

User Manual

Kit Contents

Multi-use Evacuation Tube



Instructions for use of SafetySuction

1. Following standard protocol, sterilize the SafetySuction instrument as any other stainless-steel instrument prior to use.

2. Attach Multi-use Evacuation Tube to SafetySuction instrument.



3. Attach Multi-use Evacuation Tube to HVE handle while the handle is positioned in its cradle.



4. When ready for use, have ultrasonic/piezo tip of choice inserted into handpiece.

5. The SafetySuction slides onto the ultrasonic/piezo handpiece easily by holding ultrasonic/piezo handpiece with one hand and sliding the SafetySuction onto the ultrasonic/ piezo handpiece with the other hand. Starting at the insert end of handpiece, slide the base end of the SafetySuction (where the hose is attached) over the top of the ultrasonic/piezo handle. While sliding the SafetySuction down the ultrasonic/piezo handle, the clasps on the underside of the SafetySuction will grab onto the ultrasonic/piezo handpiece.





6. The proprietary slide/clasp attachment allows positioning and maneuverability of the SafetySuction both forward and backward to expose the ultrasonic tip for complete access and aerosol containment.



Cleaning and Maintenance of SafetySuction and Multi-use Evacuation Tube

SafetySuction care

- 1. Post procedure, disconnect SafetySuction from Multi-use Evacuation Tube and ultrasonic/piezo handle.
- 2. Under running water, use cleaning brush provided to clean and remove any debris inside of SafetySuction.
- 3. SafetySuction may be placed into ultrasonic directly or in instrument cassette that is to be placed in ultrasonic.
- 4. SafetySuction is 100% autoclavable.

Multi-use Evacuation Tube Care

- 1. Post procedure, flush HVE line through Multi-use Evacuation Tube with Dental Vacuum Line cleaner according to manufacturer's directions.
- 2. Multi-use Evacuation Tube is **NOT** autoclavable.



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Cleaning and Sterilization

Device(s): Surgical Aspirator, Stainless Steel & Titanium

WADNINGS	
WARNINGS	It remains the responsibility of the end-user to ensure that the
	processing, as actually performed using equipment, materials and
	personnel in the processing facility, achieves the desired result.
Limitations on Reprocessing:	Instrument life is dependent upon a number of factors, such as method and
	duration of use, handling, care, and maintenance; therefore, end of
	functional life is determined not by a set number of uses, but by performing
	pre-operative inspection to verify continued functional performance and
	efficacy of the instrument.
INSTRUCTIONS	
Preparation for Cleaning:	Remove any packaging from the device.
Cleaning:	Equipment: Detergent, brush, running water
	Method:
	1. Rinse the devices with warm running water and using cleaning
	brush, scrub for 1 minute or until all visual soil is removed from
	the interior of the aspirator
	2. Prepare enzyme based cleaning solution according to
	manufacturer's concentrations with warm water
	3. Soak the instruments in the prepared solution for 15 minutes with
	sonication
	4. Rinse instruments thoroughly with clean water to remove
	detergent
	*Automated cleaning methods have not been validated and therefore should not be used unless validated by the end-user.
Drying:	Allow instruments to air dry or dry with a clean towel
	All instruments should be inspected to ensure proper function and condition.
Maintenance, Inspection and Testing:	Instruments with broken, cracked, chipped or worn parts should not be used.
	If any of these conditions occur, the instruments should be repaired or
	replaced immediately. For repairs, return to Quality Aspirators.
Packaging/Storage:	Standard packaging material may be used or instruments may be loaded into
r ackaging/storage:	general purpose sterilization trays. Once cleaned and sterilized,
	instruments should remain in sterilization packaging and stored in a clean,
	dry environment.
Sterilization:	Using a steam-type autoclave on pre-vacuum cycle,
	Option 1: Sterilize instruments for a minimum of 4 minutes at 134-136°C
	with a minimum dry time of 6 minutes.
	Option 2: Sterilize instruments for a minimum of 32 minutes at 121-123°C
	with a minimum dry time of 8 minutes.
	Either of these options will provide a 10 ⁻⁶ sterility assurance level
Manufacturer Contact:	Quality Aspirators
Manufacturer Contact.	
	Phone: 1-972-298-2669 Fax: 1-972-298-6592

Quality Aspirators utilizes 3rd party Biomedical Test facilities to Test and Validate cleaning and sterilization procedures according to national and international requirements. Quality Aspirators makes no changes or revisions to this procedure without referencing those changes back to the parent validation document from the biomedical test facility or conducting new validation testing.

The signature and date below indicate controlled validation of this Cleaning and Sterilization procedure.

Validation Date:

2/20/2020

Signature:

Country Gonzales Quality Manager

UI-300-10000-03 ECN 2001

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