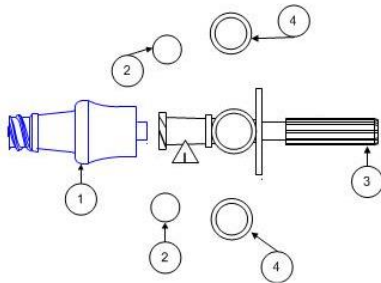


PRODUCT TECHNICAL DATASHEET

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human connections

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)	Medical Device Safety Service GmbH (MDSS), Schiffgraben 41, 30175 Hannover, Germany						
ASSEMBLY SITE	ICU Medical de Mexico, S.A. de C., Avenida Cuarza No. 250, Colonia Rancho Santa Clara, Delegacion Manaedero Ensenada, Baja California, Mexico 22790						
CLASSIFICATION CODE	GMDN Code: 60539						
INTENDED USE	The ChemoClave is a needle - free CSTD that mechanically prohibits the transfer of environmental contaminants, including bacterial and airborne contaminants into the system, and the escape of drug or vapor concentrations outside the system during drug preparation and administration, thereby minimizing exposure of individuals, healthcare personnel and the environment to hazardous drugs.						
ITEM DESCRIPTION	Universal Vented Vial Spike, CLAVE®						
PRIMING VOLUME (ml)	0.108	LENGTH (cm)	8.4	WEIGHT (g)	4.7	CASE QTY	50
							
ITEM SPECIFIC DATA	MRI Compatibility			No metal components			
	Chemical Compatibility			Lipids & Common Chemotherapeutics			
	Luer Compatibility			ISO 80369-7 Compliant. Clave compatible with male luers > 1,55mm Internal diameter			
	Sterilization and Shelf Life			Radiation; 5-Year Expiration			
CLAVE PERFORMANCE DATA	Backpressure Rating			60 psig / 3103 mmHG (Unactivated)			
	Microbial Ingress and Disinfection Compatibility			Microbial barrier for seven days utilizing a 70% IPA disinfection			
	Extended Use			600 repeat activations			

PRODUCT TECHNICAL DATASHEET

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LIST OF COMPONENTS	1	CLAVE	SILICONE SILICONE LUBRICANT ACRYLIC POLYESTER (PBT) (NON-FLUID PATH)
	2	.2UM FILTER MEMBRANE	PTFE (NON-FLUID PATH)
	3	NON LOCKING VIAL ADAPTER W/CAP	POLYCARBONATE SILICONE LUBRICANT POLYPROPYLENE (NON-FLUID PATH)
	4	VIAL ADAPTOR, ULTRASONIC RING	ABS (NON-FLUID PATH)
MATERIAL COMPLIANCE AND BIOCOMPATIBILITY	This product is made of non-latex, and non-DEHP components. Product has met the requirements of ISO 10993-1 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 is according to 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.		
PRODUCTION AND ENVIRONMENT CONTROLS	<ul style="list-style-type: none"> 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:	252.1002	
	Notified Body:	NSAI National Standards Authority of Ireland.	
	MDD Device Classification:	Class I Sterile	