

EU Declaration of Conformity



Document No.: GI 162 / 05

Manufacturer and Address:	SIRONA Dental Systems GmbH Fabrikstraße 31 64625 Bensheim Germany
Product Category:	Reprocessing Device
Medical Device Name:	DAC Universal
Product Identification (Ref.-Number):	Type MK IV REF see page 2
Classification according to Annex IX (93/42/EEC):	Class II b

We declare under our sole responsibility the compliance of the medical device concerned with the requirements of the Council Directive 93/42/EEC.

Any modification to the product, not authorized by us, will invalidate this declaration.

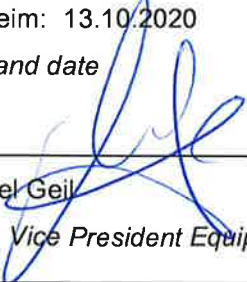

Valid from manufacturing date: 2020-10-13

The conformity of the full quality assurance system (Directive 93/42/EEC, Annex II excluding 4) is certified by:

TÜV SÜD Product Service GmbH
Ridlerstraße 65
80339 München
Germany

The identification number of the notified body for implementation of the procedure is 0123.

Bensheim: 13.10.2020
Place and date

	
Michael Geil Group Vice President Equipment and Instruments	Wolfram Müller Regulatory Compliance Manager

The declaration certifies the compliance according to Annex II of Directive 93/42/EEC. Conditions of guarantee and liability are dealt within our General Conditions of Sale.

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Name	REF
DAC Universal	6754860
DAC Universal D	6727916
DAC Universal S	6762160