

Product P/N	9500/710	Mod. 984A
Description	Micro tracheal	Rev. 06

9500/710

Micro tracheal



PRODUCT	Inlet/Outlet Connectors: 15mm female connector on patient side.		
DESCRIPTION	Bidirectional HME.		
	Approx. dimensions: 38.5mm x 28.5mm height.		
	Weight: 3.5gm (approx.).		
MANUFACTURER	GVS Filter Technology UK		
NAME	NFC House		
	Vickers Industrial Estate		
	Mellishaw Lane, Morecambe		
	Lancashire LA3 3EN - United Kingdom		
	Information		
	Tel. +44 (0) 1524 847600		
	e-mail: gvsuk@gvs.com		
INTENDED USE /	For use within Anaesthesia, Respiratory and Critical Care clinical areas. Indicated for use with		
APPLICATION	patients whose upper airways are being bypassed by an artificial tracheal airway or receiving		
	artificial ventilator support.		
CLASS OF THE	Disposable medical device - Class IIa		
PRODUCT	Rule 2 Annex IX 93/42 / EEC		
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	Rule 2 Annex VIII MDR 2017/745		
MATERIALS	Filter media: Corrugated Cellulose HME paper		
	Frame/Housing Polymer: Transparent Clear Polypropylene (PP)		
	Colour: Transparent Clear		
	Regulatory Documentation Required:		
	- Biocompatibility according ISO 10993-1		
	- ROHS		
	- BSE/TSE		
	- DEHP plasticizer Free and latex free		
	- Aging		
	- RĔAČH		
	- Conflict minerals		
PRODUCT	Appearance/Visual		
CHARACTERISTICS	As shown on drawing.		
	Discrete al/Manhaminal		
	Physical/Mechanical		
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	Interfaces (ex: Input / Output connectors): 15mm female connector on patient side. Operating temperature Range: N/A Storage temperature Range: 5 °C to 40 °C Bidirectional HME. Biological Pyrogenicity: <0.3 Eu/ml Biocompatibility to ISO10993 Category - Surface device Contact - Skin Contact Duration - <24hrs Functional Air Flow Rate: 30l/min, 60l/min, 90l/min. Pressure Drop: Flow Resistance @ 30L/min in accordance with EN ISO 9360-1: Max. 25Pa Flow Resistance @ 60L/min in accordance with EN ISO 9360-1: Max. 73Pa Flow Resistance @ 90L/min in accordance with EN ISO 9360-1: Max. 133Pa (REP: 2083/20 with 10% of safety margin added to Max.) Internal Volume: 13ml (approx.) Operating Lifetime: Refer to Instructions for Use. Shelf Lifetime: 5 years from the date of manufacture. Moisture Loss @ 500ml Tidal Volume in accordance with EN ISO 9360-1: Max. 19.0mg/l (REP: 2084/20 with +0.5ml factor of safety) Moisture Output @ 500ml Tidal Volume in accordance with EN ISO 9360-1: Min. 21.4mg/l (REP: 2084/20 with -0.5ml factor of safety) Gas leakage in accordance with EN9360: Max. 0.0 ml/min (REP:1162/17)	
INSTRUCTIONS /	Cleanliness Device assembled within Class 8 Cleanroom. Multi-language IFU available.	
WARNINGS		
PRODUCT SHELF LIFE	5 years from the date of manufacture. Expiration date and date of manufacture are detailed on the product labelling.	
STERILIZATION	Sterile version of product available (Ethylene oxide - Max 55°C)	
APPLICABLE STANDARDS AND REGULATIONS	Product Certification required: - CE mark - FDA	
	Applicable Standards and Technical Regulations: Biological evaluation of Medical Devices - Part 1: Evaluation and Testing - ISO 10993-1.	



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	Medical devices- Application of risk management to medical devices - BS EN ISO 14971.	
	Medical devices – symbols to be used with medical device labels, labelling and information to be supplied - Part1: General requirements - ISO 15223-1.	
	Anaesthetic and respiratory equipment – conical connectors – part 1: Cones and sockets – ISO 5356-1.	
	Sterilization of health care products – Ethylene oxide sterilization – ISO 11135-1.	
	Sterilization of medical devices – Microbiological Methods – Part 1: Estimation of population of Microorganisms on products – ISO 11737-1.	
PACKAGING AND LABELING	Number of pcs per bag is determined by the sales order. The first barcode label is applied to the outside of the bags. The second barcode label is applied onto the outside of the box. Each bag is labelled with the following traceability information: Vauntity Product description Product Date Lot Number (OL and 5-digit batch number to trace back to raw materials used) Operator Code Different lots in one box are separately closed and separately labelled. Bulk products will be packed in double PE bags.	
CERTIFICATE OF COMPLIANCE	With each shipment, GVS UK Customer Service will send the CofC to the Customer, based on the lot numbers and date of manufacture. Conformity declaration is printed on every invoice and Certificate is according to UNI EN 10204 type 2.1. The Quality management system is in compliance with ISO 13485.	
DRAWING	The attached drawing is part of this product specification and must not be duplicated or made accessible to a third party without written permission from GVS Filter Technology UK Ltd. Approximate dimensions for reference only	
	38.5mm	



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ACCEPTABLE	AQL:	0.65 with sampling Plan: ISO2859.		
QUALITY LEVEL				
VISUAL REQUIREMENTS	Magr Light Timir	al acceptance requirements apply when inspected und ification: Unaided eye at a distance of approximately 35-4 type: Lighting level must be reasonable for visual detection gs: Maximum inspection period per item is 25 seconds. letailed defect list, refer to product control plan.	0cm.	nditions:
		Acceptance Requirement	AQL	Sampling Plan
	:	Black particle contamination	0.65	
		Damaged/broken item	0.65	-
	3	Blocked connector/luer	0.65	
	4	Weld marks	0.65	
	į	Short fill moulding	0.65	
		Rough surface or edges	0.65	ISO 2859 Part 1 General Inspection
	7	Pronounced injection gate	0.65	Level 1
	8	Deformation/distortion	0.65	
	9	Crack	0.65	
	1	O Oil/grease	0.65	
	1	1 Wrong colour	0.65	
		2 Weld fault	0.65	
GENERAL SAFETY AND PERFORMANCE REQUIREMENTS	regul	ial characteristic: Product characteristic which can affect ations, fit, function, performance or subsequent processing ial Characteristic # 01:	•	mpliance with
NEWOINEIVIEW 13	Flow	Resistance @ 30L/min in accordance with EN ISO 9360-1	,	
		Resistance @ 60L/min in accordance with EN ISO 9360-1	•	
	Spec	Resistance @ 90L/min in accordance with EN ISO 9360-1 ial Characteristic # 02: Moisture Output @ 500ml Tidal \ 0360-1		cordance with EN
		ture Loss @ 500ml Tidal Volume in accordance with EN IS	20.0000.4	



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Special Characteristic # 03: Conical connectors compliant in accordance with EN5356.

Special Characteristic # 04: Gas Leakage compliant in accordance with EN9360.

This material specification describes the properties of product above indicated. This document contains general requirements, material description, drawing references, defect specification, biological material requirements.

REVISIONS AND APPROVALS:

DATE R	REV.	REASON FOR	ISSUED AND CONTROLLED	APPROVED BY:
		CHANGE	BY:	(NAME/FUNCTION/SIGNATURE)
			(NAME/FUNCTION/SIGNATURE)	
30/07/2021	3	Functional characteristics updated.	Kinga Gawdzik – Engineering Support Technician	Andrew Pearce – Quality Manager

CUSTOMER APPROVAL:

We accept this	material specification as a part of the agreed terms of delivery.
Company Nam	ne:
Approved by:	
	NAME/FUNCTION
	SIGNATURE
	DATE
	COMPANY STAMP

 $Please send \ back \ this \ document \ signed \ for \ approval. \ If \ we \ will \ not \ receive \ this \ specification \ signed, \ we \ consider \ the \ first \ order \ placed \ as \ implicit \ approval.$