

AB Tip İnceleme Sertifikası EU Type-Examination Certificate

Belge No / Certificate No

: 77-20-02

Belgelendirme Tarihi - Bir Sonraki Belge Tarihi /

Certification Date / Certificate Validity Date : 18.11.2020-18.11.2025 Belge Geçerlilik Tarihi / Document Validity Period : 5 yıl / 5 years

Firma Unvanı ve Adresi /

Company Name and Address

: Fit Pharm Technologies GmbH

Benzstraße 13 48619 Heek Nordrhein-

Westfalen Deutschland

Ürün Adı /Modeller / Product Name / Models

Direktifi / Directive

Modülü/Kategori / Module / Category

: FIT F261

: 2016/425 REGULATION

: MNA M-2020-00445

: B MODÜLÜ/ KATEGORİ III

MODULE B / CATEGORY III

Test Rapor No/ları / Test Report No

Ürün Tipi / Product Type:

 EN 149:2001+ A1:2009 Solunumla ilgili koruyucu cihazlar - Parçacıklara karşı koruma amaçlı filtreli yarım maskeler/ Respiratory protective devices - Filtering half masks to protect against particles

Ürünün Malzeme Bilgisi / Product Material Information: FIT F261 model ürünleri kumaş, kulak kayışı, burun klipsi ve filtre katmanı kullanılarak imal edilmiştir./ FIT F261 model products are manufactured using fabric, ear strap, nose clip, filter layer.

Volkan AKIN 18.11.2020 Karar Verici / Approver Okan AKEL 18.11.2020 Şirket Müdürü / General manager









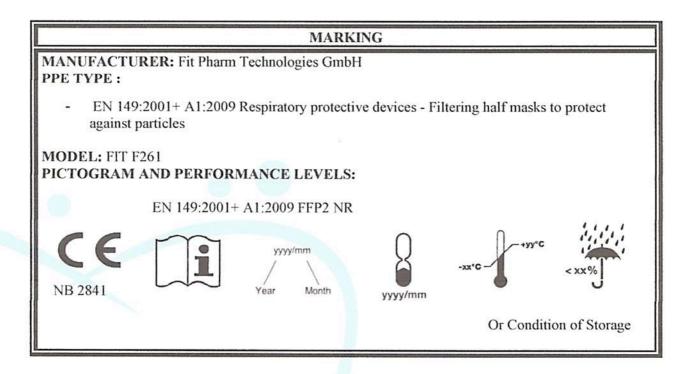
ATTACHMENTS (77-20-02)

To certify the PPE product at Category III level, C2 or D module is accompanied by applying one of the conformity assessment methods along with the EU Type Examination (Module B).

Model: FIT F261

PPE SPECIFICATION	PERFORMANCE LEVELS
Classification	FFP2
Reusable / Single Shift Use	NR

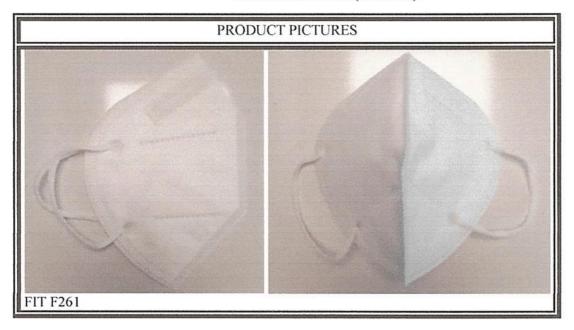
PPE produced as a single unit to fit an individual user, all the necessary instructions for manufacturing such PPE on the basis of the approved basic model:



MNA LABORATORIES SAN. TIC. LTD. ŞTİ declares that the above-mentioned product meets the requirements of the directive according to the EU Directive 2016/425, the safety of the product is covered by the conditions and use specified in this certificate and in the technical file.



ATTACHMENTS (77-20-02)



DOCUMENTS IN THE TECHNICAL

- Basic Health Safety Requirements
- Risk Assessment
- Test Reports
- Technical Report



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

MODÜL C2 - ÜRETİMİN DÂHİLÎ KONTROLÜ VE ÜRÜNÜN RASTGELE ARALIKLARLA DENETİMLİ MUAYENESİNE DAYALI TİPE UYGUNLUK

Belge No / Certificate No

: 77-20-02-01

Belgelendirme Tarihi - Bir Sonraki Belge Tarihi /

Certification Date / Certificate Validity Date

: 29.07.2021-29.07.2022

Belge Geçerlilik Tarihi / Document Validity Period: 1 yıl / 1 year

Firma Unvanı ve Adresi /

Company Name and Address

: Fit Pharm Technologies GmbH

Benzstraße 13 48619 Heek Nordrhein-Westfalen

Deutschland

: FIT F261

Ürün Adı /Modeller / Product Name / Models

Direktifi / Directive

: 2016/425 REGULATION

Modülü/Kategori / Module / Category

: C2 MODÜLÜ/ KATEGORİ III MODULE C2 / CATEGORY III

Test Rapor No/ları / Test Report No : M-2021-01236

Ürün Tipi / Product Type:

 EN 149:2001+ A1:2009 Solunumla ilgili koruyucu cihazlar - Parçacıklara karşı koruma amaçlı filtreli yarım maskeler/ Respiratory protective devices - Filtering half masks to protect against particles

Ürünün Malzeme Bilgisi / Product Material Information: FIT F261 model ürünleri kumaş, elastik kayış, burun klipsi, filtre katmanı kullanılarak imal edilmiştir./ FIT F261 model products are manufactured using fabric, elastic strap, nose clip, filter layer.

Erhan ÜSTÜNEL 29,07,2021 Karar **Ferici** Approver

Okan AKEL 29.07.2021 Şirket Müdürü / General manager









CONFORMITY TO TYPE BASED ON INTERNAL PRODUCTION CONTROL PLUS SUPERVISED PRODUCT

CHECK AT RANDOM INTERVALS (MODULE C2, ANNEX VII) (77-20-02-01)

Report No : 77-20-02-01

Report Date : 29.07.2021

Application No : 77-20-02-01

1. COMPANY INFORMATION:

Fit Pharm Technologies GmbH

Benzstraße 13 48619 Heek Nordrhein-Westfalen Deutschland

Tel: +491605319004

E-mail: s.babitzki@farm-innovation-team.de

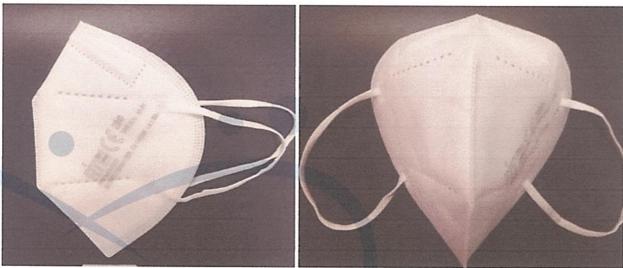
2. PPE INFORMATION:

Disposable and non-sterile half mask made of particulate protection fitler material.

3. PPE TYPE IDENTIFICATION

EN 149:2001+A1:2009 Respiratory protective devices - Filtering half masks to protect against particles -Requirements, testing, marking

4. PPE PICTURES



FIT F261

5. PPE DIMENSIONS:

FIT F261 model has been found to be produced using standard sizes.

6. PPE PRODUCT MATERIAL INFORMATION:

The mask is made of elastic strap, nonwoven fabric on the outer and inner layers and fitler material on the middle layer.

7. ESSENTIAL HEALTH AND SAFETY REQUIREMENTS

- A visual inspection was made according to EN 149:2001 +A1:2009 for ergonomics.
- Protection levels and degrees are defined by the manufacturer.
- Suitable construction materials were determined by visual inspection according to EN 149:2001 +A1:2009.



CONFORMITY TO TYPE BASED ON INTERNAL PRODUCTION CONTROL PLUS SUPERVISED PRODUCT

Notified Body Number: 2841

CHECK AT RANDOM INTERVALS (MODULE C2, ANNEX VII) (77-20-02-01)

8. ANALYSIS AND EVALUATIONS:

EN 149:2001 +A1:2009

TESTS	PARAMETER	PERFORMANCE LEVELS			RESULTS	PERFORMANC E LEVELS	EVALUATION N
		FFP1	FFP2	FFP3			
Part 7.3 Visual inspection	Shall also the marking supplied by the manuf			mation	Appropriate	-	PASS
Banned Azo Dyes	< 30 mg/kg				Not applicable	-	Not applicable
Part 7.4 Packaging	for sale packaged in are protected agains	e filtering half mask shall be offered e packaged in such a way that they otected against mechanical damage ntamination before use.			Appropriate	-	PASS
Part 7.5 Material	When conditioned in 8.3.2 the particle filte collapse.				Appropriate	-	PASS
Part 7.6 Cleaning and disinfecting	After cleaning and disinfecting the re-usable particle filtering half mask shall satisfy the penetration requirement of the relevant class.			sfy the	Not applicable	-	Not applicable
Part 7.7 Practical performance		nents should be made by garding any of the criteria			Appropriate	-	PASS
Part 7.8 Finish of parts		ts of the device likely to come into			Appropriate	-	PASS

TESTS PARAME	PARAMETER	PERFORMANCE LEVELS		RESULTS	PERFORMANCE LEVELS	EVALUATION	
		FFP1	FFP2	FFP3			
Part 7.9.1 Total inward leakage	At least 46 out of the 50 individual exercise result	<25	<11	<5	See the table below	FFP2	PASS
	At least 8 out of the 10 individual wearer arithmetic means	<22	<8	<2	See the table below	FFP2	PASS

Total Inward Leakage (%)											
	Exercise 1	Exercise 2	Exercise 3	Exercise 4	Exercise 5	Average					
Subject 1 (As received)	7,0	6,0	5,2	7,2	5,5	6,2					
Subject 2 (As received)	6,7	4,3	4,8	5,5	5,4	5,3					
Subject 3 (As received)	6,4	7,6	4,9	7,2	7,6	6,7					
Subject 4 (As received)	7,8	8,0	6,9	7,3	7,6	7,5					
Subject 5 (As received)	6,1	7,3	6,7	4,4	6,2	6,1					
Subject 6 (After temperature conditioning)	7,6	8,3	7,8	8,5	7,7	8,0					



CONFORMITY TO TYPE BASED ON INTERNAL PRODUCTION CONTROL PLUS SUPERVISED PRODUCT

Notified Body Number: 2841

CHECK AT RANDOM INTERVALS

(MODULE C2, ANNEX VII) (77-20-02-01)

Subject 7 (After temperature conditioning)	6,4	6,6	6,3	5,3	6,2	6,2
Subject 8 (After temperature conditioning)	7,7	7,9	7,6	7,8	8,0	7,8
Subject 9 (After temperature conditioning)	5,1	7,6	7,6	7,2	7,8	7,1
Subject 10 (After temperature conditioning)	3,8	4,1	3,5	4,5	3,2	3,8

Subject facial dimensions

Subject	Face Length (mm)	Face Width (mm)	Face Depth (mm)	Mouth Width (mm)
1	133	132	132	65
2	125	144	116	67
3	126	135	124	75
4	123	133	134	74
5	117	135	122	73
6	122	142	133	66
7	113	132	114	75
8	135	123	123	65
9	122	135	133	74
10	135	142	125	83

TESTS PA	PARAMETER PERFORMANCE RESULT LEVELS			7 Lin Chivirute					RESULTS	PERFORMANCE LEVELS	EVALUATION
		FFP1	FFP2	FFP3							
Part 7.9.2 Penetration of filter	Sodium chloride, 95 L/min %, max	% 20	% 6	%1	See the table below	FFP2	PASS				
material	Paraffin oil, 95 L/min %, max	% 20	% 6	%1	See the table below	FFP2	PASS				

Penetration of filter material	Sodium Chloride (%)	Paraffin Oil (%)	
As received	1,4	1,5	
As received	1,3	1,5	
As received	1,4	1,6	
After the simulated wearing treatment	1,4	1,4	
After the simulated wearing treatment	1,6	1,5	
After the simulated wearing treatment	1,5	1,7	
Mechanical strength and temperature conditioning (120 mg)	4,7	4,8	
Mechanical strength and temperature conditioning (120 mg)	5,3	5,6	
Mechanical strength and temperature conditioning (120 mg)	4,6	4,8	

TESTS PARAMETER	PARAMETER	PERFO LEVELS	RMAN	CE	RESULTS	PERFORMANCE LEVELS	EVALUATION	
	FFP1	FFP2	FFP3					
Part 7.10 Compatibility with skin	Materials shall no cause irritation or health				Appropriate	-	PASS	
Part 7.11 Flammibility	Mask shall not bu for more than 5 s	n or not to continue to burn			Flame not seen	- 1 /	PASS	
Part 7.12	Shall not exceed a	n average of % 1			0,80	- \	PASS	



CONFORMITY TO TYPE BASED ON INTERNAL PRODUCTION CONTROL PLUS SUPERVISED PRODUCT

CHECK AT RANDOM INTERVALS

(MODULE C2,	VIVIER VIII	(77-20-02-01)
(IVIODULE CZ,	AININEA VIII)	(//-20-02-01)

Carbondioxide		0,84		
content of the inhalation air		0,85		
Part 7.13 Head harness	It can be donned and removed easily	Appropriate	-	PASS
Part 7.14 Field of vision	The field of vision shall acceptable in practical performance test.	Appropriate	-	PASS
Part 7.15 Exhalation valve(s)	It shall withstand axially a tensile force of 10 N apply for 10 s. If fitted, shall continue to operate correctly after a continuous exhalation flow of 300 L/min over a period of 30 s.	Not applicable	-	Not applicable

TESTS PARAMETER	PARAMETER	PERFORMANCE LEVELS			RESULTS	PERFORMANCE	EVALUATION
		FFP1	FFP2	FFP3		LEVELS	
Part 7.16 Breathing	Inhalation 30L/min	0,6 mbar	0,7 mbar	1,0 mbar	See the table below	FFP2	PASS
Resistance	Inhalation 95L/min	2,1 mbar	2,4 mbar	3,0 mbar	See the table below	FFP2	PASS
	Exhalation 160L/min	3,0 mbar	3,0 mbar	3,0 mbar	See the table below	FFP2	PASS

Breathing Resistance (mbar)	Inhalation 30L/min	Inhalation 95L/min	
As received	0,4	1,5	
As received	0,4	1,5	
As received	0,3	1,4	
After temperature conditioning	0,4	1,4	
After temperature conditioning	0,3	1,4	
After temperature conditioning	0,3	1,5	
After the simulated wearing treatment	0,4	1,4	
After the simulated wearing treatment	0,3	1,4	
After the simulated wearing treatment	0,3	1,4	
After the flow conditioning	-	-	
After the flow conditioning	-	-	
After the flow conditioning	-	-	

Breathing Resistance 160L/min (mbar)	Facing directly ahead	Facing vertically upwards	Facing vertically downwards	Lying on the left side	Lying on the right side
As received	2,4	2,4	2,4	2,5	2,4
As received	2,5	2,4	2,4	2,5	2,4
As received	2,5	2,4	2,5	2,5	2,5
After temperature conditioning	2,5	2,5	2,4	2,4	2,4
After temperature conditioning	2,4	2,4	2,4	2,4	2,5
After temperature conditioning	2,4	2,4	2,5	2,4	2,4
After the simulated wearing treatment	2,5	2,4	2,4	2,4	2,5
After the simulated wearing treatment	2,4	2,4	2,5	2,4	2,4
After the simulated wearing treatment	2,4	2,4	2,4	2,4	2,4



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CHECK AT RANDOM INTERVALS

(MODULE C2, ANNEX VII) (77-20-02-01)

After the flow conditioning	-	-	-	1-	-
After the flow conditioning	-	-	-	-	-
After the flow conditioning	-	-	-	-	-

TESTS	PARAMETER	PERFORMANCE LEVELS		RESULTS	PERFORMANCE LEVELS	EVALUATION	
		FFP1	FFP2	FFP3			
Part 7.17	After clogging the	4	5	7	Not applicable	-	Not applicable
Clogging	inhalation	mbar	mbar	mbar			
	resistances shall						
	not exceed.						
	(valved)						
	The exhalation resistance shall not exceed				Not applicable	-	Not applicable
	3 mbar at 160 L/ min continuous flow.			3553453		****	
	(valved)						
	After clogging the	3	4	5	Not applicable	-	Not applicable
	inhalation and	mbar	mbar	mbar			**************************************
	exhalation						
	resistances shall						
	not exceed.						
	(valveless)						
Part 7.18	All demountable parts (if fitted) shall be			nall be	Not applicable		Not applicable
Demountable	readily connected	and s	ecured	were	76 67		
part	possible by hand.						

9. DECISION

Analysis and examinations FIT F261 model coded personal protective equipment; Respiratory Protective Devices EN 149:2001 +A1:2009- Filtered Half Masks for Protection Against Particles - Properties, Experiments and Marking standards are evaluated. The homogeneity of the production was monitored at the performance levels determined as a result of the technical evaluations made within the scope of MODULE C2.

10. ATTACHMENTS

- Basic Health Safety Requirements
- Risk Assessment
- Test Reports (M-2021-01236)
- User Instruction

CONTROLLER : ERHAN ÜSTÜNEL

SIGNATURE :

DATE : 29.07.2021