PRODUCT TECHNICAL DATASHEET

ITEM REF:	1042661061022	REVISION:	05

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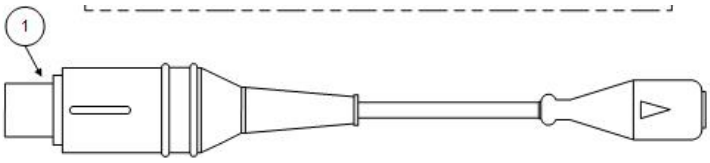
LEGAL MANUFACTURER	**ICU Medical, Inc.** 951 Calle Amanecer, San Clemente, CA 92673, USA						
IMPORTER	**ICU Medical B.V.** Hofspoor 3, 3994 VZ Houten, The Netherlands						
EU AUTHORISED REPRESENTATIVE (EUAR)	**ICU Medical B.V. Hofspoor 3, 3994 VZ Houten, The Netherlands**						
ASSEMBLY SITE	**ICU Medical (Utah), Inc.,** 4455 Atherton Drive, Salt Lake City, UT 84123						
CLASSIFICATION CODE	**GMDN** Code: 47487						
INTENDED USE	To supply power to the Transpac™ blood pressure transducer and to transmit signals from the transducer to the IBP monitor.						
ITEM DESCRIPTION	Transpac® IV 15' Cable for Use with Disposable Transducer Reusable: Do Not Discard						
PRIMING VOLUME (ml)	0	**LENGTH (cm)**	457.2	**WEIGHT (g)**	98.2	**CASE QTY**	1



LIST OF COMPONENTS *(Latex and DEHP Free)*	1	TRANSPAC IV CABLE	POLYPROPYLENE POLYESTER

MATERIAL COMPLIANCE AND BIOCOMPATIBILITY	Product has met the requirements of ISO 10993-1 and related standards for biocompatibility.
PRECAUTIONS	Use aseptic techniques. Single-use only – Do not resterilize. Sterile, Non-Pyrogenic fluid pathway in unopened, undamaged package.
LABELS AND DIRECTIONS FOR USE	Label and/or DFU contain information for proper use including any warnings or contraindications. Labels are applied on the individual package and on the outside of the cardboard box. Each sales package contains a Direction for Use where applicable.
PACKING AND PACKAGING	The non-unit packaging is biodegradable as defined by Directive 94/62/EC. Packaging met the requirements of BS EN ISO 11607. The packaging for this product is Latex Free. This item is blister packed or pouched individually using medical grade materials.
TRACEABILITY	Lot number provides full traceability of all components and manufacturing processes.
STORAGE	Store in a dry and clean place. Product should be retained in packaging until ready for use.
DISPOSAL	The user must dispose of the device according to hospital disposal policy.

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PRODUCTION AND ENVIRONMENT CONTROLS	• Product is manufactured in an environmentally controlled room. Floor, surfaces, and environment are monitored according to quality procedures. • Each component is inspected during acceptance (example: visual; dimensional; functional) to verify conformity with the requirements according to quality procedures. • Production and release specific tests are performed according to quality procedures. • Limulus amebocyte lysate (LAL) testing is performed on production samples to verify conformity of the endotoxin charge according to quality procedures. • Bioburden tests are performed on production samples to verify conformity of the microbial charge according to quality procedures.	
QUALITY SYSTEM AND PRODUCT CERTIFICATION	**Quality System complies to:**	ISO 13485:2016
	Product Certification:	The product is manufactured in compliance to Council Directive MDD 93/42/EEC as amended.
	CE Certificate Number:	N/A
	Notified Body:	NSAI National Standards Authority of Ireland.
	MDD Device Classification:	Class I Sterile