







**mna**  
LABORATUVARLARI

Notified Body Number: 2841

**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**

: 45-21-02-02

**Belgelendirme Tarihi - Bir Sonraki Belge Tarihi /**

**Certification Date / Certificate Validity Date**

: 15.09.2022-15.09.2023

**Belge Geçerlilik Tarihi / Document Validity Period**

süresi boyunca geçerlidir./ Valid for the duration of the certificate with code 45080333 (15.09.2022-15.09.2023)

**Firma Unvanı ve Adresi /**

**Company Name and Address**

: MY TİCARET VE MEDİKAL A.Ş.

Ömerli Mah. General Şükrü Koraltı Cad. No: 33  
Arnavutköy/ İSTANBUL

: MUMU / MM.P2.19

: 2016/425 REGULATION

: C2 MODÜLÜ/ KATEGORİ III  
MODULE C2 / CATEGORY III

**Marka / Model / Brand / Model**

**Direktifi / Directive**

**Modülü/Kategori / Module / Category**

**Teknik Değerlendirme Rapor No/**

**Technical Evaluation Report No**

**Ürün Tipi / Product Type:**

: 45080333

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

**Ürünün Malzeme Bilgisi / Product Material Information:** MUMU / MM.P2.19 model ürünleri kumaş, elastik kayış, burun klipsi ve filtre katmanı kullanılarak imal edilmiştir./ MUMU / MM.P2.19 model products are manufactured using fabric, elastic strap, nose clip, filter layer.

*Volkan AKIN*

15.09.2022

*Karar Verici / Approver*

*Okan AKEL*

15.09.2022

*Sirket Müdürü / General manager*



MNA Laboratuvarları San. Tic.Ltd .Şti

Adres: Küçükbağkalköy Mahallesi Yenidoğan Cad.No:21 Ataşehir/ İstanbul

Tel: 0216 574 07 08 Faks: 0216 575 13 31 [www.mnalab.com](http://www.mnalab.com)

# AB Tip İnceleme Sertifikası EU Type-Examination Certificate

**Belge No / Certificate No**

: 45-21-02

**Belgelendirme Tarihi - Bir Sonraki Belge Tarihi /**

**Certification Date / Certificate Validity Date** : 21.12.2021-28.04.2026.

**Belge Geçerlilik Tarihi / Document Validity Period** : 229-21-01 (28.04.2021-28.04.2026) kodlu sertifikanın süresi boyunca geçerlidir./ *Valid for the duration of the certificate with code 229-21-01 (28.04.2021-28.04.2026)*

**Firma Unvanı ve Adresi /**

**Company Name and Address**

: MY TİCARET VE MEDİKAL A.Ş.

Ömerli Mah. General Sükrü Koraltı Cad. No: 33 Arnavutköy/  
İSTANBUL

: MUMU / MM.P2.19

**Ürün Adı /Modeller / Product Name / Models**

: 2016/425 REGULATION

**Modülü/Kategori / Module / Category**

: B MODÜLÜ/ KATEGORİ III  
*MODULE B / CATEGORY III*

**Teknik Değerlendirme Raporu /**

**Technical Evaluation Report**

: Sertifika Kodu: 229-21-01 (28.04.2021)

**Ürün Tipi / Product Type:**

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

**Ürünün Malzeme Bilgisi / Product Material Information:** MUMU / MM.P2.19 model ürünleri kaplamalı kumaş kullanımlararak imal edilmiştir./ *MUMU /MM.P2.19 model products are manufactured using coated fabric.*

**Volkan AKIN**

**21.12.2021**

**Karar Verici / Approver**

**Okan AKEL**

**21.12.2021**

**Şirket Müdürü / General manager**



MNA Laboratuvarları San. Tic.Ltd .Şti

Adres: Küçükbağkalköy Mahallesi Yenidoğan Cad.No:21 Ataşehir/ İstanbul  
Tel: 0216 574 07 08 Faks: 0216 575 13 31 [www.mnalab.com](http://www.mnalab.com)

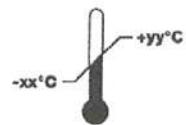
**ATTACHMENTS (45-21-02)**

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).

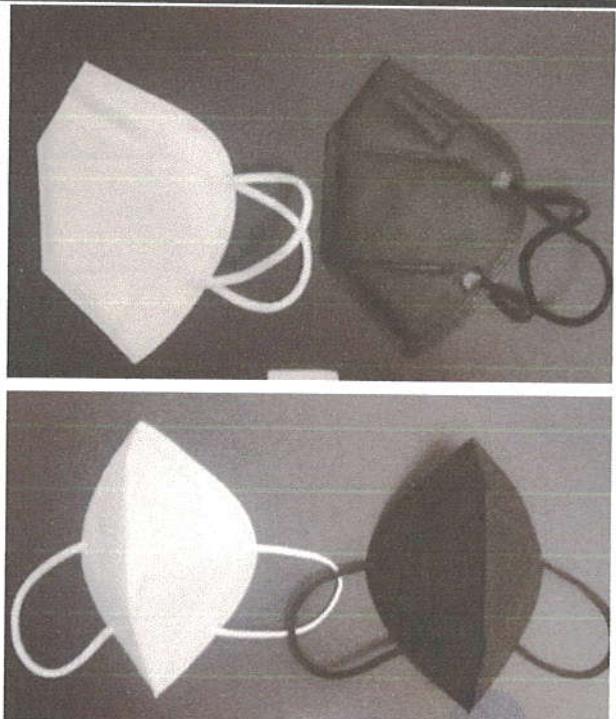
**Model :** MUMU / MM.P2.19

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:

MARKING	
<b>MANUFACTURER:</b> MY TİCARET VE MEDİKAL A.Ş.	
<b>PPE TYPE :</b>	
<ul style="list-style-type: none"> <li>- EN 149:2001+ A1:2009 Respiratory protective devices - Filtering half masks to protect against particles</li> </ul>	
<b>MODEL:</b> MUMU / MM.P2.19	
<b>PICTOGRAM AND PERFORMANCE LEVELS:</b>	
EN 149:2001+ A1:2009 FFP2 NR     	
NB 2841	
Or Condition of Storage	

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 (45-21-02)****PRODUCT PICTURES**

MUMU / MM.P2.19

**DOCUMENTS IN THE TECHNICAL FILE**

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

MNA Laboratuvarları San. Tic.Ltd .Şti

Adres: Küçükbakkalköy Mahallesi Yenidoğan Cad.No:21 Ataşehir/ İstanbul  
Tel: 0216 574 07 08 Faks: 0216 575 13 31 [www.mnalab.com](http://www.mnalab.com)



**MNA LABORATORIES**  
**TEST REPORT**

Report No: M-2021-01234	Date: 04.08.2021	Page 1 of 4	Rev:
<b>Purpose of Analysis</b>	: SPECIAL REQUEST	<b>Brand</b>	: -
<b>Sample Type</b>	: PROTECTIVE MASK	<b>Model</b>	: HP2.01 FFP2
<b>Sample Send Org.</b>	: HONNES SAĞLIK VE END.ÜRÜNLERİ A.Ş	<b>Sampler</b>	: CUSTOMER
<b>Manufacturer Name</b>	: HONNES SAĞLIK VE END.ÜRÜNLERİ A.Ş		
<b>Analysis Date</b>	: 30.06.2021		
<b>Sample Quantity</b>	: 80 pieces		
<b>Other informations</b>	: BLACK AND WHITE. MODULE C2.		

TESTS	LIMIT	RESULTS
<b>EN ISO 14362-1 / EN ISO 17234-1 Banned Azo Dyes</b>	< 30 mg/kg	< 5 mg/kg
<b>EN 149+ A1 Part 7.3 Visual inspection</b>	Shall also the marking and the information supplied by the manufacturer	Appropriate
<b>EN 149+ A1 Part 7.4 Packaging</b>	Particle filtering half mask shall be offered for sale packaged in such a way that they are protected against mechanical damage and contamination before use.	Appropriate
<b>EN 149+ A1 Part 7.5 Material</b>	When conditioned in accordance 8.3.1 & 8.3.2 the particle filter half mask shall not collapse.	Appropriate
<b>EN 149+ A1 Part 7.6 Cleaning and disinfecting</b>	After cleaning and disinfecting the re-usable particle filtering half mask shall satisfy the penetration requirement of the relevant class.	Not applicable
<b>EN 149+ A1 Part 7.7 Practical performance</b>	No negative comments should be made by the test subject regarding any of the criteria evaluated.	Appropriate
<b>EN 149+ A1 Part 7.8 Finish of parts</b>	Parts of the device likely to come into contact with the wearer shall have no sharp edge or burrs.	Appropriate
<b>EN 149+ A1 Part 7.9.1 Total inward leakage</b>	At least 46 out of the 50 individual exercise result: FFP1<25 FFP2<11 FFP3<5 At least 8 out of the 10 individual wearer arithmetic means: FFP1<22 FFP2<8 FFP3<2	See below table

Total Inward Leakage (%) EN 149+ A1 Part 7.9.1						
	Exercise 1	Exercise 2	Exercise 3	Exercise 4	Exercise 5	Average
Subject 1 (As received)	6.7	5.7	5.7	6.9	5.2	6.0
Subject 2 (As received)	6.4	5.7	6.9	5.2	5.1	5.9
Subject 3 (As received)	6.1	6.2	4.6	6.9	7.3	6.2
Subject 4 (As received)	7.5	6.9	6.6	7.0	7.3	7.1
Subject 5 (As received)	5.8	7.0	6.4	4.1	5.9	5.8
Subject 6 (After temperature conditioning)	6.3	6.2	6.3	6.4	7.4	6.5
Subject 7 (After temperature conditioning)	6.1	6.3	6.0	5.0	5.9	5.9
Subject 8 (After temperature conditioning)	7.4	6.4	7.3	7.5	5.9	6.9
Subject 9 (After temperature conditioning)	5.7	6.2	7.3	6.9	6.2	6.5
Subject 10 (After temperature conditioning)	5.8	6.2	6.3	6.5	6.6	6.3



**MNA LABORATORIES**  
**TEST REPORT**

Report No: M-2021-01234	Date: 04.08.2021	Page 2 of 4	Rev:
Purpose of Analysis	: SPECIAL REQUEST	Brand	:-
Sample Type	: PROTECTIVE MASK	Model	: HP2.01FFP2
Sample Send Org.	: HONNES SAĞLIK VE END.ÜRÜNLERİ A.Ş	Sampler	: CUSTOMER
Manufacturer Name	: HONNES SAĞLIK VE END.ÜRÜNLERİ A.Ş		
Analysis Date	: 30.06.2021		
Sample Quantity	: 80 pieces		
Other informations	: BLACK AND WHITE. MODULE C2.		

**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	LIMIT	RESULTS
EN 149+ A1 Part 7.9.2 Penetration of filter material	Sodium chloride, 95 L/min% FFP1≤20 FFP2≤6 FFP3≤1 Paraffin oil, 95 L/min% FFP1≤20 FFP2≤6 FFP3≤1	See below table

Penetration of filter material EN 149+ A1 Part 7.9.2	Sodium Chloride (%)	Paraffin Oil (%)
As received	4.4	4.4
As received	4.0	4.4
As received	4.3	4.3
After the simulated wearing treatment	4.3	4.0
After the simulated wearing treatment	4.2	4.7
After the simulated wearing treatment	4.6	4.8
Mechanical strength and temperature conditioning (120 mg)	5.5	5.3
Mechanical strength and temperature conditioning (120 mg)	5.4	5.6
Mechanical strength and temperature conditioning (120 mg)	5.5	5.4

TESTS	LIMIT	RESULTS
EN 149+ A1 Part 7.10 Compatibility with skin	Materials shall not be known to be likely to cause irritation or any other adverse effect to health	Appropriate
EN 149+ A1 Part 7.11 Flammability	Mask shall not burn or not to continue to burn for more than 5 s	Flame not seen
EN 149+ A1 Part 7.12 Carbon dioxide content of the inhalation air	Shall not exceed an average of % 1	0,66 0,77 0,75
EN 149+ A1 Part 7.13 Head harness	It can be donned and removed easily	Appropriate



**MNA LABORATORIES**  
**TEST REPORT**

Report No: M-2021-01234	Date: 04.08.2021	Page 3 of 4	Rev:
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<b>Sample Type</b>	: PROTECTIVE MASK	<b>Model</b>	: HP2.01FFP2
<b>Sample Send Org.</b>	: HONNES SAĞLIK VE END.ÜRÜNLERİ A.Ş	<b>Sampler</b>	: CUSTOMER
<b>Manufacturer Name</b>	: HONNES SAĞLIK VE END.ÜRÜNLERİ A.Ş		
<b>Analysis Date</b>	: 30.06.2021		
<b>Sample Quantity</b>	: 80 pieces		
<b>Other informations</b>	: BLACK AND WHITE. MODULE C2.		

<b>EN 149+ A1 Part 7.14</b> <b>Field of vision</b>	The field of vision shall acceptable in practical performance test.	Appropriate
<b>EN 149+ A1 Part 7.15</b> <b>Exhalation valve(s)</b>	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
<b>EN 149+ A1 Part 7.16</b> <b>Breathing Resistance</b>	Inhalation 30L/min FFP1≤0,6mbar FFP2≤0,7mbar FFP3≤1,0mbar Inhalation 95L/min FFP1≤2,1mbar FFP2≤2,4mbar FFP3≤3,0mbar Exhalation 160L/min FFP1≤3,0mbar FFP2≤3,0mbar FFP3≤3,0mbar	See below table

<b>Breathing Resistance (mbar)</b> <b>EN 149+ A1 Part 7.16</b>	<b>Inhalation 30L/min (mbar)</b>	<b>Inhalation 95L/min (mbar)</b>
As received	0.5	1.7
As received	0.5	1.7
As received	0.4	1.6
After temperature conditioning	0.5	1.7
After temperature conditioning	0.5	1.6
After temperature conditioning	0.4	1.6
After the simulated wearing treatment	0.4	1.7
After the simulated wearing treatment	0.4	1.7
After the simulated wearing treatment	0.5	1.7
After the flow conditioning	-	-
After the flow conditioning	-	-
After the flow conditioning	-	-

<b>Breathing Resistance 160L/min (mbar)</b> <b>EN 149+ A1 Part 7.16</b>	<b>Facing directly ahead</b>	<b>Facing vertically upwards</b>	<b>Facing vertically downwards</b>	<b>Lying on the left side</b>	<b>Lying on the right side</b>
As received	2,7	2,8	2,7	2,8	2,7
As received	2,8	2,7	2,8	2,7	2,8
As received	2,8	2,8	2,7	2,7	2,7
After temperature conditioning	2,8	2,7	2,7	2,8	2,8
After temperature conditioning	2,8	2,7	2,8	2,7	2,7
After temperature conditioning	2,7	2,8	2,7	2,7	2,7
After the simulated wearing treatment	2,7	2,7	2,7	2,7	2,8
After the simulated wearing treatment	2,8	2,7	2,8	2,7	2,8
After the simulated wearing treatment	2,7	2,8	2,7	2,8	2,7
After the flow conditioning	-	-	-	-	-
After the flow conditioning	-	-	-	-	-
After the flow conditioning	-	-	-	-	-



MNA LABORATORIES  
TEST REPORT

Report No: M-2021-01234	Date: 04.08.2021	Page 4 of 4	Rev:
Purpose of Analysis	: SPECIAL REQUEST	Brand	: -
Sample Type	: PROTECTIVE MASK	Model	: HP2.01 FFP2
Sample Send Org.	: HONNES SAĞLIK VE END.ÜRÜNLERİ A.Ş	Sampler	: CUSTOMER
Manufacturer Name	: HONNES SAĞLIK VE END.ÜRÜNLERİ A.Ş		
Analysis Date	: 30.06.2021		
Sample Quantity	: 80 pieces		
Other informations	: BLACK AND WHITE. MODULE C2.		

Operating as an experimental laboratory, MNA Laboratories have been accredited by TURKAK with AB-1183-T and TS\_EN\_ISO / IEC\_17025: 2017 standard. Turkish Accreditation Agency (TÜRKAK) signed a multilateral agreement with the European Accreditation Association (EA) on the recognition of test reports and a mutual recognition agreement with the International Laboratory Accreditation Association (ILAC).

\* Analysis is under accreditation.

Note :

1. No part of this analysis report can be used alone or separately, and may not be partially copied or reproduced, used to third parties and as a means of advertising without the written permission of the laboratory.
  2. Analysis results are valid for the above mentioned sample sent by MNA Laboratory company / institution / person. It may not represent the whole.
  3. Unsigned and unsealed reports are invalid.
  4. This analysis report cannot be used in judicial-administrative procedures and for advertising purposes.
  5. Results are valid for the sample as received.
  6. The decision rule is the rule that determines how measurement uncertainty is taken into account when specifying the PASS density to a specified specification. According to the TLM-052 Decision Rule Implementation instruction, the Decision Rule Implementation Method selected in agreement with CUSTOMER is clearly stated in the report.
  7. Limit Values are determined by taking from analysis methods.
  8. The laboratory is not responsible if the information provided by the CUSTOMER affects the validity of the results.
  9. Test and / or measurement results, expanded measurement uncertainties (if any) and test methods are given in the following pages, which are the supplementary part of this certificate.
  10. Water Repellency Determination Hydrostatic Pressure Determination TS ISO 811 (Hydrostatic Pressure Tester E / N: 53) Analysis, Seam Strength EN ISO 13965-2 (Strength Test Device E / N: 50) Analysis and resistance to liquid chemical permeation TS EN 659 - A1 Part 3.18 (Liquid Chemical Transfer Device E / N: 107) Analysis is carried out in the conditioning room and ISO 139 PART 3.2 conditions ( $23 \pm 2^\circ\text{C}$  temperature and  $50 \pm 4\%$  relative humidity) are applied for ambient conditions.
  11. List of phthalates analyzed is below.
- Di-iso-nonyl phthalate (DINP), CAS number: 28553-12-0 or 68515-48-0  
Di-(2-ethylhexyl) phthalate (DEHP), CAS number: 117-81-7  
Di-n-octyl phthalate (DNOP), CAS number: 117-84-0  
Di-iso-decyl phthalate (DIDP), CAS number: 26761-40-0 or 68515-49-1  
Butyl benzyl phthalate (BBP), CAS number: 85-68-7  
Di-butyl phthalate (DBP), CAS number: 84-74-2

Selin GERGİN  
Sampling and Reporting  
Officer

Erhan ÜSTÜNEL  
PPE Laboratory Responsible

Confirmed  
04.08.2021  
Volkan AKIN  
Laboratory Manager



### **EC DECLARATION OF CONFORMITY**

<b>Manufacturer</b>	: My Ticaret ve Medikal A.Ş
<b>Manufacturer's Address</b>	: Ömerli Mahallesi General Şükrü Koraltı Caddesi No: 33 Arnavutköy/İstanbul/Turkey
<b>Phone</b>	: +90 212 438 20 64
<b>E-mail</b>	: info@mymedikal.com.tr
<b>Product Description (PPER):</b>	: Filtering Half Masks to Protect Against Particles. FFP2 NR
<b>Product Name / Models</b>	: MUMU /MM. P2.19
<b>Product Type</b>	: EN 149:2001 +A1:2009 Respiratory Protective Devices- Filtering Half Masks to Protect Against Particles.
<b>Product Material Information</b>	: MUMU / MM. P2.19 model products are manufactured using coated fabric.
<b>Device Classification (PPER)</b>	: Category III (Type B and Type C)
<b>EU Type-Examination Certificate Number (PPER) For B Module</b>	: 45-21-02
<b>C2 Module Certificate Number (PPER):</b>	: 45-21-02-01
<b>Standard Reference (PPER)</b>	: EN 149:2001 +A1:2009

We, My Ticaret ve Medikal A.Ş. herewith declared that above mentioned device:

- Is in conformity with the provisions of Regulation (EU) 2016/425 on personnel protective equipment.
- Is subject to the conformity assessment procedure Module C2 Set out in Annex VII of Regulation (EU) 2016/425, under the surveillance of the notified body MNA Laboratuvarları San. Tic.Ltd.Şti , Küçükbakkalköy Mah. Yenidoğan Cad. No:21 Ataşehir TURKEY (Notified Body number 2841)

This EU declaration of conformity is issued under the sole responsibility of the manufacturer, My Ticaret ve Medikal A.Ş



MY TICARET VE MEDIKAL A.Ş

Ömerli Mahallesi General Şükriü Koraltı Caddesi No:33 Arnavutköy/İstanbul/Turkey

EU DoC Issuance Date

: 05.01.2022

Signed for and Behalf of My Ticaret ve Medikal A.Ş

Name: MURAT YILDIZ

Designation: General Manager

MY TICARET VE  
MEDIKAL A.Ş. / İŞLETİM SİRKETİ  
Ömerli Mah. Gen. Şükriü Koraltı Cad  
No: 33 Arnavutköy / İSTANBUL  
Buyukçekmece V.D: 626 040 4605  
Tel: 0212 438 20 64 Fax: 0212 438 20 65  
[www.my-medikal.com.tr](http://www.my-medikal.com.tr)



MY Medikal

## MY TICARET VE MEDIKAL A.S.

Ömerli Mahallesi General Şükrü Koraltı Caddesi No:33 Arnavutkoy -Istanbul Turkey

Tel: +902124382064 Fax: +902124382065

Website: [www.mymedikal.com.tr](http://www.mymedikal.com.tr).

### DECLARATION OF CONFORMITY

We, MY TICARET VE MEDIKAL A.S. herewith declare under our sole responsibility that below mentioned product(s) is fully compliance with the provisions of Regulations (EU) 2016/425 in Personal Protective Equipment.

Manufacturer Address	MY TICARET VE MEDIKAL A.S. Ömerli mah General Şükrü Koraltı Cd no:33, 3455 Arnavutkoy/Istanbul, Turkey
Single Registration Number (SRN)	TR-MF-000018372
Brand	Mumu/MM.P2.19
Product Description	Filtering Half Masks to Protect Against Particles. FFP2 NR
Intended Purpose	This is designed to be used as half-face masks that filter single-shift particles for protection against solid and liquid aerosols.
Product Material Information	MUMU/ MM.P2.19 model products are manufactured using coated fabric
Device Classification (PPER)	Category III (Type B and Type C)
Conformity Assessment Procedure (PPER)	Annex VII (Module C2)
EU Type-Examination Certificate (PPER)/ Module B	45-21-02
Module C2 Certificate Number (PPER)	45-21-02-01
EU Type- Examination Certificate Issued by (PPER)	MNA Laboratuvarları San. Tic.Ltd.Sti, Kucukbakkalkoy Mah. Yenidogan Cad. No:21 Atasehir TURKEY
Notified Body Number (PPER)	2841
Applicable Standard (PPER)	EN 149:2001+A1:2009

#### Authorized Signatory:

Approver : MURAT YILDIZ  
Title : General Manager/CEO  
Signature :   
: **MY TICARET VE  
MEDIKAL ANONİM ŞİRKETİ  
Ömerli Mah. General Şükrü Koraltı Cad  
No:33 Arnavutköy/İSTANBUL  
Büyükdere Mah. V.D. 526 040 4605  
Tel: 0212 438 20 65 Fax: 0212 438 20 65  
E-mail: [mymedikal.com](mailto:mymedikal.com)**  
Approval Date : 30.06.2022  
Place of Approval : Istanbul, Turkey

**DECLARATION LETTER ON THE COMPONENTS AND MATERIALS OF FFP2 MASK**

Date: 16 September 2022

**To Whom It May Concern,**

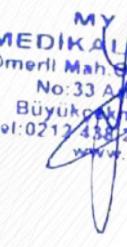
We, **My Ticaret Ve Medikal A.S.**, hereby declare the following components and materials of FFP2 mask (MM.P2.19).

COMPONENT	MATERIAL	CLASS
Outer Layer	Spunbond	55 g/m <sup>2</sup>
First Layer	Meltblown	25 g/m <sup>2</sup>
Second Layer	Meltblown	25 g/m <sup>2</sup>
Third Layer	Hot air	50 g/m <sup>2</sup>
Outer Layer	Spunbond	30 g/m <sup>2</sup>
Ear Loop	75% nylon and 25% spandex	Product Size = 19 cm Product Width = 4 mm Product Thickness = 2 mm Stretch Rate = ½
Nose wire	Plastic with double iron inside	Wire Size = 9 cm Wire Width = 5 mm

According to the components and parts mentioned above, this product does not contain latex.

For and on behalf of

**My Ticaret Ve Medikal A.S.**

<b>Approved by:</b> Name: MURAT YILDIZ Position: General Manager/CEO Date: 16.09.2022	 <p>MY TİCARET VE MEDİKAL ANONİM ŞİRKETİ Ömerli Mah. General Sükrü Koraltı Cad. No:33 Arnavutköy/İSTANBUL Büyükoğlumce V.D.626 040 4605 Tel:0212 438 20 64 Fax:0212 438 20 65 <a href="http://www.mymedikal.com">www.mymedikal.com</a></p>
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