

# **GSI TYMPSTAR™** Middle Ear Analyzer

## PRODUCT SPECIFICATIONS

The following specifications apply to both Version 1 and Version 2 of the GSI TympStar. > **Bold arrowed** type indicates the additional capabilities of Version 2.

#### **GENERAL SPECIFICATIONS**

W x D x H: 52 cm x 38 cm x 32 cm

Weight: 7.5 Kg

Shipping Weight: 13.4 Kg

Power Consumption: 120 Watts maximum

Test Types: Tympanometry, Acoustic Reflex Threshold, Reflex Decay, Eustachian Tube Function (Intact & Perforated) Protocols: Diagnostic, Screening, User-defined

> Special Tests: Two-component Tympanometry, Multiple Frequency Tympanometry, Acoustic Reflex Latency Test, Reflex Sensitization

Display: Internal or External VGA Monitor Interface: RS232, parallel and keyboard output Printout: Internal or External Deskjet or Laserjet Printer

# **PROBE TONE**

226 Hz (85 dB SPL ± 1.5 dB)  $> 678 \text{ Hz} (85 \text{ dB SPL} \pm 3.0 \text{ dB})$  $> 1000 \text{ Hz} (75 \text{ dB SPL} \pm 3.0 \text{ dB})$ 

Accuracy: ± 1%

Harmonic Distortion: Less than 5%

## **ADMITTANCE MEASUREMENTS**

Range: 226 Hz (-1.0 to +7.0) > 678 Hz (-5.0 to +25) > 1000 Hz (-5.0 to +30)

Sensitivity Scale: Auto Scales to Appropriate Range, Manual selection also possible in Reflex Modes only Accuracy (226 Hz):

Tymp Mode:  $\pm$  5% of reading or  $\pm$  0.1 ml, whichever is greater

Reflex Mode:  $\pm$  5% of reading or  $\pm$  0.2 ml, whichever is greater

#### **PRESSURE MEASUREMENTS**

(load volume of 0.2 to 7.0 ml)

Normal = +200 to -400 daPa Range: Wide = +400 to -600 daPa

Accuracy: ± 10% of reading or ± 10 daPa, whichever

Sweep Rate: 12.5, 50.0 and 600/200 daPa/sec,

> 200 daPa/sec.

Sweep Accuracy: 10% of nominal rate

Maximum limits (in 0.5cc cavity): -800 daPa & +600 daPa

# REFLEX MEASUREMENTS

Stimuli: > 250, 500, 1k, 2k, 4k, BBN, LBN, HBN, > Click (100 microseconds pulse), External Input, Non-acoustic

Frequency Accuracy: ± 3%

Harmonic Distortion (THD): Less than 5% (measured

acoustically)

**GRASON-STADLER** 

Noise Signals: (3 dB bandwidths) Low Band: 125 -1,600 Hz High Band: 1,600 -4,000 Hz Broad Band: 125 -4,000 Hz Intensity Range: 35 to 120 dB HL

Step Size: 5 dB, >1 dB and 2 dB Calibration Accuracy: ± 3 dB Step Accuracy: ± 0.5 dB

ON/OFF Ratio: 70 dB minimum

#### **ENVIRONMENTAL**

Temperature:

Storage: -40°C to +75°C

Operating: +15°C to +35°C Humidity: 90% at 35°C (non-condensing)

#### **ACCESSORIES SUPPLIED**

Probe assembly (including contralateral insert phone) Eartips (1 pkg. each standard, special, screening) Printer paper, 2 rolls thermal, 1 roll self-adhesive Calibration test cavity, Cleaning kit, Probe mount kit (shoulder, clip, wrist band), Quick User Guide, Reference Instruction Manual, Remote (RS232 link) **Dust Cover** 

#### **OPTIONAL ACCESSORIES**

| Deskjet printer       | 1700-9613 |
|-----------------------|-----------|
| External VGA monitor  | 1700-9614 |
| External keyboard     | 1700-9615 |
| Isolation Transformer | 1700-9617 |
| Service Manual        | 2000-0110 |
| Conversion Kit        | 2000-9650 |
| (V1 to V2)            |           |

#### **LANGUAGE KITS**

| GSI TympStar Version 1 |            |
|------------------------|------------|
| ´ English              | 2000-9645E |
| French                 | 2000-9645F |
| German                 | 2000-9645G |
| Spanish                | 2000-9645S |
| Italian                | 2000-9645I |
| GSI TympStar Version 2 |            |
| Énglish                | 2000-9646E |
| French                 | 2000-9646F |
| German                 | 2000-9646G |
| Spanish                | 2000-9646S |
| Italian                | 2000-9646I |
|                        |            |

# **QUALITY SYSTEM**

Manufactured, designed, developed and marketed by VIASYS Healthcare Inc. NeuroCare Group under ISÓ 13485, ISO 9001 certified quality system.

# COMPLIANCE / REGULATORY STANDARDS

Designed, tested and manufactured to meet the following domestic (USA), Canadian, European and International Standards:

UL 2601-1 American Standards for Medical Electrical

IEC 601-1, EN 60601-1 International Standards for Medical Electrical Equipment.

CSA C22.2 # 601-1-M90

**NICOLET VASCULAR** 

Medical Device Directive (MDD) (ID No.: 0344) to comply with "EC Directive" 93/42/EEC

ANSI \$3.39, ANSI \$3.6, IEC 645-1, IEC 1027, ISO 389

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NICOLET BIOMEDICAL

**TOENNIES**