MODULE D CERTIFICATE

Issued To

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

Has been found to conform to the requirements of Annex VIII (Module D) of the PPE Regulation (EU) 2016/425 – Conformity to type based on quality assurance of the production process, for the manufacture and distribution of Five fingered latex powder free surgical gloves and medispo powder free, polyisoprene surgical gloves in a range of colours and sizes.

Standards	**Module B Certificates***
EN ISO 374-1:2016+A1:2018	2777/18178-XX/E00-00
EN ISO 374-4:2019	2777/23740-XX/E00-00
EN ISO 374-5:2016	PPE-2425-23213-Cert
EN ISO 21420:2020	
EN 421:2010 (Radioactive contamination only)	

* Where 'XX' denotes the Module B certificate issue number

Date of first issue: 18th December 2023 **Expiry Date:** 17th December 2026
Latest Issue: 16th January 2026

Signed on behalf of SATRA Technology Europe Ltd – Notified Body Number: 2777

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

Certificate Validity

Subject to annual surveillance and ongoing conformity.
The certificate holder is responsible for ensuring that the notified body is advised of changes to any aspect of the overall process utilised in the manufacture of the product; failure to do so could invalidate the certificate in respect of product manufactured following the introduction of such changes.

Date	Comments	Satra Reference No.
18th December 2023	Initial audit from the 8th and 9th Nov 2023	STE4810Z2S7
26th June 2024	Addition of certificate PPE-2425-23213-Cert to the Module D certificate	STE2007937
28th November 2024	Year 1 surveillance audit from the 13th & 14th Nov 2024	STE2013276
16th January 2026	Year 2 surveillance audit from the 25th & 26th Nov 2025	STE2032503

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TERMS AND CONDITIONS

The following conditions apply in addition to SATRA's standard terms and conditions of business and those given in the current certification agreement.

Please note:

1. Products covered by this certificate are classified by Regulation (EU) 2016/425 as category III Personal Protective Equipment and as such CE Marking is reliant on current compliance with Annex VIII Module D of this Regulation.
2. Full details of the scope of the certification and product(s) certified are contained within the manufacturer's technical documentation.
3. Where a translation of this certificate exists, the English language version shall be considered as the authoritative text.
4. Certification is limited to production undertaken at the sites listed in the Module B certificates' technical file.
5. Ongoing manufactured product shall be consistent with the product(s) certified and listed on the relevant module B certificate.
6. The Manufacturer shall inform SATRA of any changes to the certified product or technical documentation.
7. Where results obtained are within the budget of uncertainty when compared to the pass requirement, classification, or performance level, then it is the responsibility of the manufacturer to ensure that the factory production control and manufacturing tolerances are such that the product placed on the market meets with the stated requirements, classifications, or performance levels.
8. This certificate shall be kept together with the relevant technical documentation in a safe place by the client named on this certificate. Production of this certificate and other documentation may be required by the authorities.
9. This certificate relates only to the condition of the testable items at the time of the certification procedure and is subject to the expiry date shown.
10. SATRA reserves the right to withdraw this certificate if it is found that a condition of manufacture, design, materials, or packaging have been changed and therefore no longer comply with the requirements of Regulation (EU) 2016/425 on Personal Protective Equipment.

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