

The management system of

Odontit S.A.

Azcuenaga 1077 4 Da,
Buenos Aires, 1115, Argentina

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

Sterile Dental Implants Systems and Autogenous bone collector device.

For placing on the market of Class III devices covered by this certificate, an EC Design Examination Certificate according to Annex II (Section 4) is required.

This certificate is valid from 19 May 2011 until 27 April 2014 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 10 February 2014
Issue 1

Certification is based on reports numbered AR/BUE 2011006
Authorised by

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