EC Declaration of Conformity

This declaration of conformity is issued under the sole responsibility of
ENRAF-NONIUS B.V., Vareseweg 127, 3047 AT Rotterdam, The Netherlands

Quality Management System according to EN ISO 13485:2016 and
Annex II (excl. section 4) of European Medical Devices Directive 93/42/EEC.
Registration No. HD 1274285-1
Notified body: TÜV Rheinland LGA Products GmbH (0197), Tillystraße 2, 90431 Nürnberg, Germany

Device name:	Endolaser 120
Description:	Low Level Laser Therapy Equipment
Article number(s):	1633900 (base unit) 1633901 (base unit with 1 probeholder) 1633902 (base unit with 2 probeholders) 1632801/1632802/1632803/1632804 (laser probes)
Classification:	IIa (according to rule 9, Annex IX of MDD 93/42/EEC)
Record of conformity:	026-400-281-41 ER
Device marking:	CE0197

We hereby declare that the above-mentioned device
complies with the European Medical Devices Directive 93/42/EEC.

This declaration of conformity is valid in combination with the test certificate of the device.
Any modifications to the product not authorized by Enraf-Nonius will invalidate this declaration.

Rotterdam, 25 February 2021

Signature:

Th. J. Doodkorte
Commercial Director