EC Design Examination Certificate

Council Directive 93/42/EEC Annex II Section 4

This is to certify that the manufacturer

Cerus Endovascular Limited

John Eccles House
Oxford Science Park
Oxford
OX4 4GP
United Kingdom

that the design of the following device(s)

Contour Neurovascular System ™
Contour 021
NEQSTENT
NEQSTENT 021

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. 523104 MR2. Changes to the approved design are subject to further approval by the Notified Body.

Basis of examination:
Contour Neurovascular System™ dated 2018-12-19
Contour Neurovascular System™ dated 2020-03-10
Contour Neurovascular System™ dated 2020-07-15
Contour Neurovascular System™ dated 2020-01-22

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

Examination report:
411_18e_Report_TFR_cerus_Contour_V3 dated 2019-09-04
0_411_18e_Report_TFR_Cerus_Contour_V5 dated 2020-03-26
411_18e_Report_TFR_Cerus_Contour_V6 dated 2020-07-18
0_411_18e_Report_TFR_Cerus_Contour_V8.docx dated 2021-03-29

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

Certificate registration no.	523104 MRA
Certificate unique ID	170775520
Effective date	2021-03-29
Expiry date	2024-05-26
Frankfurt am Main	2021-03-29

DQS Medizinprodukte GmbH

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