

Product P/N	2000/39	Mod. 984A
		Rev. 06
Description	Vent Filter	

2000/39

Vent Filter



PRODUCT DESCRIPTION	Inlet Outlet Connectors: 1/8th NPT thread 18mm, 3.5mm diameter vent hole. Approx. Dimensions: 54mm diameter x 27mm height. Weight: 11g (approx.). Bidirectional Filter.		
MANUFACTURER NAME	GVS Filter Technology UK NFC House Vickers Industrial Estate Mellishaw Lane, Morecambe Lancashire LA3 3EN - United Kingdom Information		
	Tel. +44 (0) 1524 847600		
INTENDED USE /	e-mail: gvsuk@gvs.com Filters, hydrophobic and non-hydrophobic for use within oxygen concentrators and other		
APPLICATION	medical equipment such as ventilators.		
CLASS OF THE			
PRODUCT	Disposable medical device - Class IIa Rule 2 Annex IX 93/42 / EEC		
PRODUCT	Rule 2 Annex VIII MDR 2017/745		
MATERIALS	Filter media: Hydrophobic Glass Microfibre Media		
	Frame/Housing Polymer: Transparent clear Polypropylene homopolymer (PP) White Polypropylene homopolymer (PP)		
	Colour: Transparent White		
	Regulatory Documentation Required:		
	- Biocompatibility according ISO 10993-1		
	- ROHS		
	- BSE/TSE		
	- DEHP plasticizer Free and latex free - Aging		
	- Aging - REACH		
	- Conflict minerals		
PRODUCT	Appearance/Visual		
CHARACTERISTICS	As shown on drawing.		



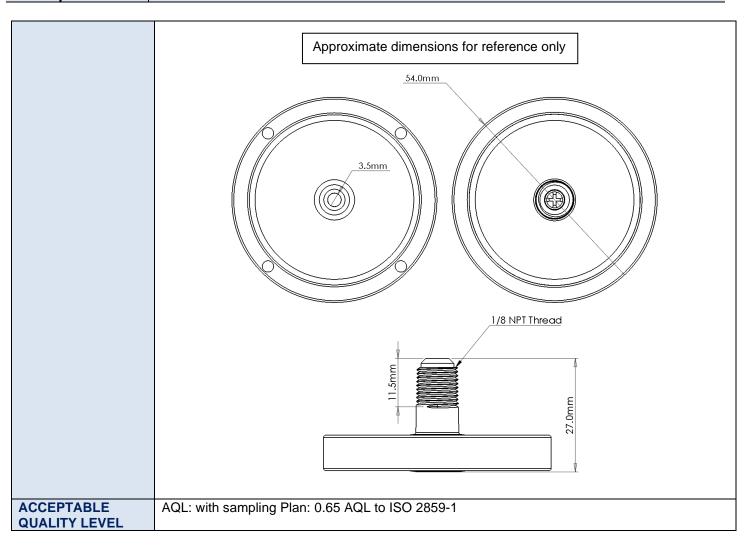
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	Physical/Mechanical Approx. dimensions: 54mm diameter x 27mm height. Weight: 11gm (approx.). Interfaces (ex: Input / Output connectors): 1/8th NPT thread 18mm and 3.5mm hole. Operating temperature Range: N/A Storage temperature Range: 5 °C to 40 °C Bidirectional Filter.	n diameter vent
	Biological Pyrogenicity: <0.3 Eu/ml Biocompatibility to ISO10993 Category – Surface device Contact – Skin Contact Duration - <24hrs	
	Functional Air Flow Rate: <i>Min. 20.5 I/min</i> @ 1PSI (REP: 2052/20 with 20% Factor of Safety applied to	o Min.)
	Filtration Efficiency: Filter Efficiency DEHS @ 0.3 microns @ 15L/min: Min. 99.9 (REP: 1383/17 with Factor of Safety applied to Min.)	99%
	Pressure Drop: N/A	
	Internal Volume: N/A	
	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/min) REP: EXT831477.	6
	Viral Filtration Efficiency in accordance with ASTM F2101-07: Min. 99.999% (Bacteriophage @ 30l/min) REP: EXT831476.	
	Cleanliness Device assembled within Clean Manufacturing Environment.	
	Testing Static Head Test 27" for 1 minute (REP 0620/15)	
INSTRUCTIONS / WARNINGS	IFU available in English language.	
PRODUCT SHELF LIFE	5 years from the date of manufacture.	
STERILIZATION	Expiration date and date of manufacture are detailed on the product labelling. N/A	



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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.
	Medical devices- Application of risk management to medical devices - BS EN IS	SO 14971.
	Medical devices – symbols to be used with medical device labels, labelling and be supplied - Part1: General requirements - ISO 15223-1.	information to
	High Efficiency Air Filters – BS EN 1822.	
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: V Quantity V Product description V Product Date V Lot Number (OL and 5-digit batch number to trace back to raw V Operator Code Different lots in one box are separately closed and separately labels products will be packed in double PE bags.	,
CERTIFICATE OF COMPLIANCE	With each shipment, GVS UK Customer Service will send the CofC to the Cust the lot numbers and date of manufacture. Conformity declaration is printed on every invoice and Certificate is according to type 2.1. The Quality management system is in compliance with ISO 9001, ISO 13485.	·
DRAWING	The attached drawing is part of this product specification and must not be dupli accessible to a third party without written permission from GVS Filter Technolog	



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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	
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	ISO 2859 Part 1 General
6	Pronounced injection gate	0.65	Inspection Level 1
7	Deformation/distortion	0.65	
8	Crack	0.65	
9	Oil/grease	0.65	
10	Wrong colour	0.65	

GENERAL SAFETY AND PERFORMANCE REQUIREMENTS

Special characteristic: Product characteristic which can affect safety or compliance with

regulations, fit, function, performance or subsequent processing of product.

Special Characteristic # 01: Air Flow rate at 1psi

Special Characteristic # 02: Filter Efficiency @ 15L/min using DEHS @ 0.3μm

Special Characteristic # 03: Bacterial Filtration Efficiency in accordance with ASTM F2101-07

Viral Filtration Efficiency in accordance with ASTM F2101-07

Special Characteristic # 04: 27" water leak test - Static head test

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



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REVISIONS AND APPROVALS:

DATE	REV.	REASON FOR CHANGE	ISSUED AND CONTROLLED BY: (NAME/FUNCTION/SIGNATURE)	APPROVED BY: (NAME/FUNCTION/SIGNATURE)
15/09/2020	1	Initial release.	Kinga Gawdzik - Engineering Support Technician	Andrew Pearce – Quality Manager

	CUSTOMER APPROVAL:
We accept this	s material specification as a part of the agreed terms of delivery.
Company Nan	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.