Silicone Tubing

Not made with natural rubber latex

Item No.	Description
T2013	.125” ID X .250” OD X 50’
T2015	.188” ID X .313” OD X 50’
T2016	.188” ID X .375” OD X 50’
T2017	.250” ID X .375” OD X 50’
T2018	.250” ID X .438” OD X 50’
T2019	.250” ID X .500” OD X 50’
T2021	.313” ID X .500” OD X 50’
T2022	.375” ID X .500” OD X 50’

Sterilization of silicone tubing is recommended by the following means:

Autoclave – Our product can be sterilized using any of the following profiles:

Prevacuum: 4 minutes at 132°C

Gravity: 10 minutes at 132°C

Sterilization instructions should match the recommendations attached.

Note: Minimum purchases may apply

Cleaning and Sterilization Recommendations for Silicone Tubing

WARNINGS	Do not exceed 135° C.
Limitations on Reprocessing	Repeated processing has an effect on the tubing. End of life is normally determined by wear and damage due to use.
INSTRUCTIONS	
Point of Use:	For best results and to prolong the life of the tubing, reprocess immediately after use. Remove excess soil with disposable cloth/paper wipe.
Containment and Transportation:	No particular requirements. It is recommended that tubing is reprocessed as soon as is reasonably practical following use.
Cleaning: Manual	Mild detergent with neutral pH (7), brush and running water. Method: 1. Rinse excess soil from tubing. 2. Apply detergent solution to all surfaces. 3. Rinse under clean running water. Standard protocols defined by the institution should be used for cleaning and disinfection.
Disinfection:	Disinfectant solution may be used in accordance with label instructions.
Drying:	Tubing should be thoroughly dried with a clean, soft cloth prior to sterilization.
Maintenance:	Discard damaged tubing.
Inspection and Function Testing:	Visually inspect for damage and wear.
Packaging	N/A
Sterilization:	Vacuum autoclave, minimum of 4 minutes at 132° C. Gravity autoclave, minimum of 10 minutes at 132° C. Do not exceed 135° C.
Storage:	No particular requirements.
Manufacturer Contact:	See brochure for telephone and address of local representative or telephone 1-800-463-7146.

Per EN ISO 17664:2004 (E); the instructions above have been provided by the medical device manufacturer as being CAPABLE of preparing a medical device for re-use. It remains the responsibility of the processor to ensure that the processing as actually performed using equipment, materials and personnel in the processing facility achieve the desired result. This requires validation and routine monotoring of the process. Likewise any deviation by the processor from the instructions provided should be properly evaluated for effectiveness and potential adverse consequences.

Southmedic

Surgical
Dynamic Tissue Systems
Respiratory
Blades & Scalpels

Southmedic Inc.,
50 Alliance Blvd.,
Barrie, ON
Canada L4M 5K3

www.southmedic.ca
705-726-9383
1-800-463-7146 Toll free in North America
contactus@southmedic.com