

The management system of

Vanetti SA

Via Principale
CH - 6672 Gordevio

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC
on medical devices, Annex II (excluding Section 4)

For the following products

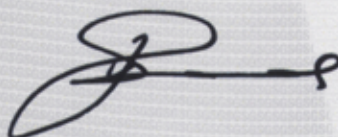
Dental rotary instruments - diamond, - for intra oral use.
Dental rotary Tungsten carbide burs - for intra oral use.
Dental rotary instruments - ceramic CSTT
Ceramic soft tissue trimmer - for intra oral use.

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 10 July 2018 until 9 July 2023
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 26 June 2021
Issue 6. Certified since 10 July 2006

Certification is based on reports numbered CH/GE 3302633

Authorised by



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