

FeNO monitoring system

Vivatmo *me*

FEATURES

Vivatmo <i>me</i>	
Measuring range	5 ppb to 300 ppb
Accuracy	±5 ppb below 50 ppb, ±10 % ≥50 ppb, ±15 % ≥ 160 ppb expressed as the upper/lower confidence limit of 95 %
Precision	±5 ppb below 50 ppb, ±10 % ≥50 ppb, ±15 % ≥ 160 ppb expressed as one standard deviation for replicate measurements with the same instrument
Lifetime, Device service life	1,000 measurements
Memory capacity	1,000 measurements



SPECIFICATIONS

Handheld	
Sensor	Chemical field-effect transistor
Display	Digital LCD display
Power source	4 AAA batteries 1.5 V, Useful life of batteries: Alkaline up to 25 attempts, lithium/iron disulfide up to 60 attempts
Weight	170 g
Dimensions	4.0 cm x 5.4 cm x 22.4 cm
Package contents	Vivatmo <i>me</i> device, package of 5 disposable mouthpieces, 4 AAA batteries, protective cap, Instructions for Use

Disposable Mouthpiece (accessory)	
Single Use	Measurement within 15 minutes after opening the pouch. Useful life limited to 5 attempts and expiration date.
Shelf life	2 years from manufacturing.

ELECTRONICAL AND SAFETY INFORMATION

Applied part:	Type BF as per EN 60601-1 for handheld and disposable mouth-piece when attached
Maximum surface temperature:	58 °C, touch time < 60 seconds
Electrical safety:	ME device with internal supply, tested as per EN 60601-1-11 IP 22 (protection against solid particles >12,5 mm and ingress of dripping water when tilted up to 15°) for basic safety but not for function
Data transfer:	Bluetooth® Smart (low energy), 2.4 GHz frequency band
Electromagnetic emissions	CISPR 11 Group 1 (battery operated)
Electromagnetic immunity	EN 61000-4-2, EN 61000-4-3 (battery operated), EN 61000-4-8

REACH REGULATION

Reporting Requirement according to Article 33 of the REACH Regulation No. 1907/2006: The pump within our product contains lead monoxide.

ENVIRONMENTAL SPECIFICATIONS

	Operation	Transportation and Storage
Temperature	+15°C to +35°C	Transportation: -20°C to +60°C Storage: +5°C to +40°C
Relative humidity (non-condensing)	15 % to 85 %	10 to 85 %
Air pressure (corresp. to 0 - 2,000 m a.s.l.)	780 hPa to 1,100 hPa	780 hPa to 1,100 hPa
Ambient NO concentration	< 200 ppb	

ELECTROMAGNETIC COMPATIBILITY (EMC)

Important information regarding electromagnetic compatibility (EMC)

This device complies with EN60601-1-2:2015 for EMC with the objective to avoid insecure product situations. This standard regulates the levels of immunity against electromagnetic interferences and the maximum electromagnetic emission values for medical equipment. This medical device manufactured by the company complies with the standard EN60601-1-2:2015 both in terms of immunity and of emissions and does therefore not need any service and maintenance regarding EMC and ESD over lifetime.

Please note that portable and mobile HF communication systems may interfere with this device even if compliant with CISPR emission requirements. Do not staple the device or use any mobile phones or other devices generating strong electrical or electromagnetic fields. This could result in malfunction of the medical device and may create a potentially insecure situation. Portable RF communication devices are not to be used closer than 30 cm next to the device.

Guidance and manufacturer's declaration – electromagnetic emissions

The Vivatmo *me* is intended for use in the electromagnetic environment specified below. The customer or the user of Vivatmo *me* should assure that it is used in such an environment.

Emissions test	Compliance	Electromagnetic environment – Guidance
RF emissions CISPR 11	Group 1	The Vivatmo <i>me</i> uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.
RF emissions CISPR 11 Harmonic emissions IEC 61000-3-2	Class B n/a	The Vivatmo <i>me</i> is suitable for use in all establishments, including domestic establishments and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.
Voltage fluctuations/ flicker emissions IEC 61000-3-3	n/a	

Guidance and manufacturer's declaration – electromagnetic immunity

The Vivatmo *me* is intended for use in the electromagnetic environment specified below. The customer or the user of the Vivatmo *me* should assure that it is used in such an environment.

Immunity test	IEC 60601 Test level	Compliance level	Electromagnetic environment – Guidance
Electrostatic discharge (ESD) IEC 61000-4-2	±8 kV contact ±15 kV air	±8 kV contact ±15 kV air	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30%.
Electrical fast transient/burst IEC 61000-4-4	n/a	n/a	n/a
Surge IEC 61000-4-5	n/a	n/a	n/a
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	n/a	n/a	n/a
Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	30 A/m	30 A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.

Notes: UT is the AC mains voltage prior to application of the test level.

Immunity test	IEC 60601 Test level	Compliance level	Electromagnetic environment – Guidance
Conducted RF IEC 61000-4-6	10 Vrms 150 kHz to 80 MHz	n/a	Portable and mobile RF communications equipment should be used no closer to any part of the Vivatmo <i>me</i> including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter. Recommended separation distance: $d = 1.2\sqrt{P}$ $d = 1.2\sqrt{P}$ 80 MHz to 800 MHz $d = 2.3\sqrt{P}$ 800 MHz to 2.7 GHz
Radiated RF IEC 61000-4-3	10 V/m 80 MHz to 2.7 GHz	10 V/m 80 MHz to 2.7 GHz	

Where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the recommended separation distance in meters (m).

Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey, should be less than the compliance level in each frequency range. Interference may occur in the vicinity of equipment marked with the following symbol:



Test specifications for immunity test against high-frequent radio-based electronic communications systems

Test frequency MHz	Modulation ^b	Immunity testing level V/m
385	Pulse modulation ^b 18 MHz	27
450	FM ± 5 kHz Deviation 1 kHz Sine	28
710		
745	Pulse modulation ^b 217 MHz	9
780		
810		
870	Pulse modulation ^b 18 MHz	28
930		
1720	Pulse modulation ^b	
1845	217 MHz	28
1970		
2450	Pulse modulation ^b 217 MHz	28
5240		
5500	Pulse modulation ^b	9
5785	217 MHz	

^b The carrier must be modulated by a square-wave signal with a duty cycle of 50%.