



DET NORSKE VERITAS

FULL PRODUCT QUALITY MANAGEMENT CERTIFICATE - EC

Certificate No. 69067-2009-CE-ARG-NA
This Certificate consists of 3 pages

This is to certify that the Quality Management System of

ODONTIT SA

Azcuénaga 1077 4° D – Ciudad Autónoma de Buenos Aires, 1115, Argentina

for production and final product inspection/testing of

Dental Devices

has been assessed with respect to

the conformity assessment procedure described in Article 11.3.a and Annex II excluding section 4 (Module H) of Council Directive 93/42/EEC on Medical Devices, as amended, and found to comply

Further details are given overleaf

Place and date:

Høvik, 10 February 2010

This Certificate is valid until:

27 July 2014

For DET NORSKE VERITAS CERTIFICATION AS
NORWAY



Notified Body No.:
0434

Marianne Spæren
Certification Manager

Aud Løken Eiklid
Technical Reviewer

Notice: The certificate is subject to terms and conditions overleaf. Any significant changes in design or construction may render this certificate invalid.

If any person suffers loss or damage which is proved to have been caused by any negligent act or omission of Det Norske Veritas, then Det Norske Veritas shall pay compensation to such person for his proved direct loss or damage. However, the compensation shall not exceed an amount equal to ten times the fee charged for the service in question, provided that the maximum compensation shall never exceed USD 300,000. In this provision "Det Norske Veritas" shall mean the Foundation Det Norske Veritas as well as all its subsidiaries, directors, officers, employees, agents and any other acting on behalf of Det Norske Veritas.



Cert. No.: 69067-2009-CE-ARG-NA
 Rev. No.:
 Project No.: PRJC-171671-2009-MSL-ARG

Jurisdiction

Application of Council Directive 93/42/EEC of 14 June 1993, adopted as 'Forskrift for Medisinsk Utstyr' by the Norwegian Ministry of Health and Care Services.

Certificate history

Revision	Description	Issue Date
	Original certificates 2004-OSL-MDD-0292 and 0293	2004-07-27
	Recertification	2010-02-10

Products covered by this Certificate

Product Description	Product	Class
Dental Implant Systems	External hex connection implant – Abutments – Prosthetic parts – Laboratory components – Ancillary equipment (eFeDeA-IF- Osseomate)	IIb
	Self tapping, external hex connection implant - Abutments – Prosthetic parts – Laboratory components – Ancillary equipment (Evolution Hexagonal- EH - Osseomate)	
	Self tapping, internal tapered connection implant - Abutments – Prosthetic parts – Laboratory components – Ancillary equipment (Evolution Swiss-ES - Osseomate)	
	Self tapping, internal Hex connection implant - Abutments – Prosthetic parts – Laboratory components – Ancillary equipment (Evolution Internal Hex- IH - Osseomate)	
	Self tapping one piece implant - Abutments – Prosthetic parts – Laboratory components – Ancillary equipment (Monoblock/Monoblock Narrow-MB/MBN - Osseomate)	
	Sell tapping mini implant - Abutments – Prosthetic parts – Laboratory components – Ancillary equipment (Evolution Ball- EB/Evolution Square-EQ - Osseomate)	
	Orthodontic implant - Abutments – Prosthetic parts – Laboratory components – Ancillary equipment (Evolution Ortodontico-EO - Osseomate)	
Autogenous Bone Collector Device	Odontit	IIa
	Osseomate	

The complete list of devices is filed with the Notified Body.



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Sites covered by this certificate

Site Name	Address
ODONTIT	Azcuénaga 1077 4° D – Ciudad Autónoma de Buenos Aires, 1115, Argentina
ODONTIT SA	Necochea 854, Ciudad Autónoma de Buenos Aires, 1158, Argentina

EU Representative

Medical Technology Promedt Consulting GmbH, Altenhofstrasse 80, D-66386 St. Ingbert, Germany

Terms and conditions

The certificate is subject to the following terms and conditions:

- Any producer (see 2001/95/EC for a precise definition) is liable for damage caused by a defect in his product(s), in accordance with directive 85/374/EEC, as amended, concerning liability of defective products.
- The certificate is only valid for the products and/or manufacturing premises listed above.
- The Manufacturer shall fulfil the obligations arising out of the quality system as approved and uphold it so that it remains adequate and efficient.
- The Manufacturer shall inform the local DNV Office of any intended updating of the quality system and DNV will assess the changes and decide if the certificate remains valid.
- Periodical audits will be held, in order to verify that the Manufacturer maintains and applies the quality system DNV reserves the right, on a spot basis or based on suspicion, to pay unannounced visits.

The following may render this Certificate invalid:

- Changes in the quality system affecting production.
- Periodical audits not held within the allowed time window.

Conformity declaration and marking of product

When meeting with the terms and conditions above, the producer may draw up an EC declaration of conformity and legally affix the CE mark followed by the Notified Body identification number of DNV.

END OF CERTIFICATE