This user guide is valid for:

**CROS device model**
Phonak CROS P-13  
CE mark applied  
2021
Your CROS device details

- If no box is checked and you do not know the model of your CROS device, please ask your hearing care professional.
- Your CROS device operates between 2.4 GHz–2.48 GHz frequency range. When flying please check if flight operator requires devices to be switched into flight mode, see chapter 11.

### CROS device models

<table>
<thead>
<tr>
<th>CROS device models</th>
<th>Battery size</th>
</tr>
</thead>
<tbody>
<tr>
<td>CROS P-13</td>
<td>13</td>
</tr>
</tbody>
</table>

### Earpieces

<table>
<thead>
<tr>
<th>Earpieces</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Dome</td>
<td></td>
</tr>
<tr>
<td>SlimTip</td>
<td></td>
</tr>
<tr>
<td>CROS Tip</td>
<td></td>
</tr>
</tbody>
</table>
Your CROS device have been developed by Phonak – a world leader in hearing solutions based in Zurich, Switzerland.

These premium products are the result of decades of research and expertise and are designed to keep you connected to the beauty of sound! We thank you for making such a great choice and wish you many years of listening pleasure.

Please read the user guide carefully to make sure that you understand and get the best out of your CROS device. Training is not required for handling of this device. A hearing care professional will help set up this CROS device according to your individual preferences during the fitting consultation.
For more information regarding features, benefits, set up, use, maintenance or repairs of your CROS device and accessories, please contact your hearing care professional or the manufacturer representative. Additional information can be found in the datasheet of your product.

Phonak – life is on
www.phonak.com
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<td>52</td>
</tr>
</tbody>
</table>
1. Quick guide

Left & right markings

Blue marking for left side.
Red marking for right side

Changing batteries

1. Remove the sticker from the new battery and wait two minutes.
2. Open the battery door.
3. Place battery in the battery door with the "+" symbol facing upwards.
**Multi-function button**

The button has several functions. It functions as a volume control in the absence of wireless transmission of sound from the CROS device to the hearing aid, a balance control when there is a wireless transmission of sound from the CROS device to the hearing aid and/or a program change depending on the CROS device programming. This is indicated in your individual instructions. If paired with a Bluetooth enabled phone, a short press will accept and a long press will reject an incoming call.

**Flight mode**

To enter flight mode press the lower part of the button for 7 seconds while closing the battery door. To exit flight mode, simply open and close the battery door again.
2. Parts of the CROS device

The following pictures show the CROS device model described in this user guide. You can identify your personal model by checking “Your CROS device details” on page 3.

The Phonak CROS device is an application for unaidable hearing loss on one side. It is placed on the impaired ear and wirelessly transmits sound to the Phonak hearing aid on the other ear.

Phonak CROS device + Phonak hearing aid = Phonak CROS system

ℹ️ Phonak CROS P only works in connection with a Phonak Audéo™ P hearing aid as the receiver.
Possible earpieces

Dome

SlimTip

CROS TIP

CROS P–13

Multi-function button

Tube

Battery door

Anchor (optional)

Speaker dummy
(without earpiece attached)
3. Left & right CROS device markings

There is a red or blue marking on the back of the CROS device and on the speaker dummy. This will tell you if the CROS device is meant to be worn on the left or the right ear.

![Diagram of a CROS device with a blue marking on the left side and a red marking on the right side.]

Blue marking for **left side.**
Red marking for **right side.**
4. On/Off

The battery door is also the on/off switch.

1. Closed battery door = CROS device is on

2. Open battery door = CROS device is off

ℹ️ When you switch on the CROS device you might hear a start-up melody in the hearing aid.
5. Batteries

1. Remove the sticker from the new battery and wait two minutes.

2. Open the battery door.

3. Place battery in the battery door with the "+" symbol facing upwards.

If it is difficult to close the battery door, check that the battery is inserted correctly and the "+" symbol is facing upwards. If the battery is not inserted correctly, the CROS device will not work and the battery door could be damaged.
Low power: You will hear two beeps when the battery is low. You will have approximately 30 minutes to change the battery (this can vary, depending on the hearing aid settings and battery). We recommend that you always have a new battery on hand.

**Replacement battery**
This CROS device requires a 13 zinc-air battery. Identify the correct battery size by checking the following table.

<table>
<thead>
<tr>
<th>Model</th>
<th>Zinc-air battery size</th>
<th>Color marking on package</th>
<th>IEC code</th>
<th>ANSI code</th>
</tr>
</thead>
<tbody>
<tr>
<td>Phonak CROS</td>
<td>13</td>
<td>orange</td>
<td>PR48</td>
<td>7000ZD</td>
</tr>
</tbody>
</table>

Please ensure you use the correct type of battery in your CROS device (zinc-air). Please also read chapter 20.2 for further information on product safety.
6. Putting on the CROS device

1. Place the CROS device behind your ear.

2. Insert the earpiece into your ear canal.

3. If there is an anchor attached to the earpiece, tuck it into the bowl of your ear to secure your CROS device.
7. Removing the CROS device

1. Pull on the bend of the tube and remove the CROS device from behind the ear.
8. Multi-function button

The multi-function button has several functions.

It functions as a volume control in the absence of wireless transmission of sound from the CROS device to the hearing aid, a balance control when there is a wireless transmission of sound from the CROS device to the hearing aid and/or a program change depending on the CROS device programming. This is indicated in your individual “CROS device instructions”. Please ask your hearing care professional for a printout.

If the CROS system is paired with a Bluetooth enabled phone, a short press on the upper or lower part of the button will accept an incoming call and a long press will reject an incoming call – refer to chapter 11.
9. Connectivity overview

The illustration below shows the connectivity options available for your CROS system.

The Bluetooth® word mark and logos are registered trademarks owned by Bluetooth SIG, Inc. and any use of such marks by Sonova AG is under license.

* The TV Connector can be connected to any audio source such as a TV, PC or hi-fi system.
** Roger wireless microphones can be connected to your CROS system as well.
10. Initial pairing

10.1 Initial pairing to Bluetooth enabled device

ℹ️ It is only necessary to perform the pairing procedure once with each device featuring Bluetooth wireless technology. After the initial pairing, your CROS system will connect automatically to the device. The initial pairing process can take up to 2 minutes.

1.
On your device (e.g. a phone), ensure that Bluetooth wireless technology is enabled and search for Bluetooth enabled devices in the connectivity setting menu.

2.
Switch on both the hearing aid and the CROS device. You now have 3 minutes to pair your CROS system with your device.
3.
Your device shows a list of Bluetooth enabled devices. Select the hearing aid from the list to pair the CROS system with the device. A beep confirms successful pairing.

ℹ️ For more information about pairing instructions for Bluetooth wireless technology, specific to some of the most popular phone manufacturers, go to: https://www.phonak.com/com/en/support.html
10.2 Connecting to the device

After your CROS system has been paired to your device, it will automatically connect again when switched on.

- The connection will be maintained as long as the device remains ON and within range.
- Your CROS system can be connected to one device at a time and paired to up to eight devices.
11. Phone calls

Your CROS system connects directly with Bluetooth enabled phones. When paired and connected to your phone, you will hear the caller’s voice directly in your hearing aid. Your voice is picked up by the hearing aid microphones and transmitted to the phone.
11.1 Making a call

Enter the phone number and press the dial button. You will hear the dialing tone through your hearing aid. Your voice is picked up by the hearing aid microphones and transmitted to the phone.

11.2 Accepting a call

When receiving a call, a calling notification will be heard in the hearing aid. The call can be accepted by a short press on the upper or lower part of the multi-function button on the hearing aid or CROS device (less than 2 seconds) or directly on your phone.
11.3 Ending a call

A call can be ended by a long press on the upper or lower part of the multi-function button on the hearing aid or CROS device (more than 2 seconds) or directly on your phone.
11.4 Rejecting a call

An incoming call can be rejected by a long press on the upper or lower part of the multi-function button on the hearing aid or CROS device (more than 2 seconds) or directly on your phone.
12. Flight mode

Your CROS system operates in the 2.4 GHz–2.48 GHz frequency range. When flying some operators require all devices to be switched into flight mode. Entering flight mode will not disable normal hearing aid functionality, and will only disable the Bluetooth connectivity functions including the wireless transmission of sound from the CROS device to the hearing aid on the other ear.

12.1 Enter flight mode

To disable the wireless function and enter flight mode in each device:

1. Open battery door.

2. Hold down the lower part of the multi-function button on the CROS device for 7 seconds while closing the battery door.

In flight mode, your CROS system cannot connect directly to your phone.
12.2 Exit flight mode

To enable the wireless function and exit flight mode in each device:

1. Open battery door.

2. Close the battery door on the CROS device again.
13. Operating, transport and storage conditions

The product is designed such that it functions without problems or restrictions if used as intended, unless otherwise noted in this user guide.

Please ensure to use, transport and store the CROS device according to the following conditions:

<table>
<thead>
<tr>
<th></th>
<th>Use</th>
<th>Transport</th>
<th>Storage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Temperature</td>
<td>+5° to +40°C</td>
<td>-20° to +60°C</td>
<td>-20° to +60°C</td>
</tr>
<tr>
<td></td>
<td>(41° to 104°F)</td>
<td>(-4° to 140°F)</td>
<td>(-4° to 140°F)</td>
</tr>
<tr>
<td>Humidity</td>
<td>30% to 85% (non condensing)</td>
<td>0% to 90% (non condensing)</td>
<td>0% to 70% (non condensing)</td>
</tr>
<tr>
<td>Atmospheric pressure</td>
<td>500 to 1060 hPa</td>
<td>500 to 1060 hPa</td>
<td>500 to 1060 hPa</td>
</tr>
</tbody>
</table>

This CROS device is classified as IP68. This means that it is water and dust resistant and designed to withstand daily life situations. It can be worn in the rain but should not be fully submerged in water or used when taking a shower, swimming or other water activities. This CROS device should never be exposed to chlorinated water, soap, salt water or other liquids with a chemical content.
14. Care and maintenance

Diligent and routine care of your CROS device contributes to outstanding performance and a long service life. To ensure a long service life, Sonova AG provides a minimum of a five year service period after phase out of the respective CROS device.

Please use the following specifications as a guideline. Further information regarding product safety, see chapter 20.2.

**General information**
Before using hair spray or applying cosmetics, you should remove your CROS device from your ear, because these products may damage it.

When you are not using your CROS device, leave the battery door open so that any moisture can evaporate. Make sure that you always completely dry your CROS device after use. Store the CROS device in a safe, dry and clean place.
Your CROS device is resistant to water, sweat and dust under the following conditions:

- The battery door is fully closed. Ensure that no foreign object such as hair is caught in the battery door when it is closed.
- After exposure to water, sweat or dust, the CROS device is cleaned and dried.
- The CROS device is used and maintained as described in this user guide.

**Use of your CROS device around water can restrict air flow to the batteries causing it to stop working. Should your CROS device stop working after coming into contact with water – refer to the trouble-shooting steps in chapter 19.**
Daily
Inspect the earpiece for earwax and moisture deposits and clean the surfaces with a lint-free cloth. Never use cleaning agents such as household detergents, soap, etc. for cleaning your hearing aid. It is not recommended to rinse with water. If you need to clean your CROS device intensively, ask your hearing care professional for advice and information on filters or drying capsules.

Weekly
Clean the earpiece with a soft, damp cloth or with a special cleaning cloth for CROS device. For more in depth maintenance instructions or for more than basic cleaning, please see your hearing care professional.
15. Exchanging the earpiece from the tube

15.1 Removing the earpiece from the tube

1.
Remove the earpiece from the speaker dummy by holding the tube in one hand and the earpiece in the other.

2.
Gently pull off the earpiece to remove.

3.
Clean the speaker dummy with a lint-free cloth.
15.2 Attaching the earpiece to the tube

1. Hold the speaker dummy in one hand and the earpiece in the other.

2. Slide the earpiece over the tube.

3. The speaker dummy and the earpiece should fit perfectly together.
16. Service and warranty

Local warranty
Please ask the hearing care professional, where you purchased your CROS device, about the terms of the local warranty.

International warranty
Sonova AG offers a one year limited international warranty, valid starting from the date of purchase. This limited warranty covers manufacturing and material defects in the CROS device itself, but not accessories such as batteries, tubes, earpieces, external receivers. The warranty only comes into force if a proof of purchase is shown.

The international warranty does not affect any legal rights that you might have under applicable national legislation governing sale of consumer goods.
Warranty limitation
This warranty does not cover damage from improper handling or care, exposure to chemicals or undue stress. Damage caused by third parties or non-authorized service centers renders the warranty null and void. This warranty does not include any services performed by a hearing care professional in their office.

Serial number (left side):  Authorized hearing care professional (stamp/signature):

Serial number (right side):

Date of purchase:
17. Compliance information

Europe:

Declaration of Conformity
Hereby Sonova AG declares that this product meets the requirements of the Medical Devices Regulation (EU) 2017/745 as well as the Radio Equipment Directive 2014/53/EU. The full text of the EU Declaration of Conformity can be obtained from the manufacturer or the local Phonak representative whose address can be taken from the list on www.phonak.com/us/en/certificates (Phonak worldwide locations).

Australia/New Zealand:

Indicates a device's compliance with applicable Radio Spectrum Management's (RSM) and Australian Communications and Media Authority (ACMA) regulatory arrangements for the legal sale in New Zealand and Australia. The compliance label R-NZ is for radio products supplied in the New Zealand market under conformity level A1.
The CROS model listed on page 2 is certified under:

**Phonak CROS P-13**

<table>
<thead>
<tr>
<th>Country</th>
<th><strong>FCC ID:</strong></th>
<th><strong>IC:</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td>USA</td>
<td>KWC-MZP</td>
<td></td>
</tr>
<tr>
<td>Canada</td>
<td>2262A-MZP</td>
<td></td>
</tr>
</tbody>
</table>

**Notice 1:**
This device complies with Part 15 of the FCC Rules and with RSS-210 of Industry Canada. Operation is subject to the following two conditions:
1) this device may not cause harmful interference, and
2) this device must accept any interference received, including interference that may cause undesired operation.
Notice 2:
Changes or modifications made to this device not expressly approved by Sonova AG may void the FCC authorization to operate this device.

Notice 3:
This device has been tested and found to comply with the limits for a Class B digital device, pursuant to Part 15 of the FCC Rules and ICES-003 of Industry Canada. These limits are designed to provide reasonable protection against harmful interference in a residential installation. This device generates, uses and can radiate radio frequency energy and, if not installed and used in accordance with the instructions, may cause harmful interference to radio communications. However, there is no guarantee that interference will not occur in a particular installation. If this device does cause harmful interference to radio or television reception, which can be determined by turning the equipment off and on, the user is encouraged to try to correct the interference by one or more of the following measures:
• Reorient or relocate the receiving antenna.
• Increase the separation between the device and receiver.
• Connect the device into an outlet on a circuit different from that to which the receiver is connected.
• Consult the dealer or an experienced radio/TV technician for help.
# Radio information of your wireless CROS device

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Specification</th>
</tr>
</thead>
<tbody>
<tr>
<td>Antenna type</td>
<td>Resonant loop antenna</td>
</tr>
<tr>
<td>Operation frequency</td>
<td>2.4 GHz – 2.48 GHz</td>
</tr>
<tr>
<td>Modulation</td>
<td>GFSK, GMSK</td>
</tr>
<tr>
<td>Radiated power</td>
<td>&lt; 2.5 mW</td>
</tr>
<tr>
<td><strong>Bluetooth</strong></td>
<td></td>
</tr>
<tr>
<td>Range</td>
<td>~1 m</td>
</tr>
<tr>
<td>Bluetooth</td>
<td>4.2 LE Dual-Mode</td>
</tr>
<tr>
<td>Profiles supported</td>
<td>HFP (Hands-free profile), A2DP</td>
</tr>
</tbody>
</table>
## Compliance with emission and immunity standards

<table>
<thead>
<tr>
<th>Emission standards</th>
<th>EN 60601–1-2:2015</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>IEC 60601–1-2:2014</td>
</tr>
<tr>
<td></td>
<td>EN 55011:2009+A1</td>
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<tr>
<td></td>
<td>CISPR11:2009/AMD1:2010</td>
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<td></td>
<td>CISPR22:1997</td>
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<td></td>
<td>CISPR32:2012</td>
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<td></td>
<td>ISO 7637-2:2011</td>
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<td></td>
<td>CISPR25:2016</td>
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<td></td>
<td>EN 55025:2017</td>
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<tr>
<td>Immunity standards</td>
<td>EN 60601-1-2:2015</td>
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<tr>
<td></td>
<td>IEC 60601-1-2:2014</td>
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<td></td>
<td>EN 61000-4-2:2009</td>
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<td></td>
<td>IEC 61000-4-2:2008</td>
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<tr>
<td></td>
<td>EN 61000-4-3:2006+A1+A2</td>
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<td>IEC 61000-4-3:2006+A1+A2</td>
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<td>EN 61000-4-4:2012</td>
</tr>
<tr>
<td></td>
<td>IEC 61000-4-4:2012</td>
</tr>
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<td></td>
<td>EN 61000-4-5:2014</td>
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<td>IEC 61000-4-5:2014</td>
</tr>
<tr>
<td></td>
<td>EN 61000-4-6:2014</td>
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<tr>
<td></td>
<td>IEC 61000-4-6:2013</td>
</tr>
<tr>
<td></td>
<td>EN 61000-4-8:2010</td>
</tr>
<tr>
<td></td>
<td>IEC 61000-4-8:2009</td>
</tr>
<tr>
<td></td>
<td>EN 61000-4-11:2004</td>
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<td></td>
<td>IEC 61000-4-11:2004</td>
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<td></td>
<td>IEC 60601-1 (§ 4.10.2):2005</td>
</tr>
<tr>
<td></td>
<td>ISO 7637-2:2011</td>
</tr>
</tbody>
</table>
18. Information and description of symbols

With the CE symbol, Sonova AG confirms that this product – including accessories – meets the requirements of the Medical Devices Regulation 2017/745 as well as the Radio Equipment Directive 2014/53/EU. The numbers after the CE symbol correspond to the code of certified institutions that were consulted under the above-mentioned regulation and directive.

This symbol indicates that the products described in these user instructions adhere to the requirements for an applied part of Type B of EN 60601-1. The surface of the hearing aid is specified as an applied part of Type B.

Indicates the medical device manufacturer, as defined in the Medical Device Regulation (EU) 2017/745
Indicates the date when the medical device was manufactured

Indicates the Authorized representative in the European Community. The EC REP is also the importer to the European Union.

This symbol indicates that it is important for the user to read and take into account the relevant information in these user guides.

This symbol indicates that it is important for the user to pay attention to the relevant warning notices in these user guides.

This symbol indicates that it is important for the user to pay attention to the relevant warning notices related to batteries in these user guides.

Important information for handling and product safety.
This symbol confers that the electromagnetic interference from the device is under limits approved by the US Federal Communications Commission.

Indicates a device's compliance with applicable Radio Spectrum Management's (RSM) and Australian Communications and Media Authority (ACMA) regulatory arrangements for the legal sale in New Zealand and Australia.

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<table>
<thead>
<tr>
<th><strong>Segurança</strong></th>
<th>The Compliance Identification Mark indicates that the device is in accordance with Brazilian conformity assessment requirements for equipment under health surveillance system. OCP indicates the certification body.</th>
</tr>
</thead>
<tbody>
<tr>
<td>🇯🇵</td>
<td>Japanese mark for certified radio equipment.</td>
</tr>
<tr>
<td>☑️</td>
<td>Indicates the manufacturer's serial number so that a specific medical device can be identified.</td>
</tr>
<tr>
<td>☑️</td>
<td>Indicates the manufacturer's catalogue number so that the medical device can be identified.</td>
</tr>
<tr>
<td>☑️</td>
<td>Indicates that the device is a medical device.</td>
</tr>
</tbody>
</table>
This symbol indicates that it is important for the user to read and take into account the relevant information in this user guide.

**IP68**

Ingress Protection Rating. IP68 rating indicates that the hearing aid is water and dust resistant. It survived continuous immersion in 1 meter of fresh water for 60 minutes and 8 hours in a dust chamber as per the IEC60529 standard.

Temperature during transportation and storage: 
-20° to +60° Celsius (−4° to +140° Fahrenheit).

Humidity during transportation:
Up to 90% (non condensing).
Humidity during storage:
0% to 70%, if not in use. See instruction in chapter 20.2 regarding drying the CROS device after use.
Atmospheric pressure during transportation and storage: 500 hPa to 1060 hPa

During transportation keep dry.

The symbol with the crossed-out garbage bin is to make you aware that this CROS device as well as the charger may not be thrown away as normal household waste. Please dispose of old or unused hearing aids and charger, at waste disposal sites intended for electronic waste, or give your hearing aid and charger to your hearing care professional for disposal. Proper disposal protects the environment and health.
## 19. Troubleshooting

<table>
<thead>
<tr>
<th>Problem</th>
<th>Causes</th>
</tr>
</thead>
<tbody>
<tr>
<td>CROS system not functioning</td>
<td>Dead battery</td>
</tr>
<tr>
<td></td>
<td>Speaker/earpiece of hearing aid blocked</td>
</tr>
<tr>
<td></td>
<td>Battery not inserted correctly</td>
</tr>
<tr>
<td>CROS device or hearing aid switched off</td>
<td>CROS system is in flight mode</td>
</tr>
<tr>
<td>Phone call function does not work</td>
<td>CROS system not paired to the phone</td>
</tr>
</tbody>
</table>

What to do

Change battery (chapter 5)

Clean speaker opening/earpiece of hearing aid

Insert battery correctly (chapter 5)

Switch CROS device on by completely closing battery door (chapter 4)

Open and close battery door (chapter 12.2)

Pair it to the phone (chapter 10)

If the problem persists, contact you hearing care professional for assistance.
20. Important safety information

Please read the information on the following pages before using your CROS device.

**Intended use**
The Phonak CROS device is placed on the unaidable ear and wirelessly transmits sound to the Phonak hearing aid on the other ear.

**Indications**
The device is indicated for unaidable hearing loss in one ear and better hearing in the other ear.

**Contraindications**
General clinical contraindications for the use of a CROS device are:

- Aidable hearing loss (on the intended CROS side)
- Acute tinnitus (in either ear)
- Anatomical deformity of the CROS ear (e.g. absence of the auricle)
The primary criteria for the referral of a patient for a medical or other specialist opinion and/or treatment are as follows:

- History of active drainage from the ear in the previous 90 days
- History of sudden or rapidly progressive hearing loss in one or both ears within the previous 90 days
- Acute or chronic dizziness
- Visible evidence of significant cerumen accumulation or a foreign body in the ear canal
- Pain or discomfort in the ear
- Abnormal appearance of the eardrum and ear canal such as:
  - Inflammation of the external auditory canal
  - Perforated eardrum
- Other abnormalities which the hearing care professional believes are of medical concern
The hearing care professional may decide that referral is not appropriate or in the best interest of the patient when the following applies:

- There is sufficient evidence that the condition has been fully investigated by a medical specialist and the possible treatment has been provided
- The condition has not worsened or changed significantly since the previous investigation and/or treatment

If the patient has given their informed and competent decision not to accept advice to seek a medical opinion, it is permissible to proceed to recommend an appropriate hearing aid system subject to the following considerations:

- The recommended hearing aid system will not have an adverse effect on the patient's health or general well-being
- The records confirm that the patient's best interest has been taken into consideration
If legally required, the patient must sign a disclaimer to confirm their rejection of the referral advice and that they made an informed decision.

A CROS device will not restore normal hearing and will not prevent or improve a hearing impairment resulting from organic conditions. Infrequent use of a CROS device does not permit a patient to attain full benefit from it. The use of a CROS device is only a part of the hearing habilitation and may need to be supplemented by auditory training and instruction in lipreading.

The CROS device is suitable for the home healthcare environment and due to its portability it may happen that it is used in professional healthcare facility environment like physician offices, dental offices etc.
**Intended patient population:**
This device is intended for patients with unaidable hearing loss in one ear and better hearing in the other ear, from 8 years of age.

**Intended user**
Intended for people with unaidable hearing loss in one ear and better hearing in the other ear, using the CROS device and their caregivers. A hearing care professional is responsible for adjusting the CROS device.

**Clinical benefit**
The CROS device itself does not provide a direct clinical benefit. The clinical benefit, which is improvement of speech understanding, is provided by the combination of the compatible hearing aid with the CROS device.
Side effects
Wearing your CROS device may result in undesirable physiological side effects or adverse reactions, which all may result from the shape and material of the housing and the earpiece, such as:

- Build-up of ear wax,
- Too much pressure,
- Sweat or moisture building up in the ear canal,
- Blisters, itching and/or rashes,
- Sensation of the ears being plugged or fullness,
- and their consequences (e.g. ear pain).

If any of these side effects occurs, consult your hearing care professional or physician for further advice.

Your CROS device is used in combination with a contralateral compatible hearing aid. Wearing a hearing aid may result in undesirable side effects related to its acoustic output. Please refer to the corresponding User Guide for more information.
Any serious incident that has occurred in relation to the CROS device, should be reported to the manufacturer representative and the competent authority of the state of residence. A serious incident is described as any incident that directly or indirectly led, might have led or might lead to any of the following:

a) the death of a patient, user or other person
b) the temporary or permanent serious deterioration of a patient's, user's or other person's state of health
c) a serious public health threat

To report an unexpected operation or event, please contact the manufacturer representative.
20.1 Hazard warnings

⚠️ Your CROS device operates between 2.4 GHz–2.48 GHz frequency range. When flying please check if flight operator requires devices to be switched into flight mode, see chapter 12.

⚠️ Changes or modifications to the CROS device that were not explicitly approved by Sonova AG are not permitted. Such changes may damage your ear or the CROS device.

⚠️ Do not use the device in explosive areas (mines or industrial areas with danger of explosions, oxygen-rich environments or areas where flammable anesthetics are handled). The device is not ATEX certified.

⚠️ WARNING: Device batteries are hazardous and can cause serious injuries if they are swallowed or placed inside any part of the body, whether the battery is used or new! Keep out of the reach of children, individuals with cognitive impairment, or pets. If you suspect that a battery was swallowed or placed inside any part of the body, immediately consult your physician without any delay!
⚠ If you feel pain in or behind your ear, if it is inflamed or if skin irritation and accelerated accumulations of earwax occur, please check with your hearing care professional or physician.

⚠ In very rare cases, the dome can remain in your ear canal when removing the hearing tube from the ear. In the unlikely case that the dome does get stuck in your ear canal, it is strongly recommended to see a physician for safe removal.

⚠ Hearing programs in the directional microphone mode reduce background noises. Please be aware that warning signals or noises coming from behind, e.g. cars, are partially or entirely suppressed.

⚠ During streaming of phone calls or music to the hearing aid, the signal from the CROS device is no longer transmitted to the hearing aid, which may result in unawareness of acoustical situations indicating danger.
⚠ This CROS device is not for children below 36 months. The usage of this device by children and individuals with cognitive impairment should be supervised at all times to ensure their safety. The CROS device is a small device and contains small parts. Do not leave children and individuals with cognitive impairment unsupervised with this CROS device. If swallowed, consult a physician or hospital immediately as the CROS device or its parts can cause choking!

⚠ The following is only applicable for persons with active implantable medical devices (i.e. pacemakers, defibrillators, etc.):

- Keep the wireless CROS device at least 15 cm (6 inches) away from the active implant. If you experience any interference, do not use the wireless CROS devices and contact the manufacturer of the active implant. Please, note that interference can also be caused by power lines, electrostatic discharge, airport metal detectors etc.
- Keep magnets (i.e. battery handling tool, EasyPhone magnet, etc.) at least 15 cm (6 inches) away from the active implant.
Use of accessories, transducers and cables other than those specified or provided by the manufacturer of this equipment could result in increased electromagnetic emissions or decreased electromagnetic immunity of this equipment and result in improper operation.

Portable RF communications equipment (including peripherals such as antenna cables and external antennas) should be used no closer than 30 cm (12 inches) to any part of the CROS devices, including cables specified by the manufacturer. Otherwise, degradation of the performance of this equipment could result.

The CROS devices should not be fitted with domes / wax protection systems when used by clients with perforated eardrums, inflamed ear canals or otherwise exposed middle ear cavities. In these cases, we recommend the use of a classic earmold. In the unlikely case that any part of this product should remain in the ear canal, it is strongly recommended to see a physician for safe removal.
⚠️ Avoid strong physical impacts to the ear when wearing the CROS device with customized earpiece. The stability of customized earpieces is designed for normal use. A strong physical impact to the ear (e.g. during sports) may cause the customized earpiece to break. This may lead to perforation of the ear canal or eardrum.

⚠️ After mechanical stress or shock to the customized earpiece, please ensure that it is intact before placing it in the ear.

⚠️ Use of this equipment adjacent to or stacked with other equipment should be avoided because it could result in improper operation. If such use is necessary, this equipment and the other equipment should be observed to verify that they are operating normally.
20.2 Information on product safety

The CROS device is water resistant and not waterproof. It is designed to withstand normal activities and occasional accidental exposure to extreme conditions. Never immerse your hearing aid in water! It is not specifically designed for extended periods of water submersion on a continual basis, that is worn in activities such as swimming or bathing. Always remove your CROS device before these activities, as the CROS device contains sensitive electronic parts.

Never wash the microphone inputs. Doing so could cause it to lose its special acoustic features.

Protect your CROS device from heat (never leave near a window or in the car). Never use a microwave or other heating devices to dry your CROS device. Ask your hearing care professional about suitable drying methods.
The dome should be changed every three months or when it becomes stiff or brittle. This is to prevent the dome from detaching from the tube spout during insertion into or removal from the ear.

When you are not using your CROS device, leave the battery door open so that any moisture can evaporate. Make sure that you always completely dry your CROS device after use. Store the CROS device in a safe, dry and clean place.

Do not drop your CROS device! Dropping onto a hard surface can damage your CROS device.

Always use new batteries for your CROS device. In case a battery is leaking, replace it immediately with a new one to avoid any skin irritation. You can return used batteries to your hearing care professional.
The batteries used in these CROS device should not exceed 1.5 Volts. Please do not use silver-zinc or Li-ion (lithium-ion) rechargeable batteries as these may cause severe damage to your CROS device. The table in chapter 5 explains exactly which type of battery your particular CROS device require.

Remove the battery if you are not using your CROS device for a long period of time.

Special medical or dental examination including radiation described below, may adversely affect the correct functioning of your CROS device. Remove and keep them outside the examination room/area before undergoing:

- Medical or dental examination with X-ray (also CT scan).
- Medical examinations with MRI/NMRI scans, generating magnetic fields.

CROS device don't need to be removed when passing security gates (airports etc.). If X-ray is used at all, it will be in very low doses, and will not affect the CROS device.

Do not use your CROS device in areas where electronic equipment is prohibited.
Your hearing care professional:

Manufacturer:
Sonova AG
Laubisführtstrasse 28
CH-8712 Stäfa
Switzerland
www.phonak.com

Australian Sponsor:
Sonova Australia Pty Ltd
12 Inglewood Place,
Norwest NSW 2153
Australia

Sonova Deutschland GmbH
Max-Eyth-Strasse 20
70736 Fellbach-Oeffingen
Germany