Oxygen Tubing – Clear, 3-Channel, Ribbed Connectors

SINGLE PATIENT USE | DISPOSABLE

Device Manufacturing Specifications

Reference Number	2001, 2002, 2004, 2014, 2025, 2030, 2035, 2040, 2050
Manufacturer	SunMed LLC/Salter Labs
FDA Product Code	Class I Medical Device
Classification – EU	Class IIa, Rule 2
CE Mark/Notified Body	CE 2797
EU Authorized Representative	Mt Promedt Consulting GmbH
Classification – Canada	Class II
EMDN Code	R03010204 Oxygen Administration Tubing
GMDN Code	12875 Oxygen Administration, Tubing
UMDNS Code	12875 Tubing, Oxygen Connecting
Made In	Mexico
Usage	Disposable, Single Patient Use
Sterile	Nonsterile
Patient Population	Neonate, Infant, Pediatric and Adult
Packaging	Individually Packaged, 20/Case, 25/Case or 50/Case
Shelf Life	5 Years

Description: Clear, 3-Channel safety oxygen supply tubing with ribbed connectors used as a conduit to delivery oxygen from the source gas to the patient.

Intended Purpose: To extend the length of oxygen tubing as accessory to nasal cannulas, oxygen masks and other breathing products.

Area of Use: Hospitals, pre-hospital, home, surgical centers, skilled nursing facilities, medical clinics.

Device Specifications

Description	Specification
Operating Temperature	5^0 C to 40^0 C
Storage Temperature	-20^0 C to 50^0 C
Internal Diameter	3/16"
Flow Rate	≤ 15 LPM
Tubing	3-Channel
End Connector	Ribbed Universal Push-On
End Connector Size	Fits 4 mm to 6 mm
Available Tubing Lengths	1' (0.3 m), 4' (1.2 m), 7' (2.1 m), 14' (4.2 m), 25' (7.6 m), 35' (10.7 m), 50' (15.2 m)

Device Material

Component	Material
Tubing	Polyvinyl Chloride, Clear
End Piece	Polyvinyl Chloride

Part Number	UOM	GTIN
2001-1	Each	00607411200007
2001-1-50	Case	10607411200004
2002-7	Each	00607411200014
2002-7-50	Case	10607411200011
2004-4	Each	00607411200038
2004-4-50	Case	10607411200035
2014-14	Each	00607411200106
2014-14-50	Case	10607411200103
2025-25	Each	00607411200168
2025-20-25	Case	10607411200165
2030-30	Each	00607411200175
2035-30-20	Case	10607411200172
2050-35	Each	00607411200182
2050-35-20	Case	10607411200189
2035-40	Each	00607411200199
2035-40-20	Case	10607411200196
2050-50	Each	00607411200236
2050-50-20	Case	10607411200233

Latex: SunMed® does not utilize latex or latex-containing material to manufacture products. To the best of our knowledge latex does not contact components or finished goods 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.

SunMed 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).

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