

EC Certificate of Conformity

The Notified Body

MEDCERT Zertifizierungs- und Prüfungsgesellschaft für die Medizin GmbH
Pilatuspool 2 – 20355 Hamburg – Germany

herewith certifies that the company

MEDICOPLAST International GmbH
Heusweilerstr. 100
66557 Illingen
Germany

has introduced, applies and maintains a quality assurance system
for the aspects of manufacture concerned with securing and maintaining sterile conditions

for the products / product categories listed in the appendix.

The compliance of this quality assurance system with the below mentioned requirements of the
Council Directive 93/42/EEC was verified by an audit:

Annex V

This certification is subject to surveillance by MEDCERT.

Effective date: 2020-06-16


Expiry date: 2024-05-27

Report No.: 1189FS19F

Process No.: QS – 1189

Certificate No.: 1189GB415200616

Hamburg, 2020-06-16


MEDCERT Certification Body
(Dr. Andreas Schich)

The certificate is only valid when provided entirely with all of its pages.
To verify the validity of this certificate, contact info@medcert.de.

MEDCERT Identification Number: 0482

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Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-237.10.15

Appendix of EC Certificate of Conformity

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List of products / product categories included in the scope of certificate

- Female-catheters
- Nelaton-catheters
- Tiemann-catheters
- Atraumatische Katheter
- Drainage bagy
- Rectal tubes
- Suction sets
- Suction catheters
- Suction cannulas
- Suction mucus extractor
- Tubes and accessoeries
 - Tube sections
 - Clamps
 - Connectors
 - Valves
- Tracheal tubes and accessories
- Tracheal cannulas and accessories
- Cord clamps
- Redon systems and accessories
- Redon drainages and accessories
- Jet catheter and accessories
- Biopsy attachment (needle guide sleeve)

– End of list –

This appendix is integral part of the above-referenced certificate.
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