

Product Procedure Specifications Sheet

Specification Sheet No	SSN-FT1390-009			
Product Name	IV Safety Cannula Pack			
Product Code	36510607 (REV 10)			
Range	365 Healthcare			
Manufacturer	Wujiang Kangxin Surgical Dressings Co. Ltd			
Packaging	180pcs/case			
Sterilisation	EO Sterile			
Issue Date	5th Aug. 2024	Version No	R01a	

Intended use	Intended to be used to assist in minor Woundcare procedures
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Pack content				
Item	Product code	Quantity	Dimensions	Material
Day Review Label	2837	1	N/A	Paper
White Waste Bag	2004	1	35x45cm	High-Density Polyethylene
Blue Tissue Backed Drape Folded 24ply	8702	1	50x50cm	PE Film & Absorbent Tissue
White Non-Woven Swab 4ply	3103	2	10x10cm	Viscose & Terylene
Cannulation Record	2832	1	N/A	Paper
Transparent Iv Dressing	5201	1	7x8.5cm	PU Film, Adhesive, Release Paper, Non-Woven Fabric
Blue Tissue Backed Drape	8802	1	50x50cm	PE Film & Absorbent Tissue
VIP Form	11401	1	N/A	Paper

Quality Certification	ISO 13485:2016		
Device Classification	Class Is	Rule	Article 22
Conformity Route	Medical Devices Regulation (EU) 2017/745 Medical Devices Regulations 2002 (SI 2002 No 618, as amended) (UK MDR 2002)		
Storage conditions	Store in a cool dry place away from heat and direct sunlight		
Shelf life	5 Years - under recommended storage conditions		
Contraindications/Warnings	None		
Markings	Product Name, REF, 365 Logo and Address, Lot Number, Manufacturing Date, Expiry Date, Medical Device, Single Use, EO Sterile, Single sterile barrier, do not use if package damaged, Keep dry and away from sunlight, CE, ECREP, Manufacturer, Distributor address		

Document Reference	Document Name	Reviewed By	Approved By	Revision Number	Issue Date
DMS 04-004-03	Procedure Pack Specification Sheet	Jack Wong	Racheal Begg	R02	03 rd April 2024

**Additional information**

Products are Latex and phthalate free
Products contain no hazardous substances
Products contain no medicinal substances

Medical Device and Reporting System: Incidents caused by this device must be reported immediately to our offices.

The information provided within this Specification Sheet is correct as of last revision but it is subject to change without notice.

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Amendment Record

This procedure is reviewed to ensure its continuing relevance to the system and process that it describes. A record of contextual additions or omissions is given below. This revision history lists all changes to this document, the date of each revision, and the reason for each revision.

Revision Number	Revision Date	Reason for Revision	Updated by
R01a	05/08/2024	Update to Mediq UK Brand	Elizabeth Ogunkolade

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