



DET NORSKE VERITAS

EC CERTIFICATE - FULL QUALITY ASSURANCE SYSTEM

Certificate No. 46275-2009-CE-IND-NA 3.0

This Certificate consists of 3 pages

This is to certify that the Quality Management System of

SMB Corporation of India

India

for design, production and final product inspection/testing of

Intra Uterine Devices and Non-Absorbable Surgical Sutures

has been assessed with respect to

the conformity assessment procedure described in Article 11.1.a and Annex II (Module H1) and 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, 27 October 2010

This Certificate is valid until:

27 February 2014

For DET NORSKE VERITAS CERTIFICATION AS
NORWAY



Aud Løken Eiklid
Certification Manager

Notified Body No.:
0434

Mariann Jeremiassen
Technical Reviewer

This Certificate has been digitally signed. See www.dnv.com/digitalsignatures for more info

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.: 46275-2009-CE-IND-NA
Rev. No.: 3.0
Project No.: PRJC-52990-2008-PRC-IND

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 certificate 2004-OSL-MDD-0093	2004-02-27
1.0	Recertification and new certificate number	2009-02-27
2.0	Revision in EU representative and scope and merging of certificate 46277-2009-CE-IND-NA Rev 1.0 into this one	2010-03-08
3.0	Editorial change to show traceability of certificates	2010-10-15

Products covered by this Certificate

Product Description	Product	Class
Intra Uterine Devices	<ul style="list-style-type: none">▪ CuT 200B▪ CuT 380A▪ Cu 375	III*
Non-Absorbable Surgical Sutures	<ul style="list-style-type: none">▪ Silk surgical suture with and without needle▪ Polyester surgical suture with needle▪ Polypropylene monofilament surgical suture with needle▪ Polyamide monofilament surgical suture with and without needle	IIb

* Design assessment is covered by a separate EC design-examination certificate with design approval no. 46275-2009-CE-IND-NA (with issue date 2nd Feb 2009 and expiry date 27th Feb 2014)

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

Sites covered by this certificate

13, 33 – 36, Prem Industrial Estate, Jogeshwari (E), Mumbai – 400 060, India

EU Representative

Obelis s.a, Boulevard General Wahis 53, 1030 Brussels, Belgium



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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