EU Quality Management System Certificate

Regulation (EU) 2017/745, Annex IX Chapter I and III

MDR 740797 R000

Manufacturer: WRP Asia Pacific Sdn Bhd

Address:
Lot 1, Jalan 3
Kawasan Perusahaan
Bandar Baru Salak Tinggi
Sepang
Selangor
43900
Malaysia

Single Registration Number: MY-MF-000004690

EU Authorised Representative: Remesco Handelsges.m.b.H.

Address:
Muthgasse 36/19
1190 Wien
Austria

Scope: See attached **Device Schedule**

On the basis of our examination of the quality system in accordance with Regulation (EU) 2017/745, Annex IX Chapter I and III, the quality system meets the requirements of the Regulation. For the placing on the market of Class III devices, and Class IIb implantable devices that are not considered well-established technologies as specified in Article 52(4) an additional Annex IX Chapter II certificate is required.

For and on behalf of BSI, a Notified Body for the above Regulation (Notified Body Number 2797):

Graeme Tunbridge, Senior Vice President Medical Devices

First Issue Date: **2023-10-20**

Current Issue Date: **2023-10-20**

Starting Validity Date: **2023-10-20**

Expiry Date: **2028-10-19**

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Device Schedule: Class IIa, Custom-made and other devices

Device(s)	Risk Classification
Sterile powder free synthetic polychloroprene surgical gloves	Class IIa
Sterile powder free synthetic nitrile surgical gloves	Class IIa
Sterile powder free synthetic polyisoprene surgical gloves	Class IIa
Sterile powder free natural rubber latex surgical gloves	Class IIa
Sterile powder free synthetic polyisoprene radiation attenuating surgical gloves	Class IIa
Sterile powder free natural rubber latex radiation attenuating surgical gloves	Class IIa
Sterile powder free nitrile examination gloves	Class Is
Sterile powder free natural rubber latex examination gloves	Class Is
For Class Is devices, the Notified Body conformity assessment is limited to the aspects relating to establishing, securing and maintaining sterile conditions.	

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Certificate History

(References to applicable Common Specifications, Harmonized Standards complied with, and the relevant test and audit reports that support any of the below certificate changes may be requested from Certificate.Verification@bsigroup.com)

Date	Reference Number	Action
Current	3335364	Issued

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Validity of this certificate is conditional on the Manufacturer's quality system being maintained to the requirements of the Regulation as demonstrated through the required surveillance activities of the Notified Body.
This certificate was issued electronically and is bound by the conditions of the contract.

NB Contact: BSI Group The Netherlands B.V., Say Building, John M. Keynesplein 9, 1066 EP, Amsterdam, Netherlands. Tel: + 31 (0) 20 346 07 80
Corporate Contact: BSI Group Assurance Limited, registered in England under number 05435540 at 389 Chiswick High Road, London, W4 4AL, UK.
A Member of the BSI Group of Companies.