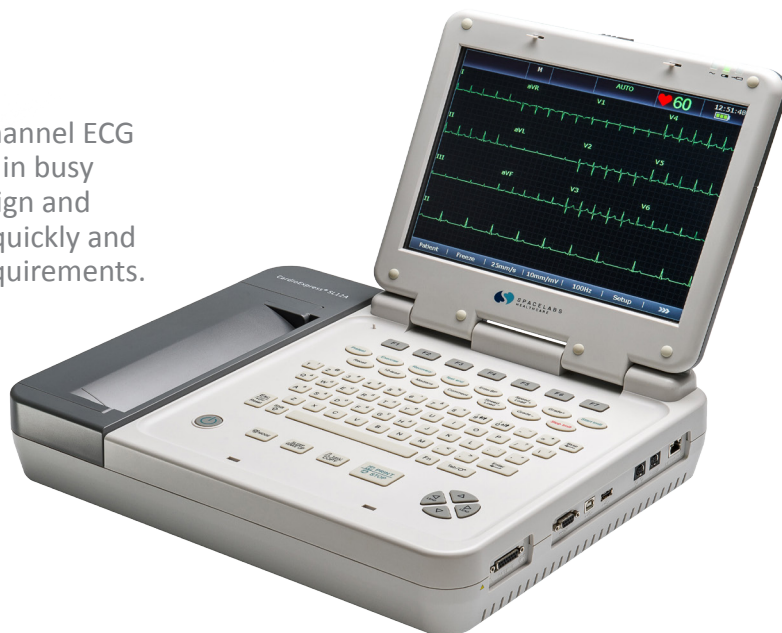


The CardioExpress SL12A is a twelve-channel ECG recorder designed for high volume use in busy departments. It features advanced design and ergonomics to enable the operator to quickly and easily control the unit for individual requirements.



FEATURES

FEATURES	
Ordering Information	SL12A-AHA SL12A-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	Internal printer for all current and stored ECG data. Print to 210mm or 216mm rolled or z-fold paper. Allows printing to compatible USB printer (<i>supported models only</i>). Supported printers: HP 2015/2035 / 1525. HP2010 / 1050 / 2000 / 2050. HP1020. HP1505.
Print Speed	5 / 6.25 / 10 / 12.5 / 25 / 50 mm/s, user selectable
Channels printed	12 (all) channels
Data storage	500 10-second 12-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, Polish, Portuguese, Russian, Spanish
PHYSICAL DIMENSIONS	
Height	120 mm
Length	420 mm
Width	330 mm
Weight	6.5 kg (without paper and battery)
Display	12.1" Color touch screen
Wi-Fi	
Wi-Fi Module	Compatible with IEEE802.11b/g <i>Note: The SL12A is not compatible with IEEE802.11n</i>



ECG ACQUISITION	
Leads	Standard Cabrera
Acquisition mode	10 second 12-lead simultaneous or sequential
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	<15 seconds
Calibration Signal Input	1mV±2%
Sample Frequency	1000Hz
Frequency response	0.05~150Hz (-3dB)
Filters	AC: -20db (at 50/60Hz Sinus) EMG filter: -3db (25/35/45Hz) or OFF 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, Low pass filter:75,100,150Hz 0.32, 0.5, 0.67Hz
Indicator LEDs	Mains power, Battery power, Battery recharging
Communication	Use of bar-code readers via USB port to import Patent ID, first name, last name.
Report	ECG data: Date and time of recording, all channels, average ECG wave template or rhythm lead, measurement and 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 5 hours normal use Recharge time (from fully discharged) 6 hours Specification 5000mAh. 14.8V Rechargeable battery
ENVIRONMENTAL REQUIREMENTS	
Storage	<ul style="list-style-type: none"> Temperature -20° to 55° C Humidity 25% - 93% (non-condensing) Atmospheric Pressure 700 to 1060 mbar (hPa)
Operating	<ul style="list-style-type: none"> Temperature 5° to 40° C Humidity 25% - 80% (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 and flicker.
Regulatory	CE marked in accordance with the Medical Device Directive 93/42/EEC. 