

Declaration of Conformity

for the electrostimulation, ultrasound therapy and suction wave device family **Soleoline**

We declare under our sole responsibility, that the product line

Soleoline
consisting of the devices

Soleo Sono
Soleo SonoStim
Soleo Galva
VacoS



GIÁM ĐỐC
Triệu Đức Thành

bears the CE mark



according to the Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4) and fulfils the essential requirements of annex I of this directive.

The device belongs to class IIa according to annex IX rule 9 of this directive.

The conformity of the quality management system is certified and controlled by TÜV SÜD Product Service GmbH, Ridlerstraße 65, 80339 Munich, Notified Body Code: 0123.

This declaration is valid from 2020-08-06 until 2024-05-26.

Stefan Leinweber
Quality Management
Medical Device Safety Officer

Neu-Ulm, 2020-08-06