IFU for Naso-Flo® nasopharyngeal airway device

TYPE AND SIZE

	Type	Code	Size (mm)
Naso-Flo® nasopharyngeal airway device	with O2 Port	NF104	4.0
		NF105	5.0
		NF106	6.0
		NF107	7.0
		NF108	8.0
		NF109	9.0
	with O2 Port, Respiratory Indicator and Filter	NF204	4.0
		NF205	5.0
		NF206	6.0
		NF207	7.0
		NF208	8.0
		NF209	9.0
	with O2 Port and Respiratory Indicator - without Filter	NF304	4.0
		NF305	5.0
		NF306	6.0
		NF307	7.0
		NF308	8.0
		NF309	9.0

WARNING/PRECAUTIONS

1. Single Use Only
2. The product should remain unopened until the point of use
3. Sterile if package is unopened, undamaged and within shelf life date
4. Do not re-sterilise
5. Do not expose to temperatures above 49°C
6. This product must be in a pre-use condition and checked prior to use
7. Do not use if the patient is suffering from a blocked nose, nasal congestion or nasal polyps
8. Do not use if the patient has a history of epistaxis or fractured nasal bone

PRE-USE CHECKS:

Do not use this product unless these checks are fully satisfactory

1. Check product is within its specified sterile shelf life
2. Visually check whole device for completeness, discolouration, damage and flaws

3. Check that there is no blockage or occlusion in the airway

DIRECTION FOR USE

These directions are general guidelines intended for use by qualified medical personnel. Any Instructions and contraindications given are not exhaustive and it is the clinician's responsibility to ensure the safe, correct use of this product.

1. Select correct size of airway.
2. Lubricate outer airway tube with a suitable water based lubricant, ensuring that the distal tip remains patent.
3. Insert airway into patient's nasal cavity and ensure that the airway is operating correctly.
4. If oxygen is required, connect the oxygen tube onto the oxygen elbow before connecting the elbow into the airway.
5. To ensure a secure connection, use a push & twist action on all fittings.
6. If the respiratory indicator is required, connect the hydrophobic filter to a suitable male sampling line before connecting to an appropriate monitor.
7. When observing the respiratory rate, oxygen should only be administered at a flow rate of less than 8 LPM.
8. Any blocked or occluded devices should be removed immediately.

STORAGE CONDITIONS

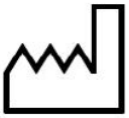
Store product inside containers or outer boxes in a clean, dry area.
Storage should be within a temperature range of 10–30^{o}C.
Do not expose to direct sunlight or UV light.

MEANING OF SYMBOLS ON PACKAGE

Do not resterilise

Manufacturer

Use-by date



Date of manufacture

Batch code

Authorised representative in the European Community

Sterilized by ethylene oxide

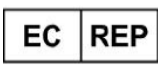
Keep away from sunlight

Do not use of package is damaged

Keep dry

Do not re-use

TIANJIN MEDIS MEDICAL DEVICE CO. LTD.
Add: 10-A Tianzhi Industrial Center, No.12 HongYuan Road, Xiqing Economic Development Area, 300385 Tianjin City, PEOPLE'S REPUBLIC OF CHINA
Tel: +86-22-83988488 Fax: 022—83988486
www.medis-medical.com



Shanghai International Holding Corp. GmbH (Europe)
Add: Eiffestrasse 80, 20537 Hamburg ,Germany
Tel: +49-40-2513175
Fax:+49-40-255726

Version 2016.08.17