

WELLS JOHNSON

Sterile Tubing

Aspiration and Infiltration Tubing

INSTRUCTIONS FOR USE

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

IFU-01070 v2
30-Sep 2026

Device Description

Wells Johnson provides a range of sterile, single-use tubing sets intended for use with Wells Johnson systems during liposuction procedures. The tubing sets vary in length, diameter, and component configuration to meet the requirements of different applications.

Each tubing set is packaged within a sterile barrier system to maintain sterility until use. Inspect the packaging before use to verify that the sterile barrier has not been compromised. Do not use the product if the packaging is damaged or opened.

Tubing Types and Wells Johnson System Compatibility

The following table identifies the tubing types and the Wells Johnson systems compatible with each tubing configuration. Use only the tubing specified for the applicable Wells Johnson system. Refer to the applicable system Instructions for Use for connection, setup, and operation instructions.

REF	Description	Compatible Wells Johnson System
24-5103-00	Standard Ultra-Limp Aspiration Tubing	Hercules and Whisperator Aspirators
24-5104-00	MB Ultra-Limp Aspiration Tubing	
24-6008-00	CB Infiltration Tubing	Infusion System (WIS)
24-6009-00	“Y” Infiltration Tubing	
24-6028-00	Large Bore Infiltration Tubing	
24-6037-00	HVP Tubing	High Volume Precision (HVP) Auto-Graft System

CAUTION: *Carefully read these instructions before using Wells Johnson Sterile Tubing. Keep these instructions in a safe place for future reference.*

WARNING: *Improper use or handling of sterile tubing may result in serious injury, infection, or other adverse patient outcomes.*

Intended Use

Wells Johnson Sterile Tubing is intended for use during liposuction procedures to facilitate the infiltration of fluids, aspiration of adipose tissue, and the transfer of adipose tissue for reinjection into targeted areas of the body.

Intended Users

Wells Johnson Sterile Tubing is intended to be used in healthcare settings by qualified and trained healthcare professionals, such as licensed surgeons and operating room personnel.

Use Environment

Wells Johnson Sterile Tubing is intended for use in healthcare settings, including operating rooms and other clinical environments where liposuction procedures are performed.

Contraindications

No known contraindications.

Expiration Date

The expiration date is printed on the sterile tubing packaging.

CAUTION: *Do not use the product after the expiration date, as sterility and performance may no longer be assured.*

Prior to Use

Carefully inspect each sterile package and verify package integrity. Do not use the device if the package is damaged, opened, or the seal is broken. If the package is compromised, dispose of the device in accordance with facility procedures.

Wells Johnson Tubing is supplied sterile and is intended for single use only. Do not reuse this device.

WARNING: *Reuse of this device may result in compromised performance, loss of sterility, risk of infection, or patient harm.*

Storage and Transport Conditions

Store and transport devices in their original sterile packaging until use. Store at controlled room temperature. Protect from excessive heat, moisture, and direct sunlight.

Disposal

Single-use tubing may present a potential biohazard after use. Handle and dispose of the tubing in accordance with accepted medical practice, facility procedures, and applicable local, state, and federal regulations.

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

Symbol	Title	Description	Standard
R_x Only	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
UDI	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.
	Do not re-sterilize	Indicates a medical device that is not to be re-sterilized	ISO 15223-1:2021, 5.2.6.
STERILE R	Sterilized using irradiation	Indicates a medical device that has been sterilized using irradiation	ISO 15223-1:2021, 5.2.4.
	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



8000 S. Kolb Road.
Tucson, AZ 85756 USA
Ph: +1 520-298-6069
eFax: +1 520-885-1189
www.wellsgrp.com