EC Certificate Full Quality Assurance System: Certificate US02/56962

The management system of

Ortho Technology, Inc.

4614 Pet Lane, Suite D-101, Lutz, FL, 33559, United States

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

Metal Brackets and Buccal Tubes, Ceramic Brackets, Composite Brackets, Orthodontic Bands and Assemblies, Orthodontic Adhesives and Cement Systems, Orthodontic Wires and Accessories, Orthodontic Burs, Elastomerics, Fixed Attachments, Auxiliaries, Intraoral Appliances

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 01 April 2019 until 15 December 2023
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 15 December 2021
Issue 10. Certified since 10 October 2002

Certification is based on reports numbered WW/MC 208138

Authorised by

Jonathan M. Hall

SGS United Kingdom Ltd, Notified Body 0120
202B Worle Parkway, Weston-super-Mare, BS22 6WA UK
t +44 (0)1934 522917 f +44 (0)1934 522137 www.sgs.com

SGS CE 02 0215