EU DECLARATION OF CONFORMITY

By the present document, the Vanetti SA-Via Principale-CH-6672 Gordevio, Switzerland assumes the responsibility that the following range of products:

Mark	Type	Code N°
Dia-Tessin	Ceramic Instrument FG	CSTT / 015 & CSTT/015L

classified as Medical Devices - Class IIa according to Directive 93/42/EEC annex IX, par. 2.1, rule 5

is in conformity with the essential requirements of the medical devices Directive 93/42/EEC as amended by Directive 2007/47/EEC.

is subject to the assessment procedure set out in Annex II (excluding point 4) of Directive 93/42/EEC under the supervision of the Notified Body stated below. Certificate number CH06/0485.

The products have been developed, manufactured and controlled within a Quality System meeting the requirements of EN ISO 13485:2016. Certificate number CH97/0206.

The main dental standards addressed are
EN 1639, EN-ISO 1797-1, EN-ISO 2157, EN-ISO 8325, EN-ISO 10993-1, EN-ISO 10993-5, EN-ISO 11737-1, EN-ISO 11737-2, EN-ISO 15223-1 and EN-ISO 17664.

These products do not need any specific certificate for clinical or technical tests in the country of origin according to the Swiss law for medical devices.

Notified body : SGS NB0120

Address : SGS United Kingdom Ltd.
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VANETTI SA
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Name: Vanetti First name: Pablo Function: CEO

Place: 6672 Gordevio Date: 10.07.2018 Signature:

Validity: 09.07.2023