EC Certificate – Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding (4)

Certificate No. MDD-202

Issued to:	IMP-Piezotehnologija d.o.o. Beograd Volgina 15, 11060 Beograd, Srbija
Place of production:	IMP-Piezotehnologija d.o.o. Beograd Volgina 15, 11060 Beograd, Srbija
Product category:	Magnetic therapy medical devices
GMDN:	/
Standard:	MDD 93/42/EEC
Annex:	II

SIQ has audited the quality system in accordance with MDD Annex II excluding (4) and found that the above-mentioned manufacturer's quality system meets the requirements of the Directive 93/42/EEC concerning medical devices Annex II. This certificate is based on

Audit report No.:
OSV 00272/2021, 2021-03-25
OSV 00100/2021, 2021-03-31
OSV 00602A/2021, 2021-05-21
OSV 00617A/2021, 2021-05-25

See also decision of NB`s commission for medical devices.

This certificate remains valid as long as the Manufacturer's quality system is subject to periodical surveillance as referred to in Directive 93/42/EEC concerning medical devices Annex II (5) and continues to meet the above requirements.

Certification date: 2021-05-25

Issue: 1/2021-05-25

Valid until: 2024-05-26

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8

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