EC Certificate Production Quality Assurance System: Certificate GB15/92944

The management system of

Technical Service Consultants Limited

Microbiology House, Fir Street, Heywood, OL10 1NW, UK

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V

For the following products

Sterile swabs for surgically invasive 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 EC Type Examination Certificate according to Annex III is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 31 May 2017 until 22 June 2021
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 20 July 2019
Issue 2. Certified since 22 June 1998

Certification is based on reports numbered GB/PC 233757

Authorised by

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

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