EC Certificate – Full Quality Assurance System
Directive 93/42/EEC on Medical Devices, Annex II excluding (4)
Certificate No. MDD-009

Issued to:	"Vitafon" Co. Ltd., Ogorodnyy pier. 23, 198097 Saint-Petersburg, Russia
Place of production:	"Vitafon" Co. Ltd., Ogorodnyy pier. 23, 198097 Saint-Petersburg, Russia
Product category:	Active therapeutic device intended to administer or exchange energy
GMDN:	61096

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 00103A/2018, 2018-03-27

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: 2010-12-03

Issue: 5/2018-03-27

Valid until: 2020-02-28

SIQ Ljubljana
Tržaška cesta 2
SI-1000 Ljubljana / Slovenija

Director of SIQ
Igor Likar