

1600HFTLC Adult High Flow Nasal Cannula with Ear Wraps and Ribbed Connector

LATEX FREE | SINGLE PATIENT USE | DISPOSABLE

Salter-Style® 1600 Adult High Flow Nasal Cannula			
Reference Number	1600HFTLC-7-25	1600HFTLC-14-25	1600HFTLC-25-10
Manufacturer	SunMed LLC/Salter Labs		
FDA Classification	Class I Medical Device		
FDA Product Code	CAT – Nasal Oxygen Cannula		
Canadian MDR	Class II, Conformity Assessment Route: Medical Device License		
Classifications – EU	Class IIa, Rule 2, MDR 2017/745, Annex VIII		
CE Mark/Notified Body	CE2797 / BSI Group		
EMDN Code	R03010203, Air/Oxygen Nasal Cannula		
GMDN Code	35201 Nasal Cannulas		
UMDNS Code	12799, Cannulae, Nasal Oxygen		
Usage	Disposable, Single Patient Multiple Use		
Sterile	No		
Patient Population	Adult		
Packaging	Individually Packaged, 25/case (for 7' and 14') Individually Packaged, 10/case (for 25')		

Device Specifications	
Description	Specification
Oxygen Tubing Length	7' (2.1 m), 14' (4.3 m), 25' (7.6 m)
Oxygen Supply Tubing	Clear, 3-Channel Safety Tubing
Oxygen Flow Rate	0 LPM to 15 LPM
Tubing End Connector	Universal, Push-On, Ribbed
Operating Temperature	5°C to 40°C
Storage Temperature	-20°C to 50°C

Product Material	
Part Description	Material
Nasal Face Piece	Plastisol, Reddish Tint
Oxygen Tubing	Polyvinyl Chloride, 3-Channel
Headset Tubing	Polyvinyl Chloride, Smooth Bore
Tubing End Connector	Polyvinyl Chloride
Bolo	Low Density Polyethylene



Intended use: Over-the-ear style nasal cannula for delivery of supplemental oxygen to nares of a spontaneously breathing patient. Headset tubing include ear cushions to help relieve pressure above the ears.

Area of use: Hospitals, medical clinics, home, surgical centers, skilled nursing facilities.

Duration of use: Discard and replace nasal cannula every 14 days or sooner if the cannula becomes soiled or damaged.

Contraindications: No known.

Caution: Keep tubing straight and free of kinks. Do not sterilize.

Warning:

- If patient develops infection, skin irritation or material sensitization consult physician.
- Patient may become hypoxic if oxygen flow is interrupted.
- Position tubing to avoid strangulation or tripping hazard.
- Do not place anything on oxygen tubing that may obstruct flow.
- Use oxygen product as prescribed.
- When using oxygen, do use open flame or heat source.

Latex: Salter Labs® does not utilize latex or latex-containing material to manufacture products. To the best of our knowledge, latex does not come in contact with components or finished good during the manufacturing process.

Phthalates: The selected materials are Non-DEHP; DEHP was not intentionally added as part of the manufacturing process.

Mercury-Lead: The selected materials do not contain lead or mercury.

Biocompatible per device classification in ISO 10993. Salter Labs manufactures product from medical grade materials in compliance with Good Manufacturing Practices (GMPs) as listed in 21 C.F.R. (U.S. Code of Federal Regulations).

Part Number	UOM	GTIN/UDI
1600HFTLC-7	Each	607411005923
1600HFTLC-7-25	Case	20607411005927
1600HFTLC-14	Each	607411006111
1600HFTLC-14-25	Case	10607411006613
1600HFTLC-25	Each	607411016128
1600HFTLC-25-10	Case	10607411006125

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