Transpak®

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CE

These instructions pertain to Class I devices

IFU-LCN SYM8600-001 Rev A

ENGLISH

Transpak®

INSTRUCTIONS FOR USE

Intended Use

The TransPak® case is intended for use as a soiled surgical instrument point of use and transport vessel.

Indications for Use

None.

Product Description

The TransPak® case is designed to reduce leakage, minimize potential for injury and exposure to potentially infectious materials or fluids and help prevent damage to soiled surgical instruments during transport.

⚠ WARNINGS

NOT INTENDED FOR USE IN AUTOCLAVES. KEEP AWAY FROM HOT SURFACES.

DIRECTIONS FOR USE

1. Wear appropriate Personal Protective Equipment (PPE), as required by facility policies, when handling soiled instruments.
2. Place soiled instruments inside TransPak®.
3. Keep instruments moist until they are cleaned. A towel moistened with water placed over the instruments may be used. Instruments that cannot be cleaned immediately may be treated with an instrument cleaner according to the device and the instrument cleaner manufacturers' written Instructions For Use (IFU). Liquids used to soak contaminated items at the point of use should be discarded before transport.
4. Close and latch the lid to prevent spillage, contamination or exposure to potential hazards.
5. Place TransPak® on the designated cart for transport to the instrument decontamination area.
6. Use caution during transport to minimize spilling and other potential hazards.
7. Transpak may be cleaned manually or in washer/decontaminator cycle suitable for processing plastics.

CLEANING

TransPak® cases are compatible with enzymatic and nonenzymatic detergents and the below list of FDA cleared liquid chemical germicides (LCG) for manual reprocessing medical devices. Always read and follow both the medical device manufacturer and LCG manufacturer's reprocessing instructions.

Product Name	Manufacturer
Glutaraldehyde solutions	
Aldahol III	Healthpoint LTD
Banicide Advanced	Pascal Company, Inc.
Cetylcide-G	Cetylite Industries, Inc.
Cidex Activated Dialdehyde	ASP/J&J
Cidex Formula 7	ASP/J&J
Cidex Plus	ASP/J&J
MedSci 3% Glutaraldehyde	MedSci, Inc.
Metricide	Metrex Research, Inc.
Metricide Plus 30 Long-Life	Metrex Research, Inc.
Metricide 28 Long-Life	Metrex Research, Inc.
Omnicide Long Life	Cottrell Limited
Omicide Plus	Cottrell Limited
Procide 14	Cottrell Limited
Sporicidin	Sporicidin International
Wavicide	Wave Energy Systems
Ortho-phthaldehyde solutions	
Cidex OPA Solution	ASP/J&J
Opaciden Solution	Ciden Technologies, LLC
Hydrogen Peroxide Solutions	
Acecide	Minntech Corp
EndoSpor Plus	Cottrell Limited
Peract 20	Minntech Corp
Resert XL	Steris Corporation
Sporox	Reckitt & Colman

WARRANTY

Symmetry Surgical warrants that this medical device is free from defects in both materials and workmanship for one (1) year from the date of purchase. **Any other express or implied warranties, including warranties of merchantability or fitness for a particular purpose, are hereby disclaimed. Suitability for use of this medical device for any particular surgical procedure should be determined by the user in conformance with the manufacturer's instructions for use. There are no warranties that extend beyond the description on the face hereof. Abuse or misuse of the product or failure to comply with the instructions for use shall void this warranty.**

Copies of the IFU are available by requesting them from Symmetry Surgical Customer Service at:
Phone: 1-800-251-3000
Email: customerservice@symmetrysurgical.com
Website: www.symmetrysurgical.com

LABELING SYMBOLS

 Symbol for Manufacturer; the name and address of the Manufacturer is next to this symbol

 Caution-Consult accompanying documents

 Lot number or batch number

 Catalog number

 Consult instructions for use

 European Representative

 **In USA contact**

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