



## 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:	Sonopuls 492
Description:	Ultrasound- and Electrotherapy Equipment (combination therapy)
Article number(s):	0498910 (base unit) 1498911 / 1498912 / 1498913
Classification:	Ila (according to rule 9, Annex IX of MDD 93/42/EEC)
Record of conformity:	026-400-253-46 ERC
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. Boodkorte  
Commercial Director