EU Declaration of Conformity

Legal Manufacturer:
Guilin HBM Health Protections, Inc.
No.1-2, Shuijing East Road, Economic and Technological Development Area, 541805 Guilin, Guangxi, China

EU Authorized Representative:
Continental Medical B.V.
Strevelsweg 700 303 Unit E7288 3083AS Rotterdam, Netherlands

Hereby declare in our own responsibility that the product:
Latex Surgical Gloves, Powder Free

Intended purpose: The surgical gloves are single use devices in sterile condition, intended to be worn on the hands of operating room personnel, to protect the patient and the surgeon against contamination and also to provide a barrier for the wearer against potentially infectious material or different chemicals used within the healthcare sector. The gloves are appropriate for use during invasive and non-invasive medical procedures requiring sterility and are designed for use in the environments within hospitals and other healthcare facilities.

Product Reference	Description
MSG7255, MSG7260, MSG7265, MSG7270, MSG7275, MSG7280, MSG7285, MSG7290	Cream Sterile Latex Surgical Gloves, Powder Free

are in conformity with Personal Protective Equipment Regulation (EU) 2016/425,
under Category III risk set out in Annex I
meeting the provisions of EN ISO 21420:2020+A1:2024, EN ISO 374-1:2016 + A1:2018, EN ISO 374- 5:2016

The products are subject to the procedure set out in:
Annex V (EU type examination) and Annex VIII (Module D) of Regulation (EU) 2016/425 issued by
Notified Body:
SATRA Technology Europe Limited
Bracetown Business Park Clonee, D15YN2P, Ireland
Notified Body Number: 2777
EU type-examination certificate No：2777/15894-04/E00-00

Shun Huang

Huang Shun / Quality Manager & PRRC

Place/date: Guilin, China, 2026-03-13

GUILIN HBM HEALTH PROTECTIONS, INC.