

# EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

**No.**

**CE 628500**

**Issued To:**

**PDI Ltd  
Aber Park  
Flint  
Flintshire  
CH6 5EX  
United Kingdom**

In respect of:

**Design, development and manufacture of non-sterile liquid impregnated disinfectant wipes for the disinfection of non-invasive medical devices.**

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex II excluding section 4. The quality assurance system meets the requirements of the directive. For the placing on the market of class III products an Annex II section 4 certificate is required.

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



Gary E Slack, Senior Vice President - Medical Devices

First Issued: **2015-11-05**

Date: **2021-03-15**

Expiry Date: **2024-05-26**

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Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.  
This certificate was issued electronically and is bound by the conditions of the contract.

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## Supplementary Information to CE 628500

Issued To:

**PDI Ltd  
Aber Park  
Flint  
Flintshire  
CH6 5EX  
United Kingdom**

Number	Device Name	Intended purpose per IFU
<b>Class IIa</b>		
MD 0108	Super Sani-Cloth Plus Disinfectant wipe - canister	-----
MD 0108	Sani-Cloth 70 Disinfectant wipe - canister	-----
MD 0108	Sani-Cloth Active Disinfectant wipe – canister, Bucket, Doy bag	-----
MD 0108	Sani-Cloth CHG 2% Disinfectant wipe - sachet	-----
MD 0108	Sani-Cloth Chlor Disinfectant wipe - canister	-----
MD 0108	Sani-Cloth AF Universal Disinfectant wipe – Canister, Bucket, Flow-wrap	-----

First Issued: **2015-11-05**Date: **2021-03-15**Expiry Date: **2024-05-26**

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Information and Contact: BSI, Say Building, John M. Keynesplein 9, 1066 EP Amsterdam, The Netherlands Tel: + 31 20 346 0780

BSI Group The Netherlands B.V. registered in The Netherlands under 33264284.

A member of BSI Group of Companies.

# EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

## List of Significant Subcontractors

Recognised as being involved in services relating to the product covered by:

Certificate No: **CE 628500**  
 Date: **2021-03-15**  
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 Aber Park  
 Flint  
 Flintshire  
 CH6 5EX  
 United Kingdom**

### Subcontractor:

### Service(s) supplied

Nex Medical Antiseptics Srl.  
 Via Arluno, 37/39  
 20010 Casorezzo Milan  
 Italy

**EU Representative**

PDI (EMEA) Ltd  
 Pywell Road  
 Willowbrook East Industrial Estate  
 Corby  
 NN17 5XJ  
 United Kingdom

**Manufacture**

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# EC Certificate - Full Quality Assurance System

## Certificate History

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 Date: **2021-03-15**  
 Issued To: **PDI Ltd  
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 CH6 5EX  
 United Kingdom**

Date	Reference Number	Action
05 November 2015	8266784	First issue.
07 February 2019	9628060	Traceable to NB 0086.
20 November 2019	3044874	Removed sterile devices from scope. Removed three subcontractors from list of critical subcontractors.
18 September 2020	3273939	Subcontractor Pluswipes Limited name change to PDI (EMEA) Ltd, Addition of supplementary information table. Addition of new product Sani-Cloth AF Universal.
22 October 2020	3219326	Renewal Addition of EU Representative Administrative correction to subcontractor's address Nice-Pak International Limited and PDI (EMEA) Ltd.,
15 March 2021	3372694	Removal of subcontractor Jonarve Limited, Flint, United Kingdom.

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 Date: **2021-03-15**  
 Issued To: **PDI Ltd  
 Aber Park  
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Date	Reference Number	Action
<b>Non-significant changes approved after the 26th May 2021 as per the Transitional Provisions of MDR Article 120.3</b>		
06 December 2021	3564554	<p>Change of legal manufacturer name and address. From:            PDI Ltd,            Aber Park            Flint            Flintshire            CH6 5EX            United Kingdom            To:            Professional Disposables International Ltd.,            Pywell Road            Willowbrook Industrial Estate            Corby            NN17 5XJ            United Kingdom</p> <p>Change of device name.            From (before): Sani-Cloth CHG 2%            To (current): Prevantics 2% CHG            Removal of sub-contractor Nice-Pak International Limited., for the service of manufacture.</p>

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Date: **2021-03-15**  
Issued To: **PDI Ltd  
Aber Park  
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Date	Reference Number	Action
06 April 2022	3661352	Change to information on the device table: Device name changed from Sani-Cloth AF universal to Sani-Cloth AF.

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6 April 2022

PDI Ltd  
Aber Park  
Flint  
Flintshire  
CH6 5EX  
United Kingdom

To whom it may concern,

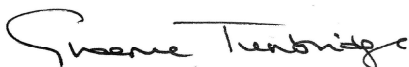
The transitional provisions specified in MDR Article 120(3) prohibit Notified Bodies from issuing new certificates or amending, modifying, supplementing any existing MDD/AIMDD certificates from 26<sup>th</sup> May 2021.

This letter is to confirm that BSI has reviewed and approved the change(s) detailed in the table below. These changes do not represent a significant change in design or intended purpose under MDR Article 120(3) and as per the guidance provided in MDCG 2020-3. The related MDD certificate specified below remains valid until the expiry date specified on the certificate.

Certificate	Directive and Annex	Reference Number	Changes approved
CE 628500	93/42/EEC Annex II excluding 4	3661352	Change to information on the device table: Device name changed from Sani-Cloth AF universal to Sani-Cloth AF.

Should you have any queries concerning your certification, or if we can be of further assistance to you, please contact your BSI Scheme Manager.

Yours sincerely,



Graeme Tunbridge  
Senior Vice President, Medical Devices