EC Design Examination Certificate

Council Directive 93/42/EEC Annex II Section 4

This is to certify that the manufacturer

MicroVention, Inc.

1311 Valencia Ave.
Tustin, CA, 92780
United States of America

that the design of the following device(s)

WEB™ Aneurysm Embolization System

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. 411133 MR2. Changes to the approved design are subject to further approval by the Notified Body.

Basis of examination: RF17-0001 Web & WDC TF 2017-03-30.docx dated 2017-05-10
RF17-0001B WEB TF dated 2018-10-12

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

Examination report: 411_18e_Report_TFR_WEB_201706 dated 2017-08-14
411_18e_Reprt_TFR_WEB_201706 dated 2019-01-22

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

Certificate registration no.	535858 MRA
Certificate unique ID	170732615
Effective date	2019-01-22
Expiry date	2022-08-13
Frankfurt am Main	2019-01-22

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.