UKCA Certificate - Production Quality Assurance

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

No. **UKCA 760030**

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

In respect of:

The manufacture of allergy prick test needles.

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.

On the basis of our examination of the quality assurance system under the requirements of Part II of the Medical Devices Regulations 2002, Annex V [as modified by Part II of Schedule 2A to The Medical Devices Regulations 2002]. The quality assurance system meets the requirements of the regulation. For the placing on the market of class IIb and class III products an Annex III certificate (modified as described above) is required.

For and on behalf of BSI, an Approved Body for the above Regulation (Approved Body Number 0086):

Graeme Tunbridge, Senior Vice President Medical Devices

First Issued: **2021-12-14** Date: **2023-05-11** Expiry Date: **2028-05-18**

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Supplementary Information to UKCA 760030

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Device code	Device name	Intended purpose per IFU
Class IIa		
MD 0106	Morrow Brown Allergy Prick Test Needle	N/A – Class IIa
Class Is, Im		
MD 0104	Sterile Transfer Sets and Accessories	N/A – Class Is, Im

First Issued: **2021-12-14** Date: **2023-05-11** Expiry Date: **2028-05-18**

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UKCA Certificate - Production Quality Assurance Certificate History

Certificate No: **UKCA 760030**

Date: **2023-05-11**

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Date	Reference Number	Action
2021-12-14	3560526	Issued. Traceable to CE 01865.
Current	3914623	Certificate Renewal

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