MED CERT®

Certification. Medical Only.

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

Transcoject GmbH
Rügenstr. 8
24539 Neumünster
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: **2021-05-25**
Expiry date: **2023-11-07**

Report No.: 0520FS26F
Process No.: QS – 0520
Certificate No.: 0520GB415210525

Hamburg, 2021-05-25

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.

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

MEDCERT Identification Number: 0482

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Appendix of EC Certificate of Conformity

Process No.: QS – 0520
Certificate No.: 0520GB415210525

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

- **Dental cannulas**
- **Syringes for single use**
- **Closure devices**
- **Application devices**
- **Connectors**

– End of list –

This appendix is integral part of the above-referenced certificate.
The certificate is only valid when provided entirely with all of its pages.
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MEDCERT Identification Number: 0482

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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:

Transcoject GmbH
Rügenstr. 8
24539 Neumünster
Germany

has introduced and maintains a quality assurance system
concerning the conformity of the medical devices with the metrological requirements

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.

This certificate is valid until 07 November 2023

Report No.: 0520FS26F
Process No.: QS – 0520
Certificate No.: 0520GB416190716

Hamburg, 16 July 2019

MEDCERT Certification Body
(Dr. Andreas Schihc)

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Appendix of EC Certificate of Conformity

Process No.: QS – 0520
Certificate No.: 0520GB416190716

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

- **Syringes for single use**

– End of list –

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