EC Design Examination Certificate

Council Directive 93/42/EEC Annex II Section 4

This is to certify that the manufacturer

DOT GmbH

Charles-Darwin-Ring 1a
18059 Rostock
Germany

that the design of the following device(s)

BONITmatrix®

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. 280145 MR2. Changes to the approved design are subject to further approval by the Notified Body.

Basis of examination: Produkthauptakte BONITmatrix dated October 2014

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

Examination report: Abschlussbericht_DOT_BONITmatrix 2014 dated 2014-11-15

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

Certificate registration no.	519887 MRA
Certificate unique ID	170608229
Effective date	2014-11-15
Expiry date	2018-11-30
Frankfurt am Main	2014-11-15

DQS Medizinprodukte GmbH

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Head of Certification Body

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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.

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