Medical Device Production Quality Assurance System Certificate GB22/00000494



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

The Scope of Registration appears on page 2 of this certificate

This certificate is valid from 27 October 2025 until 27 October 2030 and remains valid subject to satisfactory surveillance audits.

Issue 2. Certified since 01 November 2022

K Snell

Authorised by Robert Snell Head of Approved Body

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Medical Device Production Quality Assurance System Certificate GB22/00000494, continued



CliniMed Ltd

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

CliniSorb, secondary wound care dressing for the management of wound odour

Issue 2

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 Foam Applicator for use with stoma pouches and other adhesive appliances

Appeel Sterile Spray for use with stoma pouches and other adhesive appliances

Certification is based on reports numbered GB/PC/240482 Previous certificate number: N/A Change in between this certificate and previous one: N/A

