



## Monkeypox Virus Inactivation Testing Report

<b>Report identifier</b>	HCM/MPx/014/v1
<b>Report date</b>	30 August 2022
<b>Testing laboratory</b>	High Containment Microbiology, UK Health Security Agency (UKHSA)

<b>Product details</b>	
<b>Product name</b>	Xpert CT/NG Swab Transport Reagent from Xpert Vaginal/Endocervical Specimen Collection Kit  N.B. Product was decanted into a secondary vessel prior to submission to laboratory for inactivation testing
<b>Product code</b>	SWAB/A-50 (Xpert Vaginal/Endocervical Specimen Collection Kit)
<b>Batch number</b>	2115212
<b>Manufacturer</b>	Cepheid
<b>Storage conditions</b>	Ambient temperature
<b>Active substances and concentrations (if known)</b>	Ammonium chloride (5-8%), Potassium carbonate (0.5-1.5%)
<b>Instructions for use</b>	Swab resuspended in tube containing Xpert CT/NG Swab Transport Reagent

Report identifier and version number: HCM/MPx/014/v1

Report date: 30 August 2022

Page 1 of 4

**UNCONTROLLED WHEN PRINTED**

<b>Experimental conditions</b>	
Period of analysis	23 – 30 August 2022
Product test concentrations	10 volumes buffer to 1 volumes test sample
Test temperature	Ambient
Treatment times tested	10 minutes; 30 minutes
Sample type tested and virus details	Monkeypox virus stock: monkeypox virus isolate UK2 (GenBank entry MT903344), in tissue culture fluid containing 5% foetal bovine serum
Description of test	<p>Triplicate samples of monkeypox virus tissue culture fluid were treated with product at the indicated test concentration for indicated contact times. Mock-treatments were carried out in triplicate using an equivalent volume of Minimum Essential Medium (MEM, Gibco) instead of product. After treatment, all samples were subjected to a filtration step to reduce cytotoxic buffer components, using Pierce Detergent Removal Resin in accordance with the manufacturer's instructions. MEM-treated samples were subjected to the same filtration procedure in parallel. All samples were immediately titrated on Vero E6 cells and plates immunostained using an anti-vaccinia virus antibody to establish virus titre. Product only controls (purified and unpurified) were additionally titrated to determine product cytotoxicity before and after filtration.</p> <p>This test is quantitative and reports the virus titre for each treatment condition in focus forming units (FFU)/mL. Reduction in virus titre following treatment is given as the difference between the mean log<sub>10</sub> FFU/mL for treated conditions and the MEM control.</p>

<b>Table of results</b>				
<b>Treatment condition</b>	<b>Mean virus titre in FFU/mL</b>	<b>Mean virus titre in log<sub>10</sub> FFU/mL [95% CI]</b>	<b>Titre reduction in log<sub>10</sub> FFU/mL [95% CI]</b>	<b>% reduction in virus titre</b>
<b>MEM-treated</b>	5.73x10 <sup>5</sup>	5.8 [5.5-6.0]	-	-
<b>10-minute treatment</b>	1.06x10 <sup>6</sup>	6.0 [5.9-6.1]	-0.3 [-0.5-0.0]	No reduction detected
<b>30-minute treatment</b>	8.71x10 <sup>5</sup>	5.9 [5.6-6.3]	-0.2 [-0.4-0.1]	No reduction detected

**Results interpretation and limitations**

Treatment with Xpert CT/NG Swab Transport Reagent for 30 minutes failed to inactivate monkeypox virus in this test. This product should not be relied upon to inactivate monkeypox virus.

This test has been performed using tissue culture fluid. The effectiveness of this treatment against monkeypox virus may vary when used to inactivate clinical samples or other types of sample matrix.

Nucleic acid stability in this product has not been examined, nor has the suitability of this product for inactivation of other pathogens been evaluated in this study.

## **Disclaimer**

UKHSA does not in any way recommend any particular product for virus inactivation; and UKHSA shall not be responsible for the choice of product or treatment for virus inactivation, and it is the responsibility of the testing laboratories to ensure that any such product or treatment implemented has undergone the necessary verification and validation; and UKHSA shall not be liable, to the greatest extent possible under any applicable law, for any claim, loss or damage arising out of or connected with use of this and related reports and choice of virus inactivation products or treatments.

UKHSA is an Executive Agency of the Department of Health and Social Care.  
Unauthorised use of the UKHSA name and/or logo is prohibited.

## **Summary of revisions**

Version 1: New document