

Lot 6070, Jalan Haji Abdul Manan 6th Miles Off Jalan Meru 41050 Klang, Selangor, Malaysia Tel: 603-33929888 (8 lines) Fax: 603-33923328 E-MAIL: maxter@tm.net.my www.maxter.com.my

# **Declaration of Conformity – UK**

Maxter Glove Manufacturing Sdn Bhd hereby confirms that the product mentioned below complies with EU Regulations and Standards and is manufactured according to ISO9001 & ISO13485 standard requirements.

# Aurelia Perform 2.5 mil Teal Powder Free Nitrile Examination Glove

Size	Product Code	Barcode	UDI Code
X-Small	92395	955-500210-2746	1955-500210-2743
Small	92396	955-500210-2753	1955-500210-2750
Medium	92397	955-500210-2760	1955-500210-2767
Large	92398	955-500210-2777	1955-500210-2774
X-Large	92399	955-500210-2784	1955-500210-2781

# **Classification of the product:**

- Class I Medical Device based on Rule 5 transient use, Annex IX of COUNCIL DIRECTIVE 93/42/EEC of 14 June 1993 concerning medical devices, which is implemented into UK law by the Medical Device Regulations 2002, as amended.
- Class I Medical Device based on Rule 5 transient use, Annex VIII of Medical Device Regulation (EU)
  2017/745
- CAT III PPE (EU) 2016/425, which is directly implemented into UK law.

# **Product mentioned above complies with:**

- Medical Device Regulation 2002 for Class I Medical Devices, implementing COUNCIL DIRECTIVE 93/42/EEC
- The General Safety and Performance requirements of Annex I, Medical Regulation (EU) 2017/745 for Class I Medical Devices and with the Article 19 requirements.
- The provisions of Personal Protective Equipment (PPE) Regulation (EU) 2016/425, including the General Safety Requirements (Annex II), Module B EU-Type Examination Certification and Module D, Conformity to type, based on Quality Assurance of the production process.
- EEC regulations concerning the conformity of materials and products that are allowed to come into contact with food. In accordance with Regulation EEC 1935/2004, Regulation EC 10/2011 & Regulation EC 2023/2006.

### **Certification:**

 Module B, UKCA Type Examination Certificate issued by Approved Body: Satra (0321) – Certificate No. AB0321/19875-01/E00-00



Lot 6070, Jalan Haji Abdul Manan 6th Miles Off Jalan Meru 41050 Klang, Selangor, Malaysia Tel: 603-33929888 (8 lines) Fax: 603-33923328 E-MAIL: maxter@tm.net.my www.maxter.com.my

- Module D, Regulation 2016/425 as brought into the UK Law, Examination Certificate issued by Approved Body: SGS UK, Approved Body UKCA0120 – Certificate No. MY21/1811030640
- ISO9001:2015
- ISO13485:2016

#### **Gloves tested according to Harmonised Standards:**

- EN374-1 chemical resistance
- EN374-5 microbiological resistance
- EN455 1,2,3, 4 medical devices
- EN21420- physical attributes or EN420 during the transition period

#### **User Information:**

- The gloves are suitable for contact with dry, fatty, alcoholic, and aqueous food for short term contact based on the outcome of the overall migration test on the food simulants.
- The product does not contain natural rubber latex. Contains accelerators which may cause allergic reactions. Please retain the packaging for reference.
- Store below 40°C/104°F in dry, clean condition and away from direct sunlight.

# Responsibility

• This Declaration of Conformity is issued under the responsibility of the Manufacturer, as indicated below:

## **Manufacturer:**

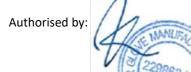
 Maxter Glove Manufacturing Sdn. Bhd., located at Lot 6070, Jalan Haji Abdul Manam, 6<sup>th</sup> Miles off Jalan Meru, 41050 Klang, Selangor, Malaysia

# **Authorised Representatives:**

- EU Representative is Supermax Healthcare Europe Limited, 38 Main Street, Swords Co. Dublin, K67 E0A2, Ireland
- UK Representative is Supermax Healthcare Limited, 12-16 Titan Drive, Fengate, PE1 5XN, Peterborough, United Kingdom



Lot 6070, Jalan Haji Abdul Manan 6th Miles Off Jalan Meru 41050 Klang, Selangor, Malaysia Tel: 603-33929888 (8 lines) Fax: 603-33923328 E-MAIL: maxter@tm.net.my www.maxter.com.my



Yap Peak Geeh Senior Manager QA & Regulatory Affairs

Date: 25-Jun-2022

\*This declaration is valid for period of 2 years from the date of issue or until any changes to regulations or products are applicable.