

Japan Food Research Laboratories

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Page 1 of 4

Date issued: August 28, 2017

REPORT

Client: Haenim Co., Ltd

15, Saryeom-ro, 21beon-gil, Seo-gu, Incheon, Republic of Korea.

Sample(s): Haenim UV Sterilizer (Model: HN-03 series)

Title: Virus Inactivation Test

Received date of sample(s): July 10, 2017

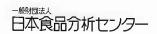
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Signed for and on behalf of JFRL

Takeko Arai

Section of Analysis Documentation

Aug. 29, 2017





Virus Inactivation Test

1. Client

Haenim Co., Ltd

2. Sample

Haenim UV Sterilizer (Model: HN-03 series)

3. Outline of the method

Influenza virus suspension was poured into a petri dish and used as a test specimen. It was placed in the sample at the position specified by the client. After the sample was allowed to operate under the conditions specified by the client, the virus infectivity titer of the test specimen was determined.

4. Results

Table 1 shows the results of the virus infectivity assay. Tables 2 and 3 show the cell and media used in the test and the test conditions, respectively. Photos 1 and 2 show the examples of the positions of the test specimens.

Table 1: Virus infectivity titer of test specimen

Test organism	Object	Position of test	log TCID ₅₀ /mL
		specimen	20g 101D30/IIIL
Influenza virus	Before operation		6.7
	After 10 minutes of	Upper row	<1.5
	operation	Lower row	<1.5

TCID₅₀: Median tissue culture infectious dose

<1.5: Not detected

Virus suspension: The culture solution was diluted 100-fold with purified water.

Table 2: Cell and media used in test

Test cell	MDCK (NBL-2) cells ATCC CC	CL-34 strain		
Cell culture medium	Eagle's MEM "Nissui" (1) containing 10 % of fetal bovine serum (Nissui Pharmaceutical Co., Ltd.)			
Cell support medium	Eagle's MEM "Nissui" (1) 10 % NaHCO ₃ L-Glutamine (30 g/L) 100 × Vitamins for MEM 10 % Albumin 0.25 % Trypsin	1000 mL 14 mL 9.8 mL 30 mL 20 mL 20 mL	Œ.	

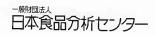




Table 3: Test conditions

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Test virus	Influenza A virus (H1N1) A/PR/8/34 ATCC VR-1469		
Virus suspension	The virus culture solution after cell incubation was centrifuged and the supernatant liquid was diluted 100-fold with purified water.		
Test specimen	A 2-mL portion of the virus suspension was poured into a petri dish (φ60 mm)		
Test procedure The test specimen was placed at the position specified by the client (just the ultraviolet ray lamp; the upper and lower rows) in the sample. The was allowed to operate for 10 minutes under the conditions specified by client.			
Determination method for virus infectivity assay	TCID ₅₀ method		





Photo 1. Example of position of test specimen: Upper row



Photo 2. Example of position of test specimen: Lower row

End of Report