

CERTIFICATE OF CONFORMITY
WITH EUROPEAN DIRECTIVE



Certificate No.: EU0707421
Order No.: 86269

We hereby certify that an examination has been carried out following the requirements of the national legislation "Regulation no. 25 of 12th January 1995 relating to medical devices pursuant to act no. 6 of 12th January 1995 relating to medical devices, transposing directive 93/42/EEC into Norwegian law to which the undersigned is subjected, confer EEA agreement, proposition no. 100 (1991-92) special appendix no. 2, volume 2A/3 A, goods, chapter XXX". We certify that the production quality system conforms to the relevant provisions of the Annex given below:

Name and address of the manufacturer: **APPLIED TRADING FZE**
P-6/164,
P.O. Box 120066
Saif Zone, SHARJAH, United Arab Emirates

Device category: Medical Devices

GMDN code: See Appendix 1 to this certificate

Models: See Appendix 1 to this certificate

Risk class as defined by the manufacturer: IIa

Standards/provisions: The audit of the quality system was based upon and assessed according to the provisions in Annex V of the EC-Directive 93/42/EEC, with the exemption of section 4.

Date of audit: 16th April 2007

Date of the end of the validity: 1st September 2012

Nemko EC notification No.: 0470

On this basis the manufacturer or the European authorised representative may draw up an EC / EEA Declaration of Conformity and affix the CE-marking as indicated below together with the Nemko EC notification number to each conforming product as long as the conformity audit and inspection procedure required by the EC directive will be fulfilled by the manufacturer and the factory. The product liability rests with the manufacturer or his representative in accordance with Council Directive 85/374/EEC.

Date of issue: 2007-08-02

Date of verification: 2007-08-02

Signature: Svein Grahnstedt
MD PhD Senior Lead Auditor

Signature: Frank Skarpsno
Senior Lead Auditor / Principal Engineer

CE 0470

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Device category: Medical Devices / Urology Products

Appendix 1: Page 1 of 1

The certificate referred to above includes the following devices/models:

SNO	PRODUCT	CLASS	STERILE	GMDN CODE
1	Male Catheter	Ila	YES	10765
2	Nelaton Catheter	Ila	YES	10734
3	Tur Set	Ila	YES	31069
4	Urethral catheter	Ila	YES	10761

Date of issue: 2007-08-02

Signature: Svein Grahnstedt

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