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Date : 4 Nov 2009

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PREMILINARY REPORT ON THE EFFICACY OF BLUOXY DISINFECTANT AGAINST SWINE INFLUENZA VIRUS (H1N1)

With reference to the above matter, the results of the test are as show as below:

The Efficacy of BluOxy Disinfectant Against Swine Influenza Virus (H1N1)

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Introduction

BluOxy is a plant-based solution developed for purpose of cleansing, sanitizing and deodorizing room ambient. The product was also reported to have ability of killing a wide range of fungi, spores and bacteria that are harmful to human. This communication reports the preliminary efficacy test result of BluOxy against Swine Influenza Virus (H1N1).

Materials and Methods

Preparation of virus pool

The virus H1N1 strain was propagated in Chicken Embryonated Eggs (CEEs) and the virus titer was determined by Hemaaglunitination test (HA).

Disinfectant

The disinfectant namely BluOxy™ was used in this experiment. Two different dilutions of disinfectant were used in this study. The concentrated stock was diluted to concentration of 2% while the solution from the small personal size bottle which is formulated at 5% was used directly in the study.

A 'carrier' dilution method

A "carrier" is used according to standard protocols for evaluating the effect of surface disinfectant (Sattar et al., 2001). Two (2) pieces of sterile stainless steel penicylinders (type 304 stainless steel) as a carrier was immersed in 7 ml of H1N1 virus pool suspension for 15 minutes. Then, the carriers (rings) were then blot dried and left in incubator for 30 minutes at 37°C (Mermert, Western Germany). Each ring was immersed in 3 ml of disinfectants for 5 minutes. One control carrier was immersed in 1 ml PBS, pH 7.2. Then, 1 ml of diluents (PBS) was added to each ring to elute the virus. These elutes were further assayed for subsequent qualitative analysis by inoculating it into 5 of CEEs.

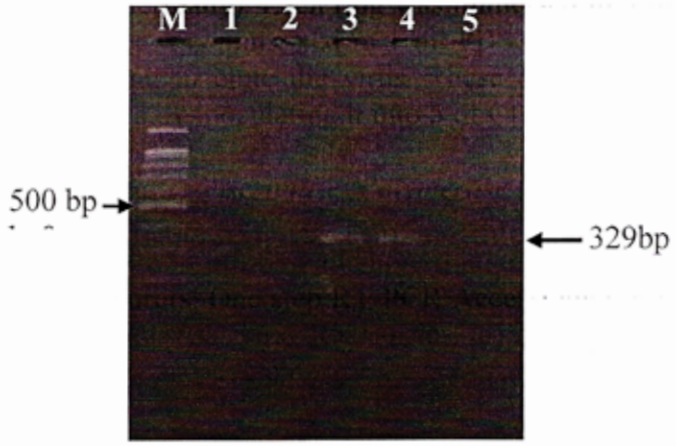
Extraction of virus and Reverse Transcriptase- Polymerase Chain Reaction (RT-PCR)

The viral RNA was extracted from infected allantoic fluid using TRIzol reagent (Invitrogen) as described by the manufacturers. One step RT-PCR AccessQuick RT-PCR Kit (Promega, USA) was employed to amplify the viral RNA using primer NP1529R/ NP1200F (M.-S. Lee et al., 2001).

Results

No amplicon was detected from BluOxy™ treated (2% and 5%) allantoic fluid. However, all the positive control shown amplicon sized at 329 bp. Negative control was also produced no amplicon. The results are illustrated in Figure 1.

Figure 1: RT- PCR product using Primer NP1529R/NP1200F (M.-S. Lee et al., 2001)



Legend: **M**: Marker, 100bp; **1**: BluOxy™ Treated at concentration of 2%; **2**: BluOxy™ Treated at 5% concentration; **3**: Positive control (Not Treated with BluOxy™); **4**: Positive Control; **5**: Negative Control

Discussion

Absence of amplicon indicated that the Swine Influenza Virus H1N1 virus was completely destroyed by the BluOxy™. This indicated that BluOxy™ posed virucidal property. However, according to the Environmental Protection Agency (EPA) and Canada General Standard Board (CGSB) guidelines for the evaluation of disinfectant, there is a requirement to conduct virus infectivity assay (Arshad et al., 2007). As such, further work needs to be conducted.

Conclusion

In conclusion, the BluOxy™ disinfectant tested in this study can be considered as an effective agent against Swine Influenza Virus (H1N1), as reflected by the absence of amplicon in BluOxy™ treated allantoic fluid.

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Sincerely,

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