EC Certificate Production Quality Assurance System: Certificate GB12/85504

The management system of

Steroplast Healthcare Ltd

Unit 2, Alpha Point, Bradnor Road, Manchester, M22 4TE, UK

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V

For the following products

Directive 93/42/EEC on Medical Devices, Annex V

Sterile hydrogel dressings, Instant Ice Pack for treatment of sprain & bruising, Sterile Wound Wash Solution and Sterile Saline Wipes, Sterile Wound Closure Strips. Sterile eyewash solution.

Annex V Sterility aspects only - Restricted to the aspects of manufacture concerned with securing and maintaining sterile conditions

Sterile: adhesive wound dressings, low adherent wound dressings, non woven swabs, first aid wound dressings and first aid kits.

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 03 July 2015 until 17 April 2020
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 09 March 2018
Issue 6. Certified since 17 April 2012

Certification is based on reports numbered GB/PC 228394

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

SGS CE 14 0215