

Attn: MIC. Co., Ltd.

## **Test Report**

### Antiviral Testing of PBM DeoSpray

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In the event the contents of this testing are to be announced, prior approval by this center is required.

Moreover, test results described in this report are for the specific sample, and do not prove the quality of the whole lot.

### **1. Purpose of the Test**

The purpose of this test is to evaluate the efficacy of viral inactivation by PBM Deo Spray using Influenza A virus.

### **2. Testing Virus**

Influenza A virus

### **3. Test Method**

#### 1) Culture and adjustment method of the sample virus

Influenza A virus

The influenza virus was inoculated into the chorioallantonic cavity of embryonated eggs and cultured in an incubator, then the chorioallantonic fluid was collected and the virus fluid was purified by density gradient centrifugation, and was determined to be the sample virus fluid.

#### 2) Test Product and Test Conditions

Supplied Test Product: PBM Deo Spray

Duration of viral activity: 0-1 minute

#### 3) Test Method

Inactivation test of the feline calicivirus was carried out according to the following procedures.

For each in vitro test, 0.9mL of the test product and 0.1mL feline calicivirus were combined, and then mixed in a vortex (infectivity titer:  $4.0 \times 10^8$  TCID<sub>50</sub>/mL). After reaction at room temperature at the prescribed duration of activity, a sampling of 0.1mL was taken and immediately diluted with 4.9mL PBS (phosphate buffered saline) to halt drug activity (sample stock solution). 10 times serial dilution was then promptly carried out by PBS, to measure viral infectivity titer of the sample. Furthermore, PBS was used in place of the sample for the of 0 minutes response time of the duration of viral activity. PBS was used.

#### 4) Virus Quantification Method

The CRFK cell culture supernatant was suction removed as a monolayer culture in a 96 hole plate beforehand, then sample undiluted solution and diluted 25  $\mu$  L virus fluid were added and allowed to stand for 1 hour at 37°C. After standing, the virus fluid was suction removed, and 100  $\mu$  L DMEM including 0.2% FBS was added to each well and cultured in a carbon dioxide gas incubator at 37°C. After culturing, cytopathogenic effect

(CPE) of each well was observed by microscope, and viral infectivity (TCID<sub>50</sub>/mL) was calculated by using Reed-Muench method.

#### 4. Test results and comments

Results are shown in Table-1.

When the virus with an initial infectivity titer of  $1.1 \times 10^5$  TCID<sub>50</sub>/mL was activated for one minute, PBM Deo Spray resulted in a reduction of more than 2.9log<sub>10</sub> infectivity titer (infectivity titer less than  $1.3 \times 10^2$  TCID<sub>50</sub>/mL).

Test supervision by Yasuhiro Nojima

End

Table-1 Antiviral Efficacy by PBM DeoSpray

Tested product	Duration of Activity (min.)		Reduction value of the infectivity titer (log 10)
	0	1	
PBM Deo Spray	$1.1 \times 10^5$	$< 1.3 \times 10^2$	$> 2.9$
Control (PBS)		***	***

Detection limit value:  $1.3 \times 10^2$

Reduction value of the infectivity titer: log<sub>10</sub>

(0 minute infectivity titer ÷ infectivity titer of 1 minute duration of activity)