EC Certificate – Production Quality Assurance
Directive 93/42/EEC on Medical Devices, Annex V
Certificate No. MDD-122

Issued to:	KIRCHNER & WILHELM GmbH + Co.KG Eberhardstrasse 56, 71679 ASPERG, Deutschland
Place of production:	KIRCHNER & WILHELM GmbH + Co.KG Eberhardstrasse 56, 71679 ASPERG, Deutschland
Product category:	Iontophoresis units
GMDN:	12-185

SIQ has audited the quality system in accordance with MDD Annex V and found that the above-mentioned manufacturer's quality system meets the requirements of the Directive 93/42/EEC concerning medical devices Annex V, including all subsequent amendments. This certificate is based on

Audit report No.:
OSV 00798/2018, 2018-07-30
OSV 00921/2018, 2018-08-20
OSV 01173/2018, 2018-11-28
OSV 01182/2018, 2018-10-29
OSV 01433/2018, 2019-01-10

Digitally signed by Ermicev Alexandr
Date: 2021.02.10 22:05:29 EET
Reason: MoldSign Signature
Location: Moldova

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 V (4) and continues to meet the above requirements.

Certification date: 2019-01-10
Issue: 1/2019-01-10
Valid until: 2024-01-10

SIQ Ljubljana
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Director of SIQ
Igor Likar