Certificate

ECM – Zertifizierungsgesellschaft für Medizinprodukte in Europa mbH,
Talbotstraße 21, 52068 Aachen, Germany

hereby declares that an examination according to DIN EN ISO/IEC 17021-1:2015 of the undermentioned quality assurance system has been carried out.

Through an audit performed on behalf of

Jecture Germany GmbH
Philipp-Reis-Straße 2, 36399 Freiensteinau, Germany

it could be demonstrated that a quality management system according to

ISO 13485:2016
EN ISO 13485:2016 + AC:2018 + A11:2021
DIN EN ISO 13485:2021
„Medical devices – Quality management systems – Requirements for regulatory purposes"

for the scope:

Production and distribution of cannulas, cannula systems and related components

has been established and implemented.

This certificate is only valid under the conditions stated in the audit report for the audit mentioned below.

Any substantial changes of the quality management system have to be notified to ecm and are subject to a separate assessment.

Audit-No.	Registered under	Valid until
0228-24-0611	**Z/25/04903E**	**18 August 2027**

Valid as of: 20 November 2025

Certification body

DAkkS Deutsche Akkreditierungsstelle D-ZM-21753-01-00

ID 0374

Annex I to certificate Z/25/04903E

This certificate covers the following sites:

Site	Scope
Jecture Germany GmbH Philipp-Reis-Straße 2, 36399 Freiensteinau Deutschland	Production and distribution of cannulas, cannula systems and related components
Jecture Poland Sp. z.o.o. ul. Gen. G. Orlicz-Dreszera 31A, Kozerki 05-825 Grodzisk Mazowiecki Poland	Production and distribution of cannulas, cannula systems and related components

Valid as of: 20 November 2025

Certification body

DAkkS
Deutsche
Akkreditierungsstelle
D-ZM-21753-01-00

ID 0374