
	EC Declaration of Conformity		Page 1 of 1
	Product:	DENTO-PREP	Version: 05.1
Doc. no.: DENTO-PREP-DoC	Author: SLP	Approved by: EHB	Date: 2024.05.27
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Identification of the Legal Manufacturer:	Rønvig Dental Manufacturing A/S Gl. Vejlevej 59 8721 Daugaard Denmark
Identification of Authorized Representative:	Rønvig Dental Manufacturing A/S Gl. Vejlevej 59 8721 Daugaard Denmark
This Declaration of Conformity is issued under the sole responsibility of the Legal Manufacturer.	
Identification of the device(s) concerned:	REF 1900 DENTO-PREP Dental Microblaster.
Risk Classification:	Class II a as per Rule 11 in Annex IX
We hereby declare that the above-mentioned device complies with the European Medical Devices Directive 93/42/EEC as modified by 2007/47/EC.	
Name and address of Notified Body:	DNV Product Assurance AS Veritasveien 1 N-1363 Høvik Norway
Applicable CE Certificate(s):	EC Quality System Certificate No. C581150 NA DNK issued by DNV, Notified Body Number 2460 in accordance with ISO 13485:2016 EN ISO 13485:2016
	EC Certificate No. 1000381165-PA-NA-DNK Rev 0.0, issued by DNV, Notified Body Number 2460, in accordance with Annex II Section 4 of this Directive.
Identification of the person authorized to sign on behalf of Legal Manufacturer:	Name: Annette Ravn Nielsen Signature:  Title: CEO Place of Issue: Daugaard, Denmark Date: 2024-10-03