

NeXus-4

User Manual



Version 2.0

CE0344

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1 Service and support

1.1 About this manual

This manual is intended for the user of the NeXus-4 system – referred to as 'product' throughout this manual. It contains general operating instructions, precautionary measures, maintenance instructions and information for use of the product. Read this manual carefully and familiarize yourself with the various controls and accessories before starting to use the product.

This product is exclusively manufactured by Twente Medical Systems International B.V. for Mind Media B.V. Distribution and service is exclusively performed by or through Mind Media B.V.

Mind Media Support can be reached via email (support@mindmedia.nl) or by phone during office hours (CET). Visit our Support section on www.mindmedia.com, because this may resolve your problem. Always provide as much information on your problem as possible, including serial numbers of the products.

Address

Mind Media B.V.

Louis Eijssenweg 2B 6049CD Herten The Netherlands

Phone +31 (0) 475410123 Website www.mindmedia.com/



In case of need for repair **ALWAYS** first contact Mind Media Support. The support staff will supply you with an RMA number in case a return is required. Never ship products back to Mind Media without this authorization and/or RMA number.

1.2 Warranty information

The product, except its cables and accessories, is warranted against failure of materials and workmanship for a period of 2 years from the date of delivery. Cables and accessories have a warranty period of 6 months.

Repairs can only be performed by the manufacturer. Warranty will terminate automatically when the product is opened by any person other than qualified personnel (authorized by Mind Media).

The warranty does not cover the following:

- failure resulting from misuse, accident, modification, unsuitable physical or operating environment, or improper maintenance
- failure caused by a product for which Mind Media is not responsible
- damage resulting from use of non-approved accessories
- any non-Mind Media products

The warranty is voided by removal or alteration of identification labels on the product or its parts. Warranty is also voided in case seals on the enclosure are broken. Mind Media does not warrant uninterrupted or error-free operation of wired or wireless data transmission.

Any technical or other support provided for a product under warranty, such as assistance with "how-to" questions and those regarding device set-up and installation, is provided without warranty.

2 Safety information

This section contains general warnings, explanation of markings, limitations of use, safety measures, and precautionary measures that are important for the safe use of the product.

2.1 Explanation of markings

This section explains the various markings and symbols used with the product.



Manual contains important safety information



Attention: read important safety information



Important information / guidance for use



Consult instructions for use



Device has type CF applied parts



Ingress protection rating



Keep dry

CE-certified (93/42/EC Annex XII), see declaration of conformity



Identification of the manufacturer



Reference number



Serial number



Contains transmitter module



Contains Bluetooth transmitter



Special EU instructions for disposal are applicable to a product on which this symbol is placed. The Maintenance section of this manual contains information on how to dispose of this equipment.

2.2 Limitations of use

Limitations of use

- Under federal law (only applicable to the USA) this product may only be sold by or on the order of a physician or licensed practitioner.
- The product may only be used under the constant supervision of or on the instructions of a physician or other authorized medical professional.

The product is NOT intended for:

- critical patient monitoring
- use in life support systems

The product is NOT to be:

- used near MRI equipment
- exposed to ionizing radiation
- used on patients undergoing electro surgery
- used in oxygen rich environments (concentration > 25 % at 1 atm)



The product is NOT:

- suitable for use in an inflammable mixture of anesthetics or agents and air, oxygen or nitrous oxide
- defibrillator proof
- suitable for sterilization

Do not use, store or transport the equipment outside the specified environmental conditions, this may damage the product.

Do not store or use in environments with Magnetic Resonance Imaging (MRI) equipment, or equipment capable of emitting diagnostic levels of ionizing radiation.

Apart from the above, there are no contra-indications. There are no known side effects from the use of this product.

2.3 Safety measures and warnings

	Warnings			
	 Do not combine the use of the product with any other electronic equipment, except those specified in this manual. Doing so may impair the product's emissions and immunity regarding EMC. 			
• The product can only be used with the accessories designated b manufacturer. The use of other accessories may impair the pro emissions and immunity regarding EMC.				
	 The accessories supplied with the device can only be used with Mind M approved devices. 			
	Sensors with their own power are not to be connected to any of the input			
	• Transmission quality decreases when there are other radio devices in the neighborhood. The wireless transmission may be interfered with by other equipment.			
	• The product should not be used adjacent to or stacked with other equipment. If this is required, then it should be observed if normal operation of the product in that configuration is confirmed.			

Warnings

- Before batteries are replaced, disconnect the patient from the device, and the device is switched off (Indicator led is off). Both batteries have to be replaced simultaneously, and have to be of the same type. Note the orientation of the batteries.
- Do not use batteries that contain lithium.
- Do not use rechargeable batteries.
- Do not immerse the product in any liquid.
- The product is to be kept dry. If operated out of office, it must be fitted in a carrying case that provides an ingress protection of at least IP02.
- Do not expose the product to direct sunlight, heat from a source of thermal radiation, excessive amounts of dust, moisture, vibrations, or mechanical shocks.
- Do not incinerate any part of the product.
- If any liquids or moisture penetrate the product or any part thereof have the product checked by the manufacturer.
- Take care in arranging patient and sensor cables to avoid risk of patient entanglement or strangulation.



- The manufacturer cannot guarantee safety and performance of the product when used in conjunction with accessories that are not manufactured or approved by the manufacturer.
- No modification of this product is allowed. The product should not be tampered with.
- Do not touch the electrical connectors that are accessible inside the battery bay if the battery cover is removed.
- Do not touch the connector pins of interface plugs or receptacles.
- Do not open the product using tools.
- The product is not to be used when it is clearly damaged or wet, or suspected to be wet inside.
- The product connectors contain nickel, avoid prolonged skin contact with patients with nickel allergy.
- Disposable electrodes, which are used for electrophysiological measurements, may be a biohazard. Handle, and when applicable dispose of these materials in accordance with accepted medical practice and any applicable local, state and federal laws and regulations.
- Reusable electrodes present a potential risk of cross-infection especially when used on abraded skin, unless they are restricted to a single patient
- To prevent contamination: store electrodes in a separate bag within the packaging.
- Except for the batteries there are no user serviceable parts within the product. Repairs can only be performed by the manufacturer.

Warnings

• When connecting the system in an IT-network:



- Simultaneous connection of other equipment to the same Bluetooth network may result in previously unidentified risks to patients, operators or third parties. Such risks must be identified, analyzed, evaluated and controlled. Subsequent changes to the Bluetooth network can introduce new risks that require additional analysis. Changes to the IT-network include: changes to its configuration, connecting additional items, disconnecting items, updates and upgrades of connected equipment.
- Clean the product only according to the cleaning instructions in this manual. Before cleaning, make sure the device is switched off. Never use any aggressive chemicals to clean the product.

2.4 Precautionary measures

Precautionary measures

• When the distance between the Bluetooth PC receiver and the device increases or when there are conducting materials in the straight line between the Bluetooth PC receiver and the device, the signal transmission is not as reliable.



- Do not use an operating cellular phone within 50 cm of the device to avoid excessive noise on the signals.
- Sharp bends or winding the cables in a loop smaller than 5 cm diameter may damage the cables.
- When the product is not in use for a longer time (more than a few days) the batteries have to be removed to prevent damage in case they start leaking.
- Dispose of batteries according to local regulations.

2.5 Disclosure of residual risk

The risk analysis process for the product has determined that there are no residual risks which need to be disclosed for the product.

2.6 Information for lay operators

Operators must convey the following information to patients in case they carry the product out of the professional's office:

- Precautions to be taken with respect to environmental temperature and EM fields, ingress of liquid
- That wireless equipment (network, phone, walkie-talkie) should be kept >4 m away from the device
- How to deal with accessories and accessory cables
- How to deal with information provided by indicators and the display
- How to replace batteries

3 Product overview

3.1 **Product components**

The product comprises the following functional components:

#	Item	Description
1.	NeXus-4 device	The data acquisition device (NeXus-4 Amplifier).
2.	Carrying bag The carrying bag to be used when the system is used in portabl measurement configurations.	
3.	Software*Device driver with application programming interface. The device is(PC Driver)supplied to you either on CD or via email (download).	
4.	Accessories	Various other electrodes, sensors and accessories may be delivered with the package. Refer to the list of supported active sensors to see which are supported by the product.
5.	Suitcase	Suitcase is used for storage of the product when not in use.
6.	Accompanying documentation	Documentation such as Labels, User Manual and certificates.

* Optional: Software may be sent to you as download by email

The device supports Mind Media approved or manufactured active sensors. A list of supported sensors can be found on the website: www.mindmedia.com

3.2 Intended use

The product is intended to be used for acquisition of (electro)-physiological signals by, or under supervision of, a physician. The user must have knowledge of current good practice in physiological measurement in science and clinical application. The product is intended to be used within a clinical or home environment and can be used stationary or ambulatory.

Electrophysiological signals (e.g. EEG, EMG or ECG) are measured via the unipolar or bipolar inputs on the device via electrode leads connected to a patient or subject. Other physiological parameters, such as respiration, body position, body movement and temperature are measured using the auxiliary input channel. These types of signals require additional sensor interface modules.



The system does **not** perform any signal interpretation or signal analysis. This is left to the researcher/physician.

 \wedge

The system is **not** intended for use in a life supporting system.

The device transfers the data to the PC by means of a wireless (Bluetooth) connection, where the signals can be viewed or stored for further processing. The device is powered by a set of batteries.

3.3 NeXus-4 views



Front View

	Description
GND	Patient Ground input
DUAL BIPOLAR INPUT (A&B)	Input for dual bipolar input. Two bipolar pairs can be connected to this input.
C & D	Auxiliary input for (active) sensors. A list of compatible sensors can be found on the website: www.mindmedia.com

3.4 User interface

On/Off Button

The device will switch on when the On/Off button is pressed shortly. When the device is transmitting data (via optical fiber or Bluetooth), the On/Off button acts as a Marker button. This will result in a signal in the digital channel of the product. In this case, the device will not shut down.

Status LED

The table below states all possible states of the LED indicators of the device

LED Indicator	Meaning	Description
OFF	System off	The LED indicator does not show LED. The device is off.
ORANGE	Startup	At startup the LED lights up orange for about 2-3 seconds. If LED does not turn green, replace batteries or contact Mind Media support, there may be a defect.
GREEN	Idle, running	LED lights up green when it is transmitting data or ready to be used.
ORANGE BLINKING (~2x per seconds)	Recording error	A recording error occurred
ORANGE/GREEN BLINKING Battery low (~1x per second)		Battery is running low.

3.5 Patient connections

Patient Ground

The patient ground should always be connected in order to keep the amplifier in range. The location of the patient ground is ideally away from your measurement electrodes.

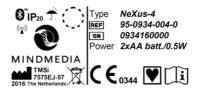
Patient Lead Connectors: Bipolar, Auxiliary

The number and type of inputs on your device depend on the configuration you have. NeXus-4 devices exist in many different configurations, varying from 2 up to 3 input channels. In general there are two types of patient connection inputs on the device: bipolar or auxiliary.

Type of input	Connector	Description	
Patient Ground	GND	Patient ground is used to keep the amplifier in range. The input is marked with 'GND'.	
Dual Bipolar input		Used for differential measurements. Leads that fit in the bipolar inputs have two bipolar cables with 2 snaps per cable going to the patient. The bipolar input uses a 6 pin connector.	
Auxiliary		Used for sensors that require (5V) power or additional sensor modules. The auxiliary input is a 5 pin connector. See list of compatible sensors on the website www.mindmedia.com.	
Important	• Note the difference in number of pins of the different inputs.		
		e orientation of the connector is marked with a red dot on both vice connector and accessories. Make sure the dots are aligned fore plugging the cable in the device.	
	• Do not rotate the connector during insertion of the cable.		

3.6 Device label

The device label can be found at the bottom of the device. It contains the REF code, Serial Number, power requirements and other properties of the device. Use the REF number and-or serial number (SN) to look up the channel specifications as listed in chapter 8



4 Instructions for use

4.1 Software

Software, that is needed to use the product, is supplied to you by email as download or by one or more CDs in the package. It is recommended to download the most up-to-date software via www.mindmedia.com. Once installed and activated, this step can be skipped.

PC requirements

Hardware		
Processor: > 1 GHz		
• RAM: > 1 GB		
• HDD: > 50 GB (> 250 GB recommended)		

• Internet connection or CD/DVD Drive

Operating system

Windows

- Windows 10 (64-bit)
- Windows 8.1 (64-bit)
- Windows 7 (32-bit / 64-bit)

Important

Disconnect all Mind Media products from the PC before installing any Mind Media software.



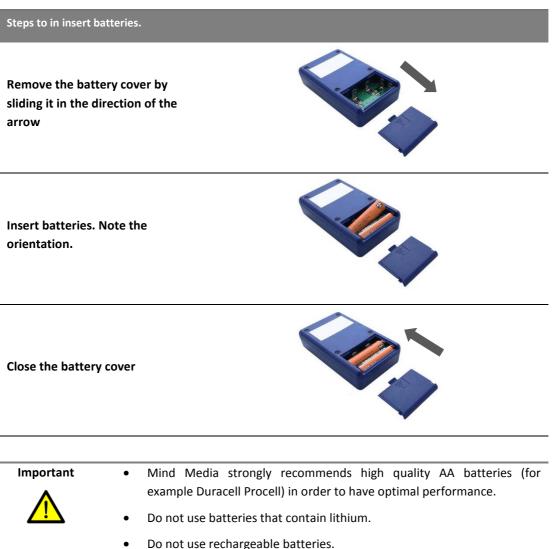
It is recommended to uninstall older versions of the driver before installing new drivers.

Mind Media PC Driver

Start the installer by executing *setup.exe*. The Mind Media PC-Driver Setup Wizard starts and guides you through the process of installation of the driver. Follow the steps on screen.

4.2 Powering the NeXus-4

The NeXus-4 can be powered using batteries. Follow the steps below to insert batteries.



Power Saving Mode

When the device is not in use (e.g. there is no active connection (pairing) with the PC), the NeXus-4 will switch off after 5 minutes to save its power.

4.3 Transfer data to PC

Bluetooth IT-network connections are supported by the product. The purpose of the IT-network connection is for device control and/or data transfer. The intended information flow is:

- Control from a PC to the device
- Raw data from the device to the PC

The supported versions of the IT network connections are:

• Bluetooth: 1.1 and higher

The following section describe the installation of the Bluetooth IT-network connection.

Please note: No hazardous situations have been identified for the product due to loss of the ITnetwork functionality.

Wireless transmission: Bluetooth interface

- 1. Insert the Bluetooth dongle in a USB port. Wait for Windows Update to install the drivers for the Bluetooth dongle.
- 2. After Windows has finished installing the Bluetooth drivers, click the Bluetooth tray icon in the right bottom corner of the Windows taskbar.
- 3. Click, 'Add a device'. If you do not see the Bluetooth tray icon, click Start > Devices and Printers > Add a device
- 4. Windows will start scanning the environment. Make sure the NeXus-4 is powered on. The NeXus-4 should automatically show up in the list.
- 5. Select the device and click '*Next*'. Windows will ask for a pairing code. The pairing code consists of the last four digits of the serial

Add a Device
Allow a Device to Connect
Show Bluetooth Devices
Send a File
Receive a File
Join a Personal Area Network
Open Settings
Remove Icon

number. The serial number can be found on the back of the NeXus-4 on the silver label next to 'SN'. Click *Next* to finish the Bluetooth setup.

6. Windows may report that drivers are being installed. Wait until Windows reports that the 'Device is ready to use'.



The Bluetooth pairing remains valid until you plug the Bluetooth dongle into a different USB port, delete the link from Windows, or pair the device with another PC.

4.4 Perform measurement

Connect Patient Leads

Connect the Patient Ground lead to the GND input of the amplifier and to the patient. Use the Mind Media Patient Ground wristband for optimal contact. Wet the band and place it around your wrist.



Mind Media recommends using the wet wristband to optimize measurement setup. This will improve the signal quality. Please read the Application Note on the reference amplifier for more background.

Connect all patient leads and patient ground. Please refer to instructions for use of the accessories and sensors for more information.

4.5 Mobility

Mind Media recommends using the carrying bag in case the patient needs to carry the device during the measurement. Slide the device in the carrying bag. This may require some force. The elastic bands hold the device in place. The carrying bag can be attached to a belt or use a shoulder band.

5 Operational principles

5.1 Bipolar input channels

The input stage for measuring bipolar electrophysiological signals is configured as an instrumentation amplifier. The difference between a 'plus' and 'minus' signal is amplified. The patient ground electrode is required to keep patient potential and amplifier potential at about the same level.

All electrode cables are shielded with the average of the 'plus' and 'minus' electrode signal (active shielding). The active shielding ensures that disturbances such as cable movement artifacts and mains interference (50/60 Hz) are reduced to a minimum.

After the first amplifier stage (gain = 20) the signals go directly to the ADC. No high pass or low pass filters that can cause signal phase shifts or filter overflows are present.

The NeXus-4 has one dual bipolar input. This connector leads to two bipolar input stages for in total two differential channels.

5.2 Auxiliary input channel

The Auxiliary input has a 5-pin connector. Signals on this connector are +5V output, -5V output, GND, +signal input and -signal input. The +5V/-5V/GND pins can be used to power an active sensor. The + and - inputs are connected to an instrumentation amplifier with a gain of 1. The output of the amplifier goes to the ADCs without any filtering.

5.3 Filtering

The ADC of the device has a digital sinc filter with a cutoff frequency of 0.2 * sample frequency. Supported sampling rates are defined in chapter 8. There is a 1^{st} order low pass filter before the ADC with a -3db point at 408Hz.

Besides above no other filtering is applied.

6 Maintenance

The product does not contain user serviceable parts. Maintenance is limited to regular cleaning. Repairs can only be performed by the manufacturer, contact support@mindmedia.com in case the product needs to be repaired. Mind Media Support staff will determine whether a repair is required and possible.

The product does not require regular servicing or re-calibration during its expected service life of 10 years. If the product is intended to be used after its expected service life, contact Mind Media to have the product inspected before continued use.

Cleaning

- Before cleaning, make sure the equipment is switched off and not in contact with a patient.
- Use only tap water, if necessary with a mild detergent, applied through a soft damp cloth.
- Do not spill fluids or submerge product in liquids.
- Never use sharp tools or aggressive chemicals for cleaning or disinfecting.
- Do not sterilize equipment.

Environmental protection

Special EU instructions for disposal are applicable to a product on which this symbol is placed. These instructions apply to all parts of the equipment.



When the product has reached End of Life, it must not be disposed of with other waste. Instead, it is the user's responsibility to dispose of their waste equipment by handing it over to a designated collection point for the recycling of waste electrical and electronic equipment.

The separate collection and recycling of your waste equipment at the time of disposal will help to conserve natural resources and ensure that it is recycled in a manner that protects human health and the environment.

For more information about where you can dispose of your waste equipment for recycling, please contact your local city office, your household waste disposal service, or Mind Media.

7 Electromagnetic guidance

Portable and mobile RF communications equipment can affect the system. The system needs special precautions regarding EMC and must be installed and put into service according to the EMC information outlined below.

Guidance and manufacturer's declaration - electromagnetic emissions The NeXus-4 is intended for use in the electromagnetic environment specified below. The customer or the user of the NeXus-4 should assure that it is used in such an environment.				
RF emissions CISPR 11	Group 1	The NeXus-4 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	Class B	The NeXus-4 is suitable for use in all		
Harmonic emissions IEC 61000-3-2	Not applicable Battery powered equipment	 establishments, including domestic establishments and those directly connected to the public low-voltage power supply network 		
Voltage fluctuations/flicker emissions 61000-3-3	Not applicable Battery powered equipment	that supplies buildings used for domestic purposes.		

Guidance and manufacturer's declaration - electromagnetic immunity

The NeXus-4 is intended for use in the electromagnetic environment specified below. The customer or the user of the NeXus-4 should assure that it is used in such an environment.

Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment - guidance
Electrostatic	±6 kV contact	± 8 kV contact	Floors should be wood, concrete, or ceramic tile. If floors are covered with synthetic
discharge (ESD) IEC 61000-4-2	±8 kV air	± 8 kV air	material, the relative humidity should be at least 30 %.
Electrical fast	±2 kV for power	Not applicable	
transient/burst	supply lines	Battery powered	
IEC 61000-4-4		equipment	
	±1 kV for	Not applicable	
	input/output lines	cabling shorter than	
		3m	
Surge	±1 kV line(s) to line(s)	Not applicable	
IEC 61000-4-5		Battery powered	
		equipment	
	±2 kV line(s) to earth	Not applicable	
		Battery powered	
		equipment	

Guidance and manufacturer's declaration - electromagnetic immunity

The NeXus-4 is intended for use in the electromagnetic environment specified below. The customer or the user of the NeXus-4 should assure that it is used in such an environment.

Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment - guidance
Voltage dips, short	<5 % U _T	Not applicable	
interruptions and	(>95 % dip in U⊤)	Battery powered	
voltage variations on	for 0,5 cycle	equipment	
power supply input lines IEC 61000-4-11	40 % U⊤ (60 % dip in U⊤) for 5 cycles	Not applicable Battery powered equipment	
	$70~\%~U_T$ (30 $\%$ dip in $U_T)$ for 25 cycles	Not applicable Battery powered equipment	
	<5 % U_T (>95 % dip in U_T) for 5 s	Not applicable Battery powered equipment	
Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	3 A/m	3 A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.

Guidance and manufacturer's declaration – electromagnetic immunity

The NeXus-4 is intended for use in the electromagnetic environment specified below. The customer or the user of the NeXus-4 should assure that it is used in such an environment.

Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment - guidance
			Portable and mobile RF communications equipment should be used no closer to any part of the NeXus-4, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter. Recommended separation distance:
Conducted RF IEC 61000-4-6	3 Vrms 150 kHz to 80 MHz	3 V	$d = 1.2\sqrt{P}$
Radiated RF IEC 61000-4-3	3 V/m 80 MHz to 2.5 GHz	3 V/m	$d = 1.2\sqrt{P} 80 \text{ MHz to } 800 \text{ MHz}$ $d = 2.3\sqrt{P} 800 \text{ MHz to } 2.5 \text{ GHz}$ where <i>P</i> is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and <i>d</i> is the recommended separation distance in meters (m). Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey, ^a should be less than the compliance level in each frequency range. ^b Interference may occur in the vicinity of equipment marked with the following symbol: $\left(\left(\underbrace{\bullet}\right)\right)$

from structures, objects, and people.

^a Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile

Guidance and manufacturer's declaration – electromagnetic immunity

The NeXus-4 is intended for use in the electromagnetic environment specified below. The customer or the user of the NeXus-4 should assure that it is used in such an environment.

Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment - guidance
radios, ama	teur radio, AM and FM r	adio broadcast ar	nd TV broadcast cannot be predicted theoretically with accuracy. To
assess the e	assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be		
considered.	considered. If the measured field strength in the location in which the NeXus-4 is used exceeds the applicable RF		
compliance	compliance level above, the NeXus-4 should be observed to verify normal operation. If abnormal performance is		
observed, additional measures may be necessary, such as re-orienting or relocating the NeXus-4.			
^b Over the fre	^b Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3 V/m.		

Recommended separation distances between portable and mobile RF communications equipment and the NeXus-4

The NeXus-4 is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of the NeXus-4 can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the NeXus-4 as recommended below, according to the maximum output power of the communications equipment.

Rated maximum output	Separation distance according to frequency of transmitter		
power of transmitter		m	
W	150 kHz to 80 MHz	80 MHz to 800 MHz	800 MHz to 2.5 GHz
	$d = 1.2\sqrt{P}$	$d = 1.2\sqrt{P}$	$d = 2.3\sqrt{P}$
0.01	0.12	0.12	0.23
0.1	0.38	0.38	0.73
1	1.2	1.2	2.3
10	3.8	3.8	7.3
100	12	12	23

For transmitters rated at a maximum output power not listed above, the recommended separation distance d in meters (m) can be estimated using the equation applicable to the frequency of the transmitter, where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer.

NOTE 1 At 80 MHz and 800 MHz, the separation distance for the higher frequency range applies.

NOTE 2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects, and people.

The NeXus-4 has no essential performance.

8 Technical specifications

General Specifications	
Туре	NeXus-4
TMS codes / REFs	95-0934-004-0
Size (device only)	112 mm x 66 mm x 28 mm (l x b x h)
Weight (g)	130 g (177 g incl. batteries)

Regulatory Specification	ıs
MDD class (Annex IX)	lla
Power source	Batteries
Electric shock protection	Applied parts: Class CF
Applied parts	 Its enclosure, also after removal of battery cover including all contacts and receptacles The patient accessories.
Accessible parts	Apart from the applied parts the NeXus-4 has no accessible parts.
Software class per IEC 62304	A
Ingress protection	Main unit: IP20

Bipolar Inputs (EEG, ECG, EOG, EMG, etc.)	
Noise	< 1.5 μV rms (@ Fs = 1024 Hz)
Gain	19.5 x
Input signal difference	-100mV to +100mV
Input common mode range	-2V to +2V
Input impedance	> 10 ¹⁰ Ω
CMRR	> 100 dB
Connector	LEMO 0B series 6 pin

Auxiliary Inputs	
Noise	< 15 µV rms (@ Fs = 512 Hz)
Gain	1 x
Input signal difference	-2V to +2V
Input common mode range	-2V to +2V
Input impedance	> 10 ¹⁰ Ω
CMRR	> 70 dB
Connector	LEMO OB series 5 pin

Filtering	
High pass	None
Low pass	Digital FIR filter in ADC;
	cutoff frequency = 0.2 * sample frequency

Battery	
Batteries	2 x AA disposable alkaline 1.5V
Power Saving	5 minutes without connection to PC
Battery low indication level	2.0 V ± 0.1V
Battery empty shut down level	1.7 V ± 0.1V

Bluetooth Co	ommunication
--------------	--------------

Bluetooth 1.1 class 2	Bluetooth [™]
Profile	Serial port profile
Range	10 meters (line of sight)
Baud rate	230400 bps

Transportation Conditions	
Temperature	-25°C to +70°C
Humidity	15% to 93%
Pressure	500 hPa to 1060 hPa

Storage Conditions	S
Temperature	0°C to +40°C
Humidity	15% to 93%
Pressure	500 hPa to 1060 hPa

Usage Conditions							
Temperature	+5°C to +40°C						
Humidity	15% to 93%						
Pressure	700 hPa to 1060 hPa						

Channel list

#	Name	Function		Resolution/bit	Range
1	A	Bipolar signal	A	0.012215 μV	-100mV to +100mV
2	В	Bipolar signal	В	0.012215 μV	-100mV to +100mV
3	С	Auxiliary sign	Auxiliary signal C		-2.0V to +2.0V
4	D	Auxiliary signal D		0.2384186 μV	-2.0V to +2.0V
5	Digi	Digital channe	Digital channel (bits)		0 to 255
		0x01 1 = but	ON/OFF ton	-	
		0x08 0 0x10 63 0x20 (a	tery state : battery empty 3: battery full lkaline)	-	
		0x40 0x80 1 = em	battery oty	-	
6	Saw	Sawtooth tes sample interv	t signal, 256- al, 1-bit step	1 (bit)	0 to 255

Bluetooth supported sample rates/channel rates

	Channel	Bytes per Bluetooth channel rate @ Fs (Hz):					
#	name	channel	1024 Hz	512 Hz	256 Hz	128 Hz	64 Hz
1	Α	3	1024	512	256	128	64
2	В	3	1024	512	256	128	64
3	С	3	128	512	256	128	64
4	D	3	128	512	256	128	64
5	Digi	1	1024	512	256	128	64
6	Saw	1	1024	512	256	128	64

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