

EC Design Examination Certificate

The Notified Body

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

herewith certifies that the design of the medical device(s):

**Vanguard Steerable Diagnostic Catheter BSA and BSB
with specifications listed in the appendix**

of the manufacturer:

**Vanguard AG
Landsberger Straße 266
12623 Berlin
Germany**

fulfills the below mentioned requirements of the **Council Directive 93/42/EEC**:

Annex II, section 4

This certificate assumes that MEDCERT has to be informed about any changes of the approved design. Changes need further approval by MEDCERT.

For the placing on the market of medical devices covered by this certificate, a separate EC certificate of conformity according to Annex II without section 4 of Council Directive 93/42/EEC is required.

Effective date: 2021-05-25

Expiry date: 2024-05-27

Report No.: 14020IA01F

Process No.: PP – 14020

Certificate No.: 14020GB411210525A

Hamburg, 2021-05-25

MEDCERT Certification Body
(Marcus Harder)

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

Form F10010009e EN / Rev. 7 / 2019.11.14



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

Appendix of EC Design Examination Certificate

Process No.: PP – 14020

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Specifications

Vanguard Steerable Diagnostic Catheter BSA						
33263	33382	33383	33384	33385	33386	33387
33388	33389	33390	33391	—	—	—

Vanguard Steerable Diagnostic Catheter BSB						
34988	34989	34990	34991	34992	—	—

– End of appendix –

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