bsi.

By Royal Charter

EC Certificate - Production Quality Assurance

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

No. **CE 01865**

Issued To: **Riverside Medical Packaging Company Ltd.**
Newmarket Drive
Derby
DE24 8SW
United Kingdom

In respect of:

The manufacture of allergy prick test needles and sterile handpieces for laser wires for ear, nose and throat surgery.

Those aspects of Annex V relating to maintaining and securing the sterility and metrology of syringes, accessories and procedure packs for compounding and single use syringe packs and accessories for clinical administration of drugs manually.

Those aspects of Annex V relating to maintaining and securing the sterility of surgical instruments and dental kits.

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex V. The quality assurance system meets the requirements of the directive. For the placing on the market of class IIb and class III products an Annex III certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 0086):

Stewart Brain, Head of Compliance & Risk - Medical Devices

First Issued: **1998-03-20** Date: **2018-05-17** Expiry Date: **2023-05-18**

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