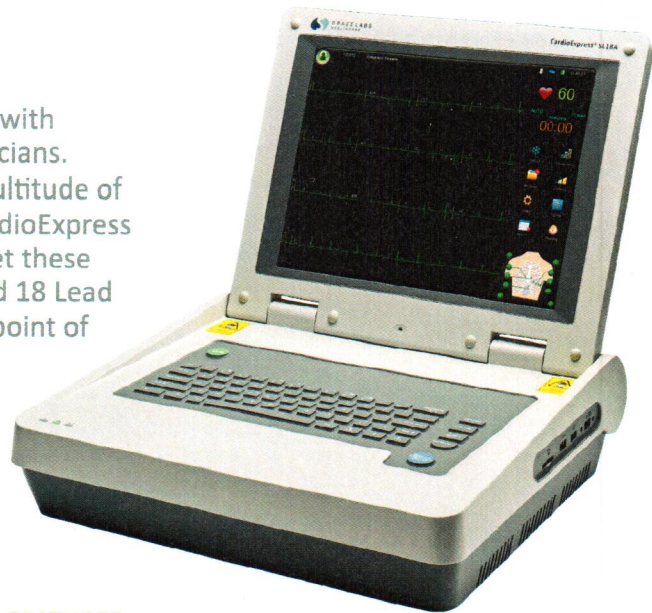


Across the world, improving health starts with providing quality data to doctors and clinicians. Today's devices are required to meet a multitude of differing needs in today's market. The CardioExpress SL18A recorder has been designed to meet these needs by including support for 12 / 15 and 18 Lead waveforms, providing quality data at the point of care.



FEATURES

FEATURES	
Ordering Information	SL18A-AHA SL18A-IEC
Recorder includes:	<ul style="list-style-type: none"> • Power cables (US or EU) • Patient cable for resting ECG (US or EU) • Chest and limb electrodes • Recorder paper • Rechargeable battery • Operations manual
Printing	<p>Internal printer for all current and stored ECG data. Print to 210 or 216mm wide roll or z-fold paper.</p> <p>Allows printing to compatible USB printer (<i>supported models only</i>). A4 or US Letter. Deskjet: HP1050 J410 Laserjet: HP1020, HP2015, HP400 M410D, CP1525N, P2-35, P2055, P1505</p>
Print Speed	5 / 6.25 / 10 / 12.5 / 25 / 50 mm/s, user selectable
Channels printed	Maximum 15 channels simultaneously
Data storage	1000, 10-second 18-channel recordings
Data export to USB port and via network port to Sentinel.	File formats: <ul style="list-style-type: none"> • PDF • SCP • FDA-XML • Dicom
System Settings	Import and export of saved system configurations settings via USB device for rapid configuration of multiple devices
Keyboard	Standard QWERTY keyboard, coloured rubber keys.
Languages	English (US), French, German, Italian, Russian, Spanish
PHYSICAL DIMENSIONS	
Height	135 mm
Length	438 mm
Width	395 mm
Weight	9.5 kg (without paper & battery)
Display	15" Color touch screen
Wi-Fi	
Wi-Fi Module	Compatible with IEEE802.11b/g & IEEE802.11n

ECG ACQUISITION

Leads	18 Standard	
Acquisition mode	18-leads simultaneously	
Sensitivity / gain	2.5, 5, 10, 20, 10/5 mm/mV, AGC	
Heart rate recognition technique	Peak-peak detection	
Heart rate recognition range	30 to 300 BPM	
Heart rate recognition accuracy	±1 BPM	
Duration to analyse 10-second 12-lead ECG, using automatic interpretation, and print-out report	<20 seconds	
Calibration Signal Input	1mV±2%	
Sample Frequency	1000Hz	Rettelse: 16kHz
Frequency response	0.05~150Hz (-3dB)	Rettelse: 0.01~300Hz(-3dB)
Filters	AC: -20db (at 50/60Hz Sinus) or OFF	EMG filter: -3db (25/35/45Hz) or OFF
<small>*The DFT filter reduces ECG baseline fluctuations, keeping the ECG on the baseline. The setting is the low limit of the frequency range.</small>	DFT filter*: 0.05, 0.15, 0.25, 0.32, 0.5, 0.67Hz	Low pass filter: 75, 100, 150, 270, 300Hz
Indicator LEDs	Mains power, Battery power, Battery recharging	
Communication	Use of bar-code readers via USB port to import Patient ID, first name, last name.	
Report	ECG data: Date and time of recording, all channels, average ECG wave template or rhythm lead, measurement & interpretation statement, histograms, trend charts, blood pressure, heart rate. Patient data: height, weight, ID, names, date of birth, race, gender, age Facility name, physician, referring physician Settings: Printing speed, sensitivity, filter, lead mark Recorder: Model type and firmware version.	

ELECTRICAL REQUIREMENTS

Power Requirement	Supports both mains and battery powered operation.	
Power supply	Auto-ranging mains power: 100V to 240V A.C., 50/60Hz. Input power 0.96A - 0.4A	
Battery	Operating time	3.5 hours normal use
	Recharge time (from fully discharged)	6 hours
	Specification	5000mAh. 14.8V Rechargeable battery

ENVIRONMENTAL REQUIREMENTS

Storage	• Temperature	-20° to 55° C
	• Humidity	15% - 95% (non-condensing)
	• Atmospheric Pressure	700 to 1060 mbar (hPa)
Operating	• Temperature	5° to 40° C
	• Humidity	15% - 95% (non-condensing)
	• Atmospheric Pressure	860 to 1060 mbar (hPa)
Water Ingress Protection	IPX0 (No protection)	

ELECTROMAGNETIC COMPATIBILITY

Emissions	IEC 60601-1-2 Group 1, Class A. RF emissions CISPR 11 IEC 61000-3-2 Class A, Harmonic emissions	
Immunity	IEC 60601-1: Internally powered ME Equipment. Class I ME equipment when connected to mains supply. Protection degree against CF electric shock with IEC or AHA patient cables. IEC 61000-3-3 Compliance, voltage fluctuations & flicker.	
Regulatory	CE marked in accordance with the Medical Device Directive 93/42/EEC.	

