

EC Certificate Full Quality Assurance System: Certificate CH06/0077

The management system of

Kenda AG

Kanalstrasse 18, 9490 Vaduz, Liechtenstein

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

Rotating instruments to finish and polish dental surfaces in restorative dentistry

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 20 November 2017 until 19 November 2022
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 1 October 2020
Issue 6. Certified since 20 November 2008

Certification is based on reports numbered CH/GE 3301545

Authorised by

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