Medical Device Production Quality Assurance System certificate
GB22/00000335

The management system of

Technical Service Consultants Limited

Microbiology House Fir Street Heywood OL10 1NW United Kingdom

has been assessed and certified as meeting the requirements of

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

For the following products
Surgically Invasive Sterile swabs for cell and secretion collection, for diagnostic purposes.

Annex V (sterility aspects only) - Restricted to the aspects of manufacture concerned with securing and maintaining sterile conditions
Sterile swabs for cell and secretion collection, for diagnostic purposes via body orifices.

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 The 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 15 August 2022 until 15 August 2027 and remains valid subject to satisfactory surveillance audits.

Issue 1. Certified since 22 June 1998.
Certification is based on reports numbered GB/PC/233757

Authorised by

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