

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

VYGON Germany GmbH
Prager Ring 100, 52070 Aachen
Germany

previous company name:
Vygon GmbH & Co. KG
Prager Ring 100, 52074 Aachen
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:

**Design and development, manufacture and distribution of
catheters and catheter sets for peripheral, central, venous,
arterial, otological, short and long-term applications,
accessories for catheter insertion and accessories for
intravenous infusion management.**

**Distribution of catheters and medical devices for infusion
management**

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.
0948-26-0114

Registered under
Z/26/04907E

Valid until
9 February 2028

Valid as of: 12 May 2026


Certification body



Deutsche
Akkreditierungsstelle
D-ZM-21753-01-00