

The management system of

Altomed Limited

Unit 2 Witney Way, Boldon Business Park, Tyne and Wear, NE35 9PE, UK

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

For the following products

Sterile and non-sterile: Symblepharon rings, Ophthalmic conformers, Ocular tumour markers

Orbital Implants
Platinum segment eyelid weight
Implantable tubes

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 25 January 2021 until 25 October 2023 and remains valid subject to satisfactory surveillance audits.

Issue 5. Certified since 23 March 1999 and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered GB/PC 09704

Authorised by

Global Medical Devices Certification Manager

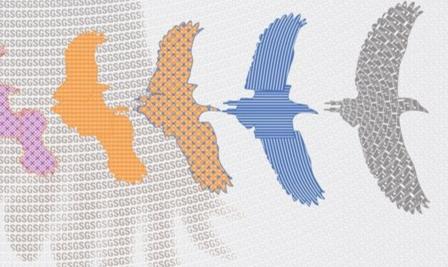
SGS Belgium NV, Notified Body 1639

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LPMD5007 - Certificate CE1639 Annex II-4_EN rev. 02

Page 1 of 1





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