



# EC Design Examination Certificate

## Council Directive 93/42/EEC Annex II Section 4

This is to certify that the manufacturer

### Acandis GmbH

Theodor-Fahrner-Str. 6  
75177 Pforzheim  
Germany

that the design of the following device(s)

#### NeuroSlider® Microcatheter

is conform to the Essential Requirements of Annex I of the Council Directive 93/42/EEC concerning medical devices.

This EC Design Examination Certificate is only valid in connection with the valid DQS Medizinprodukte GmbH Certificate No. 376850 MR2. Changes to the approved design are subject to further approval by the Notified Body.

**Basis of examination:** Produktakte B Zusammenfassung TD NeuroSlider 2018-02-07

Further basis for the examination is referenced in the examination report and relating documents mentioned below.

**Examination report:** 411\_18d\_Bericht\_NeuroSlider\_V1 dated 2018-03-16

The results of the examination are contained in the above mentioned report and the relating documents mentioned within.

Certificate registration no. 376850 MRA

Certificate unique ID 170710589

Effective date 2018-03-16

Expiry date 2023-03-15

Frankfurt am Main 2018-03-16

### DQS Medizinprodukte GmbH

Sigrid Uhlemann  
Managing Director

Dr. Thomas Feldmann  
Head of Certification Body

August-Schanz-Straße 21, 60433 Frankfurt am Main,  
Tel. +49 (0) 69 95427-300, [medical.devices@dqs-med.de](mailto:medical.devices@dqs-med.de)

DQS Medizinprodukte GmbH is a Notified Body according to Council Directive 93/42/EEC concerning medical devices with the Identification Number 0297.