

# PRODUCT SPECIFICATION

Product P/N	2200/56	Mod. 984A Rev. 06
Description	Insufflation Filter	

**2200/56**

***Insufflation Filter***



<b>PRODUCT DESCRIPTION</b>	Inlet/Outlet Connectors: 6mm hose barbed both sides. Approx. dimensions: 65mm diameter x 54.1mm height. Bidirectional filter. Weight 17gm (approx.).
<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>	To be used in conjunction with insufflators for intra-abdominal insufflation.
<b>CLASS OF THE PRODUCT</b>	Disposable medical device - Class IIa Rule 2 Annex IX 93/42 / EEC Rule 2 Annex VIII MDR 2017/745
<b>MATERIALS</b>	<b>Filter media: <i>Hydrophobic Glass Microfibre Media</i></b>  <b>Frame/Housing Polymer: <i>Clear Styrene-Butadiene Copolymer (SBC)</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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	<p><b>Physical/Mechanical</b>  <i>Approx. dimensions: 65mm diameter x 54.1mm height.</i>  <i>Weight: 17gm (approx.).</i>  <i>Interfaces (ex: Input / Output connectors): 6mm hose barbed both sides.</i>  <i>Operating temperature Range: N/A</i>  <i>Storage temperature Range: 5 °C to 40 °C</i>  <b>Bidirectional Filter.</b></p> <p><b>Biological</b>  <i>Pyrogenicity: &lt;0.3 EU/ml</i>  <b>Biocompatibility to ISO10993</b>  <i>Category – Surface device</i>  <i>Contact – Skin</i>  <i>Contact Duration - &lt;24hrs</i></p> <p><b>Functional</b>  <i>Air Flow Rate: <b>Min. 28.4l/min @ 5PSI</b> (Ref.2200/05, REP 0614/15 with 20% Factor of Safety applied to Min.)</i>  <i>Filtration Efficiency:</i>  <i>Filter Efficiency @ 15L/min using DEHS @0.3µm: <b>Min.99.9%</b> (Ref. 2200/15 REP 1218/17 with Factor of Safety)</i>  <i>Pressure Drop: N/A</i>  <i>Internal Volume: N/A</i>  <i>Bacterial Filtration Efficiency in accordance with ASTM F2101-07: <b>Min. 99.999%</b> (Staphylococcus aureus @ 30L /minute) Ref. 2200/05 REP: EXT846022-S01.</i>  <i>Viral Filtration Efficiency in accordance with ASTM F2101-07: <b>Min. 99.999%</b> (Bacteriophage @ 30L/ minute) Ref. 2200/05 REP: EXT846021-S01.</i>  <i>Operating Lifetime: <b>Refer to Instructions for Use.</b></i>  <i>Shelf Lifetime: <b>5 years from the date of manufacture.</b></i></p> <p><b>Cleanliness</b>  <i>Device assembled within Class 8 Cleanroom.</i></p> <p><b>Testing</b>  <i>Burst test: <b>Min.15.7psi</b> (Ref.2200/05, REP 0618/15 with 20% Factor of Safety applied to Min.)</i>  <i>Leakage: <b>Static Head Test 27" of water for 1 minute</b> (Ref.2200/05, REP: 0622/15)</i></p>
INSTRUCTIONS / WARNINGS	Multi-language IFU available.
PRODUCT SHELF LIFE	<i>5 years from the date of manufacture.</i> <i>Expiration date and date of manufacture are detailed on the product labelling.</i>
STERILIZATION	Sterile version of product available (Ethylene oxide - Max 55°C)
APPLICABLE STANDARDS AND REGULATIONS	<b>Product Certification required:</b> <ul style="list-style-type: none"> <li>- CE mark</li> <li>- FDA</li> </ul>

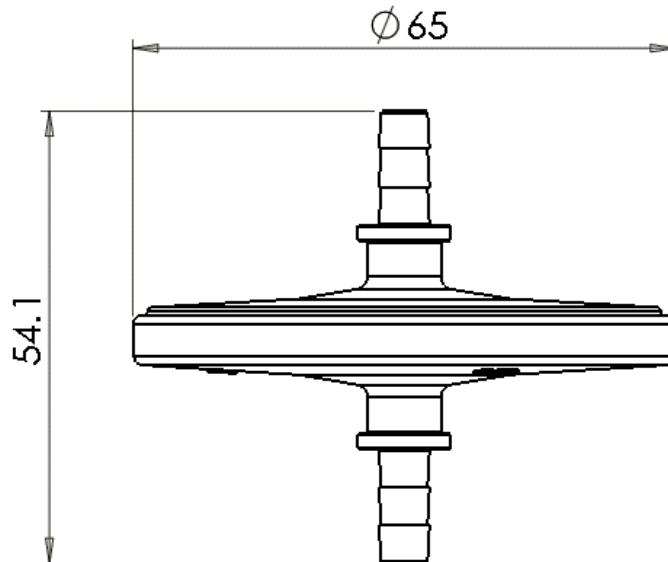
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	<p><b>Applicable Standards and Technical Regulations:</b></p> <p><i>Biological evaluation of Medical Devices - Part 1: Evaluation and Testing - ISO 10993-1.</i></p> <p><i>Medical devices- Application of risk management to medical devices - BS EN ISO 14971.</i></p> <p><i>Medical devices – symbols to be used with medical device labels, labelling and information to be supplied – Part 1: General requirements - ISO 15223-1.</i></p> <p><i>Sterilization of health care products – Ethylene oxide sterilization – ISO 11135-1.</i></p> <p><i>Sterilization of medical devices – Microbiological Methods – Part 1: Estimation of population of Microorganisms on products – ISO 11737-1.</i></p> <p><i>High Efficiency Air Filters – BS EN 1822.</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 packaged 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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Approximate dimensions for reference only

## ACCEPTABLE QUALITY LEVEL

AQL: with sampling Plan: 0.65 AQL to ISO 2859-1

## 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	Short fill moulding	0.65	
5	Rough surface or edges	0.65	
6	Pronounced injection gate	0.65	
7	Deformation/distortion	0.65	

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	8	Crack	0.65	
	9	Oil/grease	0.65	
	10	Wrong colour	0.65	
	11	Weld fault	0.65	
	12	Weld marks	0.65	
<b>GENERAL SAFETY AND PERFORMANCE REQUIREMENTS</b>	<b>Special characteristic:</b> <i>Product characteristic which can affect safety or compliance with regulations, fit, function, performance or subsequent processing of product.</i>			
	<b>Special Characteristic # 01:</b> <i>Air Flow rate at 5psi</i>			
	<b>Special Characteristic # 02:</b> <i>Filter Efficiency @ 15L/min using DEHS @ 0.3µm in accordance with High Efficiency Air Filters – BS EN 1822.</i>			
	<b>Special Characteristic # 03:</b> <i>Burst Pressure</i>			
	<b>Special Characteristic # 04:</b> <i>27” water leak test – Static head test</i>			
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)
09/09/2021	2	Sterilization section updated.	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.*