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WELLS JOHNSON

Sorenson Canister and Sorenson Liner 2000 mL

REF: 20-5157-00 – Sorenson Canister

REF: 20-5158-00 – Sorenson Liner

INSTRUCTIONS FOR USE

CAUTION: *Federal law restricts this device to sale by or on the order of a physician.*

IFU-00834 v2
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Intended Use

The Sorenson Canister with Sorenson Liner is intended for use during liposuction procedures with an external vacuum source to collect and temporarily contain adipose tissue.

Intended Users

The Sorenson Canister with Sorenson Liner is intended to be used in healthcare settings by qualified and trained healthcare professionals, such as licensed surgeons and operating room personnel.

Use Environment

The Sorenson Canister with Sorenson Liner is intended for use in healthcare settings, including operating rooms and other clinical environments where liposuction procedures are performed.

WARNING: *Sorenson Liner is non-sterile and for single use only. Do not reuse. The hard-shell outer Sorenson Canister is reusable.*

CAUTION: *Do not use the product after the expiration date.*

WARNING: *This product is not designed to be kept under vacuum when not in use. Prolonged static vacuum may increase the risk of cracking or implosion.*

Prior to Use

Before each use, inspect the canister, and liner for damage, including cracks or other defects that may have occurred during shipping, storage, or handling. Do not use if any damage is identified.

Sorenson Canister 2000 mL System Setup and Operation

1. Place the reusable canister in the canister holder on the top panel of the Hercules or Whisperator Aspirator.
2. Insert the Sorenson Liner into the canister. Make sure the liner lid properly seals the canister, and the canister Tee connector is securely tightened.
3. Connect the tubing attached to the liner lid to one end of the canister Tee connector.
4. Connect the 12-inch silicone tubing from the overflow trap to the other Tee connector.
5. Connect the sterile aspiration tubing to the port on the liner lid labeled **PATIENT**.

WARNING: *Aspiration tubing is sterile and for single use only. Do not reuse. Reuse may result in cross-contamination.*

CAUTION: *Do not fill the canister beyond 1900 mL.*

CAUTION: *Prevent aspirated material from entering the overflow trap.*

Overflow Condition:

If aspirated material enters the overflow trap, perform the following steps:

1. Turn the Hercules or Whisperator Aspirator OFF immediately.

2. Check liner positioning.
3. Check all tubing connections.
4. Check tubing for kinks or obstruction.
5. Check all tubing and accessory connections to ensure the system is correctly assembled.
6. Inspect the canister for cracks or other damage.
7. Confirm the vacuum pathway is correctly assembled.
8. Restart only after the cause has been corrected.



Disposal

The liner is for single use only. Do not reuse. Dispose of the liner and aspirated material in accordance with facility protocols.

Cleaning and Disinfection

Clean and disinfect the reusable outer canister after each use. Recommended cleaning agents include warm water with soap, ethanol, isopropanol, or bleach. Follow the cleaning agent manufacturer's instructions for use, including recommended concentration, contact time, and rinsing requirements.

CAUTION: *Do not use strong alkaline, acetone-based, or phenol-based cleaners, as they may cause chemical degradation of the material and result in product failure.*

Part Numbers for Sorenson Canister 2000 mL System

Item	Part Number
Sorenson Canister Holder	20-5159-00
Sorenson Canister Outer Reusable 2000 mL	20-5157-00
Sorenson Liner Single-Use, Non-Sterile 2000 mL (case of 10)	20-5158-00
MB Ultra-Limp Aspiration Tubing, 9 ft, Sterile, Single-Use (case of 10)	24-5104-00

Symbols Used

Symbol	Title	Description	Standard
	Use-by date	Indicates the date after which the medical device is not to be used	ISO 15223-1:2021, 5.1.4
	Catalog number	Indicates the manufacturer's catalog number so that the medical device can be identified	ISO 15223-1:2021, 5.1.6
	Batch Code	Indicates the manufacturer's batch code so that the batch or lot can be identified	ISO 15223-1:2021, 5.1.5
	Consult instructions for use or consult electronic instructions for use	Indicates the need for the user to consult the instructions for use	ISO 15223-1:2021, 5.4.3
	Prescription Only	Federal law restricts this device to sale by or on the order of a physician	FDA-recognized prescription-device designation under 21 CFR 801.109
	Unique Device Identifier	Indicates a carrier that contains unique device identifier information	ISO 15223-1:2021, 5.7.10
	Do not re-use	Indicates a medical device that is intended for one single use only	ISO 15223-1:2021, 5.4.2
	Non-sterile	Indicates a medical device that has not been subjected to a sterilization process	ISO 15223-1:2021, 5.2.7
	Do not use if package is damaged and consult instructions for use	Indicates that a medical device that should not be used if the package has been damaged or opened and that the user should consult the instructions for use for additional information	ISO 15223-1:2021, 5.2.8

Customer Service and Technical Support

Report any product malfunction, adverse event, or complaint to Wells Johnson Customer Service or Technical Support:

Phone: +1 520-298-6069 or +1 800-528-1597

Email: customerservice@wellsgrp.com



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