



# INSTRUCTIONS FOR USE

## Endotracheal Tube Reinforced (Plain/Cuffed)

Oral/Nasal Tracheal Tube with Radio-Opaque Line

### Device Description:

A Reinforced Endotracheal Tube is a sterile, single-use airway management device incorporating a stainless-steel spiral reinforcement within a transparent PVC tube to resist kinking and maintain lumen patency during patient positioning or head/neck movement.

### Intended Use:

The Endotracheal Tube is intended for airway management and delivery of oxygen and anesthetic gases in patients requiring assisted or controlled ventilation.

### Materials Used:

Medical grade Polyvinyl Chloride (PVC), Polypropylene (PP), Stainless Steel (spiral reinforcement), Acrylonitrile Butadiene Styrene (ABS – connector), Silicone (cuff, if applicable), and approved color additives.

### Indications:

- Respiratory Failure
- Airway Obstruction
- Loss of Consciousness
- Aspiration Trauma.

### Contra Indication:

- Severe airway trauma
- Difficult airway anatomy
- Cervical spine injuries
- Severe laryngeal edema

### Instructions for use:

- Select the appropriate tube size based on patient anatomy.
- Carefully inspect package for integrity and expiry and then remove product from package.
- Peel & open the pouch.
- Fully deflate the cuff.(*cuffed only*)

- Lubricate with water soluble lubricant as required.
- Intubate the trachea then inflate cuff(*cuffed only*) using minimum amount of air required to provide an effective seal.
- Listen for air passing around the cuff to determine if an effective seal has been made.
- Check for correct intubation and ongoing airway patency adjusting as necessary.
- Secure the tube (preferably using a device which incorporates a bite block) and attach the tube to the ventilation equipment

### Warnings:

- The product should not be reprocessed.
- Improper transport and handling may cause structural and/or functional damage to device or packaging.
- Do not clean or resterilise.
- Do not expose to heat or direct sunlight.
- The product should be used immediately after opening the packaging.
- **MAIS INDIA disclaims any responsibility for possible consequences resulting from improper use.**

### Cautions:

- Do not use if package is damaged.
- Discard after single use.

**Target Age Group:** Applicable for neonatal, pediatric, and adult patients depending on selected tube size.

**Duration of Contact of Device:** Short term (Use up to 72 Hours for best performance).

**Storage conditions:** Store in Cool & Dry Place. Keep away from direct sunlight.

**Device Life:** 05 Years.



Phthalates Free



Keep Dry



Non-Pyrogenic



Catalogue Number



Caution



Latex Free



Lot No./ Batch No.



Do Not Use If Package is Damaged



Do Not Reuse



Manufacturer



Sterilized By Ethylene Oxide



Do Not Resterilize



Date of Manufacturing



Single Sterile Barrier System



Medical Device



Use By / Expiry Date



Unique Device Identifier



Country of manufacture



Consult Instructions for Use



Keep Away From Sunlight



Country of manufacture

### Manufactured by:

**Mais India Medical Devices Pvt. Ltd.**  
525-P, Sector-37, Pace City-II,  
Gurgaon, Haryana-122001 (INDIA)  
Mfg. Lic. No.:MFG/MD/2019/000037

**ISO & CE**  
Certified Company

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	Q.C						
	Pro.						
CHD BY	Mkt.			<b>COLOR OF PRINT:</b> Blue 293c		DRW. No.: A09-2/7-00-IFU-000	
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			SCALE-1:1	TOLERANCE : ±2 mm	<b>REVISION DETAILS:</b>		
<b>MAIS INDIA MEDICAL DEVICES PVT. LTD.</b> 525-P SECTOR-37, PACE CITY-II, GURGAON, HARYANA-122001 (INDIA)							