Medical Device Full Quality Assurance System Certificate
GB22/00000250

The management system of

Haag Streit UK Ltd

Unit C Woodside Industrial Estate Dunmow Road Bishop's Stortford Hertfordshire CM23 5RG
United Kingdom

has been assessed and certified as meeting the requirements of

Part II of The Medical Devices Regulations 2002, Annex II excluding section 4 [as modified by Part 2 of Schedule 2A to The Medical Devices 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

Certification is based on reports numbered GB/PC/230587
Previous certificate number: N/A
Change in between this certificate and previous one: Change of company address

This certificate is valid from 15 April 2024 until 06 July 2027 and remains valid subject to satisfactory surveillance audits.

Issue 2. Certified since 13 March 1995
Certified activities performed by additional sites are listed on subsequent pages.

L. Henderson

Authorised by
Lynn Henderson

SGS United Kingdom Ltd Approved Body 0120
Rossmore Business Park, Ellesmere Port, Cheshire, CH65 3EN, UK
t +44 (0)151 350-6666 - www.sgs.com

Medical Device Full Quality Assurance System Certificate
GB22/00000250, continued

Haag Streit UK Ltd

Part II of The Medical Devices Regulations 2002, Annex II excluding section 4 [as modified by Part 2 of Schedule 2A to The Medical Devices Regulations 2002]

Issue 2

Sites

Haag Streit UK Ltd
Unit C Woodside Industrial Estate Dunmow Road Bishop's Stortford Hertfordshire CM23 5RG United Kingdom

HAAG STREIT UK LIMITED
Suites 2 & 4 2 Woodside Industrial Estate Dunmow Road Bishop's Stortford Hertfordshire CM23 5RG United Kingdom