X8 Patient Monitor Version 1.0

Data Sheet





X8 Specification	X8 Specification			
Physical Specifications				
Dimension	236±2 mm (W)) × 236±2 mm (H) × 147	7±2 mm (D)	
NA 10/ 11/	< 2.4 kg			
Max Weight	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 , 51	00 mAh		
0 " "	2550 mAh	≥ 4 h		
Operating Time	5100 mAh	≥ 8 h		
	2550 mAh	≤ 3.5 h, 90% charge		
Charge Time	5100 mAh	≤ 6.5 h, 90% charge		
Display				
Display screen	8 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			
	Continuous real-time recording			
	8-second real-time recording			
	20-second real-time recording			
	Time recording			
	Alarm recording			
	Trend graph recording			
Recording Types	Trend table recording			
recording types	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			
Data Storage				
Internal Temporary	Trond graph/trand table ravious		3 hrs, at 1 s resolution	
Memory	Trend graph/trend table review 120 hrs, at 1 min resolution			



	Alarm/Monitoring Event data	Up to 200 sets			
	NIBP Measurement Review	1200 sets			
	Arrhythmia events	Up to 200 sets			
	12-lead Diagnosis Review	Up to 50 sets			
	A single piece of patient data maximally contains the following information:				
	Trend graph and trend table	240 hours, at 1 min resolution			
Non-volatile Memory	NIBP measurement review	1200 sets			
(internal or external	Alarm review	200 sets			
storage device)	Arrhythmia event	200 sets			
	12-lead diagnosis review	50 sets			
	Waveforms	48 hours			
Wi-Fi					
IEEE	802.11b/g/n				
Frequency Band	2.4 GHz 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 Transmission				
Data Export	Ethernet / USB				
Data Management	Patientcare Viewer				
Central Monitoring System	MFM-CMS				
HIS/EMR connection	HL7				
THO/LIVITY CONNECTION	MFM-CMS / GW1 Gateway Software				
ECG					
	3-Electrode: I, II, III				
Lead Mode	5-Electrode: I, II, III, aVR, aVL, aVF, V				
Load Mode	6-Electrode: I, II, III, aVR, aVL, aVF, and leads responding to Va, Vb				
	10-Elctrode: I, II, III, aVR, aVL, aVF, V1, V2, V3, V4, V5, V6				
Lead naming style	AHA, IEC				
Display Sensitivity	1.25 mm/mV (×0.125), 2.5 mm/mV (×0.25), 5 mm/mV (×0.5),10 mm/mV (×1), 20				
(Gain Selection)	mm/mV (×2), 40 mm/mV (×4), AUTO gain				
Sweep speed	6.25 mm/s, 12.5 mm/s, 25 mm/s, 50 mm/s				
	Diagnosis: 0.05 Hz to 150 Hz				
	Surgery 1: 0.05 Hz to 40 Hz				
D	Monitor: 0.5 Hz to 40 Hz				
Bandwidth (-3 dB)	Surgery: 1 Hz to 20 Hz				
	Enhanced: 2 Hz ~18 Hz				
	Customized: High-pass Filter and Low-pass Filter				
	<u> </u>	•			



	Diagnosis: > 95 dB					
	Monitor: > 105 dB					
CMRR	Surgery: > 105 dB Enhanced: > 105 dB					
CIVIRR	Surgery 1: > 105 dB (wher	Notch is turned on				
	Customized: > 105 dB (Lo	•				
	`	. ,				
	> 95 dB (Low-pass Filter > 40 Hz)					
Lluma Ciltan	In diagnosis, Surgery 1, monitor, surgery, enhanced modes: 50Hz/60 Hz					
Hum Filter	(Hum filter can be turned on or off manually)					
Recovery time after defibrill	ation <5 s					
	Cut mode: 300 W					
ESU Protection	Coagulation mode: 100 W					
	Restore time: ≤10 s					
Pace pulse detecting lead	one among I, II, III, AVR, A	VL, AVF, V1, V2, V3,V4, V	5, V6			
Heart Rate						
Range						
rango	Ped: 15 bpm to 350 bpm					
Accuracy	±1% or ±1 bpm, whicheve	r is greater				
Resolution	1 bpm					
PVC						
Range	Adult: 0 to 300 PVCs/ min					
rvarige	Ped/Neo: 0 to 350 PVCs/ min					
Resolution	1 PVCs/min					
ST value						
Range	-2.0 mV to +2.0 mV					
Accuracy	±0.02 mV or 10% (-0.8 mV to +0.8 mV), whichever is greater. Beyond this range: not specified.					
Resolution	0.01 mV					
Arrhythmia analyses						
Asystole	Sustain VT	V-Fib/V-Tach	ExtremeTachy			
ExtremeBrady	V-Tach	Vent Brady	Tachy			
Brady	Wide QRS Tachy	Non-Sustain VT	Afib			
Vent Rhythm	Acc. Vent Rhythm	Pause	Pauses/min High			
PVCs High	R on T	PVC Bigeminy	PVC Trigeminy			
Pacer not Pacing	Pacer not Capture Missed Beat VEB					
PVC	Couplet Run PVCs Multiform PVCs		Multiform PVCs			
IPVC	Irr Rhythm PAC Bigeminy PAC Trigeminy					
Low Voltage(Limb)						
J ()	l .		L			



12-lead ECG synchroniz	ation analy	sis		
Average parameters of heart beat				
Heart rate (bpm)	Heart rate (bpm)			
Time limit of P wave (ms)				
PR interval (ms)				
QRS interval (ms)				
QT/QTC (ms)				
P-QRS-T AXIS				
RESP				
Method	Trans-thora	ncic impedance: R-F(RA-LL), R-L (RA-LA)		
Measurement lead	Options are lead I and II			
Managina Danas	Adult	0 rpm to 120 rpm		
Measuring Range	Ped/Neo	0 rpm to 150 rpm		
Resolution	1 rpm			
		6 rpm to 120 rpm: ±2 rpm		
Accuracy	Adult	0 rpm to 5 rpm: not specified		
Accuracy	Ped/Neo	6 rpm to 150 rpm: ±2 rpm		
	Ped/Neo	0 rpm to 5 rpm: not specified		
Gain Selection	×0.25, ×0.5, ×1, ×2, ×3, ×4, ×5			
Sweep	6.25 mm/s, 12.5 mm/s, 25.0 mm/s, 50.0 mm/s			
Apnea Alarm Time	10 s, 15 s, 20 s, 25 s, 30 s, 35 s, 40 s			
NIBP				
Method	Oscillometr	у		
Mode	Manual, Auto, Continuous			
Measuring Interval in Auto Mode	1/2/3/4/5/10/15/30/60/90/120/180/240/360/480 min			
Continuous	5 min, interval is 5 s			
Measuring Type	SYS, DIA, MAP, PR			
Measuring Range	Adult Mode	SYS: 25 mmHg to 290 mmHg DIA: 10 mmHg to 250 mmHg MAP: 15 mmHg to 260 mmHg		
	Pediatric Mode	SYS: 25 mmHg to 240 mmHg DIA: 10 mmHg to 200 mmHg MAP: 15 mmHg to 215 mmHg		
	Neonatal Mode	SYS: 25 mmHg to 140 mmHg DIA: 10 mmHg to 115 mmHg MAP: 15 mmHg to 125 mmHg		
Cuff Pressure Measuring Range	0 mmHg to 300 mmHg			



Pressure Resolution	1 mmHg			
Maximum Mean Error	±5 mmHg			
Maximum Standard	120 mining			
Deviation	8 mmHg			
	Adult/			
Maximum Measuring	Pediatric	120 :	S	
Period	Neonatal 90 s			
Typical Measuring Period	20 s to 35 s (depend on HR/motion disturbance)			
	Adult 297 mmHg±3 mmHg			
Overpressure Protection	Pediatric	245 mmHg±3 mmHg		
'	Neonatal	147 mmHg±3 mmHg		
PR				
Measuring range	40 bpm to 2	240 bp	om	
Accuracy	±3 bpm or 3	3.5%,	whichever is greater	
SpO ₂				
Measuring Range	0% to 100%	6		
Resolution	1%			
Data update period	1 s			
			±2% (70% to 100% SpO ₂)	
	Adult/Pediatric		Undefined (0% to 69% SpO ₂)	
Accuracy			` '	
	Neonatal		$\pm 3\%$ (70% to 100% SpO ₂) Undefined (0% to 69% SpO ₂)	
PI (Perfusion Index)			Official Control Contr	
Measuring Range	0-10			
Resolution	1			
Pulse Rate	'			
Measuring Range	25 bpm to 300 bpm			
Accuracy	±2 bpm			
TEMP	±£ bpiii			
Channel	1	1		
Sensor type	YSI-10K and YSI-2.252K			
Technique	Thermal resistance			
Measure Parameter	T1, T2			
Position	Skin, Oral, Rectum			
Unit	°C , °F			
Measuring Range	0°C to 50°C (32 °F to 122 °F)			
Resolution	0.1°C (0.1°F)			
1 COOLULIOIT				
Accuracy	Accuracy (not including sensor): ±0.1°C Sensor accuracy: ≤ ±0.2°C			
Transient Response Time	·			
Transient Nesponse Time	= 50 5			



CO ₂					
Intended patient	Adult, Pe	diatric, Neonatal			
Measure Parameters	EtCO ₂ , F	EtCO ₂ , FiCO ₂ , AwRR			
Unit	mmHg, %	, kPa			
Managina Danga	CO ₂ 0 mmHg to 150 mmHg (0% to 20%)				
Measuring Range	AwRR	2 rpm to 150 rpm			
	EtCO ₂	1 mmHg			
Resolution	FiCO ₂	1 mmHg			
	AwRR	1 rpm			
		±2 mmHg, 0 mmHg to 40 mmHg			
		±5% of reading	, 41 mmHg to 70 mmHg	Respiratory rate≤60	
		±8% of reading	, 71 mmHg to 100 mmHg	rpm	
Accuracy	EtCO ₂	±10% of reading	g, 101 mmHg to 150 mmHg		
Accuracy		±12% of reading	g or ±4 mmHg, whichever is	Respiratory rate>60 rpm	
	AwRR	<u> </u>			
Sample Gas Flowrate	50 ml/mir	50 ml/min, 70 ml/min or 100 ml/min(default), accuracy: ±15 ml/min			
Warm-up time			0 s, Reach the design accura		
Response time	<4 s				
Barometric pressure	A 1 1				
compensation	Automatic				
Zero Calibration	Support				
Calibration	Support				
Apnea alarm delay	10 s, 15 s	10 s, 15 s, 20 s, 25 s, 30 s, 35 s, 40 s			
Safety Specifications					
	IEC 60601-1: 2005+A1 :2012; IEC 60601-1-2: 2014;				
Compliant with Standards	EN 60601-1: 2006+A1 :2013; EN 60601-1-2: 2015;				
	IEC 60601-2-49: 2011				
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)		
'	Transport and Storage		-20°C to +55°C (-4 °F ~131 °F)		
Humidity	Working		15%RH to 95%RH (non-condensing)		
,	Transport and Storage		15%RH to 95%RH (non-condensing)		
Altitude	Working		86 kPa to 106 kPa		
	Transport and Storage 70 kPa to 106 kPa				

^{*} Specifications are subject to change without prior notice

