

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 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 II without section 4

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.: 0520GB410191114

Hamburg, 14 November 2019



MEDCERT Certification Body
(Dr. Andreas Schich)

The certificate is only valid when provided entirely with all of its pages.
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Appendix of EC Certificate of Conformity

Process No.: QS – 0520

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

- **Syringes for single use**
- **Application devices**
- **Cannulas**
- **Syringe ampoules**

– End of list –

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