Medical Device Full Quality Assurance System Certificate GB22/00000250

The management system of

Haag Streit UK Ltd

Edinburgh Way Harlow Essex CM20 2TT United Kingdom

has been assessed and certified as meeting the requirements of

Part II of The Medical Device Regulations 2002 Annex II excluding Section 4 [as modified by Part 2 of Schedule 2A to The Medical Device Regulations 2002]

For the following products
Tonometers.
Tonosafe sterile disposable prism.

Where the above scope includes class III medical device(s), a valid Design Examination Certificate according to Annex II (Section 4) [as modified by Part 2 of Schedule 2A of The MDR 2002] is a mandatory requirement for each device in addition to this certificate to place that device on the market

This certificate is valid from 06 July 2022 until 06 July 2027 and remains valid subject to satisfactory surveillance audits

Issue 1. Certified since 13 March 1995.
Certification is based on reports numbered GB/PC/230587

Authorised by

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