EC Certificate Full Quality Assurance System: Certificate CH12/0875

The management system of

Bien-Air Dental SA

Länggasse 60,
CH - 2504 Bienne

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

Air and electrical motors, Straight and contra-angle handpieces, Turbines, Air and electrical hoses and couplings, and Electronic consoles for: Dental applications, Oral and Maxillofacial Surgery, and Podiatry applications.

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 09 December 2016 until 30 November 2021
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 30 November 2019
Issue 7. Certified since 30 November 2007

Certification is based on reports numbered CH/GE 3301306

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

SGS CE 02 0215