

Esenyurt Firuzköy Bulvarı No:29 34325 Avcılar İstanbul/ TÜRKİYE

TEST REPORT
DENEY RAPORU

20015811ing

06-20

EKOTEKS

Customer name: GLOBICHEM KİMYA İTH.İHR.SAN. TİC. LTD. ŞTİ.

Address: BARAKFAKİH MAH. 7.CAD. NO:2/3 KESTEL/BURSA

Buyer name:

Contact Person: TALHA UĞUR

Order No:

Article No:

Name and identity of test item: White mask.

The date of receipt of test item: 20.05.2020

Re-submitted/re-confirmation

date:

Seal

Date of test: 20.05.2020-01.06.2020

Remarks:

Sampling: The results given in this report belong to the received sample by vendor.

End-Use: Mask

Care Label: Not specified.

Number of pages of the report: 4

DateCustomer RepresentativeHead of Testing Laboratory01.06.2020Servin YURTSEVENSevim A. RAZAK01.06.202001.06.2020

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REQUIRED TESTS	RESULT	COMMENTS		
MICROBIOLOGICAL TESTS				
Bacterial Filtration Efficiency-BFE (1)	P	Type I		
Microbial Cleanliness(Bioburden) (2)	P			
PHYSICAL PROPERTIES				
Breathability(Differential Pressure) (3)	P			

P: Pass

F: Fail

R: Refer to retailer technologist.

(1)Test results were evaluated according to EN 14683:2019+AC:2019 Annex-B/Table- 1)limit values

⁽²⁾Test results were evaluated according to EN ISO 11737-1:2018 limit values

(3)Test results were evaluated according to EN 14683:2019+AC:2019 Annex -C/Table- 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 Annex B Medical Face Masks,Requirements and Test Methods (*)

A specimen of the mask material is clamped between a 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	Staphylococcus aureus ATCC 6538
Bacterial concentration (cfu/ ml)	5x10 ⁵ cfu/ ml
incubation conditions	24 hour, 35°C ± 2°C
Positive control sample average	2.3x10 ³ cfu/ ml
of number of Bacteria (C)	

RESULTS			
Number of Test Sample	Test Sample (T) Number of Bacteria (cfu/ml)	Bacterial Filtration Efficiency (% B)	Requirement BFE (%)
1	74	%96.3	Type I ≥95
2	71	%96.5	Type II ≥98
3	71	%96.5	
4	76	%96.2	
5	77	%96.2	

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: Ref: EN ISO 11737-1:2018 (*)

The sample is put in extraciton liquid after shaking well, inoculated on the agar. After incubation at 30 \pm 1 $^{\circ}$ C for 72 hours, growth microorganisms are counted on the agar.

	<u>RESULTS</u>	REQUIREMENT
Microbial cleanliness (cfu/g)	12 kob/g	≤30 cfu/g Type I and Type II mask

BREATHABILITY (Differential Pressure)

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

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 .

	RESULT	REQUIREMENT
Differential Pressure) (Pa/cm²)	31.5 Pa/cm²(*)	< 40 Pa/cm ² Type I and Type II mask

^{*}average results are given