

# **TEST REPORT**

LAB NO.: 2300703/ 1 - 2 DATE: 17/04/2023

NAME OF CUSTOMER : M/S. Allwin Rotoplast (Mr.Salim Pissuwala)

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**REFERENCE**: Letter Ref: Nil dated March,29 2023

Kind Attention: Mr. Srimant Anne

DATE OF RECEIPT : 30/03/2023

**DATE OF INITIATION** : 30/03/2023 & 10/04/2023

**DATE OF COMPLETION** : 05/04/2023 & 17/04/2023

SAMPLE DESCRIPTION : Plastic sample labeled as-

Sr No.	Description
1	Sample AW - Treated (Red colour)
Untreated – Lab Control	



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<sup>•</sup> Samples are not drawn by the laboratory • Result relate only to the samples tested

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## **Test Microorganism Information:**

MS2 Bacteriophage (MS2) is an RNA virus of the family Leviviridae. Escherichia coli 15597 are the hosts for bacteriophages. Due to its environmental resistance, MS2 bacteriophages are used as a surrogate virus (particularly in place of Picornaviruses such as Poliovirus and human Norovirus) in water quality and Antimicrobial studies.

Virus: MS2 Bacteriophage

Permissive Host Cell: Escherichia coli ATCC 15597

# **Experimental Details:**

Test Carrier : Test Sample (50 mm x 50 mm); Pre-sterilized by ETO gas

Control Carrier : Sample non coated and sterilized by autoclaving (50 mm x 50 mm)

LDPE cover : LDPE film pre sterilized 40 mm x 40 mm

Virus : MS2 Bacteriophage; Inoculum volume 0.4 ml

Permissive Host Cell : Escherichia coli ATCC 15597

Contact Period : 2 hours and 24 hours

Neutralizer : DE broth

Medium : Trypticase soya agar

Incubation for survivors : 37°C for 3 days

#### Validation and Records:

## **Neutralizer Validation and Records:**

Validation Test							
Test Organism	Exptl. Condition Control (A) (PFU/ ml)	Neutralizer Toxicity Control (B) (PFU/ ml)	Dilution-neutralization Control (C) (PFU/ ml)				
MS2 Bacteriophage	52	54	56				

## Where -

A=No. of PFU/ml of Test organism in Experimental condition validation B=No. of PFU/ml of Test organism in Neutralizer Toxicity validation

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#### **Test Procedure:**

Pre-sterilized samples were loaded with diluted viral suspension to 10<sup>6</sup> PFU/ ml. Virus suspension 0.4 ml was added to 50 mm x 50 mm of Test substrate. It was covered with 40 mm x 40 mm LDPE film. Following exposure time, Virus was eluted and neutralized by serial tenfold dilution and assayed to determined surviving Viruses in comparison with Control without test product in sq. cms. Virus assay was quantitative as Plaque forming unit (PFU) visible as area of Clearance.

## Results:

#### A. Contact duration of 2 hours

Quantitative Assessment of Antiviral Activity –ISO 21702: 2019							
Untreated: Average no	Log = 4.79						
Untreated: Average no	Log = 4.85						
Sample Identification	Average No. of Plaques recovered from Treated (At)	Log of Plaques recovered from Treated (At)	Antiviral Activity(R) (Log Ut- At)	Virus Reduction Percentage			
Sample AW – Treated (Red colour)	4800	3.68	1.17	93.23			

#### B. Contact duration of 24 hours

Quantitative Assessment of Antiviral Activity –ISO 21702: 2019							
Untreated: Average r	Log = 4.79						
Untreated: Average r	Log = 4.89						
Sample Identification	Average No. of Plaques recovered from Treated (At)	Log of Plaques recovered from Treated (At)	Antiviral Activity(R) (Log U <sub>t</sub> - A <sub>t</sub> )	Virus Reduction Percentage			
Sample AW – Treated (Red colour)	150	2.17	2.72	99.81			

## Where:

R = Antiviral activity

 $U_0$  = Log of PFU recovered from Untreated specimen immediately after inoculation, in PFU/ cm<sup>2</sup>

 $\mbox{U}_{\mbox{t}}$  = Log of PFU recovered from Untreated specimen after2/24 hrs. after inoculation, in PFU/ cm²

 $A_t$  = Log of PFU recovered from Treated specimen after 2/24 hrs. after inoculation, in PFU/ cm<sup>2</sup>

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