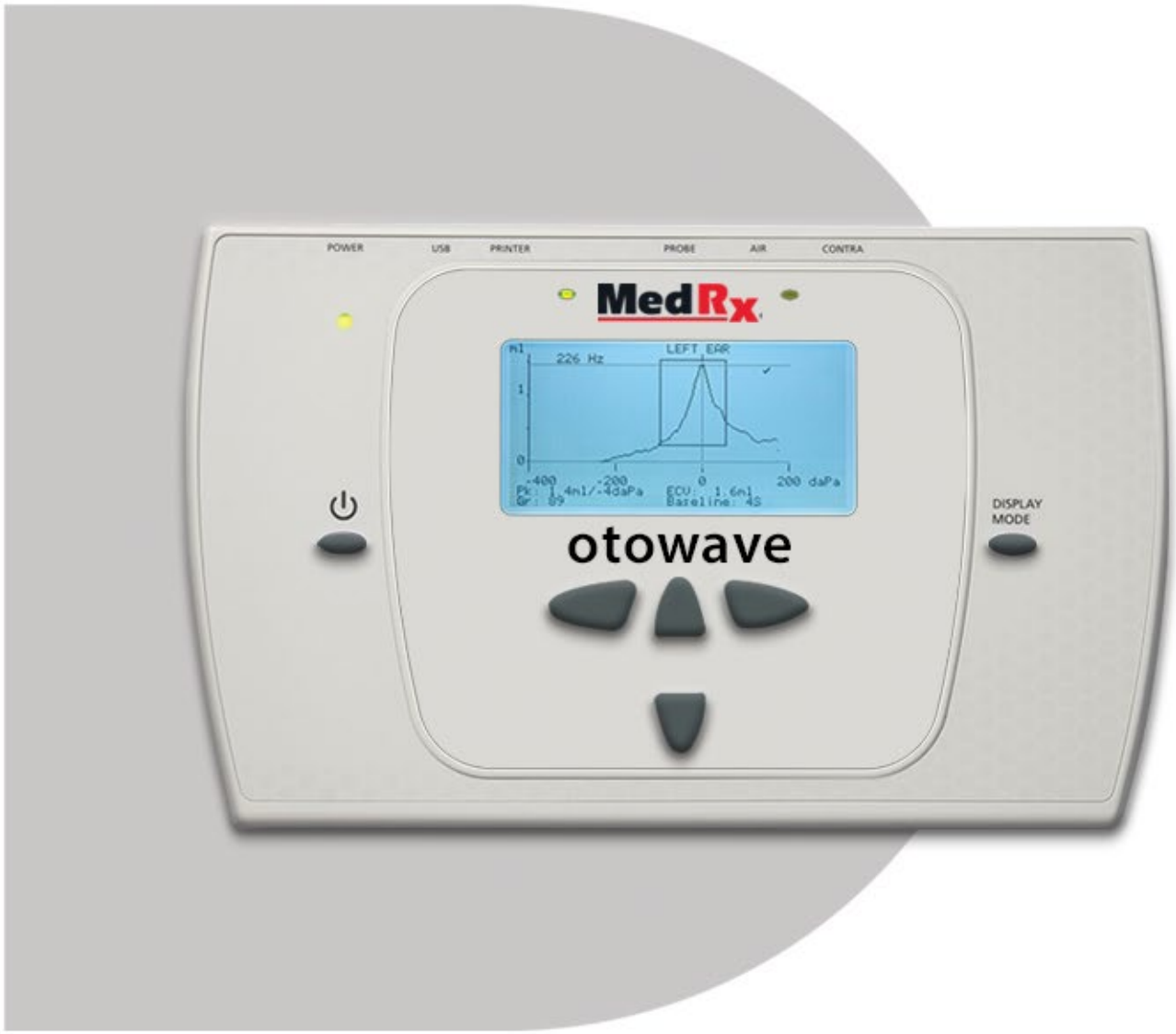

MedRx

Otowave 302 Operating Manual

READ THIS OPERATING MANUAL BEFORE ATTEMPTING TO USE THE INSTRUMENT.

This manual is valid for the Otowave 302 (applies from firmware version 1.0.0.085300 onwards – see System Information screen).

CE
0123
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INTRODUCTION

1.1. THANK YOU

Thank you for purchasing the MedRx Otowave 302, a desktop controlled impedance meter that will give many years of reliable service if treated with care.

1.2. INTENDED APPLICATIONS

This instrument is designed for use by trained personnel only, such as audiologists, ENT surgeons, doctors, general practitioners, hearing aid dispensers, child health professionals and hearing healthcare professionals with a similar level of education. It is not recommended to use the equipment without the necessary knowledge and training.

The Otowave 302 is to be used to obtain information on medical and functional conditions of the middle and outer ear as well as to assess hearing functions throughout acoustic reflex testing. The Otowave 302 can be used on all ages and performs the following kinds of measurements:

- **Tympanometry:** To measure the compliance of the tympanic membrane and middle ear at 226 Hz over a range of pressures.
- **Acoustic Reflex Testing:** The Otowave measures both ipsilateral and contralateral acoustical reflexes. The tests are performed either at ambient or peak pressure, based on the outcome of the tympanometry.

1.3. CONTRAINDICATIONS

Always visually inspect the outer ear and the external auditory canal for abnormalities before testing. Testing should not be performed on patients in the following items is applicable.

- Acute external auditory canal trauma
- Discomfort (e.g. severe otitis externa)
- Occlusion of the external auditory canal
- Discharging ear
- Recent stapedectomy or middle ear surgery
- Presence of tinnitus, hyperacusis or other sensitivity to loud sounds may contraindicate testing when high intensity stimuli are used

1.4. STANDARD AND OPTIONAL COMPONENTS

STANDARD COMPONENTS			
Otowave 302 Tympanometer	8528962	Contralateral reflex transducer (probe tip & earpiece lead)	8502177[1]
Approved power supply adapter UES12LCP	8512734	Set of disposable ear-tips	8029344[1]
USB with Software (MedRx Studio software and Noah impedance module) and Training Manuals	8514532	4 in 1 cavity assembly (0.2 ml/0.5 ml/2.0 ml/5.0 ml)	8011362
Cable USB a to USB B (2.0 m)	8011241	Carrying case	8507857
Calibration certificate			

OPTIONAL COMPONENTS			
Sanibel MPT-II Thermal Printer	8503007	Probe tip	8002592[1]
Thermal Printer paper for Sanibel MPT-II	8029305	Seal (in probe tip)	8002009[1]
Additional sets of ear tips			

OTHER COMPONENTS TO REORDER			
Printer cable – Otowave to Sanibel MPT-II	8004419	Power supply for printer	
Probe tip for contralateral reflex transducer	8001118[1]	Earpiece lead for contralateral reflex transducer	8004447
Otowave probe assembly	8502005[1]	Seal (in probe tip)	8002009[1]
Ear tip Otowave 3-5mm (pack of 25)	8012963	Ear tip Otowave 4-7mm (pack of 25)	8012965
Ear tip Otowave 7mm (pack of 25)	8013001	Ear tip Otowave 8mm (pack of 25)	8013003
Ear tip Otowave 9mm (pack of 25)	8002020	Ear tip Otowave 10mm (pack of 25)	8002021
Ear tip Otowave 11mm (pack of 25)	8002022	Ear tip Otowave 12mm (pack of 25)	8002023
Ear tip Otowave 13mm (pack of 25)	8002024	Ear tip Otowave 14mm (pack of 25)	8002025
Ear tip Otowave 15mm (pack of 25)	8002026	Ear tip Otowave 19mm (pack of 25)	8002027

[1] Applied part as according to IEC 60601-1

1.5. GUARANTEE

All the manufacture instruments are guaranteed against faulty materials and manufacture. The instrument will be repaired free of charge for a period of three years from the date of dispatch if returned, postage paid, to the MedRx service department. Return postage is free of charge for US customers and chargeable for overseas customers.

WARNING

The following exceptions apply:

- The pressure pump and transducers may go out of calibration due to rough handling or impact (dropping)
- The lifetime of probe, probe seals and ear tips is dependent upon conditions of use. These parts are only guaranteed against faulty materials or manufacture.

1.6. WARNINGS

Throughout this manual, the following meanings of warnings and cautions apply:

WARNING

The WARNING label identifies conditions or practices that may present danger to the patient and/or user.

CAUTION

The CAUTION label identifies conditions or practices that could result in damage to the equipment.

2. UNPACKING AND INSTALLATION

2.1. GENERAL

Please check the contents of the shipping carton against the delivery note to make sure that all items ordered have been included. If anything is missing, please contact the distributor who supplied the Tympanometer or MedRx, Inc. if purchased directly.

Please retain the carton and packaging as the instrument will need calibrating on an annual basis and should be returned to MedRx, Inc. in its original shipping carton.

CAUTION

For supply in US only: Federal Law restricts this device to sale by or on the order of a licensed medical professional.

2.2. MARKINGS

The following markings can be found:

Symbol	Explanation
	**Definition:** Identifies the control by means of which the instrument is switched on from (or returned to) a standby condition. A long press to turn enter standby. A short press to wake the device from standby.
	Type B applied parts. According to IEC 60601-1. Patient applied parts that are not conductive and can be immediately released from the patient.
	Refer to instruction manual.
	WEEE (EU-directive) This symbol indicates that when the end-user wishes to discard this product, it must be sent to separate collection facilities for recovery and recycling. Failing to do so may endanger the environment.
	Serial number.
	Date of manufacture.
	Manufacturer.
	The output from the power supply AC adapter is Direct Current.
	Do not re-use. Ear-tips and similar are for single use only.
	Keep dry.
	Transport and storage humidity range.
	Transport and storage temperature range.
	Medical device
	Logo

2.3. SAFETY INSTRUCTIONS

2.3.1. GENERAL

The following safety precautions must be observed at all times. General safety precautions must be followed when operating electrical equipment. Failure to observe these precautions could result in damage to the equipment and injury to the operator or patient.

The employer should instruct each employee in the recognition and avoidance of unsafe conditions and the regulations applicable to his or her work environment to control or eliminate any hazards or other exposure to illness or injury.

MedRx Inc. is aware that safety rules within individual organizations vary. If a conflict exists between the instructions in this manual and the rules of the organization using this instrument, the more stringent rules should take precedence.

The Otowave 302 is intended to be used by hearing healthcare professionals (i.e. ENT doctors, audiologists), nurses or technicians who have been trained in the proper use of the device.

2.3.2. CAUTIONS – GENERAL

CAUTION

If the system is not functioning properly, do not operate it until all necessary repairs are made and the unit is tested and calibrated for proper functioning in accordance with MedRx, Inc. specifications.

Do not drop or in any other way cause undue impact to this device. If the instrument is damaged, return it to the manufacturer for repair and/or calibration. Do not use the instrument if any damage is suspected.

This product and its components will perform reliably only when operated and maintained in accordance with the instructions contained in this manual, accompanying labels, and/or inserts. A defective product should not be used. Make sure all connections to external accessories are secured properly. Parts which may be broken or missing or are visibly worn, distorted, or contaminated should be replaced immediately with clean, genuine replacement parts manufactured by or available from MedRx, Inc .

Equipment is not user repairable. Repairs must be performed by an authorized service representative only. No modifications of the equipment are allowed by anyone other than a qualified MedRx, Inc. representative. Modification of the equipment could be hazardous.

No parts of the equipment can be serviced or maintained while in use with the patient.

Connect only accessories purchased from MedRx, Inc . to the Otowave 302. Only accessories which have been stated by MedRx, Inc. to be compatible are allowed to be connected to the device or cradle.

To comply with the standards IEC 60601-1 for safety and IEC 60601-1-2 for electromagnetic compatibility (EMC) the Tympanometer is designed to be used only with the medically-approved power supply adapter supplied, which is specified as part of the equipment. **Do not use any other type of power supply adapter with this instrument.**

The output from the power supply adapter is fitted with electronic circuit protection. In case of overload the adapter will shut down and the indicator will be extinguished. When the fault is cleared the adapter will operate as normal.

The input to the power supply adapter is protected with a non-replaceable fuse. If this fails the adapter will not operate.

The power supply adapter is the power supply disconnect device and therefore the Tympanometer should be positioned such that easy access to the power supply adapter is possible.

2.3.3. ENVIRONMENTAL FACTORS

CAUTION

Use and store the instrument indoors only. It is recommended that the instrument be operated within an ambient temperature range of 15 °C / 59 °F to 35 °C / 95 °F and in relative humidity between 30 % and 90 % (non-condensing).

Do not use the device in the presence of fluid that can come into contact with any of the electronic components or wiring. Should the user suspect fluids have contacted the system components or accessories, the unit should not be used until deemed safe by an authorized service technician.

As with all instruments of this nature the measurements taken will be influenced by significant changes in altitude & pressure. The Otowave 302 tympanometer must be re-calibrated (for volume measurement only) at the intended operating elevation if it is to be used at elevations greater than 800m above mean sea level. This applies to volume measurements up to 2.0ml maximum. Please refer to the service manual for more information.

2.3.4. ELECTRICAL AND ELECTROSTATIC SAFETY

CAUTION

Before performing any service to the insert earphones you must uncouple the Otowave 302 transducers from the patient.

WARNING

Do not touch the contacts on the back of the instrument and the patient at the same time. The consequence could be a too high leakage current to the patient.

Do not open the case of the instrument. Refer servicing to qualified personnel.

This equipment is intended to be connected to other equipment thus forming a Medical Electrical System.

External equipment intended for connection to signal input, signal output or other connectors must comply with the standard IEC 60601-1 for medical electrical equipment.

If in doubt, contact a qualified medical technician or your local representative. When the instrument is connected to a PC, or other similar items, beware of not touching the PC and patient simultaneously.

Do not use any additional multiple socket-outlet or extension cord. Use only FW7660M/05 Power Supply.

2.3.5. ELECTROMAGNETIC COMPATIBILITY (EMC)

CAUTION

Although the instrument fulfills the relevant EMC requirements, precautions should be taken to avoid unnecessary exposure to electromagnetic fields, e.g., from mobile phones, etc. If the device is used adjacent to other equipment it must be observed that no mutual disturbance appears. Please also refer to the appendix regarding EMC.

2.3.6. EXPLOSION HAZARDS

WARNING

Risk of explosion.

Do not use in the presence of flammable anesthetics or other gases.

Do NOT use in the presence of flammable gaseous mixtures. Users should consider the possibility of explosions or fire when using this device in close proximity to flammable anesthetic gases.

Do NOT use the Otowave 302 in a highly oxygen-enriched environment, such as a hyperbaric chamber, oxygen tent, etc.

2.3.7. MEASURING SECURITY

To guarantee that the Otowave 302 works properly, the instrument should be checked and calibrated at least once a year. The transducers supplied with the tympanometer are specifically calibrated with it; if these transducers are changed recalibration will be required.

The service and calibration must be performed by an authorized service technician. If these checks are not performed, EU Medical Device Regulation (MDR) and other regulations may be violated and warranties may be void.

The use of non-calibrated devices can lead to incorrect test results and is not advisable.

2.3.8. IT-NETWORK

CAUTION

Please note that connecting the device to a PC implies connecting the device to an IT-network. The connection to an IT-network may result in previously un-identified risks which must be identified, analysed, evaluated and mitigated by the responsible organisation.

Any change to the IT-network (network configuration, (dis)connection of items, update or upgrade of equipment) may introduce new risks that require additional analysis.

2.3.9. MISCELLANEOUS

Please note: DO NOT connect the Otowave 302 hardware to the computer before the software has been installed!

Storage in temperatures below 0°C /32°F and above 50°C /122°F may cause permanent damage to the instrument and its accessories.

Do not place the instrument next to a heat source of any kind.

Great care should be exercised when handling transducers, as rough handling, for example dropping onto a hard surface may break or damage the parts.

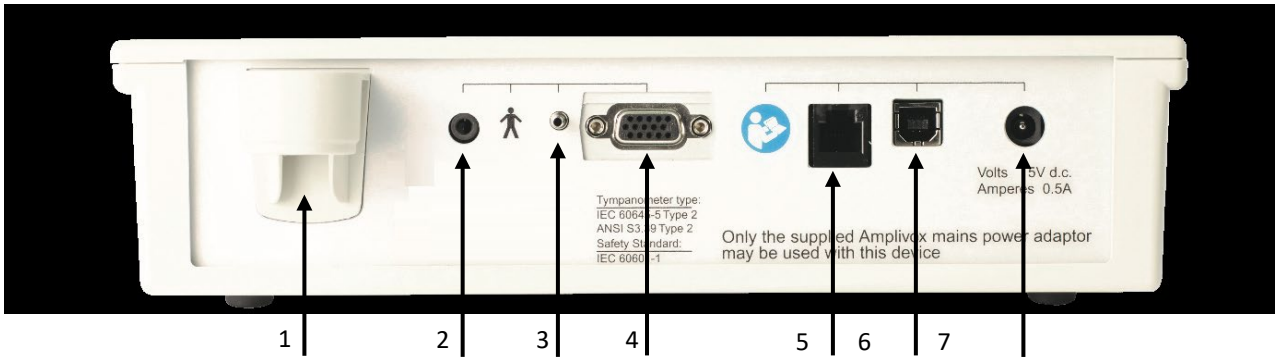
Within the European Union it is illegal to dispose of electrical and electronic waste as unsorted municipal waste. Electrical and electronic waste may contain hazardous substances and therefore have to be disposed of separately. Such products will be marked with the crossed-out wheelie-bin image shown to the left. User cooperation is important in order to ensure a high level of reuse and recycling of electrical and electronic waste. Failure to recycle such waste products in an appropriate way may endanger the environment and consequently the health of human beings.
Outside the European Union, local regulations should be followed when disposing of the product after its end of life.

2.3.10. USE OF EQUIPMENT AFTER TRANSPORT AND STORAGE

Please make sure that the instrument is functioning correctly before use. If the instrument has been stored in a cold environment (even for short period of time), please allow the instrument to become acclimatized. This can take long time depending on the conditions (such as environmental humidity). You can reduce the condensation by storing the instrument in its original packaging. If the instrument is stored under warmer conditions than the actual use conditions no special precaution is required before use. Always ensure proper operation of the instrument by following routine check procedures for audiometric equipment.

2.4. CONNECTIONS

All connections are made to the rear panel of the tympanometer as shown below.

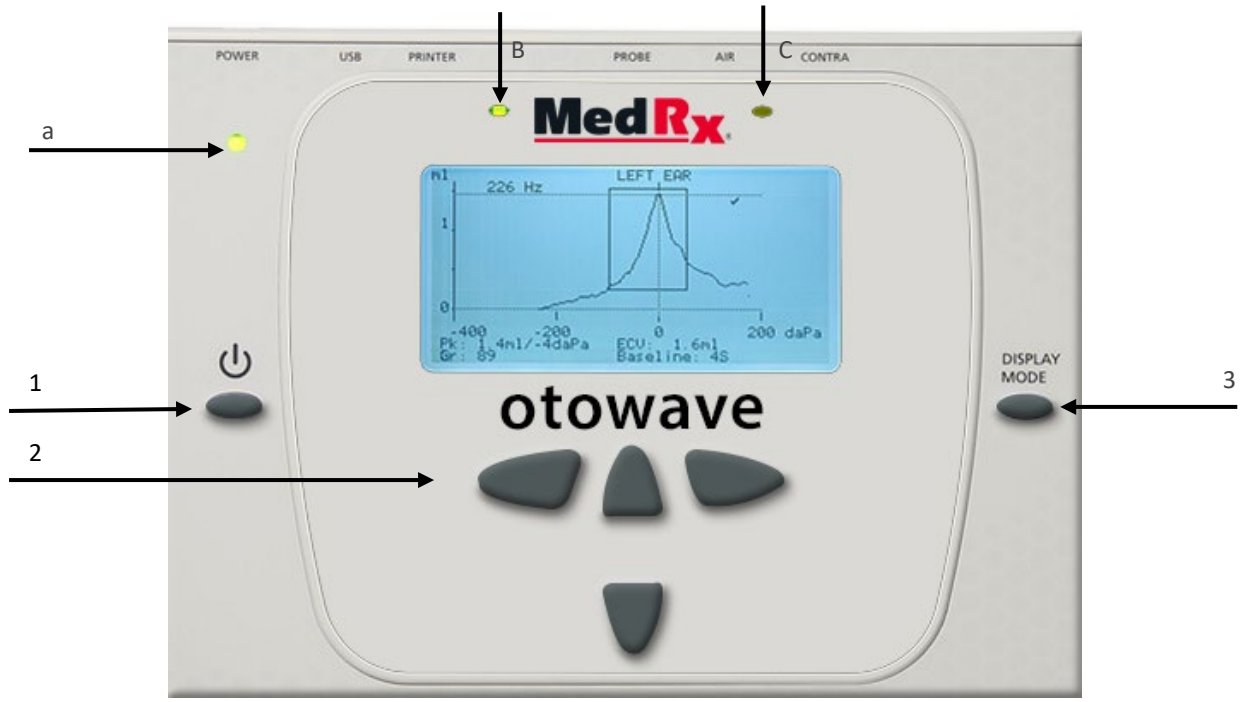


1	**Probe holder**	Cavity to store probe while not in use	
2	**Contra phone**	Contralateral transducer	3.5mm jack
3	**Air channel**	Connection of probe (pressure)	Push-fit connector
4	**Probe**	Connection of probe (electrical)	15-way D connector
5	**Printer**	Connection of external printer	RJ12 socket (6-way)
6	**USB**	Connection for computer (via USB port)	USB Connector, Type B
7	**Power**	Power supply AC/DC Adapter	2.5mm power jack

Please note: Only connect the accessories supplied with the instrument or supplied by MedRx, Inc or an MedRx, Inc distributor. These parts have been tested for use with the MedRx, Inc Otowave 302 Tympanometer for compliance with the standards IEC 60601-1 and IEC 60601-1-2. The use of accessories other than those specified may compromise compliance with these standards.

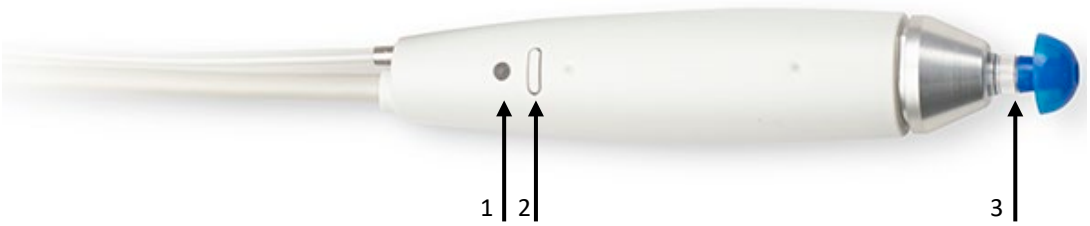
2.5. CONTROLS AND INDICATORS (BASE UNIT)

The Otowave consists of an LCD screen, three button groups in total to operate the instrument and three status LEDs.



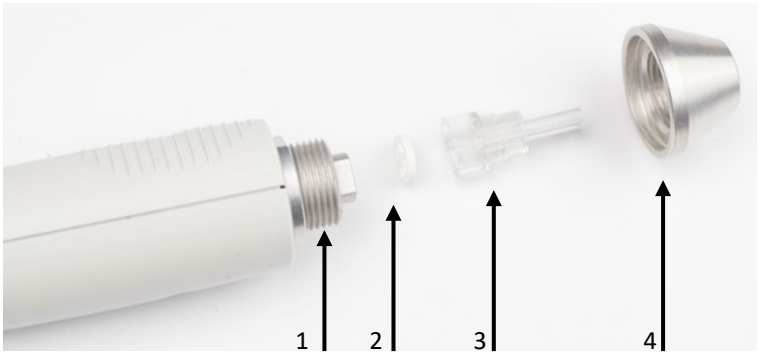
a	**Power indicator (LED a)**	Lights up as soon as the instrument is powered through the power supply adapter (also when the instrument is switched off).
b	**Indicator LED b**	Indicates if testing is in process or not.
c	**Indicator LED c**	Indicates if testing is in process or not.
1	**On/Off switch**	Short press to switch on the device, long press to switch it off.
2	**Navigation keys**	• Press the up ▲ and down ▼ navigation keys to scroll through the menus or set values • Press the right navigation key ► to accept a menu choice or go to the next step. • Press the left navigation key ◄ to cancel an operation or go back to the previous step. • The function of the left and right keys is usually shown on the bottom line of the display.
3	**Display Mode**	Quick view of the test settings currently used or change of baseline mode.

2.6. CONTROLS AND INDICATORS (PROBE)



1	**Indicator light**	Indicates if testing is in process or not.
2	**Function button**	Quick view of the test settings currently used or change of baseline mode.
3	**Probe tip with ear tip**	

2.6.1. THE PROBE HEAD



1	**Boss and Nut**	Connection on probe body for attaching nose cone
2	**Seal rubber**	Gasket used to ensure air flow
3	**Probe tip**	Transparent probe tip housing the seal rubber
4	**Nose cone**	Top part of probe to securely fasten probe tip and seal rubber

To remove the probe tip, unscrew the nose cone and remove the probe tip from the boss. A small seal will be found in the base of the probe tip. This should be examined and replaced if it is blocked or damaged. Do not remove the nut securing the boss to the probe body.

When replacing the probe tip, ensure that the seal is correctly located with the flat side aligned with the flat side within the base of the probe tip. Push the probe tip over the boss and replace the nose cone. Make sure that the nose cone is screwed home firmly but do not over-tighten. Do not use any tools to tighten the nose cone.

After replacing the tip, a Daily Check should be carried out (refer to chapter 4.5).

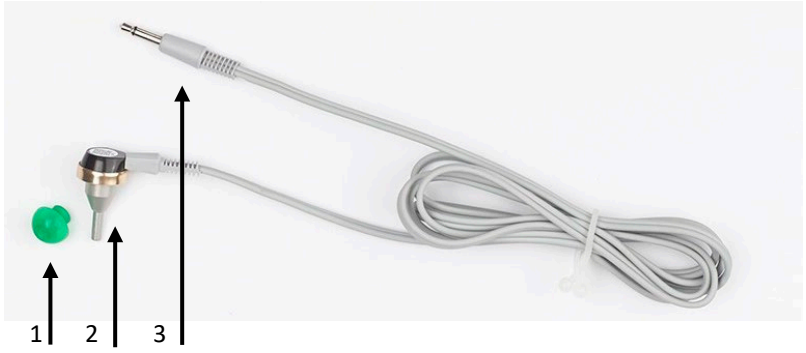
Video available on how to clean the probe tip.

2.7. LIGHT INDICATORS

The indicators on the Otowave and the probe show the status of the system.

STATUS	LED B	LED C	PROBE
Otowave turned off	Off	Off	Off
Idle, test completed or test cancelled	On	Off	Flashing (fast)
Insert probe or remove probe (refer to display for details)	Flashing (fast)	Flashing (fast)	Color alternating (Green / Yellow)
Ensure probe is held steady while an ear seal is obtained	Off	Flashing (slow)	Yellow flashing (slow)
Testing - tympanogram and/or reflex measurement	Flashing (slow)	Off	Green flashing (slow)

2.8. CONTRALATERAL TRANSDUCER



1	**Ear tip**	Ear tip to be placed on probe tip of contra phone
2	**Probe tip**	Probe tip screwed onto contra phone
3	**Plug**	Connector to **CONTRA** socket on Otowave

The contralateral transducer is used when it is required to provide a reflex stimulus to the opposite ear to that being tested with the main probe assembly. For use it should be connected to the **CONTRA** socket on the base unit and fitted with a new ear tip.

The contralateral probe tip may be replaced if necessary (e.g. if damaged). To remove the contralateral probe tip, carefully unscrew it from the body of the transducer. Carefully fit the replacement part and make sure that it is screwed home firmly but do not over-tighten. Do not use any tools to tighten the contralateral probe tip.

2.9. CHOOSING THE CORRECT EAR TIP SIZE

Video available on how to choose the correct ear tip.

The probe tip must be fitted with a new ear tip before it is presented to a patient's ear canal. If a contralateral reflex stimulus is to be applied, fit a new ear tip to the contralateral transducer before presenting it to the patient's opposite ear canal.

Please note: Never insert the probe or the contralateral transducer into a patient's ear canal without a suitable ear tip fitted.

The ear tip size is chosen based on the diameter of the external ear canal and should suit the patient's ear but also provide a comfortable pressure seal.

Ensure that the ear tip is pushed all the way down on the probe tip and that there is no gab between probe tip and ear tip.

The small holes through the Otowave probe tip must be kept clear. If these become blocked a warning message will be displayed. The probe tip must be cleaned or removed and replaced.

2.10. HARDWARE INSTALLATION

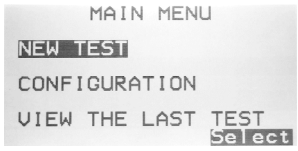
The instrument is shipped with the probe attached to the Otowave. The instrument is designed for continuous operation and is powered by a power supply adapter. Connect the output lead from the adapter into the **POWER** input socket on the back of the Otowave. When power, the indicator on the adapter and the **Power indicator (LED a)** on the Otowave will both illuminate green, showing that the instrument is ready for use.

If a contralateral transducer was purchased for contralateral reflex testing, connect the transducer to the **CONTRA socket (2)** on the base unit and fitted.

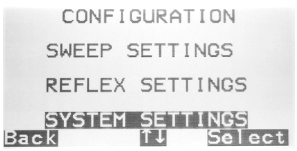
2.11. INITIAL SETTINGS

2.11.1. OPERATING LANGUAGE

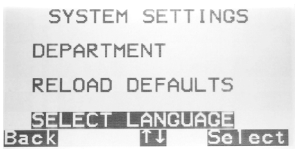
The instrument will be set in English by default. To change the operating language (English, German, French, Spanish, Portuguese or Italian) ensure you start from the **MAIN MENU.**



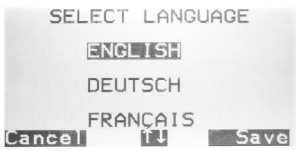
MAIN MENU
Select down ▼ 1x.
Confirm with ►.



CONFIGURATION
Select down ▼ 2x.
Confirm with ►.



SYSTEM SETTINGS
Select down ▼ 8x.
Confirm with ►.



SELECT LANGUAGE
Select language with ▲ / ▼.
Store with ►.

2.11.2. DATE AND TIME

The Otowave is equipped with a real-time clock. Before use, please set the date & time to local values in order to ensure that test data and calibration status are correctly identified.

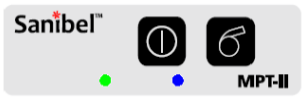
2.12. THE MPT-II PRINTER

2.12.1. INSTALLING THE MPT-II PRINTER

The Sanibel MPT-II thermal printer is available as an option for use with the Otowave 302 and is connected using the cable supplied. The printer may be specified when ordering and only this printer should be used. It will be correctly configured for use.

1. Open the lid by pushing on the sides, insert paper roll as shown, and close the lid.
2. Insert the battery.

2.12.2. SWITCHING THE PRINTER ON AND OFF



Push POWER BUTTON for two seconds in order to power ON or OFF. One short beep will be heard at power ON, two short beeps at power OFF.

The green power indicator will be lit if the printer is powered by battery.

2.12.3. USING THE PRINTER

Printer self-test: While printer is powered **OFF**, press and hold **PAPER FEED** button, then press and hold **POWER BUTTON** simultaneously. When beep is heard after approx. 3 seconds, release both buttons, and a test page will print with information on current status and character samples.

Paper feed: When powered, press **PAPER FEED** button. Paper will feed as long as the button is pressed.

Connecting process:

- Connect the printer via the cable with the device
- Power on printer
- Select print option in Otowave

Please note: Do not have several printers powered on and within range while searching.

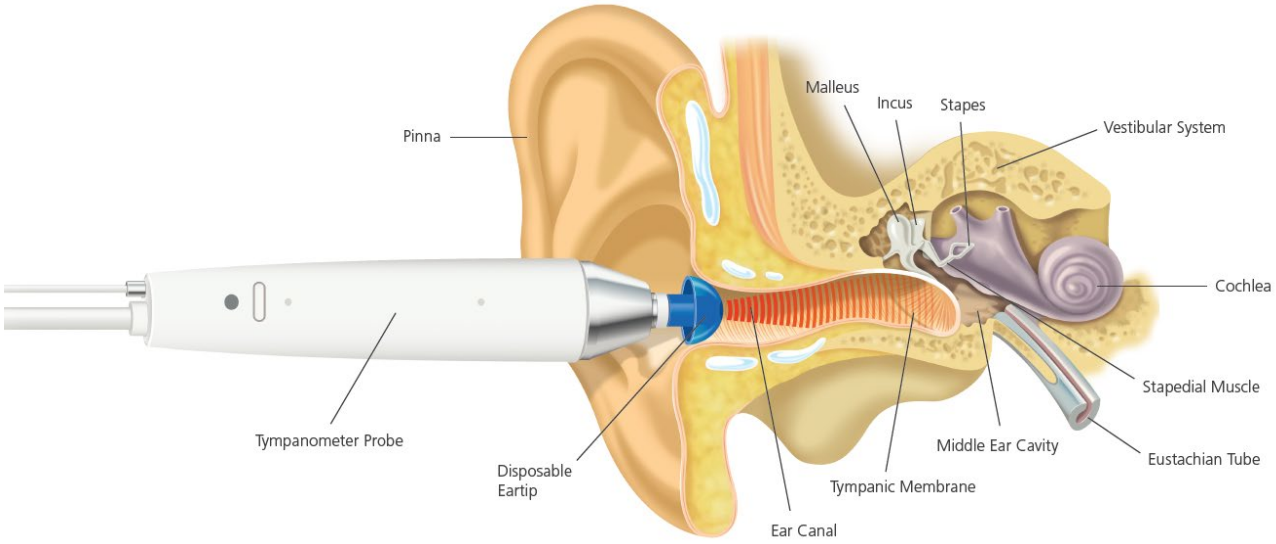
3. PRINCIPLES OF OPERATION

3.1. OTOSCOPIC EXAMINATION

A suitably-qualified health care professional should perform a thorough otoscopic examination to establish that the condition of the ear is suitable for the test options selected and that no contraindications are present. The latter would include obstruction of the external ear canal due to excessive wax and/or hairs, both of which would need to be removed. This is required to ensure that the probe tone delivered by the probe are able to reach the ear drum and are not reflected by cerumen or debris and thereby alter the test result.

3.2. PRINCIPALS OF ADMITTANCE MEASUREMENT

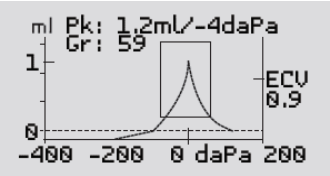
The Otowave 302 measures the admittance of the tympanic membrane and middle ear by playing a continuous tone into the ear canal at either 226 Hz. The level of this tone is calibrated to give 85 dB SP) into a 2 ml cavity. The sound level this produces in the ear canal is measured using a microphone and the admittance calculated from the result.



In line with normal audiometric practice admittance is displayed as an equivalent volume of air in ml (for 226Hz). The residual ear canal volume between the probe and the tympanic membrane is always displayed in ml.

3.3. TYMPANOGRAM

Tympanometry is part of the objective impedance test battery and provides information about the middle ear mobility and pressure in the middle ear system.



To record the tympanogram, the admittance is measured while the air pressure in the ear canal is varied from +200 daPa to -400 daPa by means of a small pump. The admittance peaks when the air pressure is the same on both sides of the tympanic membrane. The change of admittance with pressure is displayed graphically.

3.4. ACOUSTIC REFLEX MEASUREMENT

Using the same principle as in tympanometry measures, it is also possible to establish whether an acoustical reflex is present. The acoustic reflex is caused by the contraction of the stapedial muscle as a response to high-intensity stimulation of the ear. The acoustic reflex is also a natural protection of the inner ear from too high sound pressure levels and thereby damage of the hearing organ.

In acoustic reflex testing, the 226Hz tone is used to measure the admittance of the ear, while a short tone at a different frequency is presented (the reflex stimulus). The level of this stimulus is increased in steps until the stapedial muscles respond causing the tympanic membrane to become stiffer, or a preset maximum level is reached. When the change in admittance exceeds a predetermined threshold this constitutes a reflex and the change in admittance at that level when the stimulus is applied is displayed as a plot against time.

The stapedial reflex is measured at the static ear canal pressure that produces the maximum membrane admittance, so reflex measurements are taken after the tympanogram is measured when the peak admittance pressure has been established.

The reflex stimulus may be produced in the ear being measured (ipsilateral mode), the opposite ear (contralateral mode) or in both ears (ipsilateral mode followed by contralateral mode). For contralateral stimulation the reflex tone is produced in a separate transducer supplied with the instrument.

4. USING THE OTOWAVE

4.1. GENERAL PRECAUTIONS

When operating the instrument, please observe the following general precautions:

CAUTION

1. Use this device only as described in this manual.
2. Use only the disposable ear tips designed for use with this instrument.
3. Always use a new ear tip for each patient to avoid cross-contamination. The ear tips are not designed for reuse.
4. Never insert the probe tip into the ear canal without affixing an ear tip as omission may damage the patient's ear canal.
5. Keep the box of ear tips outside the reach of the patient.
6. Be sure to insert the probe tip in a way which will ensure a tight fit without causing any harm to the patient. Use of a correct and clean ear tip is mandatory.
7. Be sure to use only stimulation intensities acceptable to the patient. When presenting contralateral stimuli using the insert phones – do not insert the phones or in any way try to conduct measurements, without a correct insert ear tip in place.
8. Clean the probe and insert phones regularly using a recognized disinfectant.
9. Clean the probe tip regularly to ensure wax or other debris stuck in the probe tip does not affect the measurement.
10. Contraindications to testing include recent stapedectomy or middle ear surgery, a discharging ear, acute external auditory canal trauma, discomfort (e.g, severe otitis externa) or occlusion of the external auditory canal. Testing should not be performed on patients with such symptoms without a medical doctor's approval.
11. The presence of tinnitus, hyperacusis or other sensitivity to loud sounds may contraindicate testing when high intensity stimuli are used.

Please note:

Careful handling of the instrument whenever in contact with a patient should be given high priority. Calm and stable positioning while testing is preferred for optimal accuracy.

1. Never clean the transducer housing with water or insert non-specified instruments into the transducer.
2. Do not drop and avoid other undue impact to this device. If the instrument is dropped or in any other way damaged, return it to the manufacturer for repair and/or calibration. Do not use the instrument if any damage is suspected.

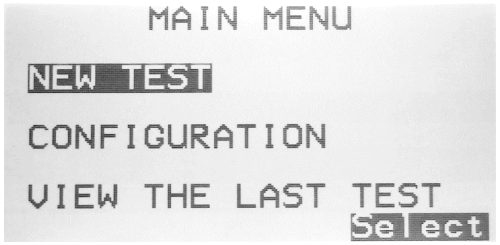
4.2. SWITCHING THE INSTRUMENT ON AND OFF

Select the on/off button for 1 s to switch the instrument on. You will see a small hourglass in the middle of the display indicating that the instrument is powering on. No warm-up time is required, although a short diagnostic routine will run for a few seconds. During this time the internal pump will operate.

To switch the instrument off again, hold the button for approx. 2 s and the instrument and LED b will turn off.

Please note: The power indicator (LED a) will light as long as the instrument is connected through the mains.

4.3. MENU DISPLAYS



When the start-up sequence is complete the **MAIN MENU** is displayed. From here, you can reach different submenus.

Use the navigation keys (up ▲ and down ▼) to scroll through the options and select submenus.

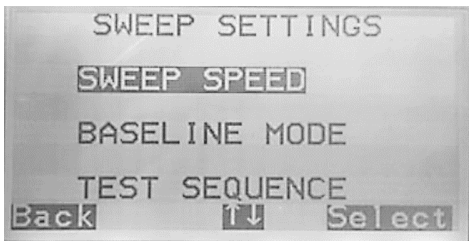
To select a submenu select the right navigation key ►.

To be brought one menu back, push the left navigation key ◄.

The following submenus can be reached from the **MAIN MENU**:

- New Test
- Configuration
- View the last Test
- Daily Check
- Data Management
- System Information

4.4. CONFIGURATION



The configuration submenu contains the following settings:

- Sweep Settings (Tympanometry only)
- Reflex Settings (Acoustic reflex only)
- System Settings (General)

4.4.1. SWEEP SETTINGS

Video available on how to use different test settings in tympanometry.

ITEM	DESCRIPTION	DEFAULT
Sweep Speed:	The rate of change of air pressure may be selected to be 100daPa/s 200daPa/s or 300daPa/s. This determines the time taken for a pressure sweep from +200 to -400 daPa (6, 3 and 2 seconds, respectively).	200 daPa/s
Baseline Mode:	The Otowave 302 can display tympanograms in a variety of graphical formats allowing the operator to choose the most appropriate for the patient under examination. **Please refer to the appendix for further information on how to use the Baseline mode**	226 Hz
Ear Seal Check:	The **STANDARD** option is adequate for most tests, although it may not always be possible to generate the extremes of pressure during a tympanogram measurement with this setting. If difficulty is experienced in using the ear tips to create a seal the alternative **EXTENDED** option may be helpful. This function checks that a range of pressures will be available before starting a test by means of a visual indication of the quality of the seal. The **EXTENDED** function is especially helpful if small ear canal volumes should not experience excessive pressure.	Standard
Defaults:	Reset the sweep settings of the selected profile to its original settings.	

4.4.2. REFLEX SETTINGS

Video available on how to add ipsilateral reflexes to test protocol.

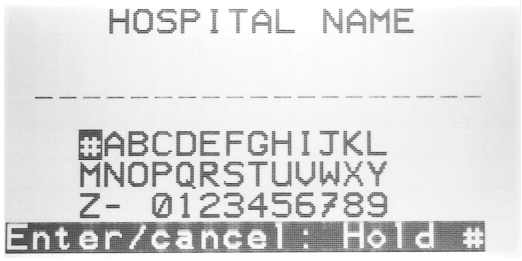
Item	Description	Default
Level Mode:	**Please note:** Depending on the **LEVEL MODE** selection, the **LEVELS** screen will contain different content. **ONE LEVEL:** Use the S keys to choose the level of reflex stimulus to apply. Only one level will be tested in the measurement. The maximum level of ipsilateral stimulus may be set to maximum 100dBHL; the maximum level of contralateral stimulus may be set to 110dBHL. **MULTILEVEL:** Use the S keys to choose the maximum level of reflex stimulus to apply and the step size between the levels of the preceding stimuli. The maximum level of ipsilateral stimulus may be set between 85dBHL and 100dBHL; the maximum level of contralateral stimulus may be set between 85dBHL and 110dBHL.	Multilevel
Sequence:	Choose the type of reflex stimulus to apply: ipsilateral only, ipsilateral followed by contralateral or contralateral only.	Ipsi
Levels:	Choose ipsilateral or contralateral and press the ► key to confirm the selection. Then use the ▲ and ▼ keys to choose the maximum level of reflex stimulus to apply and the step size between the levels of the preceding stimuli. The maximum level of ipsilateral stimulus may be set between 85dBHL and 100dBHL; the maximum level of contralateral stimulus may be set between 85dBHL and 110dBHL.	95 dB 5 dB steps
Frequencies:	Use the ▼ key to scroll through the frequencies available for each of the ipsilateral and contralateral stimuli (500Hz, 1000Hz, 2000Hz & 4000Hz), and then the ▲ key to select (✓) or deselect (-) the frequencies at which the reflex stimulus is to be applied. Then press ► to save the entire selection.	1kHz ipsi
Selection:	Choose the circumstances when a reflex measurement is to be made (always, never, only if an admittance peak is found, or only after confirmation is made at the start of the test sequence). In cases where an admittance peak has not been established a pressure of 0daPa is used.	Only if peak found
Threshold:	Choose the change in admittance required to signify that a reflex response has been detected (0.01ml to 0.5ml).	0.03 ml
Auto-Stop:	By default the reflex test at each frequency will stop at the lowest level of stimulus that produces a response. By setting Reflex Auto-Stop to **NO** the Otowave will test for a reflex at all selected levels.	Yes
Filter:	Choose either 2Hz or 1.5Hz. 2Hz is suitable for most circumstances.	2 Hz

	If a smoother reflex plot is required for better interpretation 1.5Hz may be chosen.	
Defaults:	Reset the sweep settings of the selected profile to its original settings.	

4.4.3. SYSTEM SETTINGS

ITEM	DESCRIPTION	DEFAULT
Set Time/Date:	Set the internal clock date and time. Use the ◄ and ► keys to select a field and the ▲ and ▼ keys to adjust the value.	
LCD Contrast:	Adjust the display contrast using the ▲ and ▼ keys.	
LCD Brightness:	Adjust the display brightness using the ▲ and ▼ keys.	
Report Cal. Dates:	Select **PRINT CAL. DATES** to show the serial number for the base unit and the transducers on the print-out provided by the Sanibel Thermal printer.	PRINT CAL. Dates
Set Date Format:	Set the format of how the date is displayed: DD/MM/YY or MM/DD/YY	DD/MM/YY
Hospital Name:	Allows the Hospital name to be entered. The name will appear at the top of the print out.	
Department:	Allows the Department name to be entered. The name will appear at the top of the print out.	
Reload Defaults:	Reset the instrument to its original settings.	
Select Language:	Change the operation language to English, German, French, Spanish, Portuguese or Italian.	
Defaults:	Reset the system settings of the selected profile to its original settings.	

Data entry of Hospital Name and Department:



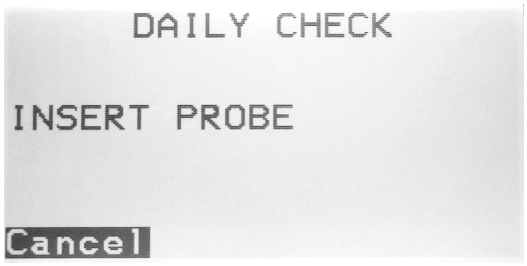
To enter the data use the ▲ ▼ ◄ and ► keys to select a character. Press and hold the ► key to enter the selected character. To delete the last character, press and hold the ◄ key.

In order to save the entry, move the selection on the **#-key** and hold the ► key. The selection will now be stored.

To cancel the entry, move the selection on the **#-key** and hold the ◄ key. You will be brought back to the **SYSTEM SETTINGS.**

4.5. DAILY CHECK

The operation of the Otowave should be checked daily using the 4-in-1-test cavity assembly supplied with the instrument.

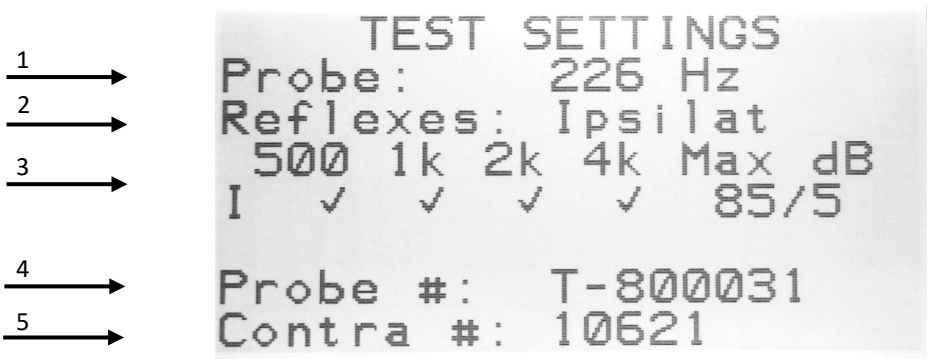


Select the **DAILY CHECK** option in the main menu and wait until **INSERT PROBE** is displayed.
Insert the probe, without an ear tip, into the hole at the 2ml end of the test cavity. Make sure that the probe is pushed fully home and is held tight against the stop. The probe must be square to the end of the test cavity.
The display should show the volume of the test cavity to within ± 0.1ml.
Remove the probe, wait until **INSERT PROBE** is displayed and repeat the test with the three remaining test cavities.
The display should show the volume of the 0.2ml, & 0.5ml test cavities to within ± 0.1ml.
The volume of the 5.0ml test cavity should be shown within ± 0.25ml.

When the checks have been completed press ◄ to return to the main menu.

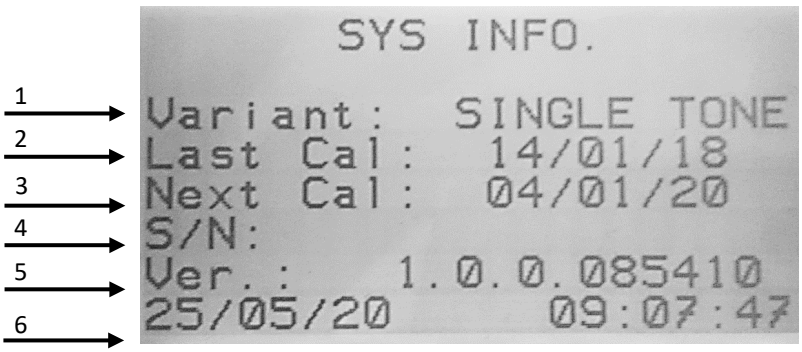
4.6. DISPLAY MODE

To view the current selected test settings ensure that the **MAIN MENU** is displayed and then press and hold the function button on the probe to display the **TEST SETTINGS** screen as shown below.



1	**Probe:**	Frequency being used for tympanometry (226)
2	**Reflexes:**	Ear sides tested (ipsilateral or contralateral).
3	**Frequencies and Level:**	Selected frequencies for reflex testing (marked with an A when selected), max. level to test and step size.
4	**Probe #:**	Serial number of probe used.
5	**Contra #:**	Serial number of contra phone calibrated to equipment.

4.7. SYSTEM INFORMATION

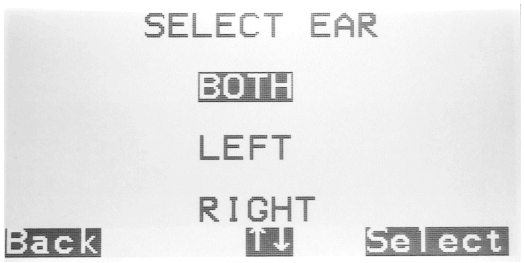


1	**Variant:**	Instrument version (Dual Tone = High Frequency option enabled)
2	**Last Cal:**	Last calibration date
3	**Next Cal:**	Next calibration date
4	**Serial No:**	Serial number of Otowave
5	**Ver.:**	Firmware version
6	**Date and Time:**	User defined date and time

4.8. PERFORMING A TEST

4.8.1. SELECTING THE EAR SIDE

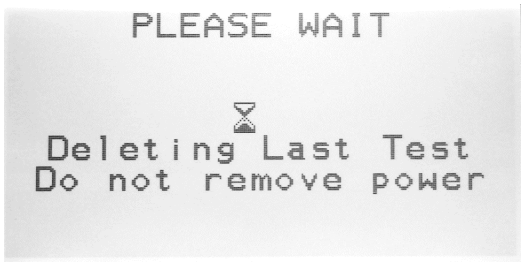
Having selected the required test settings a typical tympanogram measurement and reflex tests are carried out as follows. Highlight **NEW TEST** and push the right navigation key ► to continue with the selection.



Select the ear(s) required for test (**BOTH** signifies **LEFT** followed by **RIGHT**).

Selecting the right navigation key ► will start the test procedure.

Press ◄ at any time to cancel the test and return to the ear selection menu.

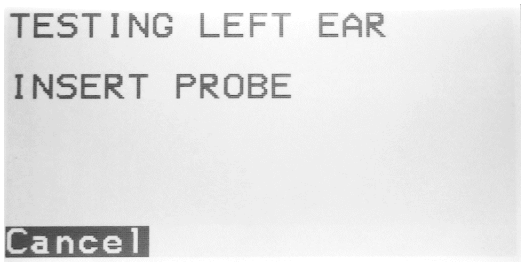


Please note: Continuing with the test procedure will delete the last recorded test results, stored in the short-term memory of the instrument.

4.8.2. PERFORMING TYMPANOMETRY

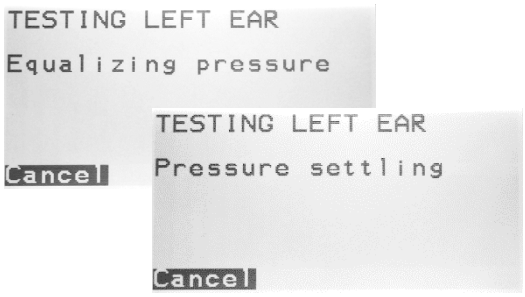
Depending on the protocol selected, the test sequence will perform both tympanometry and acoustic reflex testing in one run without removing the probe.

Please note: The protocol selected by default runs a tympanogram together with ipsilateral reflexes.



An instruction on the screen will guide you through the test sequence. The test will start automatically by inserting the probe into the patient's ear.

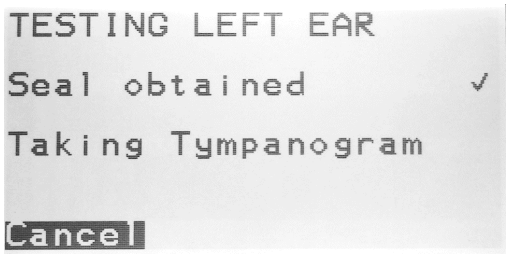
Insert the probe into the test ear. While the probe placement is evaluated, indicator LED b and c on the device will light up alternating as well as the indicator light on the probe will change it's color from yellow to green by turns to indicate the test start.



Once an adequate seal is detected the tympanogram measurement is made. This takes about 3 seconds. It is important not to move the probe and to ask the patient to remain very still during the test.

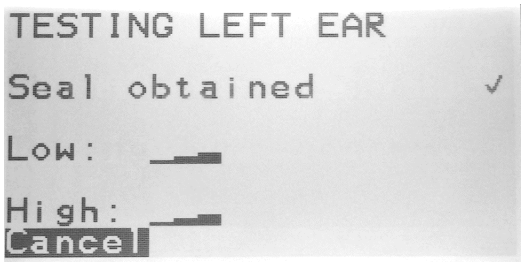
TROUBLESHOOTING:
In some cases no pressure can be build up and the test sequence will stay in the **EQUALIZE PRESSURE SCREEN**. This situation can occur for the several reasons. Solution to this issue can be found in the **TROUBLESHOOTING SECTION**.

STANDARD EAR SEAL CHECK



The default **STANDARD** option only shows if a seal could be obtained.

EXTENDED EAR SEAL CHECK



The **EXTENDED EAR SEAL CHECK** shows a number of bars indicating the robustness of the seal.

The probe should be adjusted in the ear until two or more bars are shown for **LOW** and **HIGH**.

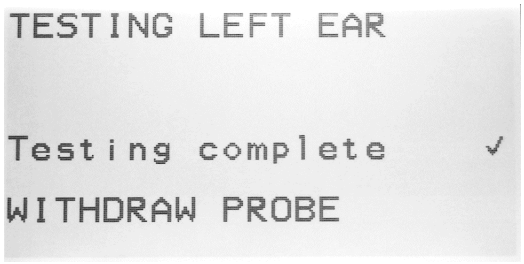
LOW: The pump is moved by a fixed distance in an attempt to reduce the pressure in the ear and is held in that position. If the measured pressure reduces by a minimum amount, and remains low (within present tolerances), the seal is considered OK.

HIGH: The process is repeated, at a pressure higher than ambient. Otherwise, the process restarts. If the higher pressure is sufficiently above the ambient pressure, and is held, the seal is good and the pump goes just above the starting pressure

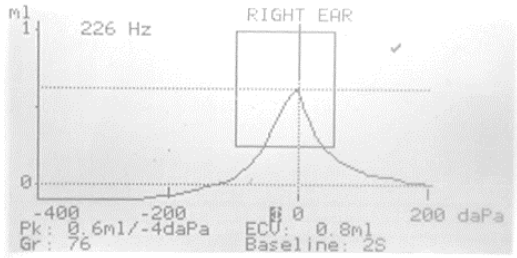
The method used for the extended ear seal check places a maximum limit on the ear canal volume of ~4.5 ml.

As soon as the pressure can be established, the tympanometry measurement will be performed.

During the actual testing phase, the indicator **LED b** on the device and the indicator light on the probe will pulse in a green light.



When the test sequence is completed, **LED b** on the device and the indicator light on the probe will continuously light in a green color. Also, the instruction on the screen will ask you to remove the probe from the patients ear.



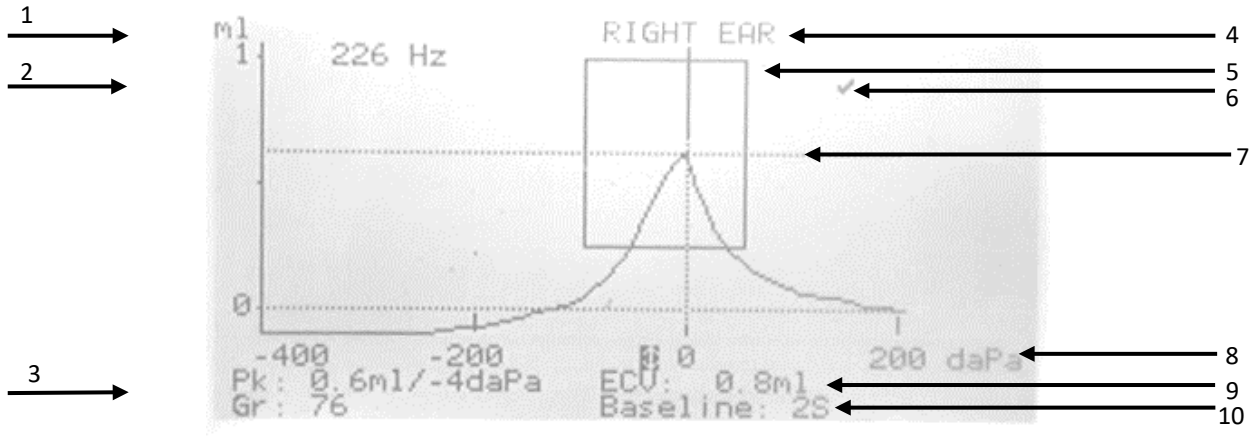
As soon as the probe is removed from the ear, the test result for the ear measured will show on the screen.

Please note: If **BOTH EARS** had been selected to be tested, the test sequence of the other ear will be continued from the test review screen. Selecting the right navigation key ► will start the test procedure on the other ear.

Selecting ◄ will cancel the procedure and return to the ear selection menu.

After the test has been successfully performed, three different actions (print, store, review) can be taken from here in order to proceed working with the data obtained. Selecting the right navigation key ► will bring you to **PROCESS RESULTS** screen from where the data can be printed, stored or reviewed again.

4.8.3. UNDERSTANDING THE TYMPANOMETRY TEST RESULT



1 y-axis label, in ml for 226 Hz.

2 y-axis, ranging from 0 to 1 ml in this example.

3 Values defining compliance curve based on cursor position **(9)** and the Baseline Mode **(5)**
- Pk (Peak): Volume in ml or mmho, pressure in daPa
- Gr (Gradient): Width of compliance curve at half of peak compliance in daPa

4 Ear side, L for left and R for right.

5 Normative box

6 Pass (a) / Refer (x) sign when tymp peak falls into normative box or not (refer).

7 Pressure cursor to be operated with up▲ and down ▼ navigation keys.

8 x-axis, default baseline offset, ranging from -400 daPa to +200daPa in this example.

9 The Ear Canal Volume (ECV) in ml measured at the Baseline Mode **(5)**.

10 Baseline Mode to show tympanogram
- **226 Hz:** Y-only compensation, 2S or 4S mode (scalar mode).

11 The test frequency [Hz]

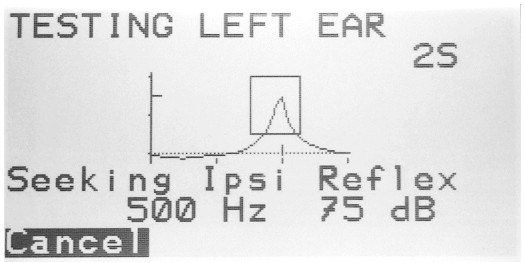
4.8.4. PERFORMING TYMPANOMETRY AND ACOUSTIC REFLEX TESTING

Video available on how to run a test.

Before starting an acoustic reflex test, a tympanometry is performed first. The tympanometry measurement is performed the same way as described in chapter 4.8.2. The tympanogram will be followed immediately by an acoustic reflex test.

Performing a tympanometry before the reflex test is recommended, so the ear canal pressure will be set to the value that gave the peak admittance during the tympanogram test.

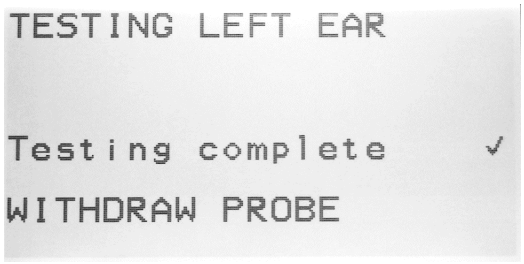
TROUBLESHOOTING:
By default, the acoustic reflex testing is only performed if a peak is found in the tympanogram. This setting can be changed in the **CONFIGURATION** menu.



The acoustic reflex test screen consists of the preview of the measured tympanogram. Underneath the graph, the tested ear side, the test frequency and intensity are listed. The reflex test starts with the lowest frequency and level selected.

The instrument will then step through the tone frequencies and levels set in the **CONFIGURATION** menu searching for a reflex response.

Please note: If contralateral testing is enabled, the ipsilateral reflexes are tested first.

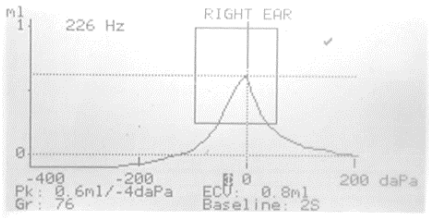


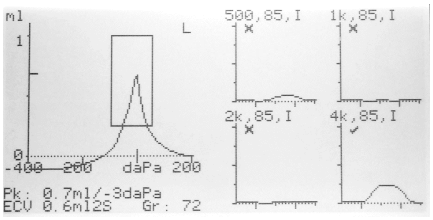
When the measurement is complete the indicator on the probe changes from flashing green to steady green. The display confirms that the test has been completed along with the instruction **WITHDRAW PROBE.**

After the test has been successfully performed, three different actions can be taken from here in order to proceed working with the data obtained. Selecting the right navigation key ► will bring you to **PROCESS RESULTS** screen from where the data can be printed, stored or reviewed again.

4.8.5. UNDERSTANDING THE ACOUSTIC REFLEX TEST RESULT

The navigation key ► and ◄ are required to navigate in the reflex result screens.





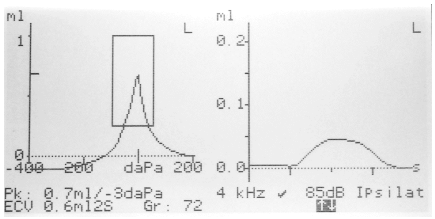
The acoustic reflex result screens will always start to show the tympanogram first.

Selecting the navigation key ► will minimize the tympanogram and show the reflex additionally (combined view).

The combined view shows a smaller tympanogram and the measured reflexes for either the threshold measured or the loudest intensity presented, if no threshold could be found.

Selecting the navigation key ◄ will show the full tympanogram view again. Selecting the right navigation key ► will show the reflex measures in more detail, depending on the **REFLEX AUTOSTART** function.

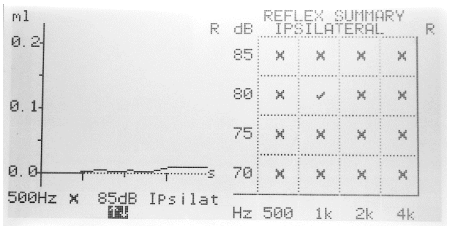
REFLEX AUTOSTART ON



If Reflex autostop has been selected the tympanogram remainson the left hand side of the screen with a larger scale trace of the reflex at the lowest selected frequency with the lowest level that gave a response.

Select the ▲ ▼ keys to move scroll through the different frequencies.

REFLEX AUTOSTART OFF



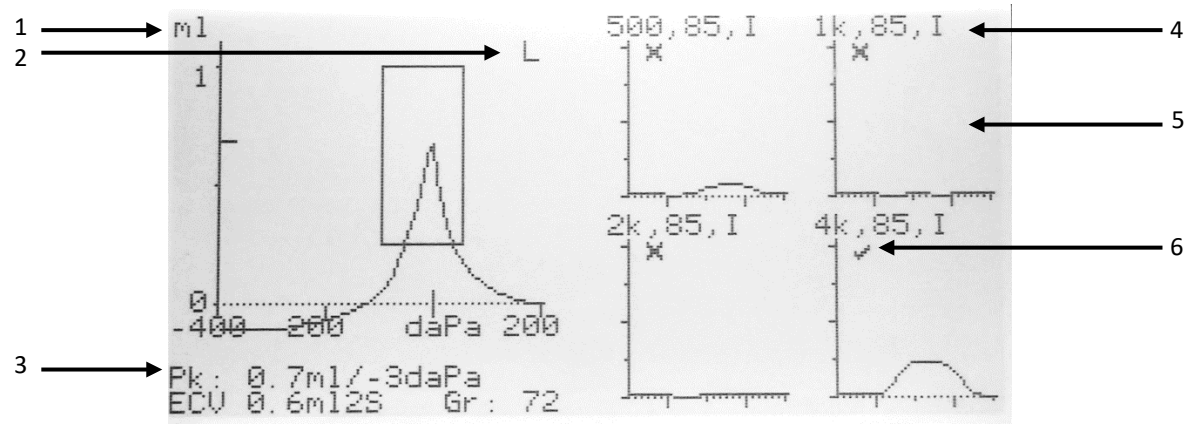
If Reflex autostop has not been selected the larger scale trace of the reflex at the lowest frequency with the lowest level that gave a response is shown on the left hand side of the screen.

A summary of the levels and frequencies at which a reflex tone was presented is shown on the right hand side of the screen along with the result of the test ("✓" if a reflex was found, otherwise "**X**"). The dash symbol "**-**" is shown if a reflex tone was not presented at the level indicated.

Please note: Depending on your firmware version, a threshold line will be shown in the reflex graph. This line is one of the criteria defining the pass/refer evaluation of the reflex result is based on.

Selecting the navigation key ◄ will show the four frequencies again. Selecting the navigation key ► will show the **PROCESS RESULTS** screen, from where the measurement outcomes can be processed.

If contralateral reflex measurements were taken pressing the ► key will display similar results for these reflexes.



1 Tympanogram on the left hand side and the four reflex traces shown on the right.

2 Ear side, L for left and R for right.

3 Values defining compliance curve based on cursor position **(9)** and the Baseline Mode **(5)**
- Pk (Peak): Volume in ml or mmho, pressure in daPa
- The Ear Canal Volume (ECV) in ml measured at the Baseline Mode **(5)**.
- Gr (Gradient): Width of compliance curve at half of peak compliance in daPa

4 Four reflex traces, containing frequency of the reflex stimulus, lowest level of tone (dBHL) at which a reflex was found and the type of reflex stimulus used (I for ipsilateral, C for contralateral).

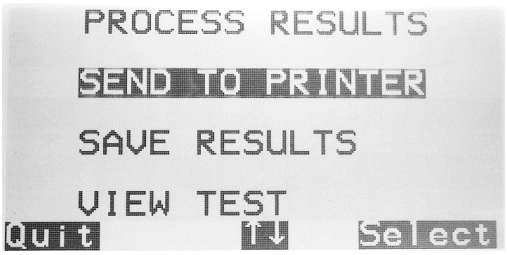
5 Grafical display of reflex. To view the reflex traces in more detail press ►. One of two displays will then be shown depending on whether or not Reflex autostop has been selected.
- **Single frequency used:** Diagrams contain the different levels tested for the specific frequency.
- **Several frequencies used:** each diagram represents 1 frequency, showing only the intensity where a reflex was detected.

6 "✓" if a reflex was found, otherwise "**X**"

4.8.6. IMMEDIATE PROCESS OF RESULTS

4.8.7. GENERAL

After a test has been finished, the data can either be sent to a printer and/or stored to the internal database of the instrument or the NOAH or the MedRx Studio software.



From the test result screen select the right navigation key ► until the **PROCESS RESULTS** screen is reached.

From here the following options are given:
- Print the current record
- Save the current record
- View the record again
- Return to the **MAIN MENU**

To be brought one menu back, push the left navigation key ◄.

4.8.8. SENDING RESULTS TO A PRINTER

The 302 can be upgraded with an option to allow connection via the supplied cable to the designated portable Sanibel MPT-II thermal printer for printing test results. Upon receipt of the printer it must be initially charged for a minimum of 15 hours prior to use.

Please note that thermal paper printouts can fade with exposure to light or heat. Consider transferring the data to a computer for permanent storage.

To print the results of the last test select **SEND TO PRINTER** from the **PROCESS RESULTS** menu on completion of the test. The printing process has to be confirmed by pressing **SELECT** again. The Otowave will then attempt to connect to the printer.

To stop the print operation (for example if a printer is not connected) press ◄ to select Cancel.

The print out consists of the three characters printed in the **NAME** field followed by the Otowave graphical displays, the analysis and the results. There is space for additional details to be handwritten by the clinician (patient's full name/age, operator & comments). Also, the name of the hospital, the department, and the calibration dates for the instrument may also be printed if required (refer to chapter 4.4.3).
After successful printing the **PROCESS RESULTS** menu is displayed.

Please note:

- When printing one test result, the print out will contain the last selected baseline mode.
- When printing several test results, the print out will contain the stored baseline mode.

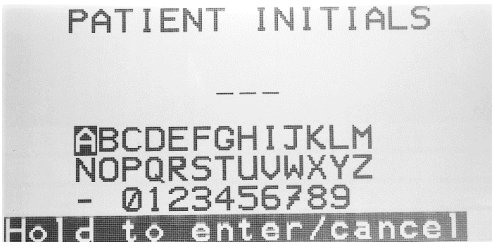
4.8.9. SAVING RESULTS TO THE INTERNAL DATABASE

Up to 36 tests can be stored in the Otowave internal database.

To save the results of a test select **SAVE RESULTS** from the **PROCESS RESULTS** menu that is displayed on completion of a test. This option can also be accessed by selecting **VIEW THE LAST TEST** from the main menu and scrolling through the results using the ► key as long as the test results have not already been saved or deleted (e.g. by starting and then aborting a new test).

A three character identifier is used for the record. This is also used as the reference for the patient's name on the printed record and for data transferred to a computer.

The identifier would typically be the patient's initials, and as the tympanometer uses a combination of this identifier and the date/time of a test to refer to stored records this same identifier may be used for different tests for the same patient.

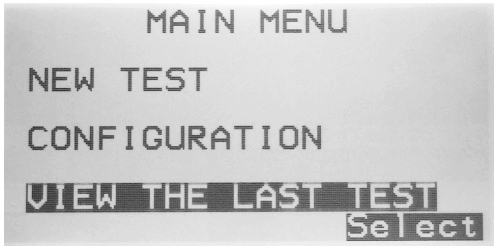


To enter the data use the ▲ ▼ ◄ and ► keys to select a character. Press and hold the ► key to enter the selected character. To delete the last character, press and hold the ◄ key.

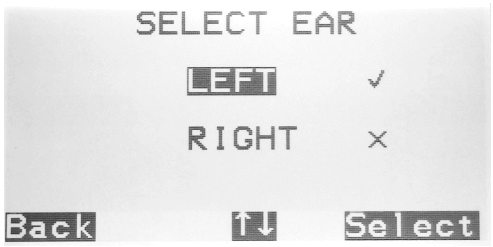
In order to save the entry ensure all three identifier characters have been entered and then hold the ► key. The selection will now be stored. You will then be brought back to the **PROCESS RESULTS** screen**.** The save record option is now removed.

To cancel the entry, delete all three characters and hold the ◄ key. You will be brought back to the **PROCESS RESULTS** screen**.**

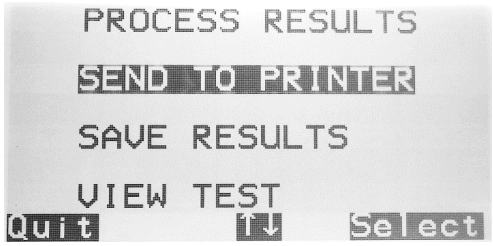
4.9. (RE)VIEW THE LAST TEST(S)



The Otowave 302 offers a short-term memory that allows you to store a test per ear until a new test has been conducted.



The symbols ✓ or **X** are used to signify whether results are available for each ear. Only test results with a ✓ can be selected for a review.



From the test result screen select the right navigation key ► until the **PROCESS RESULTS** screen is reached.

From here the following options are given:

- Print the current record
- Save the current record
- View the record again
- Return to the **MAIN MENU**

Please refer to chapter 0 for further steps on how to proceed from the PROCESS RESULTS screen.

Please note that the last test could have already been stored and still show in the short-term memory. In this case the function **SAVE** is disabled.

4.10. DATA MANAGEMENT

4.10.1. GENERAL

Video available on how to process test data.

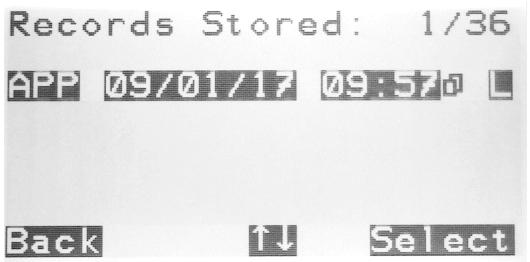
Up to 36 tests can be stored in the Otowave's internal database.

Records stored in the database of the Otowave can be listed, viewed, printed or deleted using the **DATA MANAGEMENT**.

From here the following options are given:

- Review all stored records (view, print, delete single records from here)
- Delete stored record
- View the record again

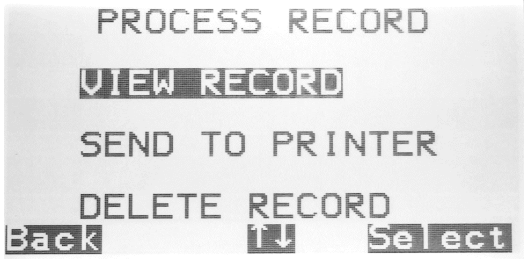
4.10.2. LIST RECORDS (VIEW, PRINT, DELETE)



LIST RECORDS shows the stored tests, 6 at a time, most recent first.

The overview contains the following information, to help identifying the test result look for:

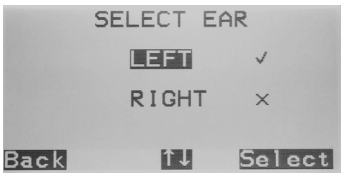
- The three-letter patient identifier entered when the test was stored
- Date and time of the test
- Whether the test has been printed ( )
- Whether the test has been sent to a computer ( )
- Whether the test is for the Left (L), Right (R) or both (2) ears

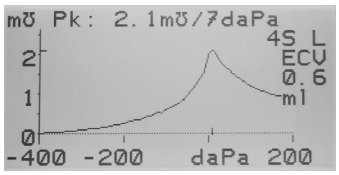


When a record is selected the **PROCESS RECORD** menu will be displayed. This accesses the following functions:

- View the selected record
- Print the selected record
- Delete a record

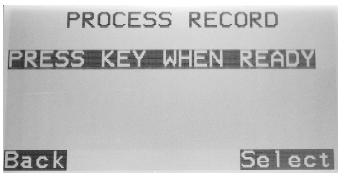
VIEW RECORD

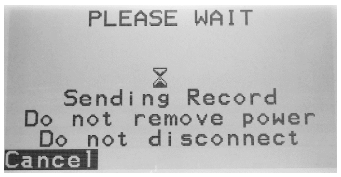




The symbols ✓ or **X** are used to signify whether results are available for each ear. Only test results with a ✓ can be selected for a review.

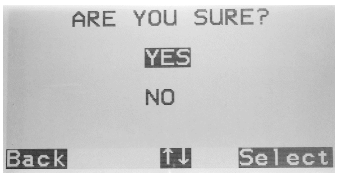
SEND TO PRINTER

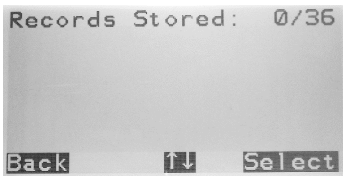




Confirm the **SENT TO PRINTER** selection as soon as the printer is ready.

DELETE RECORD





When select **DELETE RECORD** a confirmation of the deletion progress is required in order to proceed. If several tests are stored in the database, you will be brought back to the **LIST VIEW** after the successful deletion.

If only one record is available in the database a note will show on the screen that the database is empty.

Please note: When a new test is "saved as last test" the baseline mode most recently viewed will be saved, although any of the other modes can be re-created when the test is loaded back into the instrument using "View the last test". The same applies to results stored in the instrument's database. This allows a different display mode and baseline offset to be used for display or printing, but it does not affect the baseline mode originally stored in the database.

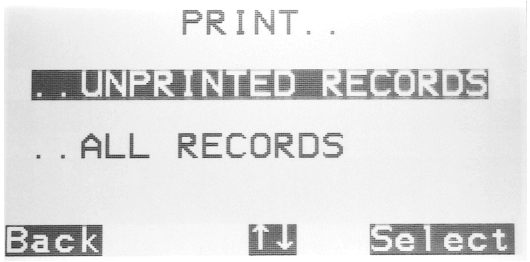
4.10.3. DELETE RECORDS

DELETE RECORDS allows a group of records to be deleted.

It is possible to delete all records stored in the database, only those records that have been printed or those records that have been sent to a computer.

4.10.4. PRINT RECORDS

PRINT RECORDS allows a group of records to be sent to the printer.



It is possible to print all stored records or just those records that have not already been printed.

Please note: If printing the entire database it is recommended that a full roll of paper is loaded into the printer.

4.10.5. DATA TRANSFER TO NOAH OR MEDRX STUDIO SOFTWARE

The Otowave 302 is supplied with software to allow connection to a computer for the transfer of test results. You must use the designated USB cable which is available from MedRx, Inc.

To transfer test results stored within the Otowave to a NOAH database, the NOAH Impedance module must be installed on to a computer. Alternatively, MedRx Studio software allows data to be transferred to a computer and subsequently viewed, annotated & printed. This software is supplied on a CD which includes this operating manual.

The computer will automatically detect the instrument when it is connected with the USB cable. Data transfer is initiated from the computer (not from the Otowave). To disconnect simply remove the cable when data transfer is completed.

Refer to the installation & operating instructions provided with the NOAH Impedance Module or MedRx Studio software for further details.

4.10.6. DATABASE FULL

When the internal memory is full, a warning will be displayed if the database is full when attempting to save a test.

- Selecting **MANAGE RECORDS** will display the **DATA MANAGEMENT** menu which provides options for printing or transferring data to a computer prior to deleting records to make space for the new test.
- **OVERWRITE OLDEST** will overwrite the oldest record in memory with the results being stored.
- **BACK** will return to the previous menu.

5. TROUBLESHOOTING

If a fault condition cannot be cleared, the operator is cautioned against repeatedly starting the instrument.

Please note: Refer to the installation & operating instructions provided with the NOAH Impedance Module or ampliSuite software for details of the data transfer operation and errors that may occur.

PROBLEM	CAUSE	SOLUTION(S)
No pressure can build up and the test sequence will stay in the **EQUALIZE PRESSURE SCREEN**.	• No seal can be obtained • Estimated volume is too high (perforated ear drum) • Wrong ear tip size chosen • Probe is blocked	• Examine the probe tip for contamination and clean or replace the probe tip • Reposition the probe • Change the ear tip
No reflex test is conducted after the tympanometry even though the reflex test is active in the **REFLEX SEQUENCE.**	In **REFLEX SELECTION** the configuration is set to **ONLY IF PEAK IS FOUND** or **NEVER MEASURE.**	Change settings in **REFLEX SELECTION** to desired option.
Last measured data cannot be found under **VIEW THE LAST TEST**.	**NEW TEST** might have been selected in between and thereby deleted the last test(s) from the short-term memory.	To be stored data should be stored immediately.
BLOCKED PROBE Indicator LED b and c flash fast.	• Probe is blocked • Probe placed against ear canal skin • Probe disconnected from base unit	• Examine the probe tip for contamination and clean or replace the probe tip • Reposition the probe • Change the ear tip • Check probe connection with base unit
WITHDRAW PROBE Indicator LED b and c flash fast.	• The probe has been moved during measurement. • Test has been started with the probe already inserted into the ear.	Reposition the probe
Volume outside range **WITHDRAW PROBE** Indicator LED b and c flash fast.	• Ear canal volume is > 5ml. • Probe is not properly inserted into the ear.	Reposition the probe
Pressure lost **WITHDRAW PROBE** Indicator LED b and c flash fast.	Ear seal has been broken while testing for seal.	Reposition the probe
Measurement timed out Indicator LED b and c flash fast.	• Occurs when the ear seal check is set to EXTENDED • Pump failed to achieve the starting pressure within 4 s. • Pressure failed to reach -400 daPa within 12 s.	Reposition the probe. Retry the test. If the problem persists, contact your MedRx, Inc service center.
VOLUME OUTSIDE RANGE Indicator LED b and c flash.	Probe not placed correctly in ear canal.	Reposition probe.
	•	•

PROBLEM	CAUSE	SOLUTION(S)
PROBE NOT CLEAR Indicator LED c steady light.	• Probe is blocked Probe placed incorrectly	• Check that the probe is not inserted into a test cavity at start-up. Please ensure the probe is not blocked or obstructed.
AIRFLOW ERROR Indicator LED c steady light.	Fault with air system and/or pump. Cannot determine pump direction.	Unknown pump fault. Restart the unit. If problem persists, contact your MedRx, Inc service center.
AIRFLOW ERROR **RESTART THE UNIT** Indicator LED c steady light.	Fault with air system and/or pump.	Restart the unit. If problem persists, contact your MedRx, Inc service center.
WARNING! CALIBRATION EXPIRED Indicator LED c steady light.	The current date is later than the next calibration date. Check that the clock is set to the correct date. If so, arrange for the instrument to be recalibrated. Tests can still be performed.	Recalibration needed before further tests are performed.
WARNING! DEVICE UNCALIBRATED. Indicator LED c steady light.	One or more default values require recalibration before further tests are performed.	Contact your MedRx, Inc service center.
WARNING! DEFAULTS RELOADED. Indicator LED c steady light.	Default configuration settings reloaded.	Default configuration settings reloaded. If the error persists, contact your MedRx, Inc service center.
Printing Error No connection can be established with the printer	• Printer is switched off or not charged • Connection between printer and base unit cannot be established.	• Restart the base unit • Restart the printer • Charge printer • Ensure the connection between printer and base unit is established.

If difficulties resolving fault conditions occur the equipment distributor (or MedRx, Inc if purchased directly) should be consulted.

6. ROUTINE MAINTENANCE

6.1. GENERAL MAINTENANCE PROCEDURES

The performance and safety of the instrument will be maintained if the following recommendations for care and maintenance are observed:

1. It is recommended that the instrument go through at least one annual service, to ensure that the acoustical, electrical and mechanical properties are correct. This should be carried out by an authorized repairer in order to guarantee proper service and repair.
2. Observe that no damage is present to the insulation of the power supply cable or the connectors and that it is not exposed to any kind of mechanical load that could involve damage.
3. To ensure that the reliability of the instrument is maintained, we recommend that the operator at short intervals, for instance once a day, performs a test on a person with known data. This person could be the operator.
4. If the surface of the instrument or parts of it is contaminated, it can be cleaned using a soft cloth moistened with a mild solution of water and detergent or similar. Always disconnect the power supply adaptor during the cleaning process and be careful that no fluid enters the inside of the instrument or accessories.
5. After each patient examination, ensure that there has been no contamination to the parts touching the patient. General precautions must be observed in order to avoid cross-contamination of disease from one patient to another. Water should be used for frequent cleaning, but in the case of severe contamination it may be necessary to use a disinfectant.

CAUTION

- Before cleaning always switch off and disconnect from the power supply
- Use a soft cloth lightly dampened with cleaning solution to clean all exposed surfaces
- Do not allow liquid to come in contact with the metal parts inside the earphones/headphones
- Do not autoclave, sterilize or immerse the instrument or accessory in any fluid
- Do not use hard or pointed objects to clean any part of the instrument or accessory
- Do not let parts that have been in contact with fluids dry before cleaning
- Rubber ear-tips or foam ear-tips are single use components

Recommended cleaning and disinfection solutions:

- Warm water with mild, nonabrasive cleaning solution (soap)
- 70% isopropyl alcohol only on hard cover surfaces

6.2. CLEANING THE OTOWAVE

CAUTION

- Use caution while cleaning.
- Before cleaning, remove the Otowave from power supply.
- Single use components such as ear-tips do not require cleaning.
- Do not allow any liquid to enter any part of the instrument or accessories.
- Do not autoclave or sterilize the instrument or any accessory.
- Do not use hard, sharp or pointed objects to clean any part of the instrument or an accessory.
- If parts have been in contact with fluids do not allow them to dry before cleaning.
- Follow local best practice and safety guidelines if available.
- Clean the instrument by wiping the outer case with a lint free cloth lightly dampened with cleaning solution. Recommended cleaning and disinfection solutions are warm water with mild, nonabrasive cleaning solution (soap) and/or Clinical wipes (for example Clinell Universal).
- If disinfection is required, use a disinfectant wipe rather than a spray product. Make sure that excess liquid from the wipe does not seep into any sensitive areas such as connectors and seams where plastic pieces connect such as under the rubber buttons on the Otowave. Follow the instructions on the disinfection product.

6.3. CLEANING THE PROBE

The probe tip should be checked before each ear insertion to ensure it is undamaged and that none of the tubes through it are blocked. It should be replaced if necessary.

The sealing washer should be replaced when the probe tip is replaced, if it shows signs of wear, or if a pressure leak is suspected.

CAUTION

Handle the probe and accessories with care. Do not allow moisture, condensation, fluids or debris to enter the probe.

6.4. DISPOSABLES

Ear tips should be replaced after a single use. This applies to ear tips used with the main probe assembly and the contralateral transducer.

Use only the disposable supplies that are supplied with your Otowave. Ear tips, ear cups and adhesive electrodes are intended for single-use only. These should be discarded after use. They cannot be disinfected.

CAUTION

In the event of re-use of the single-use disposables, you enhance the risk of cross contamination!

6.5. COMPONENTS/REPLACEMENT PARTS

Some reusable components are subject to wear with use over time. We recommend that you keep theses replacement parts available (as appropriate for your Otowave device configuration).

6.6. REPAIR

The manufacturer is only considered to be responsible for the validity of the CE marking, effects on safety, reliability and performance of the equipment if:

- assembly operations, extensions, readjustments, modifications or repairs are carried out by authorised persons
- a 1-year service interval is maintained
- the electrical installation of the relevant room complies with the appropriate requirements, and
- the equipment is used by authorised personnel in accordance with the documentation supplied by manufacturer

It is important that the customer contact MedRx, Inc. or designated distributor for technical support before shipping the Otowave.

This should be done every time an instrument is returned to manufacturer.

When packing the instrument for shipping, please use the original shipping carton and packing materials. Place the instrument parts in plastic bags before packing to prevent dirt ingress.

6.7. WARRANTY

The manufacturer gives the purchaser the following warranty;

If within three years from the date of dispatch, any defect in respect of material or workmanship within our control is discovered, we will make good the defect without charge, subject to the following conditions;

- Notice of the fault is given to manufacturer within the warranty period.
- The instrument is forwarded, postage paid, to the manufacturer at the above address or as otherwise directed.
- Return postage is free of charge for US and chargeable for overseas customers.
- The responsibility of the manufacturer under this warranty is strictly limited to making good the defect in the instrument itself.
- No attempt has been made to effect a repair or adjust the calibration or alter the instrument from the original build standard.
- Defects caused by abnormal conditions of use, accident or neglect are expressly excluded.

If any product requires service during the applicable warranty period, the purchaser should communicate directly with the local service centre to determine the appropriate repair facility. Repair or replacement will be carried out at the manufacturer's expense, subject to the terms of this warranty. The product requiring service should be returned promptly, properly packed, and postage prepaid. Loss or damage in return shipment to the manufacturer shall be at purchaser's risk.

In no event shall manufacturer be liable for any incidental, indirect or consequential damages in connection with the purchase or use of any the manufacturer product.

This shall apply solely to the original purchaser. This warranty shall not apply to any subsequent owner or holder of the product. Furthermore, this warranty shall not apply to, and the manufacturer shall not be responsible for, any loss arising in connection with the purchase or use of any the manufacturer product that has been:

- repaired by anyone other than an authorised manufacturer service representative;
- altered in any way so as, in manufacturer opinion, to affect its stability or reliability;
- subject to misuse or negligence or accident, or that has had the serial or lot number altered; defaced or removed; or
- improperly maintained or used in any manner other than in accordance with the instructions provided by the manufacturer.

This warranty is in lieu of all other warranties, express or implied, and of all other obligations or liabilities of the manufacturer. The manufacturer does not give or grant, directly or indirectly, the authority to any representative or other person to assume on behalf of the manufacturer any other liability in connection with the sale of the manufacturer products.

THE MANUFACTURER DISCLAIMS ALL OTHER WARRANTIES, EXPRESSED OR IMPLIED, INCLUDING ANY WARRANTY OF MERCHANTABILITY OR FOR FUNCTION OF FITNESS FOR A PARTICULAR PURPOSE OR APPLICATION.

6.8. CALIBRATION AND RETURN OF THE INSTRUMENT

MedRx recommends that the Otowave 302 is calibrated annually. A warning message will be displayed at power up if the instrument was calibrated more than twelve months ago. The date of the last calibration is displayed on the **SYS INFO** screen.

Please contact MedRx or the designated distributor for details of calibration services.

7. TECHNICAL SPECIFICATIONS

7.1. STANDARD AND REGULATORY

Medical CE mark	The CE mark indicates that manufacturer meets the requirements of Annex II of the Medical Device Regulation 2017/745. TÜV Product Service, Identification No. 123, has approved the quality system.	
Class	The Otowave 302 Tympanometer is classified as a Class IIa device under Annex IX (Section 1) of the EU Medical Devices Regulation.	
Standards and Conformance	**Safety:**	IEC 60601-1 (plus ES, CSA & EN deviations), Class II, Type B applied part
	EMC:	IEC 60601-1-2
	Performance:	IEC 60645-5, Type 2 Tympanometer ANSI 3.39, Type 2
Physical	**Display:**	256 x 64 pixels / 8 lines of 21 characters
	Dimensions (base unit):	270 x 60 x 165 mm / 10.63 x 2.36 x 6.49 inch (excluding connections)
	Weight (base unit):	760 g / 1.68 lbs
	Dimensions (probe):	130 x 25 mm / 5.11 x 0.98 inch
	Weight (probe):	115 g / 0.25 lbs
	Interconnection: (probe to base)	1.5 m combined electrical cable and air tube
Power Supply	**Power supplypower:**	100-240Vac; 50-60Hz; 205-110 mA
	Input rating:	5Vdc; 0.5 A
	Warm-up period:	None at room temperature
	Idle current:	70 mA
	Current while testing:	230 mA
Environmental	**Operating temperature:**	+15°C to +35°C / + 59°F to +95°F
	Operating humidity:	30 % to 90 % RH (non-condensing)
	Operating atmospheric pressure:	980 mb to 1040 mb
	Transport: storage temperature:	-20°C to +70°C / -4°F to +94°F
	Transport and storage humidity:	10 % to 90 % RH (non-condensing)
	Transport and storage atmospheric pressure:	900 mb to 1100 mb

7.2. GENERAL

Time and Date	**Stamps:**	Time and date stamp applied to all recordings, and to the last calibration date
Languages:		English, German, French, Spanish, Portuguese, Italian
Database	**No. of records stored:**	36
	Data storage:	Any recording can be stored once the tympanogram is viewed. Patient Initials (A-Z, 0-9, "-") must be entered before storage.
	Data held:	Patient Initials, Tympanogram and Reflex graphs and analysis for Left Ear and/or Right Ear, Time and Date of recording, which ears were tested, whether or not the record has been printed and/or sent to a computer, parameters used for analysis, 128 bit Globally Unique Identifier (GUID)
	Data presentation:	Records listed in reverse chronological order (latest first), with indication of data stored as described above
Printing	**Supported printer:**	Sanibel MPT-II
	Interface:	Cable supplied
	Information printed:	Tympanogram, Tympanogram analysis parameters, Reflex graphs, Reflex analysis parameters, Serial Number of device, Last and Next Due Calibration dates; space for patient & clinician's details to be entered.
PC Interface	**Serial Interface:**	USB Version 1.1
	Information sent:	Patient header, full left & right ear data.

7.3. TYMPANOMETRY

Probe Tone	**Frequency:**	226 Hz ±2%
	Level:	85 dB SPL ±2dB and 79 dB SPL ±2 dB over ECV range
Pressure	**Range:**	+200 daPa to -400 daPa ±10daPa or ±10 % (whichever is larger) over range 0.1 ml to 6 ml
	Limits (safety cutout):	+600 and -800 daPa
Sweep	**Speed:**	Selectable: 100, 200 or 300 daPa/sec
Analysis		Admittance peak level (in ml or mʊ) pressure at peak; Gradient in daPa (for 226Hz); Ear Canal Volume (ECV) @ 200 daPa or -400 daPa
	Number of samples stored	100 per tympanogram

7.4. ACOUSTIC REFLEX TESTING

Ipsilateral	**Test Frequencies:**	500 Hz, 1 kHz, 2 kHz & 4 kHz (±2 %)
	Level:	Configurable over range: 500Hz, 1kHz, 2kHz & 4kHz (+/-2%) 70dBHL to100dBHL (+/-3dB) (2kHz level is restricted to maximum 95dBHL for ear canal volumes greater than ~3.5ml) (4kHz level is restricted to maximum 85dBHL for ear canal volumes greater than ~3.5ml & maximum 95dBHL for all ear canal volumes)
	Number of reflex levels presented below the selected maximum and step size(s) available:	100dBHL max, with 5dB or 10dB steps 95/90/85dBHL max, with 5dB steps
Contralateral	**Test Frequencies:**	500Hz, 1kHz, 2kHz & 4kHz (±2%)
	Level:	Configurable over range: 500Hz, 1kHz, 2kHz & 4kHz (+/-2%) 70dBHL to110dBHL (+/-3dB) (1kHz level is restricted to minimum 75dBHL for ear canal volumes less than ~0.2ml) (2kHz level is restricted to maximum 105dBHL for ear canal volumes greater than ~3.5ml) (4kHz level is restricted to maximum 100dBHL for ear canal volumes greater than ~3.5ml & maximum 105dBHL for ear canal volumes greater than ~1.5ml)
	Number of reflex levels presented below the selected maximum and step size(s) available:	110/105/100dBHL max, with 5dB or 10dB steps 95/90/85dBHL max, with 5dB steps
General	**THD:**	< 5 %
	Reflex analysis	Reflex pass/fail at each level tested; maximum amplitude of each reflex; nominal pressure used for the reflex test (computer display only)
	Pressure used for reflex measurement	Pressure at tympanogram peak (if found) or at 0daPa
	Reflex stimulus control	Stimulus presented at all levels, or Stimulus ceases when a reflex is found
	Reflex detection threshold and accuracy	0.01ml to 0.5ml ±0.01ml (configurable in 0.01ml steps)
	Reflex tone duration	0.6 seconds

8. EMC GUIDANCE & MANUFACTURER'S DECLARATION

CAUTION

- This instrument is suitable in hospital environments except for near active HF surgical equipment and RF shielded rooms of systems for magnetic resonance imaging, where the intensity of electromagnetic disturbance is high
- Use of this instrument adjacent to or stacked with other equipment should be avoided because it could result in improper operation. If such use is necessary, this instrument and the other equipment should be observed to verify that they are operating normally
- 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. The list of accessories, transducers and cables can be found in this appendix.
- 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 this instrument, including cables specified by the manufacturer. Otherwise, degradation of the performance of this equipment could result

NOTICE

- ESSENTIAL PERFORMANCE for this instrument is defined by the manufacturer as:
  This instrument does not have an ESSENTIAL PERFORMANCE Absence or loss of ESSENTIAL PERFORMANCE cannot lead to any unacceptable immediate risk
- Final diagnosis shall always be based on clinical knowledge There are no deviations from the collateral standard and allowances uses
- This instrument is in compliance with IEC60601-1-2:2014, emission class B group 1
  NOTICE: There are no deviations from the collateral standard and allowances uses
  NOTICE: All necessary instruction for maintaining compliance with regard to EMC can be found in the general maintenance section in this instruction. No further steps required.

Guidance and manufacturer's declaration – electromagnetic emissions		
The Otowave 302 Tympanometer is intended for use in the electromagnetic environment specified below. The customer or user of the Otowave 302 Tympanometer should assure that it is used in such an environment.		
Emissions test	**Compliance**	**Electromagnetic environment – guidance**
RF emissions CISPR 11	Group 1	The Otowave 302 Tympanometer uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause interference in nearby electronic equipment.
RF emissions CISPR 11	Class A	The Otowave 302 Tympanometer is suitable for use in all establishments, other than domestic and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.
Harmonic emissions IEC 61000-3-2	Class A	
Voltage fluctuations/flicker emissions IEC 61000-3-3	Complies	

Guidance and manufacturer's declaration – electromagnetic immunity (1)			
The Otowave 302 Tympanometer is intended for use in the electromagnetic environment specified below. The customer or user of the Otowave 302 Tympanometer should assure that it is used in such an environment.			
Immunity test	**IEC 60601 test level**	**Compliance level**	**Electromagnetic environment – guidance**
Electrostatic Discharge (ESD) IEC 61000-4-2	±6 kV contact ±8 kV air	±6 kV contact ±8 kV air	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30%
Electrical fast transient/burst IEC 61000-4-4	±2 kV for power supply lines ±1 kV for input/output lines	±2 kV for power supply lines ±1 kV for input/output lines	Power supply quality should be that of a typical commercial or hospital environment
Surge IEC 61000-4-5	±1 kV differential mode ±2 kV common mode	±1 kV differential mode ±2 kV common mode	Power supply quality should be that of a typical commercial or hospital environment

Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment – guidance
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	<5% U_T (>95% dip in U_T) for 0.5 cycle 40% U_T (60% dip in U_T) for 5 cycles 70% U_T (30% dip in U_T) for 25 cycles <5% U_T **(>95% DIP IN U_T) FOR 5 SEC**	<5% U_T (>95% dip in U_T) for 0.5 cycle 40% U_T (60% dip in U_T) for 5 cycles 70% U_T (30% dip in U_T) for 25 cycles <5% U_T (>95% dip in U_T) for 5 sec	Power supply quality should be that of a typical commercial or hospital environment. If the user of the Otowave 302 Tympanometer requires continued operation during power supply interruptions, it is recommended that the Otowave 302 Tympanometer be powered from an uninterruptible power supply or a battery
Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	3 A/m	3 A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.
NOTE U_T is the a.c. power supply voltage prior to the application of the test level.			

Guidance and manufacturer's declaration – electromagnetic immunity (2)			
The Otowave 302 Tympanometer is intended for use in the electromagnetic environment specified below. The customer or user of the Otowave 302 Tympanometer should assure that it is used in such an environment.			
Immunity test	**IEC 60601 test level**	**Compliance level**	**Electromagnetic environment – guidance**
Conducted RF IEC 61000-4-6	3 Vrms 150kHz to 80MHz	3 Vrms	Portable and mobile RF communications equipment should be used no closer to any part of the Otowave 302 Tympanometer, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter. Recommended separation distance d = 1.2√P d = 1.2√P (80MHz to 800MHz0 d = 2.3√P (800MHz to 2.5GHz) where P is the maximum output power rating of the transmitter in Watts (W) according to the transmitter manufacturer and d is the recommended separation distance in meters (m). Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey, [a] should be less than the compliance level in each frequency range. [b] Interference may occur in the vicinity of equipment marked with the following symbol:
Radiated RF IEC 61000-4-3	3 V/m 80MHz to 2.5GHz	3 V/m	
NOTE 1 At 80MHz and 800MHz, the higher frequency range applies. NOTE 2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.			

Guidance and manufacturer's declaration – electromagnetic immunity (2)	
a	Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio, AM and FM radio broadcast and TV broadcast cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which the Otowave 302 Tympanometer is used exceeds the applicable RF compliance level above, the Otowave 302 Tympanometer should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the Otowave 302 Tympanometer.
b	Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3 V/m.

Recommended separation distances between portable and mobile RF communications equipment and the Otowave 302 Tympanometer

The Otowave 302 Tympanometer is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of the Otowave 302 Tympanometer can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the Otowave 302 Tympanometer as recommended below, according to the maximum output power of the communications equipment.

Rated maximum output power of transmitter W	**Separation distance according to frequency of transmitter m**		
	150 kHz to 80 MHz $d = 1.2\sqrt{P}$	**80 MHz to 800 MHz** $d = 1.2\sqrt{P}$	**800 MHz to 2.5 GHz** $d = 2.3\sqrt{P}$
0.01	0.12	0.12	0.23
0.1	0.38	0.38	0.73
1	1.2	1.2	2.3
10	3.8	3.8	7.3
100	12	12	23

For transmitters rated at a maximum output power not listed above, the recommended separation distance d in metres (m) can be estimated using the equation applicable to the frequency of the transmitter, where P is the maximum output power rating of the transmitter in Watts (W) according to the transmitter manufacturer.

NOTE 1 At 80MHz and 800MHz, the separation distance for the higher frequency range applies.

NOTE 2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

9. SAFETY PRECAUTIONS TO TAKE WHEN CONNECTING OTOWAVE 302

Please note that if connections are made to standard equipment like printers and networks, special precautions must be taken in order to maintain medical safety. Please follow instructions given in this chapter.

Diagram 1: Otowave 302 used with the medically-approved power supply adapter

Diagram 2: Otowave 302 used with the medically-approved power supply adapter and printer

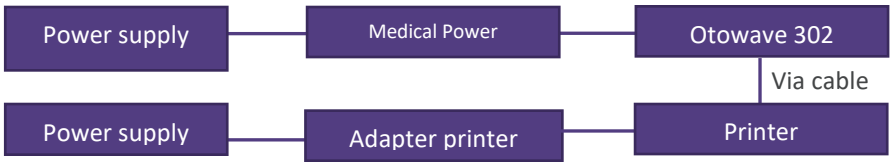
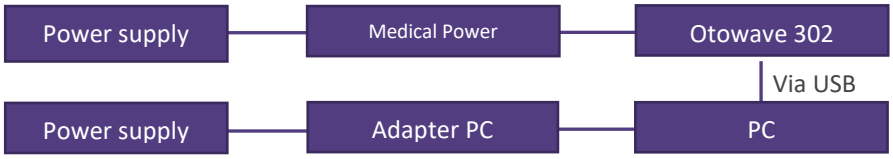


Diagram 3: Otowave 302 used with the medically-approved power supply adapter and PC



APPENDIX A –BASELINE MODE

GENERAL

The Otowave 302 can display tympanograms in a variety of graphical formats allowing the operator to choose the most appropriate for the patient under examination.

This is achieved by altering the **DISPLAY MODE** and the meatus compensation (or **BASELINE OFFSET**). The **DISPLAY MODE** determines how the tympanogram trace is derived from the raw data, and the baseline offset selects the pressure to which the meatus compensation is referenced (either -400daPa or +200daPa). **DISPLAY MODE** and **BASELINE OFFSET** are collectively referred to as **BASELINE OFFSET** in the instrument menus and the accompanying documentation.

The tympanogram is initially presented using default settings for display mode and baseline offset. Additionally, whenever a tympanogram is shown it may be re-displayed using any of the alternative **DISPLAY MODES** and **BASELINE OFFSETS** available described in this section.

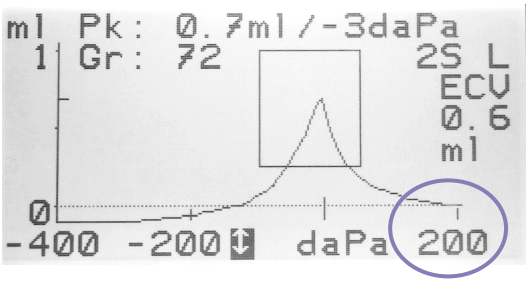
Switching between **DISPLAY MODES** and **BASELINE OFFSETS** is carried out using either the display mode key on the front panel or the function button on the probe.

Press and briefly hold the key or button to cycle round Scalar, Vector and Component display modes. Short button presses will circle around the current selected mode, long button presses will access another display mode.

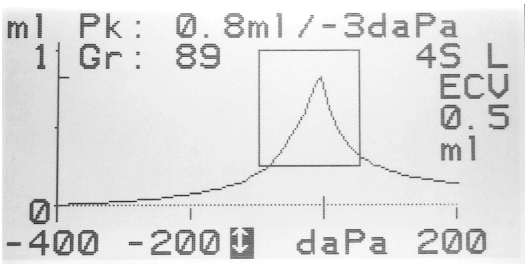
226 HZ TYMPANOMETRY – Y COMPENSATION ONLY

Please note that only the Scalar display mode is available for 226Hz probe frequency.

2S MODE



4S MODE



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