

ATTESTATION OF CONFORMITY

Certificate Nr: MDD-267

In conformance to the European Economic Commission 93/42/EEC Medical Devices Directive on harmonisation of laws, regulations and administrative documentation of Member States on Medical Devices and European Economic Commission directive 93/68/EEC amending Medical Devices Directive dated 22 July 1993,

the products manufactured by

**EN ECZA DEPOSU İLAÇ MEDİKAL ÖZEL SAĞLIK HİZMETLERİ
İNŞAAT TAAHHÜT TİCARET A.Ş.**

at the following address

Karaduvar Mahallesi Serbest Bölge 7.Caddesi No:21 33020 Akdeniz / MERSİN / TURKEY

EN 14683:2019+AC:2019 Medical Face Masks

Brand Name : ENMED

Model : ENM-313

Type IIR

are tested according to the following initial type tests by the manufacturer

Technical standard EN 14683:2019+AC:2019 Medical face masks - Requirements and test methods

For the assessment of conformity, the following documents were also applied to:

Results of laboratory tests Ekoteks Laboratuvar Testing Laboratory Bacterial Filtration Efficiency, Microbial Cleanliness, Differential Pressure and Splash Resistance Pressure tests.

UNIVERSAL CERTIFICATION has evaluated production, design, intended use, risk evaluation according to safety purpose, product itself and add-on components (if exists) and product technical drawings of the medical face masks manufactured and designed for use during the medical operations or similar medical situations with same requirements which require restriction of infectious materials to be spread to patients. With this certificate, it is approved that the product fulfils all essential requirements and the related rules of 93/42/EEC Medical Devices Directive (MDD) Class I are applied. The information on the packaging for the above listed products covers the necessary information stated in Annex I, §13, of the Medical Devices Directive (93/42/EEC) or Annex I, §23, of the Medical Device Regulation (EU) 2017/745. This information includes; reference to EN 14683 standard, type of mask (as indicated in Table 1) and other relevant information given in EN ISO 15223-1:2016 and EN 1041:2008+A1:2013. It is considered to be suitable to attach a CE mark, as seen below, on your products in accordance with the information given in this certificate with publishing an EU Declaration of Conformity.

This certificate is issued on 25/09/2020 and valid until 24/09/2021 with the conditions that no change has been made with the product references and no change in the production process or not suspended or withdrawn for any reason.

ISTANBUL -25/09/2020



Verify the validity with the QR Code

Suat KAÇMAZ
UNIVERSAL CERTIFICATION
Director

EU DECLARATION OF CONFORMITY

MANUFACTURER

EN ECZA DEPOSU İLAÇ MEDİKAL ÖZEL SAĞLIK HİZMETLERİ İNŞAAT TAAHHÜT
TİCARET A.Ş.

Karaduvar Mahallesi Serbest Bölge 7.Caddesi No:21 33020 Akdeniz / MERSİN / TURKEY

PRODUCT DESCRIPTION

Layered and molded medical device classified in the Class I - Medical Device to be used as protection against inhalation of viruses, bacteria, other microorganisms, allergens from the environment

Brand Name : ENMED

Model : ENM-313

Type IIR

The Producer / the Manufacturer declares on his sole responsibility that the product above is, under conditions of normal use and conditions defined by the Producer / the Manufacturer, safe and meets all the necessary legal conditions and requirements. The product, a medical device that is intended for single use and solely in accordance with the Producer's / the Manufacturer's instructions.

The Conformity is assessed especially with the following provisions:

- Government Regulation no. 93/42/EEC Medical devices establishing technical requirements for medical devices, in effective wording
- Technical standard EN 14683:2019+AC:2019 Medical face masks - Requirements and test methods
- Other relevant harmonized legislation
- Other relevant local, national and community standards
- For the assessment of conformity, the following documents were also applied to:
- Tests for irritation and delayed-type hypersensitivity
- Results of laboratory tests Ekoteks Laboratuvar Testing Laboratory Bacterial filtration efficiency
- Results of laboratory tests Ekoteks Laboratuvar Testing Laboratory Microbial Cleanliness
- Results of laboratory tests Ekoteks Laboratuvar Testing Laboratory Differential Pressure
- Results of laboratory tests Ekoteks Laboratuvar Testing Laboratory Splash Resistance Pressure

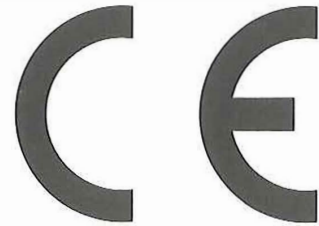
MARKING, LABELLING

Annex I, §13, of the Medical Devices Directive (93/42/EEC) or Annex I, §23, of the Medical Device Regulation (EU) 2017/745 specifies the information that should be specified on the packaging in which the medical face mask is supplied. The following information shall be supplied:

type of mask (as indicated in Table 1). EN ISO 15223-1:2016 and EN 1041:2008+A1:2013 should be considered

MEASURES TO ENSURE CONFORMITY

The Producer / the Manufacturer declares that he has taken all necessary measures to ensure the conformity of products placed on the market with technical documentation and basic requirements for this type of product.



General Manager

25/09/2020

EN ECZA DEPOSU İLAÇ MEDİKAL
ÖZEL SAĞLIK HİZM. A.Ş.

EN ECZA DEPOSU İLAÇ MEDİKAL ÖZEL SAĞLIK HİZMETLERİ

İNŞAAT TAAHHÜT TİCARET ANONİM ŞİRKETİ

Adres Adı: Karaduvar Mah. 313. Sokak, Karaduvar Apt. No: 5/B Merkez MERSİN

Adres No: 12943/2476 İstiklal V.D. : 3350707675

MERSİS NO: 0335010767500001 TİC. SİCİL NO: 51525

Tel: 0324 357 56 44 Fax: 0324 357 58 44



ENMED® ENM - 313 3 PLY UTRASONIC SURGICAL MASK (TYPE II-R)

The ENMED ENM - 313 3 PLY UTRASONIC SURGICAL MASK (TYPE II-R) is intended to be worn by medical personnel, patients and people to protect from transfer of microorganisms, body fluid and particulate material transfer. Type II-R Face masks include a splash resistant layer to protect against blood and other bodily fluids. Type IIR face masks are tested in the direction of exhalation (inside to outside) and take into account the efficiency of bacterial filtration.

KEY FEATURES

- The products are for single use only.
- Latex Free
- Hypoallergenic
- Fiberglass free
- Fluid Resistant
- Three Ply construction
- 3 pleats of folds to allow the user to expand the mask so it covers the area from the nose to the chin.
- Mask allows itself to be secured with an ear loop when placed behind the ears.
- Reduces exposure to blood and body fluids.
- Minimizes Doctor and patient contamination to exhaled microorganisms.

COUNTER INDICATION

None in particular.

PROPERTIES

Bacterial Filtration Efficiency (BFE): >98%

Breathing resistance: <40 Pa/cm²

Splash Resistance of ≥16 kPa

EN 14683 classification: Type IIR facemasks

INTENDED USE

3 ply mask is intended to be worn by

Medical personnel, patients and people to protect from transfer of microorganisms, body fluid and particulate material transfer.

DONNING INSTRUCTIONS

- The mask should be worn so that it covers the mouth and nose completely and there is no gap between the face and the mask.
- Touching the mask while wearing should be avoided, and hands should be washed with soapy water after touching a used mask. In the absence of water, antibacterial gel should be used.
- When the mask gets wet or moist, it should be replaced with a new dry mask. Moisture can cause diaper rash and irritation in that area.
- Torn, punctured or damaged masks should not be used.
- The mask is personal and Damp, wet products should never be used.

STORAGE CONDITIONS

- Unopened package can be used for 5 years from the date of manufacture.
- Store in a dry and cool place, away from intense sources of heat.

WARNINGS AND LIMITATIONS

- Do not use the product other than its intended use.
- Never use wet and moist products.
- The products are for single use only.
- Do not use it again when you remove it.
- It is personal. Never use someone else's mask.
- Our products have been tested. However, We recommend it not to be used on sensitive skin and if allergic to any materials used.

TECHNICAL DATA

MATERIAL	Multilayer SMS Fabric
COLOR	Blue, Green, White
HEADBAND	Latex free elastic ear loops, white, 17 cm
SIZES	Universal (17cm x 9cm)
PACKAGING	Optional packaging of 1/4/10/50/100
Nose Clips	Optional Aluminum Nose Clips
COO	TURKEY

PRODUCT OVERVIEW

A surgical mask is intended to prevent the release of potential contaminants from the user into their immediate environment which can also protect the wearer from large droplets, sprays and splashes of bodily fluids. It is easily identifiable by the presence of two ear loops to secure the mask to the face for quick donning.

Procedure masks are used on hospital floors, isolation units, and labor and delivery units, among other areas of the hospital for generally to prevent clinicians, patients and visitors from spreading germs by talking, coughing, or sneezing.

These masks may also be used in the emergency department and the intensive care unit. However, beware that some specifications are non-sterile and designed for clean environments which not suitable for use in the operating room.

Our disposable surgical face masks contain 3 layers: a mix of non-woven fabric and a middle melt-blown material that acts as a filter and have a >99% particle filtration efficiency (PFE). They provide physical barrier that protects both wearers and others from the transfer of respiratory secretions, fluids or other debris and avoid the contamination of bacteria or viruses present in mucus and saliva.

WASHING INSTRUCTIONS

Our products do not have a washing feature.

REGULATORY INFORMATION

Product CE marked as per 93/42/EEC Directive on Medical Devices.

Class of the device: I.

Product compliant with EN14683

HOW TO PROPERLY WEAR A FACE MASK



ENM - 313 3 PLY UTRASONIC SURGICAL MASK (TYPE II-R)

Product	3 PLY UTRASONIC SURGICAL MASK (TYPE II-R)
Material	Outer layer: 20 Gsm Spunbound (175mm)
	Filter layer: 25 Gsm Meltblown (BFE >%99) (175mm)
	Inner layer: 30 Gsm Spunbound (200mm)
Size	17cm x 9 cm
Color	Blue, Green, White
Nose Clips	Optional Aluminum Nose Clips
Standard	1- BFE >%99
	2- PFE >%95
	Polypropylene inner and outer facing assures comfortable on the inside and protection on the outside;
Certificate	CE, ISO, FDA



**EKOTEKS LABORATUVAR ve GÖZETİM
HİZMETLERİ A. Ş.**
Esenyurt Firuzköy Bulvarı No:29 34325 Avcılar
İstanbul/ TÜRKİYE



TEST REPORT
DENEY RAPORU

AB-0583-T
20041685
11-20

Müşterinin adı:

EN ECZA DEPOSU A.Ş.

Adresi:

Atatürk mah. 31043 Sk. Kubat Apt. No:8/B Mezitli/ MERSİN

Alıcı firma:

-

İlgili kişi:

ENGİN ŞAHİN

İstek numarası:

-

Model numarası:

-

Numunenin adı ve tarif:

Mavi dokusuz yüzey medikal maske. (Müşteri tarafından belirtilmiştir; Mavi)

Numunenin kabul tarihi:

06.11.2020

**İlave numune ve/veya ilave
bilgi geliş tarihi:**

-

Deneyin yapıldığı tarih:

06.11.2020-17.11.2020

Açıklamalar:

-

Numune alımı:

Bu raporda verilen sonuçlar müşteri tarafından gönderilen numuneye aittir.

Numunenin son kullanımı:

-

Yıkama talimatı:

Belirtilmedi.

Raporun sayfa sayısı:

5

Türk Akreditasyon Kurumu (TÜRKAK) deney raporlarının tanınması konusunda Avrupa Akreditasyon Birliği (EA) ve Uluslararası Laboratuvar Akreditasyon Birliği (ILAC) ile karşılıklı tanınma antlaşmasını imzalamıştır.

Deney laboratuvarı olarak faaliyet gösteren EKOTEKS LABORATUVAR ve GÖZETİM HİZMETLERİ A.Ş. TÜRKAK'tan AB-0583-T akreditasyon dosya numarası ile ISO 17025:2017 standardına göre akredite edilmiştir.

Deney ve/veya ölçüm sonuçları, genişletilmiş ölçüm belirsizlikleri (olması halinde) ve deney metodları bu sertifikanın tamamlayıcı kısmı olan takip eden sayfalarda verilmiştir.



Mühür

EKOTEKS

Tarih
17.11.2020

Müşteri Temsilcisi
Özlem ULLU

Laboratuvar Müdürü
Sevim A. RAZAK
17.11.2020

*Bu rapor laboratuvarın yazılı izni olmadan kısmen kopyalanıp çoğaltılamaz.
İmzasız ve mühürsüz raporlar geçersizdir.*

**EKOTEKS LABORATUVAR ve GÖZETİM
HİZMETLERİ A.Ş.**

AB-0583-T

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İSTENEN TESTLER	SONUÇ	AÇIKLAMA
MİKROBİYOLOJİK TESTLER		
Bakteri Filtrasyon Etkinlik Testi-BFE	P	TİP IIR
Mikrobiyal Temizlik (Biyoyük)	P	
Kan Sıçrama Direnci	P	
FİZİKSEL TESTLER		
Nefes Alabilirlik(Basınç Farkı)	P	
P: Geçer F: Kalır R: Alıcı firmanın teknik kişisine başvurunuz Test sonuçları EN 14683:2019+AC :2019 EK-B/Tablo 1) limit değerlerine göre değerlendirilmiştir.		

NOT: Aksi belirtilmediği takdirde testler ile ilgili kayıtlar 5 yıl, orjinal numuneler 3 ay saklanır. Müşteri tarafından talep edildiğinde, testlere ait ölçüm belirsizliği raporlanır fakat "Geçer/Kalır" değerlendirmesinde ölçüm belirsizliği değeri dikkate alınmaz. Raporlanan belirsizlik, genişletilmiş belirsizlik olup standart belirsizlik kapsam faktörü k=2 kullanılarak elde edilmiştir. Güvenilirlik düzeyi % 95'tir. Bu raporda (*) işaretli deneyler akreditasyon kapsamına dahil değildir.

Gen.fl36-1/03



Bu rapor, laboratuvarın yazılı izni olmadan kısmen kopyalanıp çoğaltılamaz
İmzasız ve mühürsüz raporlar geçersizdir.

TEST SONUÇLARI

Tıbbi yüz maskeleri – Gereklilikler ve Deney Yöntemleri EN 14683:2019+AC:2019 (TS EN 14683+AC:2019)

BAKTERİ FİLTASYON VERİMLİLİK TAYİNİ TESTİ- BFV

Test Metodu: EN 14683:2019+AC :2019 (TS EN 14683+AC:2019) EK-B

Örnek, aerosol ve mikrobiyal yük örnekleme haznesi arasına sıkıştırılır. Vakum sistemi yardımıyla bakteri içeren aerosol, filtreden geçirilir. Örneğin bakteri filtrasyon etkinliği, örnekten geçen koloni oluşturan birimlerin sayısının bakteri yüklü, aerosolde mevcut olan koloni oluşturan birimlerin sayısının yüzdesi olarak ifade edilir.

Deney Akış Hızı	28,3 L/dk
Toplam Deney Akış Süresi	2 dakika
Numune Ölçüleri	20x20 cm ²
Test Kondüsyon	(21 ± 5) °C and (85 ± 5) % bağıl nem, 4 saat
Test Mikroorganizması	<i>Staphylococcus aureus</i> ATCC 6538
Bakteri konsantrasyonu (kob/ ml)	5x10 ⁵ kob/ ml
Inkübasyon süresi, sıcaklık	37 ± 2 °C, 20 - 52 h
Pozitif Kontrol Numune Bakteri Sayısı Ortalaması (C)	2,3 x10 ³ kob/ ml
Ortalama Partikül Boyutu (MPS)	3 µm

SONUÇLAR			
Deney Numunesi Sayısı	Deney Numunesi Bakteri Sayısı(kob) (T)	Bakteri Filtrasyon Verimliliği (%B)	İstenen Değer BFV (%)
1	43	98,1%	Tip I ≥95 Tip II ≥98
2	38	98,3%	
3	45	98,0%	
4	41	98,2%	
5	31	98,7%	

kob: koloni oluşturan birim

$$B = (C - T) / C \times 100$$

%B: Bakteri Filtrasyon Verimliliği

C: Kontrol numunede üreyen bakteri sayısının ortalaması

T: Deney numunesinde üreyen bakteri sayısı

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MİKROBİYAL TEMİZLİK (BİYOYÜK)

Test Metodu: EN 14683:2019+AC :2019 (TS EN 14683+AC:2019) EK-D
EN ISO 11737-1:2018 /TS EN ISO 11737-1 :2018

5 numune çalışılır. Numune tartılır ve test çözeltisi içerisine atılarak iyice çalkalanır (250 rpm de 5 dk) ve uygun besiyerlerine ekilir. Toplam aerobik bakteriler için 30±1°C'de 72 saat, küf ve maya için ise 20-25 °C'de 7 gün inkübasyon sonrası agarda oluşan mikroorganizmalar sayılır ve toplam sonuç verilir. Ortalama sonuç verilir.

	<u>SONUÇ</u>	<u>İSTENEN</u>
Mikrobiyal Temizlik (kob/gr)	18 kob/gr	≤30 kob/gr

*kob: Koloni oluşturan birim

SIÇRAMA DİRENCİ

Test Metodu: EN 14683:2019+AC :2019 (Madde 5.2.4) (*) tıbbi yüz maskesinin sıvı sıçramalarına nüfuz etmesine karşı direnci, Tablo 1'de Tip IIR için verilen minimum değere uygun olmalıdır .

ISO 22609 :2004 (*) Giysilerin enfekte edici ajanlara karşı koruma - Tıbbi yüz maskeleri - Sentetik kanın nüfuz etmesine karşı direnç için test yöntemi (sabit hacim, yatay olarak yansıtılmış)

Test Kondüsyon koşulu ve süresi: (21 ± 5) °C ve (85 ± 5) % bağıl nem, 4 saat

6 farklı deney numunesi alınır.

<u>NUMUNE</u>	<u>SIÇRAMA DİRENCİ BASINCI (kPa)</u>	<u>SONUÇ</u>	<u>İSTENEN</u>
1	>21.3 kPa	GEÇER	≥16 kPa
2	>21.3 kPa	GEÇER	
3	>21.3 kPa	GEÇER	
4	>21.3 kPa	GEÇER	
5	>21.3 kPa	GEÇER	
6	>21.3 kPa	GEÇER	
Ortalama Sonuç	>21.3 kPa	GEÇER	

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BASINÇ FARKI (NEFES ALABİLİRLİK)

Test Metodu: EN 14683:2019+AC :2019 (TS EN 14683+AC:2019) EK-C

Test Kondüsyon koşulu ve süresi: (21 ± 5) °C ve (85 ± 5) % bağıl nem, 4 saat
2,5 cm çaplı 5 farklı deney numunesi alınır.

8 l/dk hava akışı uygulanır.

Fark Basınç Manometresi üzerinden okunan basınç farkı değeri Pa (Pascal) olarak kayit edilir.

NUMUNE	BASINÇ PARKI SONUÇ	İSTENEN
1	42,5 Pa/cm ²	< 60 Pa/cm ²
2	44,7 Pa/cm ²	
3	44,0 Pa/cm ²	
4	42,8 Pa/cm ²	
5	44,5 Pa/cm ²	
Ortalama Sonuç	43,7 Pa/cm ²	

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TEST REPORT
DENEY RAPORU

Customer name: EN ECZA DEPOSU A.Ş.
Address: Atatürk mah. 31043 Sk. Kubat Apt. No:8/B Mezitli/ MERSİN
Buyer name: -
Contact Person: ENGİN ŞAHİN
Order No: -
Article No: -
Name and identity of test item: Blue non-woven medical mask.(Claimed to be; Blue)
The date of receipt of test item: 06.11.2020
Re-submitted/re-confirmation date: -
Date of test: 06.11.2020-17.11.2020
Remarks: -
Sampling: The results given in this report belong to the received sample by vendor.
End-Use: -
Care Label: Not specified.
Number of pages of the report: 5

The Turkish Accreditation Agency (TURKAK) is signatory to the multilateral agreements of the European co-operation for the Accreditation (EA) and of the International Laboratory Accreditation (ILAC) for the Mutual recognition of test reports.

EKOTEKS LABORATUVAR ve GÖZETİM HİZMETLERİ A.Ş. accredited by TÜRKAK under registration number [AB-0583-T] for ISO 17025:2017 as test laboratory.

The test and/or measurement results, the uncertainties (if applicable) with confidence probability and test methods are given on the following pages which are part of this report.



Date
17.11.2020

Customer Representative
Özlem ULUS

Head of Testing Laboratory
Sevim A. RAZAK
17.11.2020

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REQUIRED TESTS	RESULT	COMMENTS
MICROBIOLOGICAL TESTS		
Bacterial Filtration Efficiency-BFE	P	TYPE IIR
Microbial Cleanliness(Bioburden)	P	
Splash Resistance	P	
PHYSICAL PROPERTIES		
Breathability(Differential Pressure)	P	
P: Pass F: Fail R: Refer to retailer technologist. Tests results were evaluated according to EN 14683:2019+AC :2019 Tablo 1 limit values.		

REMARK: Original samples are kept for 3 months and all technical records are kept for 5 years unless otherwise specified.If requested, measurement uncertainty will be reported. But unless otherwise specified, measurement uncertainty is not considered while stating compliance with specification or limit values The reported uncertainty is based on a standard uncertainty multiplied by a coverage factor k=2, providing a level of confidence of approximately 95 %. Tests marked (*) in this report are not included in the accreditation schedule.



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TEST RESULTS

BACTERIAL FILTRATION EFFICIENCY (BFE)

Test Metod: (Bacterial Filtration Efficiency Testing –BFE /Ref: EN 14683:2019+AC:2019 Medical Face Masks,Requirements and Test Methods

A specimen of the mask material is clamped between an impactor and an aerosol chamber. An aerosol of *Staphylococcus aureus* is introduced into the aerosol chamber and drawn through the mask material and the impactor under vacuum. The bacterial filtration efficiency of the mask is given by the number of colony forming units passing through the medical face mask material expressed as a percentage of the number of colony forming units present in the challenge aerosol.

Test Flow Rate	28,3 L/min
Test Flow Time	2 minute
Sample Sizes	5 pieces mask
Microorganism	<i>Staphylococcus aureus</i> ATCC 6538
Bacterial concentration (cfu/ ml)	5x10 ⁵ cfu/ ml
incubation conditions	24 hour, 35°C ± 2°C
Positive control sample average of number of Bacteria (C)	2,3 x10 ³ cfu/ ml

RESULTS			
Number of Test Sample	Test Sample (T) Number of Bacteria (cfu/ml)	Bacterial Filtration Efficiency (% B)	Requirement BFE (%)
1	43	98,1%	Type I ≥95 Type II ≥98
2	38	98,3%	
3	45	98,0%	
4	41	98,2%	
5	31	98,7%	

cfu: Colony-forming unit

$$B = (C - T) / C \times 100$$

%B: Bacterial Filtration Efficiency

C: is the mean of the total plate counts for the two positive control runs

T: is the total plate count for the test specimen

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TEST RESULTS

MICROBIAL CLEANLINESS (Bioburden)

Test Metod: EN ISO 11737-1:2018

The sample is put in extraciton liquid after shaking well, inoculated on the agar.
After incubation at 30 ± 1 ° C for 72 hours, growth microorganisms are counted on the agar.

	<u>RESULTS</u>	<u>REQUIREMENT</u>
Microbial cleanliness (cfu/g)	18 cfu/g	≤ 30 cfu/g

*cfu= Colony forming unit.

SPLASH RESİSTANCE

Test Metod: EN 14683:2019+AC :2019 (Clause 5.2.4) the resistance of the medical face mask to penetration of splashes of liquid shall conform to the minimum value given for Type IIR in Table 1

ISO 22609 :2004 Clothing for protection against infectious agents — Medical face masks — Test method for resistance against penetration by synthetic blood (fixed volume, horizontally projected)

Test Condition (21 ± 5) °C ve (85 ± 5) % relative humidity, 4 hrs

6 different samples were taken

	<u>SPLASH RESİSTANCE PRESSURE (kPa)</u>	<u>RESULTS</u>	<u>REQUIREMENT</u>
1	>21.3 kPa	PASS	≥ 16 kPa
2	>21.3 kPa	PASS	
3	>21.3 kPa	PASS	
4	>21.3 kPa	PASS	
5	>21.3 kPa	PASS	
6	>21.3 kPa	PASS	
Average Result	>21.3 kPa	PASS	

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TEST SONUÇLARI

BREATHABILITY (Differential Pressure)

Test Method: EN 14683:2019+AC :2019 (TS EN 14683+AC:2019) Annex-C

Test Condition (21 ± 5) °C ve (85 ± 5) % relative humidity, 4 hrs

Test area is 25 mm in diameter , 5 different sample was taken

Adjusted airflow is 8 l/min. The differential pressure is read directly using a differential pressure manometer .

SAMPLE	DIFFERENTIAL PRESSURE RESULT	REQUIREMENT
1	42,5 Pa/cm ²	< 60 Pa/cm ²
2	44,7 Pa/cm ²	
3	44,0 Pa/cm ²	
4	42,8 Pa/cm ²	
5	44,5 Pa/cm ²	
Average Result	43,7 Pa/cm ²	