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Document No.: TR-25c-02 KiaNonoBioVista laboratory Accredited Laboratory of Medical Devices Testing



# Bacterial filtration efficiency test report

## **Customer Name and Address Information:**

Vira Teb Termeh Negar Company

Address: No. 36, Chamran Boulevard, Hashtgerd, Alborz, Iran

Phone number: +98-26-44353594, E-mail: Virateb.termeh@gmail.com, web: viratebtn.com

### Test Place Name and Address Information:

KiaNonoBioVista laboratory

Address: Unit 10, Floor 4, No. 58, Rose building, Taherian Street, Sadeghiyeh Square, Tehran,

Phone number: +98-21-440240223, E-mail: info@kiananobio.com, web: kiananobio.com

## Reference Standards and Conformity Methods:

- ASTM F2101-19 "Evaluation the bacterial filtration efficiency (BFE) of medical face mask materials, using a biological aerosol of Staphylococcus aurous"
- INSO 6138-1399 "single-use face masks for use in health care-requirements and test methods"
- BS EN 14683:2019+AC-Annex B "Medical face masks Requirements and test methods"
- ASTM F2100-11:2018 "Standard specification for performance of materials used in medical face masks"

#### Time Schedule:

Sample receiving date:

29/08/2022

Start test date:

29/08/2022

End test date:

05/09/2022

Issue date:

06/09/2022

## Sample Identification:

Name:

Disposable Protective Clothing (Tamam Ghad) Vira Teb Termeh Negar Company - Iran

Manufacturer: Sterilization method:

N010606-28-01

Case Lab number:

0106-0020

Sample Lab code: Manufacturing Date:

16/08/2022

Expiry date:

16/08/2025

LOT Number

REF Number:

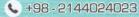
EP45

### **Test Summary Report:**

Test type	Test method	Test results		
Medical Mask bacterial filtration efficiency	ASTM F2101 BS EN 14683: 2019	Normal BFE ≥98.703%		

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### cceptance criteria:

The test conditions will be satisfied if no bacterial challenge counted during the collection of the negative control.

- The test conditions will be satisfied if approximately 1700-3000 CFU counted during the collection of positive controls.
- The mean particle size (MPS) in the bacterial challenge should be maintained at 3.0±0.3 μm.
- The sample will be approved if bacteria filtration efficiency comply with the criteria given below:

Table 1- The pharmaceutical criteria for bacteria filtration efficiency (BFE %)

	Example of applications	BFE %				
level 1	For general purpose medical procedures, where the wearer is not at risk of blood or body fluid splash or to protect staff and/or the patient from droplet exposure to microorganisms  (e.g. patient with upper respiratory tract infection visits general practitioner)					
level 2	For use in emergency departments, dentistry, changing dressings on small wounds where minimal blood droplet exposure may possible occur (e.g. endoscopy procedures)	≥98				
level 3	For all surgical procedures, major trauma first aid or in any area where the health care worker is at risk of blood or body fluid splash (e.g. orthopedic, cardiovascular procedures)	≥98				

#### Results:

The percent bacteria filtration efficiency and standard deviation are shown in table2.

Table2- microorganisms counted (CFU) in samples and controls

			Samples			Positive Controls		Negative
	No.1	No.2	No.3	No.4	No.5	No.1	No. 2	Control
Cascade 1	7	6	9	8	7	385	372	0 CFU
Cascade 2	6	7	6	4	7	372	340	0 CFU
Cascade 3	5	5	5	5	4	363	338	0 CFU
Cascade 4	3	4	6	3	3	340	322	0 CFU
Cascade 5	1	2	0	1	3	323	305	0 CFU
Cascade 6	1	0	0	1	0	288	262	0 CFU
Total	23	24	26	22	24	2071	1939	0
BFE %	98.853	98.803	98.703	98.903	98.803		-	-
Average	98.813					20	0	
Mean particle	e size (MPS	S): 3.3µm	Test side: Either			Testing area: 39.5cm2		

Expanded uncertainty=±0.31 % Reading \*Expanded uncertainty: An interval about the measurement result that encompasses 95% of probability density function of measurement result.

#### Conclusion:

On the basis of the results, interpreted according to ASTM F2101 the test sample "Disposable Protective Clothing (Tamam Ghad) Model EP45 - Vira Teb Termeh Negar Company " had a Normal Bacteria Filtration Efficiency and satisfied the requirements of the test.

#### Record filing:

The study program and all raw data will be retained in Kiananobiovista Lab archives for a period of 10 years from the issue of the final report. An original copy of the report is available at Kiananobiovista Lab.

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