands HME | 7331791-HME-0-000-0003-XJ |
| Provox Life FreeHands HME | 7331791-HME-0-000-0008-XZ |
| Removal Aid | 7331791-HME-0-000-0005-XQ |
| Provox FreeHands Support | 7331791-HME-A-000-0000-EU |

| Justification: | Function: | Electronic signature justification: | Signed: Date (yyyy-mm-dd) - Time (hh:mm): |
|----------------|-----------|-------------------------------------|---|
| Issued: | QA | Sara Dahl - X-SARDAH | 2021-11-10 - 13:01 |
| Reviewed: | QA | John Wennborg - JOHWEN | 2021-11-10 - 15:05 |
| Approved: | DD | Diana Tieger - DIATIE | 2021-11-11 - 16:15 |
| Released: | QA | Sara Dahl - X-SARDAH | 2021-12-10 - 09:53 |

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Provox® FreeHands Support™



Product description:

The Provox FreeHands Support system consists of two parts; Provox FreeHands Support and Provox FreeHands Support Adhesive.

The reusable Provox FreeHands Support is available in three different variants; 'Flat', 'Medium' and 'Deep'. The three variants, as well as the possibility to adjust the metal ring and plastic parts, enable personal fit. Provox FreeHands Support Adhesive is a single-use adhesive that is attached to the skin, beneath the Provox Adhesive.

Product Information

| | | | |
|--|--|-----------------|----|
| Document ID: | PF078-01-TechInfo | Edition: | 05 |
| Manufacturer: | Atos Medical AB Kraftgatan 8 SE-242 35 Hörby,Sweden | | |
| Classification: (EU) 2017/745 | Class 1, Rule 1 | | |
| Intended Use: | Provox FreeHands Support provides support to the Provox adhesive when using a Provox hands-free speaking valve after total laryngectomy. The device is a single patient use device with a single use adhesive. | | |
| Use specifications: | <p><i>Intended medical indication:</i> Product for rehabilitation for patients breathing through a tracheostoma after a total laryngectomy.</p> <p><i>Intended patient population:</i> Patients of any age. Cognitive ability, by a clinician judged as sufficient. Manual dexterity, by a clinician judged as sufficient.</p> <p><i>Intended usage:</i> The FreeHands Support is a reusable single patient device whereas the FreeHands Support Adhesive is a (disposable) single use device. The product is intended to be available to patients over-the-counter. Should only be used when awake (i.e., not during sleep). Intended for laryngectomized patients using Provox hands free speaking valves. The product should only be used by patients with sufficient manual dexterity to handle attachment/removal of the device.</p> <p><i>Intended part of the body/type of tissue applied to or interacted with:</i> The device will contact intact skin.</p> <p><i>Intended user profile:</i> Patient (Nurse, physician or other caregiver only for fitting and instructions).</p> <p><i>Intended conditions of use:</i> Environment of use: (Primarily) Home use, secondary hospital use. Frequency of use: Daily use, not during sleep. Location: Ambient temperature -20°C to +45°C Mobility: The product should only be used by patients with sufficient manual dexterity to handle attachment/removal of the device."</p> | | |
| Contraindications: | Do not use Provox FreeHands Support on breached skin. Provox FreeHands Support is NOT intended to be in place during MRI examination (Magnetic Resonance Imaging), or during the period of Radiation Therapy treatment. | | |
| CE Mark: | Yes. Devices are CE-marked. | | |
| GMDN code: | 62155 (Tracheostomy base plate, reusable) | | |
| Sterilization: | Non-sterile | | |
| Raw material: | Provox FreeHands Support consists of a ring of stainless steel and a plastic part (Polycarbonate, PC). | | |

Product Information

| | |
|-------------------------------------|--|
| Latex information: | Not manufactured with natural rubber latex |
| Biological origin: | The device is not manufactured with materials derived from human or animal source. |
| Handling and storage: | Store the product dry and away from sunlight at room temperature. Excursions permitted between 2°C - 42°C. |
| Waste handling and disposal: | Waste handling and disposal should be carried out in agreement with medical practice and applicable national laws and legislations. Used product may be a potential biohazard. |
| Hazardous components: | None |
| Expiration date: | 3 years after manufacturing. |
| Packaging: | Provox FreeHands Support is packed in a cardboard box together with instructions for use. |

Product Information

Devices under Basic UDI-DI: 7331791-HME-A-000-0000-EU

| REF | Name | UDI-DI |
|------|--------------------------------------|----------------|
| 8020 | Provox FreeHands Support Starter Set | 07331791009266 |
| 8021 | Provox FreeHands Support Flat | 07331791009273 |
| 8022 | Provox FreeHands Support Medium | 07331791009280 |
| 8023 | Provox FreeHands Support Deep | 07331791009297 |

Atos Medical AB compatible products:

| Range | BASIC UDI-DI |
|-----------------------------------|---------------------------|
| Provox FreeHands Support Adhesive | 7331791-HME-A-000-0004-F8 |
| Provox FreeHands FlexiVoice | 7331791-HME-0-000-0007-XW |

| Justification: | Function: | Electronic signature justification: | Signed: Date (yyyy-mm-dd) - Time (hh:mm): |
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| Issued: | DD | Pontus Eklund - X-PONEKL | 2020-04-20 - 15:59 |
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| Approved: | DD | Fredrik Calais - FRECAL | 2020-04-21 - 14:55 |
| Released: | DD | Pontus Eklund - X-PONEKL | 2020-10-28 - 16:38 |

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Product Information

Provox® FreeHands Support™ Adhesive



Product description:

The Provox FreeHands Support system consists of two parts; Provox FreeHands Support and Provox FreeHands Support Adhesive.

The reusable Provox FreeHands Support is available in three different variants; 'Flat', 'Medium' and 'Deep'. The three variants, as well as the possibility to adjust the metal ring and plastic parts, enable personal fit. Provox FreeHands Support Adhesive is a single-use adhesive that is attached to the skin, beneath the Provox Adhesive.

Product Information

| | | | |
|--|--|-----------------|----|
| Document ID: | PF078-02-TechInfo | Edition: | 05 |
| Manufacturer: | Atos Medical AB Kraftgatan 8, P.O. Box 183, SE-242 22 Hörby, Sweden | | |
| Classification: (EU) 2017/745 | Class I, Rule 1 | | |
| Intended Use: | Provox FreeHands Support provides support to the Provox Adhesive when using a Provox hands-free speaking valve after total laryngectomy. The device is a single patient use device with a single use adhesive. | | |
| CE Mark: | Yes, the devices are CE marked. | | |
| GMDN code: | 62175 (Stomal appliance skin-adherent patch, non-sterile) | | |
| Sterilization: | Non-sterile | | |
| Raw material: | Provox FreeHands Support Adhesive consists of a plastic plate (Polycarbonate, PC) and an adhesive tape (Acrylate and Polyester film). | | |
| Latex information: | Not manufactured with natural rubber latex | | |
| Biological origin: | The device is not manufactured with materials derived from human or animal source. | | |
| Handling and storage: | Store the product dry and away from sunlight at room temperature. Excursions permitted between 2°C - 42°C. | | |
| Waste handling and disposal: | Waste handling and disposal should be carried out in agreement with medical practice and applicable national laws and legislations. Used product may be a potential biohazard. | | |
| Hazardous components: | None | | |
| Expiration date: | 3 years after manufacturing. | | |
| Packaging: | Provox FreeHands Support Adhesive is packed in a blister package made of PET film and with a re-sealable top film (PP). It is then packed in a cardboard box. | | |

Product Information

Devices under Basic UDI-DI: 7331791-HME-A-000-0004-F8

| REF | Name | UDI-DI |
|------|--|----------------|
| 8024 | Provox FreeHands Support Adhesive (15pc) | 07331791009303 |

Atos Medical AB compatible products:

| Range | BASIC UDI-DI |
|--------------------------|---------------------------|
| Provox FreeHands Support | 7331791-HME-A-000-0000-EU |