

Saint-Gobain Life Sciences – Bioprocess Solutions



• Ultra-flexible bend radius

- Available in 25-foot standard lengths
- Sterilizable via autoclave or gamma
- Temperature range from -80°F (-62°C) to 500°F (260°C)
- Custom color-coding available
- Meets all USP Class VI, EP 3.1.9 FDA and ISO criteria
- High-pressure and vacuum rating

Single-use Applications

- Biopharmaceutical manufacturing
- Pharmaceutical processing and production
- Vacuum pump applications
- Bulk transfer
- Load cell
- Bioreactor process lines
- Production fermentation

Connections

- Sanitary, non-metallic fittings
- Radially crimped 316L SS or non-metallic fittings
- Adapter connections



Sani-Tech[®] STHT[®]-WR Wire-Reinforced Silicone Hose

Sani-Tech® STHT®-WR Wire-Reinforced Silicone Hose

Sani-Tech[®] STHT[®]-WR platinum-cured silicone, wire-reinforced hose, constructed of multi-ply reinforcement and 316L SS helical wire, provides unsurpassed flexibility while maintaining high-pressure ratings with full vacuum capabilities. Sani-Tech STHT-WR is manufactured with an ultra-pure, biopharmaceutical-grade silicone to ensure your process integrity.

Characteristics

The manufacturing process is carefully controlled from receiving through production. In all our production sites our tubing is produced and double bag packaged in an ISO 7 clean room. Inspection and lot traceability are readily accessible as batch numbers are assigned. All packages are identified by external labeling on both the bag and the highquality, crush-proof box.

Saint-Gobain Performance Plastics' manufacturing facility has the ability to create a variety of unique color-coding systems for your particular application needs. STHT*-WR available with custom coded coloring.

Biocompatibility

Sani-Tech[®] STHT[®]-WR is manufactured from the finest grade of silicone materials and is fully characterized, validated and tested to a variety of specifications including USP Class VI criteria, ISO 10993 and European Pharmacopoeia 3.1.9. Sani-Tech STHT-WR platinum-cured braid-reinforced silicone hose has a masterfile with the U.S. Food and Drug Administration. For additional compliance data, please review characterization information on the back page.

Sani-Tech® STHT®-WR Hose Inventory Sizes

				Maximum	Minimum	Minimum		
				Working	Burst	Bend		Vacuum
				Pressure	Pressure	Radius	Weight	Rating
	ID	OD	Wall Inches	PSI (bar)	PSI (bar)	Inches	Per Foot	IN. HG
Part Number	Inches (mm)	Inches (mm)	(mm)	at 68°F	at 68°F	(mm)	lb. (kg/m	(mmHg)
STHT-WR-0500	.500 (12.7)	.920 (23.4)	0.21 (5.3)	150 (10.3)	600 (41.4)	1.5 (38.1)	0.25 (0.4)	29.9 (760)
STHT-WR-0750	.750 (19.1)	1.190 (30.2)	0.22 (5.6)	125 (8.6)	500 (34.5)	2.5 (63.5)	0.36 (0.5)	29.9 (760)
STHT-WR-1000	1.000 (25.4)	1.450 (36.8)	0.225 (5.7)	125 (8.6)	500 (34.5)	4 (101.6)	0.48 (0.7)	29.9 (760)
STHT-WR-1500	1.500 (38.1)	2.160 (54.9)	0.33 (8.4)	125 (8.6)	500 (34.5)	5.5 (139.7)	1 (1.5)	29.9 (760)
STHT-WR-2000	2.000 (50.8)	2.690 (68.3)	0.345 (8.7)	100 (6.9)	400 (27.6)	7 (177.8)	1.25 (1.9)	29.9 (760)

Typical Physical Properties

Property	ASTM Method	Value or Rating
Durometer Hardness Shore A, 15 Sec	D2240	65
Tensile Strength psi (MPa)	D412	1291 (8.9)
Ultimate Elongation, 100%	D412	693
Tear Resistance Ib-f/inch (kN/m)	D624	329 (.037)
Specific Gravity	D792	1.19
Tensile Modulus @ 100% Elongation, psi (MPa)	D412	351 (2.42)

Unless otherwise noted, all tests were conducted at room temperature (73°F). Values shown were determined on 0.075" thick extruded strip or 0.075" thick molded ASTM plagues or molded ASTM durometer buttons.

Sterilization Methods

Autoclavable	
Radiation	Up to 2.5 Mrad (25 Kilogray)
Gas	ethylene oxide

NOTE: STHT® hose will not deteriorate with repeated autoclaving. This method of sterilization is strongly recommended. STHT silicones should not be considered for be continuous steam applications.

WARNING: Do not use STHT® silicone hose in hot oil or acid applications

Characterization

The bio-compatibility of STHT[®] platinum-cured silicone hose, manufactured with Sani-Tech[®] silicone, has been tested and complies with the parameters set forth in the following test protocols:

- USP XXIV (88) biological reactivity, in vivo
- Intracutaneous test, systemic injection test, implantation test
- USP XXIV (87) biological reactivity, in vitro
- L929 MEM elution, AGAR diffusion
- European Pharmacopoeia 3.1.9



Saint-Gobain Performance Plastics

3910 Terry Diane St. Beaverton, MI 48612 Tel: (888) 387-0067 Tel: (989) 435-9533 Fax: (989) 435-2355 IMPORTANT: It is the user's responsibility to ensure the suitability and safety of Saint-Gobain Performance Plastics products for all intended uses and that the materials to be used comply with all applicable medical regulatory requirements. Saint-Gobain Performance Plastics assumes no responsibility for any product failures that occur due to misuse of the materials it provides arising out of the design, fabrication or application of the products into which the materials are incorporated.

WARRANTY: For a period of 12 months from the date of first sale, Saint-Gobain Performance Plastics warrants this product to be free of defects in materials and workmanship. Our only obligation will be to replace any portion proving defective, or at our option, to refund the purchase price thereof.

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