EC Certificate of Conformity

Directive 93/42/EEC on Medical Devices, Annex IV

Certificate No. MDD-130

Issued to:	REPULS Lichtmedizintechnik GmbH Lemböckgasse 61 / Top 1, 1230 Wien, Austria
Place of production:	SVI Austria GmbH, Frauentaler Str. 100, 8530 Deutschlandsberg, Austria
Place of production:	Sementis Engineering GmbH, Resselstrasse 16, 2120 Wolkersdorf, Austria
Product category:	Electronic therapy device
UMDNS:	11-503

SIQ has performed examinations and tests in accordance with MDD Annex IV (4) and found that the device(s) or batches of devices conform(s) with the technical documentation. This certificate is based on

Audit report No.:
OSV 00012A/2019, 2019-02-28
OSV 00229A/2019, 2019-03-22
OSV 000764/2019, 2019-06-21

See also decision of NB`s commission for medical devices.
See also a Detailed list of product names, models and types with batch numbers.

Certification date: 2019-06-28

Issue: 1/2019-06-28

SIQ Ljubljana
Tržaška cesta 2
SI-1000 Ljubljana / Slovenija

Director of SIQ
Igor Likar

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