Vyntus® CPX

Technical Specifications



Ergospirometry (physiological measurement)		
Parameter	Measurement range	Accuracy
Ventilation (V'E)	0 to 300 L/min	2 % or 0.50 L/min
O2 uptake (V'O2)	0 to 7 L/min	3 % or 0.05 L/min
CO2 production (V'CO2)	0 to 7 L/min	3 % or 0.05 L/min
RER	0.6 to 2.0	4 % or 0.04

Volume sensor (physiological measurement)		
Parameter	Measurement range	Accuracy
Volume	0 to 10 L	2 % or 0.05 L
Flow	0 to 15 L/s	3 % or 0.07 L/s
Resolution	3 mL	
Resistance	< 0.1 kPa/L/s at 15 L/s	
ATS	Compliant	

O2 analyser	
Туре	Fully digital, high speed analyser, based on electrochemical principle
Measurement range	0 to 100 Vol%
Accuracy	0.05 Vol% or 0.2 %
Resolution	0.01 Vol%
Typ. rise time (T10-90)*	75 ms
Operating life	Typical 2 years

CO2 analyser		
Type Fully digital, high speed analyser based on principle of infrared absorption		
Measurement range	0 to 15 Vol%	
Accuracy	0.05 Vol% or 1%	
Resolution	0.01 Vol%	
Typ. rise time (T10-90)*	75 ms	

Ambient measurement		
Ambient unit	Measurement range	Accuracy
Temperature sensor	-40 to +85 °C (-40 to +185 °F)	0.5 °C at 20 °C (68 °F)
Humidity sensor	10 to 95 % relative humidity	4 % relative humidity
Vyntus CPX	Measurement range	Accuracy
Air pressure sensor	30 to 110 kPa (300 hPa to 1100 hPa)	0.25 kPa (2.5 hPa) at 0 to 65 °C (32 to 149 °F)

Notes

^{*} Rise time after digital filtering. During every gas calibration the filter coefficients are specifically optimized for the sensors.



SpO2 sensors	
Displayed oxygen saturation range (SpO2)	0 to 100 %
Displayed pulse rate range	18 to 321 beats per minute (BPM)
SpO2 accuracy (arms*)	± 3 digits @ 70 to 100 %

25 - 80 L/min

Canopy module specifications

Dilution flow range

Power supply	
Vyntus CPX with Vyntus CART 3.0	
Mains voltage	115 V; 230 V, AC 50/60 Hz
Power input	690 VA
Electrical safety	Protection class I
Vyntus CPX without cart	
Туре	Magic Power MPM-X125
Mains input voltage	100 - 240 V, AC 47 - 63 Hz
Power consumption	1.5 - 1 A
Output voltage	24 V DC
Output	120 VA / 5 A
Electrical safety	Protection class I

Moisture protection	
Vyntus CPX with Vyntus CART 3.0	no IP protection
Vyntus CPX without cart	IP 20

Ambient conditions	
Temperature	+10 °C to +34 °C (+50 °F to 93.2 °F)
Relative humidity	20 to 80 % RH,
	non-condensing
Ambient pressure	700 to 1060 hPa (525 to 795 mmHg)
Altitude	≤ 3000 m

Classification of applied parts	
Type applied part	В

Category according to MDD 93/42/EEC (2007)	
Complete system	Active class IIa medical product

Pressure reducer	
Туре	Test gas pressure reducer
Inlet pressure	200 bar (2901 psi)
Outlet pressure	2 bar fixed (29 psi)
1 outlet	G 1/4 internal screw thread

D' (186 ' L)		
Dimensions / Weight		
Vyntus CPX module	22 W 20 F D 42 F	11/42// 44.2// 5.2//
Dimensions	33 cm W x 28.5 cm D x 13.5 cm	H (13" x 11.2" x 5.3")
Weight	4.08 kg (8.9 lbs)	
Vyntus CPX with Vyntus CART 3.0		
Dimensions total	61 - 82.5 cm W x 62 cm D x 172 (24 - 32.5" x 24.4" x 67.7 - 77.6	- 197 cm (depending on monitor assembly) ")
Weight total	82 kg (180.8 lbs) inclusive CardioSoft ECG, KISS suction unit and 2 x 19" TFT monitors	
Castors	4 twin swivel castors Ø 10 cm (3.9	9"), lockable, conductive
Shelves	Number	2
	Platform area	45 cm W x 36 cm D (17.7" x 14.2")
	Load capacity	20 kg (44 lbs) distributed load
Keyboard drawer	Keyboard platform area	40 cm W x 20 cm D (15.7" x 7.9")
with mousepad (left / right)	Load capacity	3 kg (6.6 lbs) distributed load
Drawer	Interior area	34 cm W x 24 cm D x 8 cm H
		(13.4" x 9.4" x 3.1")
	Platform area	45 cm W x 36 cm D (17.7" x 14.2")
	Load capacity shelf	15 kg (33 lbs) distributed load
	Load capacity drawer	3 kg (6.6 lbs) distributed load
Computer holder with two tension belts	Platform area	24 cm W x 3.7 cm D (9.4" x 1.5")
'	Load capacity	5 kg (11 lbs)
Dual monitor mount	Max. monitor width	49.4 cm (19.45")
	Load capacity for each monitor	14 kg (30.9 lbs)
34'' curved monitor as dual monitor	Dimensions	82.5 cm W x 7.3 cm D x 37.2 cm H
application		(32.5" x 3.0" x 14.7")
	Weight	8 kg (17.6 lbs)
/yntus CPX Canopy option		
Vyntus Canopy module	Dimensions	17.5 cm W x 25.5 cm D x 11 cm H
		(6.9" x 10" x 4.3")
	Weight	2.62 kg (5.8 lbs)
Holder for hood	Dimensions	40 cm W x 53 cm D x 28 cm H
		(15.7" x 20.9" x 11")
	Weight	1.2 kg (2.6 lbs)
Canopy tube	Length	183 cm (72")
Foil	Dimensions	150 cm W x 160 cm D (59.1" x 63.0"),
		50 μm thickness
	Material	LDPE, biocompatible

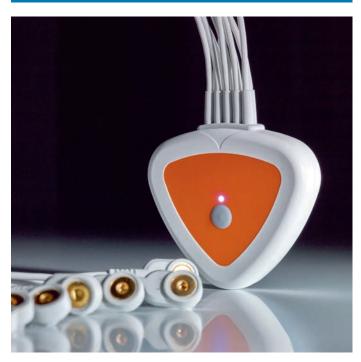
Standards, directives and market clearances		
Standards	EN 60601-1, EN 60601-1-2, EN 62304, EN 62366, EN ISO 14971, EN ISO 10993-1	
Directives	93/42/EEC amended by 2007/47/EC, RoHS 2011/65/EU compliant	
Market clearances	CE for Vyntus CPX and Vyntus CPX Canopy option, FDA 510(k) clearance for	
	Vyntus CPX only	

Notes

^{*} \pm 1 Arms represents approximately 68 % of measurements.

Vyntus® ECG

Technical Specifications





ECG Amplifier, 10 electrodes, 12 lea	ds	
Application	ECG recording	
ECG leads	according to Einthoven, Wilson,	Goldberger
Bandwidth	0.05 - 150 Hz digital	
Sampling rate per channel	500 Hz	
Pacemaker detection	4000 Hz	
Resolution	< 2.5 μV/bit	
Connection to the PC	Wireless, Bluetooth®	
Bluetooth®	Bluetooth® module	Panasonic PAN 1322
	Bluetooth® standard	V2.1, class 2 device
	Output power	2.5 dBm, max. 4.5 dBm
	Frequency range	2400 - 2483.5 MHz
	FCC identifier	T7VEBMU
	IC identifier	216QEBMU
	Bluetooth® QD ID	B021246
Connection of the electrodes	4 mm snap, gilded	

Dimensions / Weight		
ECG housing	Dimensions	9.5 cm W x 8.5 cm D x 2 cm H (3.7" x 3.3" x 0.8")
	Weight	0.2 kg (0.4 lbs)
Length of electrode cables	V1 - V6	60 cm (23.6")
	LL	45 cm (17.7")
	LA / RL	70 cm (27.6")
	RA	80 cm (31.5")

Power supply		
Supply voltage	1.5 V DC AA battery or 1.2 V DC AA NiMH rechargeable battery, at least 2500 mAh	
Power input (max.)	500 mA	
Service life with fully charged battery	at least 6 hours	

Moisture protection		
Type applied part	IP 20	

Ambient conditions	
Temperature	+10 °C to +34 °C (+50 °F to 93.2 °F)
Relative humidity	20 to 80 % RH, non-condensing
Ambient pressure	700 to 1060 hPa (525 to 795 mmHg)

Classification of applied parts	
Type applied part	CF, defibrillator proof

Category according to MDD 93/42/EEC (2007)	
Complete system	Active class IIa medical product

Standards, directives and market clearances	
Standards	EN 60601-1, EN 60601-1-2, EN 60601-2-25, EN 62304, EN 62366, EN ISO 14971, EN ISO 10993-1
Directives	93/42/EEC amended by 2007/47/EC, RoHS 2011/65/EU compliant
Market clearances	CE, FDA 510(k) clearance

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