

LUXAMED®

LUXAMED® Reflexhammer

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Marteau reflex LUXAMED®

LUXAMED® Martello reflex

Martillo LUXAMED® Reflex

Młotek refleksyjny LUXAMED®

Martelo reflexos Luxamed®

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LUXAMED®



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- ENGLISH -



To ensure proper use of this LUXAMED® product, please read this user manual carefully and retain for future reference.

LUXAMED® Reflex hammer

Intended use of instrument:

The LUXAMED® reflex hammers are only to be used for testing the muscle- or tendon reflexes and skin reflexes on healthy skin. The needle, on the base of the handle, is used to test for the skin sensitivity. Use the neurological sensor tip carefully to avoid any injuries or perforation at the scope of application. These instruments are only to be used by authorized, trained personnel. Keep out of the reach of children. Use the device in accordance with its designated use and for its intended purpose only.

Contraindication:

none

Warning:



Please make sure that, prior to any use, the rubber inserts are fixed correctly at the head of the hammer. NEVER use without rubber inserts.



Use the neurological sensor tip carefully to avoid any injuries or perforation at the scope of application due to improper use.



All serious incidents that have occurred with the product must be reported to the manufacturer and the competent authority of the Member State in which the user and / or patient is established.

Application:

Using the reflex hammer, tap gently on the corresponding reflex zone. Make sure of a careful application. With the caused reflex contraction a diagnostic can be established. By lightly pressing or moving the top of the needle back and forth on the skin, the skin sensitivity can be diagnosed.

Cleaning:

Always clean the reflex hammer right after use. The reflex hammer can be cleaned with a damp cloth (use ph-neutral or slightly alkaline cleaning materials).

Disinfection:

Disinfection or disinfecting cleaning of the outer surfaces of the reflex hammer simply by wipe disinfection. For this, isopropanol can be used.

Sterilisation:

No sterilisation necessary, not approved for sterilisation.

Disposal:

This product and its accessories can be disposed of alongby with household waste.

Storage, transport and operating conditions:

Ambient temperature between -10° and +40°

Rel. humidity between 10% and 90%

Symbol key

	Manufacturer
	Date of manufacture
	Attention to the user manual
	Caution
	In Conformity with EU guidelines
	It is a medical device
	Conforms with UK regulations
	Authorised agent for medical products in Switzerland: Inspira GmbH Thunstrasse 64 CH-3110 Münsingen, Switzerland

Warranty Conditions:

We give 2 years warranty from the date of purchase. The purchase date has to be proven by the invoice. Within the warranty period defects are eliminated free of charge.

After repairs the warranty period is not extended on the whole unit, but only to the replaced components.

Excluded from the warranty are parts subject to normal wear and tear (e.g. cuff), transport damages and any damage caused by improper handling (e.g. non-compliance with the instructions for use). Damages due to disassembly by unauthorized persons are also excluded from warranty. No claims for damages against us are substantiated by the warranty.

In the case of justified warranty claims the device has to be sent along with the original invoice to:



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UK Responsible Person

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