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EAR CERTIFICATE

(Article 11 of the European Medical Devices Regulation - MDR (EU) 2017/745)

Date of Issuance	May 17, 2022
Renewal Date	May 16, 2023
Certificate No.	EC-EAR-2022-0059
Manufacturer	LIENTEH TECHNOLOGY SDN BHD
Manufacturer Address	Lot 6483, Jalan Sungai Puloh KU5, Kawasan Perindustrian Sungai Puloh, 42100 Klang, Selangor Darul Ehsan, Malaysia
Device Class(es)	Class I, Rule 5
Device(s) with TF-ID	1. Non-Sterile Nitrile Examination Gloves Powder-Free TF-LT-01

Conclusion

Europecert declares that the aforementioned manufacturer has fulfilled the essential requriement of appointing an European Authorized Respresentatitve in accordance with Article 11 of the MDR (EU) 2017/745 and MEDDEV 2.5/10 requirements.

europecert
Alsstrasse 97
41063 Mönchengladbach - Germany
Tel: +49 (0) 2161 990 883 1
support@europecert.eu

Joe Raj Kumar
- General Manager -
Europecert

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