Taglia XS - monouso/non-sterile

DOCUMENTAZIONE





Taglia XS - monouso / non-sterile









Taglia XS - monouso / non-sterile

CARATTERISTICHE

Mascherina FFP2 monouso taglia XS, in tessuto-non-tessuto (TNT), dotata di elastici auricolari che garantiscono grande comodità e aderenza a una vasta gamma di conformazioni del volto.

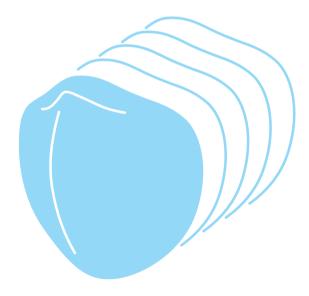
Adatta per la protezione delle vie respiratorie da aerosol e particelle non-oleose, polvere, smog, allergeni e altri agenti contaminanti.

Una volta usate, le mascherine monouso debbono essere immediatamente smaltite all'interno del bidone della raccolta indifferenziata, reggendola dagli elastici auricolari ed evitando contatti con la superficie. Seguire le norme locali vigenti per lo smaltimento dei rifiuti indifferenziati.

Certificazione: le mascherine sono dotate di attestato di certificazione CE e rispettano i requisiti delle norme tecniche armonizzate EN 149:2001+A1:2009.

Confezionamento: Busta 1 pz. / Box 25 pz.

DETTAGLI TECNICI



FFP2 NR (monouso)

Composizione materiale: 49% TNT, 19% Hot air cotton, 32% meltblown

Formato: Box (20 pezzi) / involucro singolo

Dimensioni: 12,0 * 10,5 cm

Standard Filtrazione: EN149:2001+A1:2009 FFP2 NR

Condizioni di conservazione: immagazzinare in luogo fresco e asciutto, a temperatura compresa tra -20° e 40° C. Non esporre a luce solare diretta, luce UV e lampade fluorescenti. Se il confezionamento e danneggiato o bagnato scartare il prodotto. Non utilizzare oltre la data di scadenza (riportata sulla

Durata d'utilizzo massimo consigliata: 8 ore

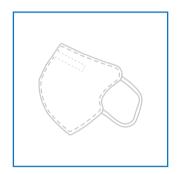
Shelf life: 2 anni dalla data di produzione

FUNZIONI

Le Mascherine FFP2 vengono utilizzate frequentemente per prevenire contagi e per prevenire la respirazione di particelle dannose in ambito edile, nell'industria chimica, oppure come dispositivo di protezione da allergeni.

Taglia XS - monouso / non-sterile

ISTRUZIONI PER L'USO









Controllare l'integrità dell'involucro prima dell'uso. Se la confezione è rotta, non utilizzare.

- 1. Aprire la confezione, controllando che la mascherina non rechi segni d'usura o macchie
- 2. Indossare la mascherina appoggiandola sul volto a partire dal naso. Agganciare gli elastici auricolari dietro le orecchie.
- 3. Modellare la mascherina adattandola alla forma del volto fino a un livello di aderenza ottimale
- 4. Modellare la mascherina sulle mani fino all'ottenimento di un'aderenza ottimale. Regolare clip nasale ed elastici finchè la mascherina non aderisce al viso correttamente.
- 5. Premere entrambi le mani sulla mascherine e soffiare vigorosamente, controllando per mancate aderenze; regolare clip nasale e d elastici finchè la mascherina non aderisce al viso correttamente.

INDICAZIONI

Questa maschera non deve essere utilizzata se l'uso previsto del prodotto è quello di proteggere l'indossatore da alte concentrazioni di agenti infettivi (lavoro svolto a stretto contatto con batteri, virus e funghi - e.g. laboratorio).

Non protegge da gas e vapori (e.g. gas anestetici, fumiganti e gas tossici).

La mascherina non è sterile. Non deve essere utilizzata in tutte le procedure che richiedano il requisito della sterilità.

Per impedire potenziali contaminazioni, non riporre la mascherina in tasca o dentro a indumenti/borse e non appoggiare la mascherina su superfici. Se, durante l'uso, dovesse rendersi necessario rimuovere la maschera, evitare contatto con mani / naso / bocca e la superficie esterna della maschera. La mascherina è monouso, non riutilizzabile o reciclabile.

IMPORTANTE

Per un uso corretto, seguire attentamente le istruzioni d'uso. Materiali in diretto contatto con la pelle possono causare una reazione allergica in certi individui ipersensibili.

Certificazioni / Test report





Certificate of approval Konformitätsbescheinigung

Product name: Filtering half mask Produktname: FFP2 Filter Halbmaske

Model: CARE0961K Modell: CARE0961K

Production batch No: EN149:2001+A1:2009 FFP2 NR Executive Standard: EN149:2001+A1:2009 FFP2 NR

Product specification: Nonwovens, Molt Blown, Hot Air Cotton Produktspezifkation: Vliesstoffe, schmelzgeblasen, Heißluftbaumwolle

Main ingredients: 49 % non-woven fabric, 19 % hot air cotton, 32 % melt blown fabric

Hauptzutaten: 49% Vliesstoff.

19% Heißluftbaumwolle, 32% schmelzgeblasener Stoff

Batch No./Date: FX21022 Produktionscharge/Datum: 2021.03.05 Period of validity: 2 Years Gültigkeitsdauer: 2 Jahre

Inspector:02 Quantity: 20 Inspektor:02 Menge: 20

Careable Biotechnology Co.,Ltd Building O, No.3, hongxing road, jiangmen City, China

FFP2 Faltbare Atemschutzmaske Máscara Plegable FFP2 Masque Pllable FFP2

Maschera Respiratoria Pieghevole FFP2
FFP2 opvouwbaar ademhalingsbeschermingsmasker

DEUTSCH AUFSETZANLEITUNG - siehe Gebrauchsanleituing

ESPAÑOL INSTRUCCIONES DE COLOCACIÓN - ver instrucciones

FRANÇAIS COMMENT METIRE LE MASQUE - voir le mode d'amploi

ITALIANO ISTRUZIONI PER LUSO - vedi istruzioni per l'uso
NEDERLANDS INSTRUCTIES VOOR HET DRAGEN - zie de gebruiksaanwijzing







Manufacturer/Hersteller/Fabricante/ Fabricant/ Produttore/ Fabrikant

Careable Biotechnology Co.,Ltd Building O, No.3, hongxing road, jiangmen City, China

Imported By: Befast International Logistics Europe Rue Colone Bourg 127-129 1140 Brussl Belgium

DEUTECH GEBRUICHEAMISSUNG PRODUKTEIGENSCHAFTEN:

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ARRIVASI:

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EU DECLARATION OF CONFORMITY

The manfacturer: Careable Biotechnology Co. Ltd.

Building O. No. 3, hongoing road, ilangmen City, China

Imported By: Befast International Logistics Europe Rue Colone Bourg 127-129 1140 Brussl Belgium

Here declares under its own responsibility that the personal protective

equipment (PPE) in category III: Model Number:CARE0961K Description: Filtering Half Mask

Product type: Respiratiry protective device. Filtering Half Mask to protect against particles

It is produced in compliance with the essential health and safety requirements according to Regulation (EU) 2016/425 of the European Parliament and of Council of 09 March 2016 on personal protective equipment regulation and with the following reference standard

and with the following reference standard

EN 149-2001 + A1:2009 Respiratory protective devices - Filtering half masks to protect against particles - Requirements, testing, marking is identical to the PPE which is the subject of EC Type certificate no.

is identical to the PPE which is the subject of EC Type certificate in CW/PPER/46/12/2020/rev 1

issued by Notified Body no. 1463: Polski Rejestr Statkó w S.A. - al. gen. Jó zefa Hallera 126,80-416 Gdansk, Poland

The PPE listed above is manufactured under a system which satisfies the requirements of Module C2 (Annex VIII) of the same regulation, certificate no. CW/PPER/S/03/2021.
Polisk Reiser's Statis on S. A. al. een. Józefa

Hallera 126,80-416 Gdansk, Poland Poland Notified Body Number 1463

Name: James Jing Position: General Manager Signature/Unterschrift: Date/Datum: 2020-06-20







CERTYFIKAT ZGODNOŚCI Z TYPEM W OPARCIU O WEWNĘTRZNĄ KONTROLĘ PRODUKCJI ORAZ NADZOROWANE KONTROLE PRODUKTU W LOSOWYCH ODSTEPACH CZASU (Moduł C2)

CONFORMITY TO TYPE CERTIFICATE BASED ON INTERNAL PRODUCTION CONTROL PLUS SUPERVISED PRODUCT CHECKS AT RANDOM INTERVALS (Module C2)

No.	CW/PPER/2	2/01/2021		Oknes objety cartyfile Period covered by the		2021-0	1-12 - 20	122-01-1
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	tering half mas ass FFP2 NR).), Model: CARE	0961K					
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Foliki Reasty Statkow S.A.

tr jednostki retylkowanej

1463

tel. (+48) (58) 346 17 00 fax (+48) (58) 346 03 92 e-mail: mailbookkgr.uk

Roczna ocena niejednorodności produkcia Annual assessment of production non-homogeneity Zastronowana materia przy dokonanie ocene

inspekcja procesu produkcyjnego i zapistw z prilit On site review of productive and test recents

Audit kentroli protessi pro-

Ocera niejednorodności produkcji pograsa ocereg jednej dużej próbki
Anada top opo-topnaprenty ocerosof by selection of a cleak, longe speciele

Goesa niejedwarodności produkcji pograna ocene politek w ciągu roku. Anadactice non-boroagenisty ouzstanii by stanownich of pampies throughout the year

2a Doong prosprovadoli (ireig, naowisko) Zwiazek z jednostka notyfikowana

2b Przedstawiciel Formy Limit, nazwiskoś

Company representative (Name)

Na podržavia prosprovadovnii oceny stalierdono, še proces produkcijni jest jednorodny

O Tak

Conclusion undrieria siegandroici

Nie było żadnych niegzodności / There were no non-conformities.

Writish jednostii rotyt kraumi

Środek ochrony osobistej jest kompatybikny z typem okredionym w certyfikacie badania typu UE.

Personal protective equipment is compatible with the type defined in the EC type-expraination certificate. Umani

1. Półmaska filtrująca bez zaworu.

2. Prihmaska filtrujaca przeznaczone do jednorazowano użytku.

3. Dokumentacja techniczna zatwierdzona w języku angielskim.

4. Produkt ten nie może buć stosowany jako maska przeciwgazowa w środowisku toksycznym. 5. Półmaska filtrująca nie jest prozonaczona do użytkowonia medycznego i chiruralcznego.

6. Półmaska filtrująca nie powinna być używana w środowisku o stężeniu tienu poniżej 19.5 %.

2. Filtering half mask without valve.

2. Filtering half mask shall not be used for more than one shift.

4. This product can not be used as a gas mask in a taxic environment. 5. Filtering half mask can not be used for medical and surgical purposes.

6. Filtering half mask should not be used in an environment with anypen contens less then 19.5%.

Załączniki

Oetina ocena z rocznego nadzonu Overall assessment of the annual syrvelllance Gdańsk, 2021-01-12

Hora Sinna Cortafilms

Fern, 10 PCW-01/PPER





CERTYFIKAT BADANIA TYPU UE (MODUŁ B) EU TYPE-EXAMINATION CERTIFICATE (MODULE B)

Nr CW/PPER/46/12/2020

ZAŚWIADCZA SIE

Le Pobli Riquist Statiów S.A. (PSS) przeprowadził procedurę bedania typu wymienionego mżej wyrobu i stwierdził jego zgodność. 2 wymaganiani ośroślowymi w zakączniku V do Roporządomie Parlamentu Europojskego i Rady (UZI 2016/455 (PPC) w sprawie środoko ochrony indystatulanie poza udoślenia dyskłowym Bady SSIGE/SWC, ze miamenti.

THIS IS TO CERTIFY

THIS I VICENIES IN THE PART OF THE PART OF

Wnioskodawca Applicant Careable Biotechnology Co., Ltd. Building O, no. 3, Hongxing road, Jiangmen City, China.

Producent Manufecturer Careable Biotechnology Co., Ltd. Building O, no. 3, Hongking road, Jiangmen City, China.

Typ wyrobu

Sprzet ochrony układu oddechowego. Półmaski filtrujące do ochrony przed cząstkami. Respiratory protective devices. Filtening holf masks to protect against particles.

Opis wyrobu

Pólmaska filtrująca, model: CARE 0961K (klasa FFP2 NR). Filtering half mask. Model: CARE 0961K (class FFP2 NR).

Anadact description

PN-EN 149+A1:2010 EN 149-2001+A1:2009

This certificati

Nei ejszy certyfikat pozoszaje wsamy do czasu uniewszmienia przy zachowania warunków uznania (patrz str. 2). This certificate remains whó unless cancelled or revoked, provided the approval conditions (see page 2) are compilal with

Data walności

2025-12-10

9

Gdańsk, 2020-12-11

tr jednosti setylikowanej vs. of notfed body 1463 Polski Rejestr Statków S.A. al. Gen. litzefa Hollera 126 80-416 Gdańsk, Poland tel (+48)/58) 345 17 00 fax (+48)/58) 345 17 00 e-mail: deligning server, \$110 J/Marker and old

Zantepora Dyrektora Pjonu Certyfikacji

Form. 8/FCW-01/PPER. 2020-02-36

Wykac dokumentacii

- Instrukcja użytkowania zatwierdzona przez PRS S.A. dnia 2020-12-03.
 - Ocena purvisa zatwierdzona przez RIS S A. dnia 2020-12-03. Dokumentacja techniczną "Półmask filtrującei, model: CARE 0961K" - zatwierdzony przez PRS S.A.
 - deia 2020,13,03 4. Report a hadari or IKF20034101 wydany przes Zhellana Academy of Science and Technology for
 - Inspection & Quarantine (Technology Center of Hangzhou Customs District/Zheijang Lead Product Technical Co., Ltd.) z akredytacją CNAS L0354 z dnia 2020-12-08. 5. Sprawoodanie z przepladu PRS S & cz CW/MoK/PPFR/283/2828 z dnia 2020-12-10.

 - I. Instruction of use approved by PRS 5.A. on 2020-12-03.
 - 2. Risk analysis approved by PRS S.A. xr 2020-12-03.
 - 3. Technical documentation "Filtering half mask, Made): CARE 0961A" approved by PRS S.A. on 2020 12 03 4. Test report No. XF20034101 issued by Zheliang Academy of Science and Technology for Inspection & Quarantine (Technology Center of Hangzhou Customs District/Zhejiong Lead Product

Technical Co., Ltd.) with CNAS accresitation no. LOSS4, dated on 2020-12-08. 5. PRS S.A. Survey Report No. CW/Mox/PPER/283/2020 dated on 2020-22-20.

(innem) podane na strenie 1) (different than alsen on ease 1)

Cereniczenia uznania 1. Dane techniczne:

Approval designations a) półmaska filtrująca z regulowanym klipsem na nos,

- b) klips na nos montowany wewnąti z półmaski filtrującej, ci półmaska filtrująca wykonana z 4 warstwowej włókniny z filtrem z tkaniny.
- di półmaska filtrująca wyposażona w zauszniki, e) półmaska filtrująca bez zaworu.
- f) wymiary: 105 mm ± 1 mm x 120 mm ± 1 mm, g) docelowa grupa użytkowa: dzieci powyżej trzeciego roku życia,
- h) kolony: półmaska filtrująca zausmki biole
- 2. Pôłmaska filtrująca przeznaczona do jednorazowego użytku. 3. Dokumentacja techniczna zatwierdzona w języku angielskim.
- 4. Produkt ten nie może być stoszware lako maska przeciwanowa w środowisku tokycznym. 5. Półmaska filtrująca nie powinna byćubywana w środowisku o steżeniu tlenu poniżei 19.5 %.
- 6. Półmaska filtrująca nie jest przeznaczona do użytkowania medycznego i chinurgicznego.

L. Specifications:

- a) filtering half mask with adjustable wase clip. b) nose clip mounted inside the filtering half mask,
- c) filtering half mask made with 4 layers non-waven fabric with melt-blown fabric filter.
- d) Altering half mask with ear loops. el filtering half-mass without valve
- () size: 105 mm ± 1 mm x 120 mm ±1 mm. a) target aroup: children over the ace of three,

b) colors filtering half mask valve

- 2. Filtering half mask shall not be used for more than one shift 3. Technical documentation approved in English.
- 4. This product can not be used as a pas mask in a toxic environment.
- 5. Filtering half mask should not be used in an environment with awaren contens less then 19.5%. 6. Filtering half mask can not be used for medical and surplical numbers.

Warunki senania Approval consistent

1 Miniepty certyfikat styleci ważność po wprowadzenia zmian lab medyfikacjiw wyrobie bez aprzednego uzgodnienia z PRS. This cortificate becomes invalid after changes or modifications to the product without prior agreement with PRS

2 Your madvaks made but unnecessary as company worship our made but wastaway delibracia products belie and wastaway to be because a backeriere typu UE zostanio prosprosestoma ocena agodnosici produktiji pod nadzonem jednostići notyfikowanej, według zafącznika Wi lub Will wymienionego wyżej rosporządzenia. The Mark of Confirmity, you nate by officed to the above time manneyd product and a remainstrater's Declaration of Confirmity manner provided the production is assessed under surrevillence of a notified body according to Asses IVI or VIV of the girn Angulation.



TEST REPORT



Report No.: JKF20034101

Applicant : Careable Biotechnology Co., Ltd

Thejiang Academy of Sea are and Technology for Inspection and Quarantine
Add: No. 398, June 1s. 3 No. 3 No.



The informa	tion are provided by clien	t(applicant):					
	Sample Name:	Filtering half	mask				
Sample Information	Style No.:	Caref9961K					
	Brands	Fusi Care					
	Applicant:	Careable Bios	echnology Co., Ltd	Ei)			
Customer	Address	Building O, n	o.3, hongxing road,	Jingmen	city (002)		
Information	Manufacturer:	Careable Bios	achnology Co., Ltd				
	Manufacturer address:	Building O, o	o.3, hongxing road,	jingmen	city (002)		
The informa	ation are confirmed by test	ing organizatio	in:				
	Date of sample received:	2020-12-02	Testing period-	2020-1	2-02 to 2020-12-08		
2200	Quantity:	70 Pieces					
Test Information	Sample description:	White mask					
Information	Basis of judgments			Filtering I	nalf masks to protect		
Test Conclusion	The items tested meet the	requirements of	EN 1492001+A12	009 FFP:	! NR		
Test Result	Please refer to next pages.						
Remark	,						
Edit:	ha-† e% 2 Ye yiwen		5	šign:	Letti . Zhao dong		

*** End of this page ***



Test Results:

Clause 7.5 Material

(EN 149/2001+A1/2009 Clause 8.2 & 8.3.1 & 8.3.2)

Requirement	Results	Rating
Miterials used shell be satisfied to withstand handling and wear over the period for which the period for behind paled much to dispipate the second. After undergoing the conditioning described in 3.3.1 more of the particle filtering from masks stall how suitedem develoated in allow of the temperor or straps. When conditioned in accordance with 8.3.1 and 8.3.2 the particle filtering half mask shall not codigate. Agy material from the filter media released by the air files through the filter shall not consistence.	Comply	Pass

Clause 7.6 Cleaning and disinfecting

(EN 149/2001+A1/2009 Clause 8.4 & 8.5 & 8.11)

Requirement	Results	Rating
If the purcle Titering half mask is designed to be re-usable, the materials used shall without the cleaning and disinfecting agents and procedures to be specified by the manufactures. With reference to 7.9.2, after cleaning and disinfecting the re-usable particle Ethering half mask shall satisfy the penetration requirement of the relevant clean.	Not applicable (Not designed to be re-usable)	N/A

Clause 7.7 Practical performance

(EN 149-2001+A1-2009 Clause 8.4)

Requirement	Results	Rating
The particle filtering half mask shall undergo practical performance tests under readstic conditions. These general tests serve the purpose of checking the equipment for imperfections that cannot be determined by the tests described elsewhere in this standard.	No imperfections	Pass

Clause 7.8 Finish of parts

(FIS 149-2001- a 1-2009 Charle 5-2.)

Requirement	Results	Rating
Parts of the device likely to come into contact with the wearer shall have no sharp	No sharp edges or	Pass
edges or burrs.	burns	50.00

Requirement	Results	Rating
For purise (therity, half masks froid in accordance with the mounteneurs' information, at Jean 46 and of the 50 individual exercise rouths (i.e. 10 subjects a 5 receives) for and insured bedage shall be not greater than 25% for FFFP, 11% for FFFP, 3% for FFFP, 3% for FFFP and, in addition, it shall so all out of the 10 individual weare withmeric means for the tool in used leakage shall be not generat than. 25% for FFFP, 3% for FFFP, 3% for FFFP 3	46 out of the 50 individual exercise \$\leq 11\% 8 out of the 10 individual wearer arithmetic means \$\leq 3\%	Pass

Table 7.9.1-A lineard leakage test data

Subject	Sample No.	Condition	Walk (%)	Head side/side (%)	Head up/down	Talk (%)	Walk (%)	Mean (%)
CQQ	: 10		6.915	7.814	7.415	10.251	7,314	7.942
WLJ	2		5.617	6.810	6.091	10.991	6.591	7.220
WG	3	As received	9.415	10.615	10.364	15.361	11.947	11.540
ZJH	4		6.954	7.941	6,519	8.915	6.487	7.363
TLB	5		5.619	6.415	7.006	10.935	7.641	7.523
ZMY	6		8.614	6.817	7.015	9.718	6.691	7.771
LJF	7	1 1	10.574	9,609	11.569	18.841	10.694	12.257
HML	8	Temperature	6.481	6.552	7.138	9.891	6.947	7,402
RK.	9	conditioned -	5.369	6.814	7.136	10.364	8.314	7.599
ZD	10	1 1	4.369	5.147	5.618	9.617	6.108	6.172

Table 7.9.1-E Facial dimensions

Subject	Face Length (mm)	Face Width (mm)	Face Depth (mm)	Mouth Width (mm)
CQQ	136	167	125	65
WLJ	132	159	110	60
WG	120	152	109	57
ZJH	122	150	104	50
TLB	125	152	101	57
ZMY.	137	150	120	60
LJF	125	135	90	55
HML	124	130	115	55
RK	112	161	146	50
ZD	116	168	115	55



Clause 7.9.2 Penetration of filter material

(EN 149/2001+A1/2009 Clause 8.11 & EN 13274-7/2019)

	stequiev meat		Buckuns	PCHUNG.
penetration of the foliarements of the folia	iker of the particle filtering I wing table.			
Classification	Sodium chloride test 95 L/min	Paraffin oil test 95 L/min	Detail refer to Table 7.9.2	Pass
FFP1	<20%	≤20%	Table 7.9.2	5555
FFP2	<6%	156%		
FFP3	<1%	<1%		

Table 7.9.2 Penetration of filter material

Acrosol	Condition	Sample No.	Penetration (%)
		- 11	0.250
	As received	12	0.078
		13	0.077
	Simulated wearing	14	0.112
Sodium chloride test		15	0.065
	treatment.	16	0.038
	Mechanical strength+ Temperature conditioned	17	0.750
		18	0.415
	Temperature constitutes	19	0.492
	As received	20	1.768
		21	3.810
		22	3.353
	02820200000	23	1.817
Paraffin oil test	Simulated wearing treatment	24	3.995
	treatment	25	2.617
	W. C. C. C.	26	5.827
	Mechanical strength+	27	5.526
	Temperature conditioned	28	5.931
	Flow conditioning	: single filter: 95.0 L/m	in

Clause 7.10 Compatibility with skin

(EN 1492001+A) 2009 Clause 8.4.6. 8.5.)

Requirement	Results	Rating
Materials that may come into contact with the wearer's skin shall not be known to be likely to cause initation or any other adverse effect to health.	No irritation or any other adverse effect to health	Pass



Clause 7.11 Flammability

(EN 149/2001+A1/2009 Chase 8.6)

Requirement	Results	Rating
When tested, the particle filtering half mask shall not burn or not to continue to burn	Detail refer to	Pass
for more than 5s after removal from the flame.	Table 7.11	Pass

Table 7.11 Flammability

Condition	Sample No	Result	
	29	Not burn	
As received	36	Not burn	
- 1	31	Not burn	
Temperature conditioned	12	Not been	

Clause 7.12 Carbon dioxide content of the inhalution sir

(EN 149/2001+A1/2009 Clause 8.7)

Requirement	Results	Rating
The carbon disolde content of the inhalation air (dead space) shall not exceed an average of 1.0 % (by volume).	Detail refer to Table 7.12	Pass

T	able 7.12 Carbon dioxde come	nt of the inhalation air	
Condition.	Sample No	1	Result (%)
	33	0.26	0920000000
As received	34	0.29	Mean value: 0.28
	35	0.28	91.28

Clause 7.13 Head harness

(EN 149:2001+A1:2009 Chuse E.4 & E.5)

Requirement	Resum	rorning
The head harness shall be designed so that the porticle filtering, half mask can be		
donned and ramoved casely.		
The head harness shall be adjustable or self-adjusting and shall be sufficiently robust	Comply	Pass
to hold the particle filtering half mask firmly in position and be capable of		15.575530
maintaining total inward leakage requirements for the device.		

Clause 7.14 Field of vision

(EN 149/2001+A1/2009 Clause 8.4)

Requirement	Results	Rating
The field of vision is acceptable if determined so in practical performance tests.	Comply	Pass

Rating

Report No.: JKF20034101 Report date: 2020-12-08

Results

Clause 7.15 Exhalation valve

(EN 149/2001+A1/2009 Clause 8.2 & 8.9 1 & 8.3.4 & 8.8.)

Requirement	Results	Rating
A particle flereing half mask may have one or more eshalation valoe(s), which shall function correctly in allorizations. If an exhalation varie is provided a shall be protected against or be resistant to dirt and mechanical damage and may be abreauted or may include any other device that may be necessary for desputicle flereing flereing the flereing with Task to comply with Task. Eshalation valve(s), if finel, shall continue to operate correctly after a continuous enhabetion flow of 100 bullion over a period of 10 s. When the chalation whe housing as manhed to the facebank, it shall withstand analyte among force of 10 N gendle for the continuous enhabets may be a smally assuming force of 10 N gendle for the continuous continuous many harmonic force of 10 N gendle for the continuous continu	Not applicable (No exhalation valve)	N/A

Clause 7.16 Breathing resistance

(EN 149:2001+A1:2009 Chuse 8.9

- 3	Maximum	permitted resists	ince (mbse)		
Classification	Inhe	lation	Exhalation	Detail refer to	Pass
	30L/min	95L/min	160L/min	Table 7.16	
FFP1	0.6	2.1	3.0		
FFP2	0.7	2.4	3.0		
FFP3	1.0	3.0	3.0		

Requirement

Test item	Condition	Sample No.	A	В	C.	D.	E
		36	0.48	0.47	0.49	0.48	0.48
Inhalation (30 L/min)	As received	37	0.49	0.48	0.47	0.48	0.48
		38	0.49	0.50	0.49	0.49	0.49
	Carlo Market Control	39	0.47	0.48	0.47	0.47	0.48
	Simulated wearing treatment	40	0.48	0.47	0.48	0.48	0.48
	ricatibent	41	0.49	0.48	0.49	0.47	0.48 0.48 0.49 0.48 0.48 0.49 0.44
	TA-40 (1955)	42	0.45	0.45	0.45	0.45	0.44
Temperature conditioned		43	0.45	0.44	0.45	0.44	0.44
	44	0.44	0.41	0.2	0.43	0.43	

		36	1.73	1.74	1.75	1.73	1.74
Inhabition (95 L/min)	As received	37	1.75	1.75	1.76	1.77	1.75
	1000,000,000	38	1.75	1,77	1:76	1.75	1.76
		39	1.72	1.73	1.72	1.71	1.72
	Simulated wearing treatment	40	1.74	1.75	1.74	1,74	1.73
(95 L/mm)	treatment	41	1.74	1.74	1.75	1.76	1.75
	Temperature	42	1.68	1.67	1.67	1.66	1.6
		43	1.67	1.66	1.67	1.66	1.65
		44	1.64	1.63	1.63	1.65	1.68
Eshulation	As received	36	2.28	2.29	2.28	2.30	2.30
		37	2.29	2.30	2.31	2.30	2.30
	150000000000000000000000000000000000000	38	2.30	2.29	2.32	2.32	2.3
	Simulated wearing	39	2.26	2.29	2.27	2.26	2.25
		.40	2.25	2.23	2.24	2.26	2.25
(160 L/min)	treatment	41	2.29	2.27	2.30	2.28	2.27
	w	42	2.14	2.13	2.13	2.14	2.15
	Temperature	43	2.13	2.14	2.15	2.16	2.15
	conditioned	44	2.11	2.10	2.11	2.12	2:12

Sample No. A

Condition

Clause 7.17 Clogging

(EN 149:2001+A1:2009 Clause 8.9 & 8.10)

Requirement	Results	Rating
Requirement Entirely and Sequirement Entirely and Tracks 1.7.1.1 Valved particle (Blering laff marks) (I.7.1.1 Valved particle (Blering laff marks) (Her clegging the histon resistances studient exceed FFF1-5mbar, FFF2-5mbar, FFF3-5mbar at 95 Limit continuous flow. The exhibition resistance shall not exceed ffFF1-5mbar at 96 Limit continuous flow. 1.7.1.2.1 Valved say gardiele (Blering half marks) (Her clegging the histonians and exhibition resistances shall not exceed FFF1-5mbar FFF2-6mbar at 95 Limit continuous flow. 1.7.1.2.1 Valved flow fifter marketist. Val types (wheld and valvedess) of particle (Blering half marks claimed to ment the chapping requirement and lad now much requirements grain in 7-92, for the	Results Optional for single shift device only	Not required

Clause 7.18 Demountable parts (EN 149-2001-A1-2009, Clause 8.2.)

Requirement	Results	Rating
All demountable parts (if fitted) shall be readily connected and secured, where	Comply	Pass

At facing directly ahead; Bt fining vertically upwards; Ct facing vertically downwards; Dt lying on the left side; Et lying on the right side



