EC Design Examination Certificate

Council Directive 93/42/EEC Annex II Section 4

This is to certify that the manufacturer

pro med instruments GmbH

Bötzinger Straße 38
79111 Freiburg
Germany

that the design of the following device(s)

DORO® Blades

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. 221096 MR2. Changes to the approved design are subject to further approval by the Notified Body.

Basis of examination: TF02_A00_Technische_Doku_DORO_v1.4 dated 2015-04-15

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

Examination report: 411_18d_Bericht_Produktprüfung_DORO+Spatel_V2 dated 2015-10-28

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

Certificate registration no.	221096 MRA
Certificate unique ID	170633721
Effective date	2015-10-28
Expiry date	2020-10-27
Frankfurt am Main	2015-10-28

Digitally signed by Cucerenco Olesea
Date: 2020.06.19 14:53:35 EEST
Reason: MoldSign Signature
Location: Moldova

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.

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