ECG100S

General Information	
Product Name	ECG100S
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Product Code	80508096
Manufacturer	Cardioline Spa
	Registered Office and Factory: Via Linz, 19-20-21 Zona Ind. Spini di Gardolo 38121 Trento Italy
	Sales Office: Via F.lli Bronzetti, 8 20129 Milan Italy
Device Description	The device is a 12-lead, fully diagnostic electrocardiograph which displays, acquires, prints and stores ECG tracings for adults and children. It also calculates the main overall ECG parameters. The device is equipped with USB connectivity. ECG exams can be exported in SCP or PDF format. The device can be supplied with the optional 12-lead Glasgow resting ECG interpretation algorithm, with specific criteria by age, sex and race. If this option is enabled, the algorithm provides full ECG interpretation in short or extended form, including infant, pediatric and acute ST elevation myocardial infarction detection. For further information on the resting ECG interpretation algorithm, see the Guidance for the physician on the application on adults and children (see accessories list). The device is battery or mains operated. The printing formats supported include: standard or Cabrera 3, 3+1, 3+3 or 6 channels in automatic mode and 3 or 6 channels rhythm strip printing.
Intended Use	 ECG100S is a high-performance, multi-channel, interpretative resting electrocardiograph. The ECG signal is acquired with a 10-wires patient cable and is displayed in real time on a LCD screen integrated in the device. The electrocardiograph can analyse and store the ECG traces, send them to an external peripheral via USB, print the 12 lead ECG in automatic or manual mode by means of a thermal printer. ECG100S is intended for assessment and diagnosis of cardiac functions. In any case the results of analysis performed by the electrocardiograph must be validated by a Physician. ECG100S is intended for use in hospitals, in medical clinics and doctor's offices of any size. The device is indicated for use to acquire, analyse, display and print electrocardiograms. The device is intended to provide the physician with an automatic interpretation of the ECG to be reviewed by a physician. The device is indicated for use in a clinical setting, by a physician or by trained personnel who are acting on the orders of a licensed physician. It is not intended as a sole means of diagnosis. The interpretations of ECG offered by the device are only significant when used in conjunction with a physician over-read as well as consideration of all other

- The device is indicated for use on adult and pediatric populations.
- The device is not intended to be used as a vital signs physiological monitor.

Technical Specifications	
ECG Acquisition	
ECG channels	12-lead (I, II, III, aVR-L-F, V1-6)
Patient Cable	Standard 15D, 10 wires patient cable
CMRR	> 100dB
Input impedance	100ΜΩ
A to D converter	24 bit, 32000 samples/second/channel
Sampling rate of the input stage	32000 samples/second/channel
Sampling rate for signal analysis	1000 samples/second/channel
A/D conversion	20 bit
Output Data Resolution	1 µV/LSB
Dynamic Range	+/- 400 mV
Bandwith	0,05-300 Hz
Pacemaker detection	Hardware detection coupled with convolution digital filtering
Defibrillation Protection	AAMI/IEC standards
Front-end performance	ANSI/AAMI IEC 60601-2-25:2011
Acquisition Mode	Automatic (12 leads), Manual (3/6 leads), Stat (12 leads)
Lead Configuration	Standard, Cabrera
Processing	
Operating system	Linux
Pace detection	Hardware detection in compliance with the requirements 60601-2-25
Lead fail detection	Independent on all leads
Heart Rate Meter	30 - 300 bpm
Baseline stabilization	Diagnostic fully digital high pass filter
AC Filter	50/60 Hz adaptive digital filter
Filters	Digital low pass filters at 25/40/150 Hz, for display and printing only
FCC Managements	All leads, average, corrected
ECG Measurements	
ECG Interpretation	Glasgow Analysis Program for Adults, Pediatric, STEMI (Optional)
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ECG Interpretation ECG Interpr. Data input	Glasgow Analysis Program for Adults, Pediatric, STEMI (Optional) Race, sex, age, drugs
ECG Interpretation ECG Interpr. Data input Storage	Glasgow Analysis Program for Adults, Pediatric, STEMI (Optional) Race, sex, age, drugs
ECG Interpretation ECG Interpr. Data input Storage Processing Options	Glasgow Analysis Program for Adults, Pediatric, STEMI (Optional) Race, sex, age, drugs Internal storage up to 100 ECGs

SCP	Yes (exported to USB drive)
PDF	Yes (exported to USB drive)
Connectivity	
USB	Yes
LAN	No
WiFi	No
Display	
Display Type	4.3" Backlit Color LCD
Display resolution	640x480
Display data	3/6/12 leads realtime
Display formats	6x2, 6x1 1st, 6x1 2nd, 6x1 3rd, 3x1 1st, 3x1 2nd, 3x1 3rd, 3x1 4th, 3x1 5th
Keyboard	
Keyboard Type	Full alphanumeric
Keyboard Technology	Polycarbonate layer, mechanical keypad
Dedicated Keys	ID, Start, Stop, Auto, Link - Function keys
Printer	
Technology	108 mm Thermal printhead
Resolution	8 dots/mm
Speed	5, 10, 25, 50 mm/s
Sensitivity/Gain	2.5, 5, 10, 20 mV/mm
Paper Type	Z-Fold thermal paper pack 100x150 mm
Auto Print	3, 3+1, 6 channels; Standard or Cabrera
Printing formats	6x2, 3x4, 3x4+1, 3x4+3
Manual Print	3/6 channels; 5,10,25,50 mm/sec
USB External Peripherals	
Barcode reader	Optional
Magnetic Card Reader	Optional
External data storage	Optional
Electrical Characteristics	
Power source	Medical grade - Mod. AFM60US18 - XP Power Limited
Power supply	Medical grade - Mod. AFM60US18 - XP Power Limited
Input Voltage power supply	100-240 Vac
Input Current power supply	1.5A
Input frequency power supply	50/60 Hz
Rated Output power supply	30 W, 18 V, 1.67 A
Protection Class power supply	I
Degree of Protection power supply	IP20

Battery Type	NiMH	
Battery Duration	more than 500 ECGs – more than 6 hours	
Battery Charging Time	4 hours to 85% full capacity	
Physical Characteristics		
Dimensions	285x204x65 mm	
Weight	1,8 Kg	
Shipping container	360x360x250 mm - 4Kg	
Operating Environmental Specifications		
Temperature	+10°C - +40°C	
Humidity	25% - 95%	
Pressure	700hPa - 1060hPa	
Storage Environmental Specifications		
Temperature	0°C - +40°C	
Humidity	25% - 95%	
Pressure	700hPa - 1060hPa	

Regulatory and Safety			
Classification according MDD 93/42/CEE			
Class	Class IIa		
Rationale	rule 10 annex IX 93/42/EEC Directive and its amendments		
Notified body	TUV (1936)		
Classification according to FDA	A regulation		
Classification:	II without exemption		
Product Code:	DPS		
Review Panel:	Cardiovascular		
Regulation Number:	870.2340		
Classification according to IEC 60601-1 - Electrical Safety			
Protection against electric shock:	IP (internal power ME) - class I on the external AC/DC		
Applied parts:	type CF – defibrillation-proof		
Protection against harmful ingress of water or particular matter:	IP20		
Method(s) of sterilization:	NA (not intended to be sterilized)		
Suitability for use in an oxygen rich environment:	No		
Mode of operation:	continuous operation		
Classification according to IEC 60601-1-2 - Electro Magnetic Compatibility			
Group	1		
Class	В		

Performances	
	EN 60604 2 25 2044
Standard	EN 60601-2-25:2011
Other classifications	
GMDN	110407 - Electrocardiographs, Multichannel, Interpretive
CND	Z12050302 - ELETTROCARDIOGRAFI PER DIAGNOSI AVANZATA
RDM (Registration number in Italy)	
Applicable Standards	
EN 980	Symbols for use in the labelling of medical devices
EN 1041	Information supplied by the manufacturer of medical devices
EN ISO 13485	Medical devices - Quality management systems - Requirements for regulatory purposes (ISO 13485:2003)
EN ISO 14971	Medical devices - Application of risk management to medical devices (ISO 14971:2007, Corrected version 2007-10-01)
EN 60601-1	Medical electrical equipment - Part 1: General requirements for basic safety and essential performance
EN 60601-1-2	Medical electrical equipment - Part 1-2: General requirements for basic safety and essential performance - Collateral standard: Electromagnetic compatibility - Requirements and tests
EN 62304	Medical device software - Software life-cycle processes
EN 60601-1-6	Medical electrical equipment - Part 1-6: General requirements for basic safety and essential performance - Collateral standard: Usability
EN 62366	Medical devices - Application of usability engineering to medical devices
EN 60601-2-25	Medical electrical equipment - Part 2-25: Particular requirements for the safety of electrocardiographs

Product codes and accessories

630301054 Peripheral ECG electrodes clamp AG/ agcl	
63030106Set of 4 peripheral ECG electric clamp Ag/Agl	
630301074 peripheral ECG electric clamp pediatric	
630301636 chest ECG electric suction type Ag/agcl	
63050025ECG patient cable IEC, 10 lead, plug 4 mm	
63050068ECG patient cable AHA, 10 lead, plug 4 mm	
650900057 Carrying case "Cardioline ECG 100+"	
66030031CDisposable electrodes ECG, snap, 50 pics	
66030034CDisposable electrodes ECG, tab, 100 pics	
66030036CDisposable electrodes ECF neonatal, 25 pics	
66030037CDisposable electrodes ECG banana, 60 pics	
63090236Set of 10 snap adapters for 4 mm plug	
66010051 Z-FOLD 100X150 PAPER	