Medical Device Production Quality Assurance System Certificate GB22/0000494

The management system of

CliniMed Ltd

Cavell House, Knaves Beech Way, Loudwater High Wycombe Buckinghamshire HP10 9QY 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]
Restricted to the aspects of manufacture concerned with securing and maintaining sterile conditions

For the following products

CliniSorb, secondary wound care dressing for the management of wound odour
Instillaquill, sterile single use extension tube for use in gynecological investigations,
LBF Sterile Wipes for the prevention of skin irritation
LBF Sterile No Sting Barrier Film 1ml & 2ml Foam Applicators for the prevention of skin irritation
LBF Sterile No Sting Barrier Film Spray (30ml and 50ml) for the prevention of skin irritation
Appeel Sterile Sachet for use with stoma pouches and other adhesive appliances
Appeel Sterile Wipe for use with stoma pouches and other adhesive appliances
Appeel Sterile Foam Applicator for use with stoma pouches and other adhesive appliances
Appeel Sterile Spray for use with stoma pouches and other adhesive appliances.

This certificate is valid from 01 November 2022 until 01 November 2027 and remains valid subject to satisfactory surveillance audits.

Issue 1. Certified since 01 November 2022
Certification is based on reports numbered GB/PC/240482

Authorised by
Lynsey Hall
Head of Approved Body 0120

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