

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



**SGS United Kingdom Ltd, Notified Body 0120**

202B Worle Parkway, Weston-super-Mare, BS22 6WA UK  
t +44 (0)1934 522917 f +44 (0)1934 522137 www.sgs.com

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Page 1 of 1

