

CASE[™] Cardiac Assessment System for Exercise Testing

Signal processing

ST measurements – Resting	ST amplitudes, slope
ST measurements – Stress	ST amplitudes, slope, integral, index, ST/HR slope, ST/HR loops, ST/HR
E, J and post-J point	Manual or computer selected
Signal processing technique	Incremental median updating using HEART Exercise program
Baseline Correction	ADS
Artifact/Baseline correction	ADS or Finite Residual Filter (FRF) algorithm
QRS detection and analysis	Based on automatic or manual lead selection
Arrhythmia detection	Automatic arrhythmia detection, documentation and annotation
Full disclosure ECG	Up to 60 minutes of full disclosure with event review both during and post acquisition
Reanalysis – Resting	Re-analyze after manual correction of median beats and measurements
Reanalysis – Stress	Post-test median measurements from E, J, post-J point selections
ECG analysis	Marquette [™] 12SL ECG Analysis Program for Adult and Pediatric (optional)
Computerized Measurements	15-lead analysis includes measurements of user-selectable additional 3 leads
Additional ECG function	Vectorcardiography

Heart Rate Meter

30 to 300 BPM ±10% or 5 BPM, whichever is greater. Heart rates outside this range will not be displayed.

Communications/storage

ECG data formats	GE Hi-Fidelity ECG, XML
MUSE [™]	MUSE Cardiology Information System Compatible (v7 or later) with bi-directional orders and ADT support
MUSE Web	Compatible for retrieval view and printing of MUSE system data
CardioSoft Web	Compatible for report viewing
Data export	PDF export of final reports (auto export and custom file name); PDF export of Full Disclosure data; Microsoft [®] Word export of configured reports; XML or Microsoft Excel [®] export of specified data
EMR connectivity	Other EMRs through MUSE Cardiology Information System (v8 or later); or GDT/ BDT Interface
DICOM	Bidirectional, DICOM modality worklist/orders

Data acquisition (via CAM Connect 14)

Technology	Type CF, Defibrillation-Proof Defibrillation protection: Per IEC 60601-2-25:2011
Lead system	15 lead
Dynamic Range	AC Differential \pm 5mV, DC offset \pm 300 mV
Common Mode Rejection	>130 dB (>100 dB with AC filter disabled)



Input Impedance	>10M Ω @ 10 Hz, defibrillator protected	Sample Rate	500
Patient Leakage	<10 µA	Recording	
Analog to Digital	Bandwidth: DC to 500 HZ;	Lead OFF	Yes
conversion	Sample rate: 2 ksps;	Detection	
	Resolution: 0.1192 µV	Communication	Digital RF 2400 – 2483 MHz,
Down Sampled	Bandwidth: 0.04 to 150 Hz;	D	0.4mW, conform to FCC Part 15.249
ECG waveform	Sample rate: 2 ksps; Resolution: 1.22 µV	Battery	Size AA \times 2, Alkaline or NiMH
		Operation Time	Up to 40 hours with Alkaline Batteries
Input to ECG Analysis	Bandwidth: 0.0., 0.56ZPD to 150Hz; Sample rate: 500sps;	Dimensions and	Weight [g]: 350
Allalysis	Resolution: 4.88 µV/LSB	weight	Size [mm]: 140 x 95 x 50
Pacemaker	Bandwidth: 23.5 Hz to 10.5 kHz;	Safety Standards	IEC 60601-1, 60601-1-2, 60601-2-25
waveform	Sample rate: 75 ksps;	Classification	Type-CF, Internally powered
	Resolution: 11.85 µV/LSB	Certification	CE
Pace Detection	Duration: 0.2 ms to 2.2 ms	Pace Detection	Sampling rate: 500 sps
	Amplitude: 2mV to 700 mV	Additional report	20, 40, 100, 150 Hz (selectable)
	Separation: 1ms or greater	filters	
Quality Indicators	Real-time electrode placement support	Battery Indicator	0%–100% (at a gap of 20%)
	with LED lead quality indicators	on HOST	
Remote control	ECG acquisition button	Receiver	
Ingress Protection	IP×4	ECG Out	0.5 to 150 Hz (Bandwidth-3db)
Level		ECG Out Gain	1000
Noise	<15 µV (-3dB) bandwidth	ECG Out Sample	500
High pass filter	0.04	Rate	
Additional report	20, 40, 100, 150 Hz (selectable)	TTL Trigger width	16-128ms
filters		TTL Trigger delay	<11ms (delay from R-wave)
Line filter	50.0 or 60.0 Hz notch filter (selectable)	Interface	USB 2.0 compliant
QRS trigger	TTL synchronization output	Communication	USB 2.0 Full Speed
			Digital RF, 2400-2483.5 MHz, 0.4 mW,

Wireless data acquisition (via GEH ECG 1200)

Acquisition unit	
Lead system	Standard 12 Lead
Patient Leads	Detachable 10 Lead wires conform to AAMI
Defibrillation protection	Protected against 360J discharge
Patient leakage current	< 10 uA
Input impedance	> 10 MOhm
CMMR	> 90dB
Frequency range	0.05 – 150Hz
Dynamic range	+/- 2.4V
Resolution	24 bits (0.286 uV/LSB)
Sample Rate Internal	8000

TTL Trigger width	16-128ms
TTL Trigger delay	<11ms (delay from R-wave)
Interface	USB 2.0 compliant
Communication	USB 2.0 Full Speed Digital RF, 2400-2483.5 MHz, 0.4 m\ conform to FCC part 15.249
Power: USB 5V	100mA max at 5V input
Dimensions and weight	Weight [g]: 80 Size [mm]: 100 x 54 x 30
	C

Physical specifications

Operating System	Microsoft Windows [®] 10, 64Bit IoT
Height (approx)	147 cm (58 in) minimum to 167.6 cm (65.98 in) maximum, fully extended
Width (approx)	62 cm (24 in) minimum to 83.8 cm (32.99 in) maximum, fully extended
Depth (approx)	75 cm (30 in) minimum to 116.8 cm (45.98 in) maximum, fully extended
Wire paper tray	31.75 cm x 23.50 cm (12.50 in x 9.25 in) Attaches to left side of CASE system
Weight	68 kg (150 lb) without monitor

Acquisition module holder tray	•	Environmental – operating requirements	
Interfaces included Acquisition module – CAM 14 Keyboard – alpha numeric - Key Switch: Membrane key switch with tactile feedback - Waterproof / Dustproof Ratings: IP 68, NEMA 4X Keypad – dedicated stress function keys Mouse Integrated Thermal Printer	Operating temp. range	10 to 40°C	
	 Key Switch: Membrane key switch with tactile feedback 	Storage temp. range	-20 to 60°C
		Relative Humidity	10 to 95% RH non-condensing
		Temperature	10 to 40°C (+50 to 104°F)
		Humidity	20 to 95% RH non-condensing
	0	Pressure	700 to w1060 hPa
	10 MB/100 MB/ 1 GB Ethernet – MUSE compatibile	Display type	
USB ports – 6 PS2 ports – 2 Serial ports – 6; COM 1-2, COM A-D, treadmill, BP, ergometer, SpO ₂ Analog port – 4; TTL (trigger) – 1; output (imaging sync) and analog device communication External DVD/CD-RW External SD	•	Monitored leads	12, 15 – Resting; 3, 6, 12, 15 – Stress
	PS2 ports – 2	Displayed leads	Number on screen 3, 6, 12 or 15
	Display format	4×2.5 , $4 \times 2.5 + 1$ rhythm, 2×6 , 6 rhythm, 3 rhythm - Resting $4 \times 2.5 + 1$ rhythm, 2×6 , 6 rhythm, 3 rhythm, 3 rhythm + medians, 3 rhythm + trend - Stress	
	Display sensitivity/gain	2.5, 5, 10, 20, 40 mm/mV - Resting 2.5, 5, 10, 20 mm/mV - Stress	

Environmental – power requirements

Power supplyAC operation onlyOperating voltage
range100-120 VAC,
47-63 Hz, 2.8 A
200-240 VAC,
47-63 Hz, 1.4 APower consumption350 W max (1200 BTU/h)
<250 W normal (850 BTU/h)
<30 W standby (100 BTU/h)</td>

CASE monitor

Display type	LCD (flat panel display)
Display resolution	LCD - 1680 × 1050
Display size	56cm (22") diagonal

1 Marquette 12SL ECG Analysis Program Physician's Guide, 2032056592-002 Revision B. 2015, GE Healthcare: Milwaukee, WI

© 2019 General Electric Company – All rights reserved.

GE Healthcare reserves the right to make changes in specifications and features shown herein, or discontinue the product described at any time without notice or obligation. CardioSoft v7 is a CE-marked medical device. CardioSoft v7 is not available in all markets and is not 510k cleared. Contact your GE Healthcare representative for the most current information. GE, the GE Monogram, CASE, Marquette, MUSE, and TONOPORT are trademarks of General Electric Company. GE Healthcare, a division of General Electric Company. Microsoft, Windows, and Excel are trademarks of Microsoft Corporation. HP is a trademark of Hewlett-Packard, Inc. Pentium is a trademark of Intel Corporation. GE Medical Systems, Inc., doing business as GE Healthcare.

