

X12

Patient Monitor

Version 1.2



Main Unit Specification

Physical Specifications

Dimension	306±2 mm (W) × 309±2 mm (H) × 151±2 mm (D)
Max Weight	< 3.5 kg Standard configurations, no battery or accessories

Power Supply

Line Voltage	100 V to 240 V~
Current	1.4 A to 0.7 A
Frequency	50 Hz/60 Hz

Battery

Capacity	2550 mAh , 5100 mAh
Operating Time	2550 mAh ≥ 4 h 5100 mAh ≥ 8 h
Charge Time	2550 mAh ≤ 3.5 h, 90% charge 5100 mAh ≤ 6.5 h, 90% charge

Display

Display screen	12.1-inch color TFT screen, touch screen available
Resolution	800 × 600
Waves	A maximum of 13 waveforms can be displayed on the same screen

Recorder

Record Width	48 mm
Paper Speed	12.5 mm/s, 25 mm/s, 50 mm/s
Channels	3
Recording Types	Continuous real-time recording 8-second real-time recording 20-second real-time recording Time recording Alarm recording Trend graph recording Trend table recording NIBP review recording Arrhythmia review recording Alarm review recording Drug calculation titration recording Hemodynamic Calculation result recording 12-lead analysis recording C.O. measurement recording ST view recording QT view recording

Data Storage

Internal Temporary Memory	
Trend graph/trend	3 hrs, at 1 s resolution

review

120 hrs, at 1 min resolution

Alarm/Monitoring

Up to 200 sets

NIBP Measurement

1200 sets

Review

Up to 200 sets

Arrhythmia events

Up to 200 sets

12-lead Diagnosis

Up to 50 sets

Review

Non-volatile Memory (internal or external storage device)

A single piece of patient data maximally contains the following information:

Trend graph and trend

240 hours, at 1 min resolution

table

1200 sets

NIBP measurement

200 sets

review

200 sets

Alarm review

50 sets

Arrhythmia event

3 electrodes/5 electrodes/6 electrodes: 48 hours

12-lead diagnosis

10 electrodes: 35 hours

review

Full disclosure

50 sets

Waveforms

3 electrodes/5 electrodes/6 electrodes: 48 hours

Table

10 electrodes: 35 hours

Wi-Fi

IEEE

802.11b/g/n

Frequency Band

2.4 GHz ISM band & 5 G ISM band

Interfaces and others

VGA output (optional)

1

USB interface

2

Nurse Call / Analog Output/ Defibrillator

Synchronization (optional)

1

Network Interface

1

Data Transmission

Data Export

Ethernet / USB / Wi-Fi (Optional)

Data Management

CMS-Lite

Central Monitoring System

MFM-CMS

HIS/EMR connection

HL7

MFM-CMS / GW1 Gateway Software

ECG

Lead Mode

3-Electrodes: I, II, III

5-Electrodes: I, II, III, aVR, aVL, aVF, V₁

6-Electrodes: I, II, III, aVR, aVL, aVF, V_a, V_b

10-Electrodes: I, II, III, aVR, aVL, aVF, V₁-V₆

AHA, IEC

Electrode Standard

300 μV

Display Sensitivity

×0.125, ×0.25, ×0.5, ×1, ×2, ×4, AUTO gain

Sweep

6.25 mm/s, 12.5 mm/s, 25 mm/s, 50 mm/s

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Bandwidth (-3 dB)	Diagnosis: 0.05 Hz to 150 Hz Diagnosis 1: 0.05 Hz to 40 Hz Monitor: 0.5 Hz to 40 Hz Surgery: 1 Hz to 20 Hz Enhanced: 2 Hz ~18 Hz Customized: High-pass Filter and Low-pass Filter	Sweep 6.25 mm/s, 12.5 mm/s, 25.0 mm/s, 50.0 mm/s
CMRR	Diagnosis: > 95 dB Diagnosis 1: > 105 dB (when Notch is turned on) Monitor: > 105 dB Surgery: > 105 dB Enhanced: > 105 dB Surgery 1: > 105 dB (when Notch is turned on) Customized: > 105 dB (Low-pass Filter < 40 Hz)	Apnea Delay 10 s, 15 s, 20 s (Default), 25 s, 30 s, 35 s, 40 s
Hum Filter	> 95 dB (Low-pass Filter > 40 Hz)	NIBP
Recovery Time After Defibrillation	In diagnosis, diagnosis 1, monitor, surgery, enhanced and customized modes: 50 Hz/60 Hz (Hum filter can be turned on or off manually)	Method Oscillometry
ESU Protection	< 5 s (measured without electrodes as IEC60601-2-27:2011, Sect. 201.8.5.5.1 requires.)	Mode Manual, Auto, Continuous, Sequence
Pace Pulse Detecting	Cut mode: 300 W Coagulation mode: 100 W Restore time: ≤10 s	Measuring Interval in Auto Mode 1/2/3/4/5/10/15/30/60/90/120/180/240/360/480 min, and User Define
Lead	one among I, II, III, aVR, aVL, aVF, V1-V6	Continuous 5 min, interval is 5 s
Heart Rate		Measuring Type SYS, DIA, MAP, PR
Range	ADU: 15 bpm to 300 bpm PED/NEO: 15 bpm to 350 bpm	Measuring Range Adult Mode SYS: 25 mmHg to 290 mmHg DIA: 10 mmHg to 250 mmHg MAP: 15 mmHg to 260 mmHg
Accuracy	±1% or ±1 bpm, whichever is greater	Pediatric Mode SYS: 25 mmHg to 240 mmHg DIA: 10 mmHg to 200 mmHg MAP: 15 mmHg to 215 mmHg
Resolution	1 bpm	Neonatal Mode SYS: 25 mmHg to 140 mmHg DIA: 10 mmHg to 115 mmHg MAP: 15 mmHg to 125 mmHg
PVC		Cuff Pressure Measuring Range 0 mmHg to 300 mmHg
Range	ADU: (0 to 300) PVCs/ min PED/NEO: (0 to 350) PVCs/ min	Pressure Resolution 1 mmHg
Resolution	1 PVCs/min	Maximum Mean Error ±5 mmHg
ST value		Maximum Standard Deviation 8 mmHg
Range	-2.0 mV to +2.0 mV	Maximum Measuring Period Adult/ Pediatric: 120 s Neonate: 90 s
Accuracy	±0.02 mV or 10% (-0.8 mV to +0.8 mV), whichever is greater. Beyond this range: not specified.	Typical Measuring Period 20 s to 35 s (depend on HR/motion disturbance)
Resolution	0.01 mV	Dual Independent Channel Overpressure Protection
Arrhythmia analysis		Adult (297±3) mmHg
		Pediatric (245±3) mmHg
		Neonatal (147±3) mmHg
12-Lead ECG Synchronization Analysis		SpO₂
Average parameters of heart beat	PR interval (ms)	Measuring Range 0% to 100%
Heart rate (bpm)	QRS interval (ms)	Resolution 1%
Time limit of P wave (ms)	QT/QTC (ms)	Data update period 1 s
P-QRS-T AXIS		Accuracy Adult/Pediatric: ±2% (70% to 100% SpO ₂) Undefined (0% to 69% SpO ₂) Neonatal: ±3% (70% to 100% SpO ₂) Undefined (0% to 69% SpO ₂)
RESP		PI (Perfusion Index)
Method	Impedance between RA-LL, RA-LA	Measuring Range 0-10, invalid SI value is -?-.
Measurement lead	Options are lead I and II. The default is Lead II.	Resolution 1
RR Measuring Range	Adult: 0 rpm to 120 rpm Ped/Neo: 0 rpm to 150 rpm	TEMP
Resolution	1 rpm	Channel 2
Accuracy	Adult: 6 rpm to 120 rpm: ±2 rpm 0 rpm to 5 rpm: not specified Ped/Neo: 6 rpm to 150 rpm: ±2 rpm 0 rpm to 5 rpm: not specified	Sensor type YSI-10K and YSI-2.252K
Gain Selection	×0.25, ×0.5, ×1, ×2, ×3, ×4, ×5	Technique Thermal resistance
		Measure Parameter T1, T2, TD
		Position Skin, oral cavity, rectum
		Unit °C, °F
		Measuring Range 0°C to 50°C (32 °F to 122 °F)
		Resolution 0.1°C (0.1 °F)
		Accuracy ±0.3 °C (±0.54 °F) [±0.1 °C (±0.18 °F), exclude sensor error]
		Transient Response Time ≤ 30 s
		PR

PR (SpO₂)		
Measuring range	EDAN: 25 bpm to 300 bpm	
Accuracy	EDAN: ± 2 bpm	
Resolution	EDAN: 1 bpm	
PR (NIBP)		
Measuring range	EDAN: 40 bpm to 240 bpm	
Accuracy	EDAN: ± 3 bpm or 3.5%, whichever is greater	
Resolution	EDAN: 1 bpm	
PR (IBP)		
Measuring range	EDAN: 20 bpm to 300 bpm	
Accuracy	EDAN: 30 bpm to 300 bpm: ± 2 bpm or $\pm 2\%$, whichever is greater; 20 bpm to 29 bpm: undefined	
Resolution	EDAN: 1 bpm	
IBP		
Channel	2	
Technique	Direct invasive measurement	
Measuring range	Art: 0 mmHg to +300 mmHg PA: -6 mmHg to +120 mmHg CVP/RAP/LAP/ICP: -10 mmHg to +40 mmHg P1/P2: -50 mmHg to +300 mmHg	
Resolution	1 mmHg	
Accuracy	$\pm 2\%$ or ± 1 mmHg, whichever is greater (not including sensor)	
Unit	kPa, mmHg, cmH ₂ O	
CO₂		
Intended patient	Adult, Pediatric, Neonatal	
Measure Parameters	EtCO ₂ , FiCO ₂ , AwRR	
Unit	mmHg, %, kPa	
Measuring Range	EtCO ₂ : 0 mmHg to 150 mmHg (0% to 20%) FiCO ₂ : 0 mmHg to 50 mmHg AwRR: 2 rpm to 150 rpm	
Resolution	EtCO ₂ : 1 mmHg FiCO ₂ : 1 mmHg AwRR: 1 rpm	
EtCO₂ Accuracy	Typical conditions: Ambient temperature: (25 \pm 3) °C Barometric pressure: (760 \pm 10) mmHg Balance gas: N ₂ Sample gas flowrate: 100 ml/min All conditions	± 2 mmHg, 0 mmHg to 40 mmHg $\pm 5\%$ of reading, 41 mmHg to 70 mmHg $\pm 8\%$ of reading, 71 mmHg to 100 mmHg $\pm 10\%$ of reading, 101 mmHg to 150 mmHg $\pm 12\%$ of reading or ± 4 mmHg, whichever is greater ± 1 rpm
AwRR Accuracy	Sample Gas Flowrate	50 ml/min, 70 ml/min or 100 ml/min (default), accuracy: ± 15 ml/min
Warm-up time	Display waveform within 20 s, Reach the design accuracy within 2 minutes.	
Response time	< 4 s (with 2 m gas sampling tube, sample gas flowrate: 100 ml/min/70 ml/min) < 5.5 s (with 2 m gas sampling tube, sample gas flowrate: 50 ml/min)	
Barometric pressure compensation	Automatic (The change of barometric pressure will not add additional errors to the measurement values.)	
Zero Calibration	Support	
Calibration	Support (It is recommended to be operated by trained personnel.)	
Apnea delay	10 s, 15 s, 20 s (Default), 25 s, 30 s, 35 s, 40 s	
C.O.		
Technique	Thermodilution Technique	

Measure Parameters	C.O., TB, TI
Measuring Range	C.O.: 0.1 L/min to 20 L/min TB: 23°C to 43°C (73.4°F to 109.4°F) TI: -1°C to 27°C (30.2°F to 80.6°F)
Resolution	C.O.: 0.1 L/min TB, TI: 0.1°C (+0.1°F)
Accuracy	C.O.: $\pm 5\%$ or ± 0.2 l/min, whichever is greater TB: ± 0.1 °C (not including sensor) TI: ± 0.1 °C (not including sensor)

Safety Specifications	
Compliant with Standards	IEC 60601-1: 2005+A1 :2012; IEC 60601-1-2: 2014; EN 60601-1: 2006+A1 :2013; EN 60601-1-2: 2015; IEC 60601-2-49: 2018
Anti-electroshock Type	Class I equipment and internal powered equipment
Anti-electroshock Degree	CF
Ingress Protection	IPX1

Environmental Specifications	
Temperature	Working: +0°C to +40°C (32°F ~104°F) When the battery is charged: +0 °C to +35 °C (32°F~95°F)
Humidity	Transport and Storage: -20°C to +55°C (-4°F ~131°F) Working: 15%RH to 95%RH (non-condensing)
Altitude	Transport and Storage: 15%RH to 95%RH (non-condensing) Working: 86 kPa to 106 kPa Transport and Storage: 70 kPa to 106 kPa

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