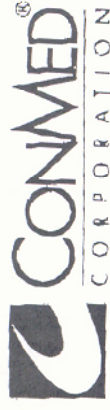


# DECLARATION OF CONFORMITY CERTIFICATE



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

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 Applicators [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 Infusers [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 27<sup>th</sup> 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  
D-30163 Hannover - Germany  
Phone: +49-511-6824440