EC Certificate Full Quality Assurance System: Certificate GB18/962169

The management system of

Technovent Limited

Unit 5, York Park, Bridgend Industrial Estate, CF31 3TB, 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 magnetic attachments for craniofacial and dental prostheses.

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 18 September 2018 until 20 March 2022
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 17 September 2019
Issue 1. Certified since 18 September 2018

Certification is based on reports numbered GB/PC 08504

Authorised by

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