

4100/30	
Oxygen Concentrator Filter	Mod. 984 Rev. 05
	4100/30 Oxygen Concentrator Filter

4100/30

Oxygen Concentrator Filter



PRODUCT DESCRIPTION	Inlet / Outlet Connectors: 22mm Male, length: 12.5mm (approx.) Approx. Dimensions: 80.2mm length x 60.9mm width x 69.4mm height Weight: 50gm (approx.) Unidirectional Filter Similar existing product: ITI N/A ITI Specify: 4100 series filters.
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 e-mail: gvsuk@gvs.com
INTENDED USE / APPLICATION	Pre-filter for use within oxygen concentrator machines, to filter the air entering the device.
PRODUCT	Disposable medical device - Class 1 Annex V 93/42/EEC Rule 1 Annex VIII MDR 2017/745
MATERIALS	Filter media: I N/A Specify: Glass Microfibre Media Frame/Housing Polymer: I N/A Specify: Bronze Tinted Acrylonitryle Butadiene Styrene (ABS) Colour: I N/A Specify: Bronze Other insert(s): I N/A Specify: Adhesive: Polyolefin Hot Melt Adhesive Regulatory Documentation Required: I Biocompatibility according ISO 10993-1 I I IMDS



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Description	Oxygen Concentrator Filter Rev. 05
	IFI ROHS
	IFI BSE/TSE
	DEHP plasticizer Free and latex free
	I⊽l Aging
	IFI REACH IFI Conflict minerals
	(*) Connict minerals
	Other:
	IFI N/A IFI Specify:
PRODUCT	Appearance/Visual
CHARACTERISTI	7 toothotto part.
	I⊽I Yes; I⊤I No
	Finishing: I Trim finishing mirror polished surfaces
	Thin roughness surfaces
	I⊓ Gross roughness surface
	I⊽l As shown on drawing
	Presence of graphics/Artwork:
	141 No, 1 Topechy.
	Flash bears admitted:
	I⊓ N/A IPI Specify: Refer to the control plan.
	Injection gate vestige:
	I⊓ N/A I Specify: Refer to the control plan.
	A TAIL TO POSITY THE STATE OF T
	Black spots admitted (size/area):
	I⊓ N/A I¬I Specify: Refer to the control plan.
	Infiltration into insert admitted:
	IFIN/A I□ Specify:
	Other:
	I⊽I N/A I□I Specify:
	Physical/Mechanical
	Dimensions (Approx.): 80.2mm length x 60.9mm width x 69.4mm height Weight: 50gm (approx.)
	Interfaces (ex: Input / Output connectors):
	IFIN/A IFI Specify: 22mm Male
	Operating temperature Range: N/A
	Storage temperature Range: 5 °C to 40 °C
	Other: ITI N/A ITI Specify: Unidirectional Filter
	Tital Coposity. Chambed on a Tital
	Chemical
	Compatibility to solvents:
	IVIN/A I⊓I Specify:
	Resistance to weathering:
	IFI N/A IFI Specify:



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Other: MN/A MSpecify: Biological Sterility: IN/A ITI Specify: Pyrogenicity: IVI N/A ITI Specify: Haemolysis: IFI N/A IFI Specify: Toxicity: IV N/A IV Specify: Biocompatibility to ISO10993 Category - Surface device Contact - Intact Skin Contact Duration - <24hr **Functional** Air Flow Rate: ITI N/A ITI Specify: 30l/min, 60l/min, 90l/min Water Break Through Pressure: IPI N/A ITI Specify: Liquid Flow Rate: IPI N/A ITI Specify: **Bubble Point Pressure:** MN/A ITI Specify: Filtration Efficiency: ITI N/A ITI Specify: Filter Efficiency @ 30L/min: Min. 95% (REP: Ref. to Test Result 22-01-10) Pressure Drop: ITI N/A ITI Specify: Flow Resistance @ 30L/min in accordance with EN ISO 9360-1: Max. 101Pa Flow Resistance @ 60L/min in accordance with EN ISO 9360-1: Max. 296Pa Flow Resistance @ 90L/min in accordance with EN ISO 9360-1: Max. 602Pa (REP: Ref. to Test Result 22-01-10) Internal Volume: IN/A ITI Specify: Operating Lifetime: IN/A I Specify: Refer to Instructions for Use. Shelf Lifetime: ITI N/A ITI Specify: 5 years from the date of manufacture. Other:

I N/A I Specify:



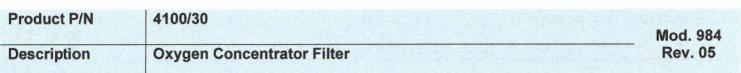
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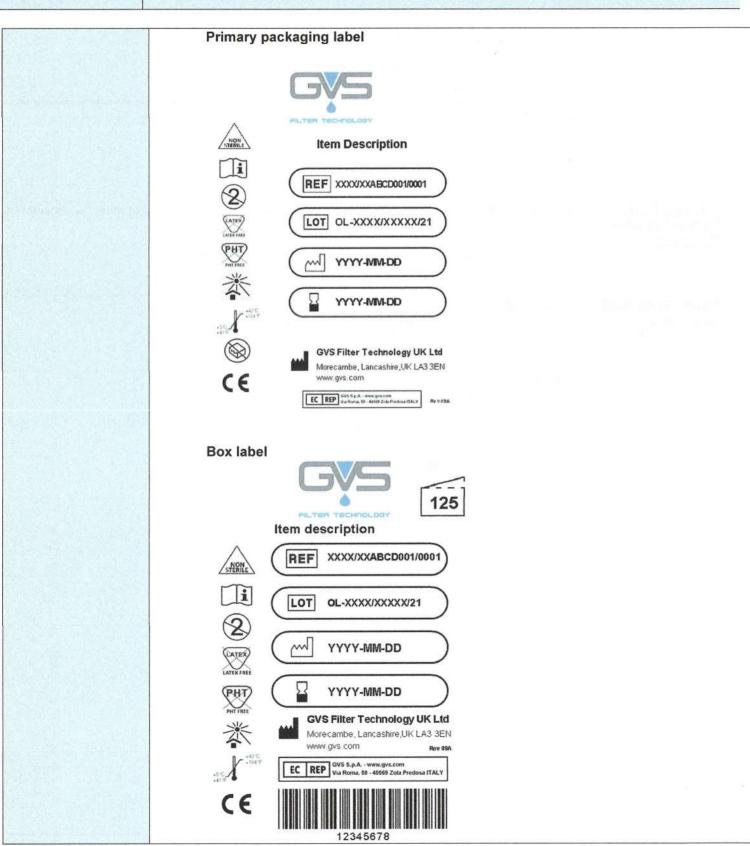
	Cleanliness Part cleanliness: IFI N/A IFI Specify: Clean-room Production required: IFI N/A IFI Specify: Class 8. Testing
	Insert embedding Test: IF N/A IF Specify: Burst test: IF N/A IF Specify: Leakage: IF N/A IF Specify: Other: IF N/A IF Specify:
INSTRUCTIONS / WARNINGS	IFU □ N/A □ Required: Specify Languages: □ English □ French □ Italian □ Spanish □ Portuguese □ German □ Chinese □ Japanese □ Korean □ Russian □ Other (Specify):
PRODUCT SHELF LIFE	5 years from the date of manufacture. Expiration date and date of manufacture are detailed on the product labelling.
STERILIZATION	IFI Not required; IFI EtO; IFIGamma; IFI Beta; IFISteam; IFI e-beam
APPLICABLE STANDARDS AND REGULATIONS	Product Certification required: I♥I CE mark I♥I FDA I□ Canadian Health I□ Brazil ANVISA I□ Argentina ANMAT I□ NIOSH I□ Japan I□ Korea I□ China I□ Russia I□ Other (Specify):
	Applicable Standards and Technical Regulations: Biological evaluation of Medical Devices - Part 1: Evaluation and Testing - ISO 10993-1. Respiratory protective devices - Method for test - Part 7: Determination of particle filter penetration - BS EN 13274-7.
	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.



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	Customer Specific Requirements: No
	Is the Product a safety product? I□I Yes I□I No
	Environment Impact: Have atmosphere emissions, liquid effluents, underground leachables been considered? For Yes For No
	Energy released (heat, vibration):
INTELECTUAL PROPERTY AND FUNDING	Intellectual Property: Patents that have been granted or applied for that must be considered.
FUNDING	Grants: Is the project (partially) funded by public institutions? I™I N/A I™I Specify:
PACKAGING AND LABELING	Packaging I♥I Primary: Refer to individual Part Numbers. I♥I Secondary: Refer to individual Part Numbers.
	Labelling: I⊽l Standard; Iℂl Customer specific:
	Filter label for 4100/30LAUA
	FILTER TECHNOLOGY
	HIGH EFFICIENCY PARTICULATE AIR (HEPA) FILTER 4100/30LAUA
	INSTALLED BY
	DATE









Product P/N	4100/30 Mod. 984
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	UDI:
	IFI N/A; IFI No; IFI Yes (Specify): 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: ✓ Quantity ✓ Product description ✓ Product Date ✓ Lot Number (OL and 5 digit batch number to trace back to raw materials used) ✓ Operator Code
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 9001, ISO 13485.
DRAWING The attached drawing is part of this product specification and must not be dup accessible to a third party without written permission from GVS Filter Technology.	
	80.2 A 69 A
	3
ACCEPTABLE QUALITY LEVEL	I⊽I AQL: with sampling Plan: 0.65 AQL to ISO 2859-1 I□I PPM: I□I Other:
VISUAL REQUIREMENTS	Visual acceptance requirements apply when inspected under below conditions:



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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	Weld marks	0.65	
5	Short fill moulding	0.65	
6	Rough surface or edges	0.65	ISO 2859 Part 1 Genera
7	Pronounced injection gate	0.65	Inspection Level 1
8	Deformation/distortion	0.65	
9	Crack	0.65	
LO	Oil/grease	0.65	
.1	Wrong colour	0.65	
.2	Weld fault	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:

Flow Resistance @ 30L/min in accordance with EN ISO 9360-1

Flow Resistance @ 60L/min in accordance with EN ISO 9360-1

Flow Resistance @ 90L/min in accordance with EN ISO 9360-1

Special Characteristic # 02: Filter Efficiency @ 30L/min in accordance with EN 13274-7.

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:

REV.	REASON FOR CHANGE	ISSUED AND CONTROLLED BY: (NAME/FUNCTION/SIGNATURE)	APPROVED BY: (NAME/FUNCTION/SIGNATURE)
1	First Issue	Kinga Gawdzik – Engineering Support Technician	Andrew Pearce – Quality Manager
	REV.	CHANGE	CHANGE BY: (NAME/FUNCTION/SIGNATURE) 1 First Issue Kinga Gawdzik – Engineering

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.

Land To the Street Persons Sent

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