

# PRODUCT SPECIFICATION

Product P/N	4444/06	<b>Mod. 984A Rev. 06</b>
Description	<b>Slimline Electrostatic with Integral Mouthpiece</b>	

**4444/06**  
**Slimline**  
**Electrostatic**  
**with Integral**  
**Mouthpiece**



<b>PRODUCT DESCRIPTION</b>	Inlet/Outlet Connectors: 22mm Female/15mm Male – Mouthpiece. Approx. dimensions: 68.6mm diameter x 93mm height. Weight: 26g (approx.). Bidirectional Filter.
<b>MANUFACTURER NAME</b>	<b>GVS Filter Technology UK</b> NFC House Vickers Industrial Estate Mellishaw Lane, Morecambe Lancashire LA3 3EN - United Kingdom  <b>Information</b> Tel. +44 (0) 1524 847600 e-mail: gvsuk@gvs.com
<b>INTENDED USE / APPLICATION</b>	Filters protect the patient's airways effectively from exogenous microbial loads, thus reducing the risk of extrinsic colonisation and infection. Used to help reduce cross contamination between patient and machine.
<b>CLASS OF THE PRODUCT</b>	Disposable medical device - Class IIa  Rule 2 Annex IX 93/42 / EEC  Rule 5 Annex VIII MDR 2017/745
<b>MATERIALS</b>	<b>Filter media: <i>Electrostatic Blended Synthetic Fiber</i></b> <b>Frame/Housing Polymer: <i>Transparent Clear Polypropylene (PP)</i></b> <b>Colour: <i>Transparent Clear</i></b>  <b>Regulatory Documentation Required:</b> <ul style="list-style-type: none"> <li>- Biocompatibility according ISO 10993-1</li> <li>- ROHS</li> <li>- BSE/TSE</li> <li>- DEHP plasticizer Free and latex free</li> <li>- Aging</li> <li>- REACH</li> <li>- Conflict minerals</li> </ul>
<b>PRODUCT CHARACTERISTICS</b>	<b>Appearance/Visual</b> As shown on drawing.

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## Physical/Mechanical

*Approx. dimensions: 68.6mm diameter x 93mm height.*

*Weight: 26g (approx.).*

Interfaces (ex: Input / Output connectors): **22mm Female/15mm Male – Mouthpiece.**

Operating temperature Range: **N/A**

Storage temperature Range: **5 °C to 40 °C**

**Bidirectional Filter.**

## Biological

Pyrogenicity: **<0.3 EU/ml**

**Biocompatibility to ISO10993**

Category – **Surface device**

Contact – **Oral cavity**

Contact Duration - **<24hrs**

## Functional

Air Flow Rate: **30l/min, 60l/min, 90l/min.**

Filtration Efficiency: *Filter Efficiency @ 30L/min using TSI 8130: **Min. 97.5%***  
(REP: 1687/19 with factor of safety applied to Min.)

Pressure Drop:

*Flow Resistance @ 30l/min in accordance with EN ISO 9360-1: **Max. 118Pa***

*Flow Resistance @ 60l/min in accordance with EN ISO 9360-1: **Max. 264Pa***

*Flow Resistance @ 90l/min in accordance with EN ISO 9360-1: **Max. 441Pa***

(REP:1688/19 with 10% of safety margin added.)

Internal Volume: **48ml (approx.)**

Operating Lifetime: **Refer to Instructions for Use.**

Shelf Lifetime: **5 years from the date of manufacture.**

*Bacterial Filtration Efficiency in accordance with ASTM F2101-07: **Min. 99.999%***  
(*Staphylococcus aureus @ 30L /minute*) REP: EXT439447.

*Viral Filtration Efficiency in accordance with ASTM F2101-07: **Min. 99.9 %***  
(*Bacteriophage @ 30L/ minute*) REP: EXT439446.

*Gas leakage in accordance with EN9360: **Max. 0.0 ml/min*** (REP: 1689/19)

## Cleanliness

Device assembled within Class 8 Cleanroom.

## Testing

**Leak test at 3PSI.**

## INSTRUCTIONS / WARNINGS

*Multi-language IFU available.*

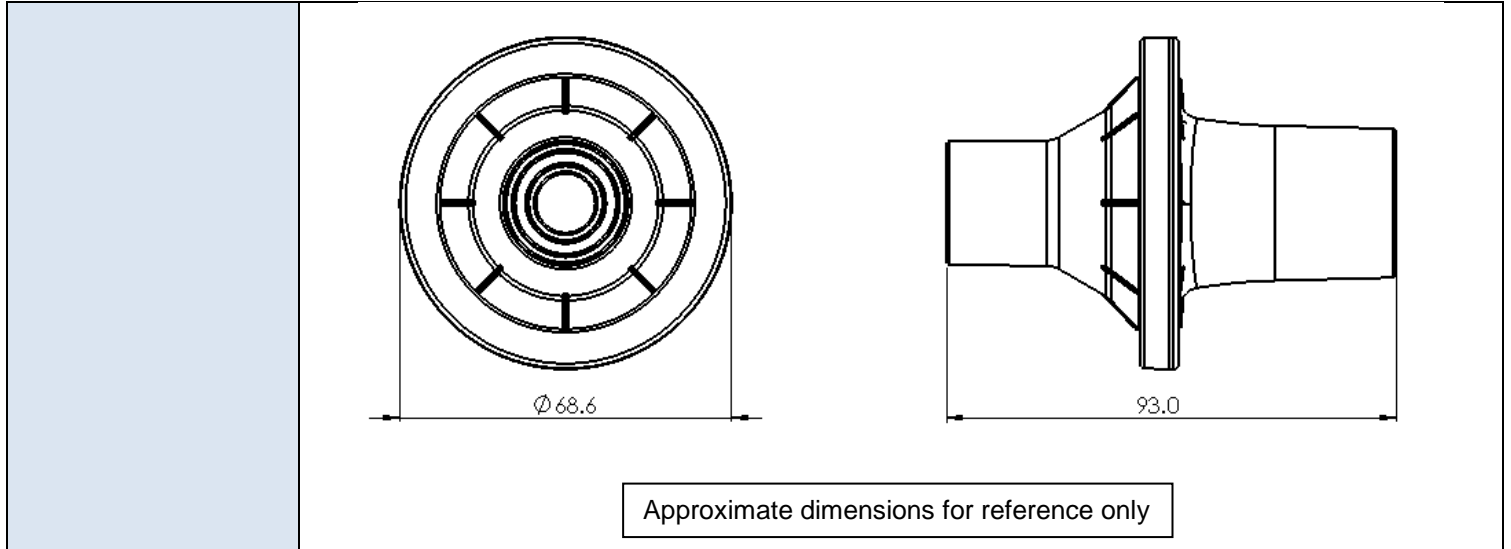
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<b>PRODUCT SHELF LIFE</b>	<p>5 years from the date of manufacture.</p> <p><i>Expiration date and date of manufacture are detailed on the product labelling.</i></p>
<b>APPLICABLE STANDARDS AND REGULATIONS</b>	<p><b>Product Certification required:</b></p> <ul style="list-style-type: none"> <li>- CE mark</li> <li>- FDA</li> </ul> <p><b>Applicable Standards and Technical Regulations:</b></p> <p><i>Biological evaluation of Medical Devices - Part 1: Evaluation and Testing - ISO 10993-1.</i></p> <p><i>Respiratory protective devices - Method for test - Part 7: Determination of particle filter penetration - BS EN 13274-7.</i></p> <p><i>Medical devices- Application of risk management to medical devices - BS EN ISO 14971.</i></p> <p><i>Medical devices – symbols to be used with medical device labels, labelling and information to be supplied - Part1: General requirements - ISO 15223-1.</i></p> <p><i>Breathing system filters for anaesthetic and respiratory use — Part 1: Salt test method to assess filtration performance - ISO 23328-1.</i></p> <p><i>Anaesthetic and respiratory equipment – conical connectors – part 1: Cones and sockets – ISO 5356-1.</i></p>
<b>PACKAGING AND LABELING</b>	<p><i>Number of pcs per bag is determined by the sales order.</i></p> <p><i>The first barcode label is applied to the outside of the bags.</i></p> <p><i>The second barcode label is applied onto the outside of the box.</i></p> <p><i>Each bag is labelled with the following traceability information:</i></p> <ul style="list-style-type: none"> <li>✓ Quantity</li> <li>✓ Product description</li> <li>✓ Product Date</li> <li>✓ Lot Number (OL and 5-digit batch number to trace back to raw materials used)</li> <li>✓ Operator Code</li> </ul> <p><i>Different lots in one box are separately closed and separately labelled.</i></p> <p><i>Bulk products will be packed in double PE bags.</i></p>
<b>CERTIFICATE OF COMPLIANCE</b>	<p><i>With each shipment, GVS UK Customer Service will send the CofC to the Customer, based on the lot numbers and date of manufacture.</i></p> <p><i>Conformity declaration is printed on every invoice and Certificate is according to UNI EN 10204 type 2.1.</i></p> <p><i>The Quality management system is in compliance with ISO 9001, ISO 13485.</i></p>
<b>DRAWING</b>	<p><i>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.</i></p>

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**ACCEPTABLE QUALITY LEVEL** AQL: 0.65 with sampling Plan: ISO2859.

**VISUAL REQUIREMENTS**

**Visual acceptance requirements apply when inspected under below conditions:**

Magnification: *Unaided eye at a distance of approximately 35-40cm.*  
 Light type: *Lighting level must be reasonable for visual detection.*  
 Timings: *Maximum inspection period per item is 25 seconds.*  
*For detailed defect list, refer to product control plan.*

Acceptance Requirement	AQL	Sampling Plan
1 Black particle contamination	0.65	ISO 2859 Part 1 General Inspection Level 1
2 Damaged/broken item	0.65	
3 Blocked connector/luer	0.65	
4 Weld marks	0.65	
5 Short fill moulding	0.65	
6 Rough surface or edges	0.65	
7 Pronounced injection gate	0.65	
8 Deformation/distortion	0.65	
9 Crack	0.65	
10 Oil/grease	0.65	

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	11	Wrong colour	0.65		
	12	Weld fault	0.65		
<b>GENERAL SAFETY AND PERFORMANCE REQUIREMENTS</b>	<p><b>Special characteristic:</b> <i>Product characteristic which can affect safety or compliance with regulations, fit, function, performance or subsequent processing of product.</i></p> <p><b>Special Characteristic # 01:</b></p> <p><i>Flow Resistance @ 30L/min in accordance with EN ISO 9360-1</i></p> <p><i>Flow Resistance @ 60L/min in accordance with EN ISO 9360-1</i></p> <p><i>Flow Resistance @ 90L/min in accordance with EN ISO 9360-1</i></p> <p><b>Special Characteristic # 02:</b> <i>Filter Efficiency @ 30L/min using TSI 8130 in accordance with EN 13274-7.</i></p> <p><b>Special Characteristic # 03:</b> <i>Bacterial Filtration Efficiency in accordance with ASTM F2101-07</i></p> <p><i>Viral Filtration Efficiency in accordance with ASTM F2101-07</i></p> <p><b>Special Characteristic # 04:</b> <i>Conical connectors compliant in accordance with EN5356</i></p>				

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	REV.	REASON FOR CHANGE	ISSUED AND CONTROLLED BY: (NAME/FUNCTION/SIGNATURE)	APPROVED BY: (NAME/FUNCTION/SIGNATURE)
08/09/2021	4	Internal volume corrected.	Kinga Gawdzik – Engineering Support Technician 	Andrew Pearce – Quality Manager 



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## CUSTOMER APPROVAL:

We accept this material specification as a part of the agreed terms of delivery.

Company Name:

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.*