CONMED CORPORATION

WITH FACILITIES LOCATED AT:
UTICA, NEW YORK USA
NEW HARTFORD, NEW YORK USA
ROME, NEW YORK USA
DENVER, COLORADO USA
EL PASO, TEXAS USA
JUAREZ, CHIHUAHUA, MEXICO

DECLARATION OF CONFORMITY CERTIFICATE

We, **CONMED Corporation, 525 French Road, Utica, NY 13501 USA**, according to EN45014, declare with sole responsibility, that our medical devices:

Aspirators [10-208] – Class IIa Medical Devices
Cables/Leads, Electrosurgical Unit [11-496] – Class IIb Medical Devices
Cannulae, Other [15-206] – Class IIa Medical Devices
Catheter, Cholangiography [16-429] – Class IIa Medical Devices
Clip Appliers [10-894] – Class IIb Medical Device
Dressing, Other [15-216] – Class IIa Medical Device
Electrodes, Defibrillation [15-033] – Class IIa Medical Devices
Electrodes, Electrosurgical, Active [16-860] – Class IIb Medical Devices
Electrodes, Electrosurgical, Active, Foot-controlled [16-206] – Class IIb Medical Devices
Electrodes, Electrosurgical, Active, Hand-controlled [11-499] – Class IIb Medical Devices
Electrodes, Transcutaneous, Electrical Nerve Stimulation [17-191] – Class IIa Medical Devices
Electrosurgical Unit Adapters, Cable [11-493] – Class IIb Medical Devices
Electrosurgical Unit, Monopolar/Bipolar [18-231] - Class IIb Medical Device
Electrosurgical Unit, Monopolar/Bipolar, Argon-Enhanced Coagulation [18-232] – Class IIb Medical Device
Elevators, Uterine [15-677] – Class IIa Medical Device
Forceps, Electrosurgical [11-502] – Class IIb Medical Devices
Needles, Pneumoperitoneal [12-750] – Class IIa Medical Devices
Pressure Infusors [13-100] – Class IIa Medical Devices
Retractors, Abdominal [13-375] – Class IIa Medical Devices
Retractors, Vaginal [13-392] – Class IIa Medical Devices
Specimen Containers [13-655] – Class IIa Medical Devices
Suction Irrigators [13-845] – Class IIa Medical Devices
Suction Tips, Electrosurgical [15-314] – Class IIa Medical Devices
Suture Units, Automatic [15-065] – Class IIb Medical Devices
Suture Units, Other [16-264] – Class IIa Medical Devices
Trocars [14-154] – Class IIa Medical Devices

meet, (where applicable), the provisions of Council Directive 93/42/EEC pertaining to medical devices which apply to them. The obligation as laid down in Annex II of the 93/42/EEC are fulfilled via Annex II, Clause 3 certification issued by **TUV Product Service, Notified Body No. 0123**.

We hereby appoint **MDSS GmbH, Burckhardtstrasse 1, D-30163 Hannover, Germany** to act as European Authorised Representative as defined in Art. 1(j) of the Medical Device Directive 93/42/EEC.

Signed this day 27th of JAN 2004 at Utica, NY USA

Ira D. Duesler, Jr.
Management Representative

This certificate was filed under No. 22320 by MDSS on 24 of Feb. 2004
MDSS has notified the authorities. The above named manufacturer receives the

Reg. No: See Certificate of CE Registration

The necessary pre-requisites for placing the CE mark on the above mentioned products and marketing them in all Member States of the European Union, have thus been fulfilled.

Signed this day 24 of Feb. 2004

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