Medical Device Production Quality Assurance System Certificate GB22/00000251

The management system of

Haag Streit UK Ltd

Unit A Cartel Business Estate 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 V [as modified by Part 2 of Schedule 2A to The Medical Device Regulations 2002]

For the following products

Annex V Metrological aspects only – Restricted to the aspects of manufacture concerned with the conformity of the devices with metrological requirements Synoptophore Prism Bars Pinhole Occluder RAF Binocular Gauge Maddox Wing Test
Annex V Sterility aspects only – Restricted to the aspects of manufacture concerned with the conformity of the devices with sterility requirements Schirmer tear test.

Where the above scope includes class IIb or class III medical device(s), a valid Type Examination Certificate according to Annex III [as modified by Part 2 of Schedule 2A to 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/08905

Authorised by

SGS United Kingdom Ltd, Approved Body 0120
Rossmore Business Park, Ellesmere Port, Cheshire, CH65 3EN, UK
t +44 (0)121 541 4743 www.sgs.com