

EU Quality Management System Certificate

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

MDR 780250 R000

Manufacturer: Professional Disposables International Ltd

Address:

Pywell Road
Willowbrook Industrial Estate
Corby
NN17 5XJ
United Kingdom

Single Registration Number: GB-MF-000035214

EU Authorised Representative: NEX Medical Antiseptics Srl

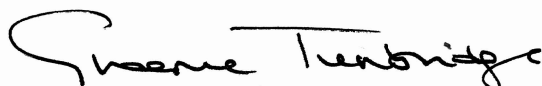
Address:

Via Per Arluno
37/39 20003 Casorezzo (MI)
Italy

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 Global Regulatory & Quality

First Issue Date: **2024-07-26**

Current Issue Date: **2024-07-26**

Starting Validity Date: **2024-07-26**

Expiry Date: **2029-07-25**

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

Device(s)	Risk Classification
Liquid impregnated disinfectant wipes for non-invasive medical devices	Class IIa



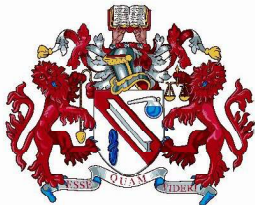
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By Royal Charter

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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	3787066	Issued



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