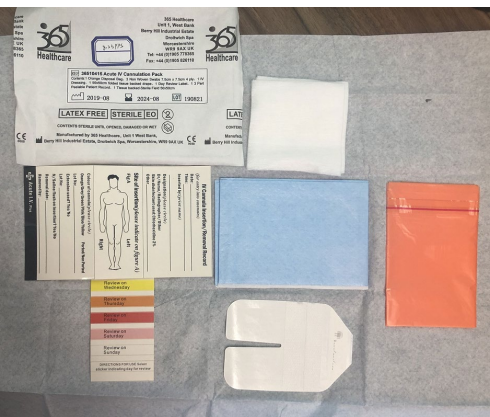


Product Procedure Specifications Sheet

Specification Sheet No	SSN-FT1390-007		
Product Name	IV Cannulation		
Product Description	36510415		
Range	365 Healthcare		
Manufacturer	Wujiang Kangxin Surgical Dressings Co. Ltd		
Material	100pcs/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
Orange waste bag	2003	1	45 x 29cm	High-density polyethylene + Adhesive
White non-woven swab	3102	3	7.7 x 7.5cm	Viscose & Terylene
IV Dressing	5201	1	7 x 8.5cm	PU Film Glassine Backing Paper Non-woven Acrylic Glue
Blue tissue backed drape folder 24 ply	8702	1	50 x 50cm	PE Film and absorbent tissue
1 day review label	2837	1	N/A	Paper
Patient record	2842	1	N/A	Paper
Blue Tissue backed Drape	8802	1	50 x 50cm	Pulp Tissue + PE Film Lamination

Quality Certification	ISO 13485		
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		

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



	use if package damaged, Keep dry and away from sunlight, CE, ECREP, Manufacturer, Distributor address
Additional information	Products are Latex and phthalate free Products contain no hazardous substances Products contain no medicinal substances
<i>Medical Device and Reporting System: Incidents caused by this device must be reported immediately to our offices.</i>	

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

365 Healthcare	Mediq House
T: +44 (0) 1332 413 700	Arundel Avenue, Castle Donington
W: https://mediq.co.uk/	Derby, DE74 2HJ

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Registered in England No. 62537

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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DMS 04-004-03	Procedure Pack Specification Sheet	Jack Wong	Racheal Begg	R02	03 rd April 2024