



EC-CERTIFICATE

Full Quality Assurance System
(Annex II of the Directive 93/42/EEC on Medical Devices)
Registration No. 7988

Elkomed AG
CH - 9524 Zuzwil

The Notified Body of QS Zürich AG hereby declares that an examination of the under mentioned full quality assurance system has been carried out following the requirements of the national legislation to which the undersigned is subjected, transposing annex II (with the exemption of section 4) of the Directive 93/42/EEC on medical devices. We certify that the full quality assurance system conforms to the relevant provisions of the aforementioned directive.

This EC-Certificate is only valid for the following products:

(The products are specified in annex „Product range“. This Certificate is only valid together with this annex)

ELKOPUR

During the validity of this Certificate, the requirements of the above mentioned regulations must be fulfilled permanently.

This is supervised by QS ZÜRICH AG

For precise and updated information concerning possible changes occurred in the certification object of the present certificate, please contact qs-zuerich@quality-service.ch



CE 1254

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Expiration date: 04.04.2017

QS Zürich AG
P.O. Box 6335
CH-8050 Zürich
qs-zuerich@quality-service.ch



A handwritten signature in blue ink, appearing to read 'C. Jovan'.

SCESm 047
www.sas.ch

Direction