PDI Dry Scrub Brush - Information for Use (IFU)

PDI Dry Scrub Brush is a single use brush/sponge with nail cleaner supplied sterile.

Uses: Disinfection for general surfaces in the Medical Area.

Directions: Remove a scrub brush from the carton tower dispenser. Peel back the easy peel label to release the brush/sponge. Fully remove from packaging and use as required. PDI Dry brush/sponge is compatible with any detergent or antiseptic cleansing solution.

Caution: None.

PPE: Not applicable.

Addresses:
Distributed by: PDI Ltd, Aber Park, Flint, Flintshire, CH6 5EX
T: +44(0)1352 736700
E: sales@pdi-hc.co.uk
Manufactured by: NEX MEDICAL ANTISEPTICS S.r.l, Via Per Arluno 37/39, 20010 Casorezzo (MI)-Italy

Storage: Store upright. Keep out of direct sunlight.

Description: This product is sterile single use. Once a brush/sponge dispensed from the packaging, it can only be used once and then must be disposed. Overuse and re-use will not provide the efficacy required for correct product use. Do not use if the packaging appears damaged.

Sterile: This product has been sterilised by Ethylene Oxide.

Re-sterilisation: There is no data available for re-sterilisation.

Composition:
Brush: PE medical grade
Sponge: polyurethane
Nailcleaner: PP medical grade
All components are LATEX free

Regulatory Classification: CE marked 93/42/EEC

	Product Code	Pack Format	Brush/sponges per Pack	Packs per Case	NHS Code
	MXP00394	Single brush sponge	25 brush/sponges	9 (225 units)	FSL1533

Microbiology & skin irritation tests	
Hemolysis test	ISO 10993-4:2008
Cytotoxicity test	ISO 10993-5:2009
Skin irritation test	ISO 10993-10/Amd 1:2006

Packaging Symbols

Here are some of the most common symbols you may find on PDI product packaging and their description:

Title of Symbol	**Description of Symbol**	**Symbol** ***(Colours may vary)***
Consult instructions for use	Indicates the need for the user to consult the Instructions for use, available on the website at www.pdihc.com/global	
CE mark	CE mark Medical Device Directive Class IIa	
Do not re-use	Indicates a medical device (wipe) that is intended for one use, or for use on a single patient during a single procedure	
Manufacturer	Indicates the medical device manufacturer, as define in EU directives 90/385/EEC, 93/42EEC and 98/79/EC	
Date of Manufacture	Indicates the date the product was manufactured	
LOT	Allocated LOT or batch number of the product	
Expiry	The expiry date of the product	
Sterile	This product is sterile and has been sterilised by Ethylene Oxide	

Authorised Representative in the European Community

EC REP

NEX MEDICAL ANTISEPTICS S.r.l.
Via Per Arluno 37/39 20010 CASOREZZO (MI) Italy

Additional Information

Printed copies of this IFU are available upon request. Email: sales@pdi-hc.co.uk
Previous revisions of IFU's are available upon request. Email: sales@pdi-hc.co.uk
Document language: English (EN)