



SUBJECT Microbiological Test of Disinfectant wipes 991919 & 991920

TEST LOCATION TÜV SÜD China
TÜV SÜD Products Testing (Shanghai) Co., Ltd.
B-3/4, No.1999 Du Hui Road, Minhang District
Shanghai 201108, P.R. China

CLIENT NAME NuMedical Pty Ltd.

CLIENT ADDRESS 9-13 Hender Avenue Magill SA 5072 Australia

TEST PERIOD 03-Aug-2020~24-Aug-2020

Prepared By

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Report Drafter

Authorized By



(Steven Zhang)
Authorized Signatory

Note: (1) General Terms & Conditions as mentioned overleaf. (2) The results relate only to the items tested.(3) The test report shall not be reproduced except in full without the written approval of the laboratory.(4) Without the agreement of the laboratory, the client is not authorized to use the test results for unapproved propaganda.

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RECEIPT DATE / TEST DATE

03-Aug-2020/ 03-Aug-2020

THE FOLLOWING SAMPLE(S) WAS/WERE SUBMITTED

BY/ ON BEHALF OF THE CLIENTS AS:

Sample Name: NuMedical Pty Ltd.
Sample Specification: /
Batch No./Date: 2020.05.20
Manufacturer: 9-13 Hender Avenue Magill SA 5072 Australia

SAMPLE NO.	DESCRIPTION	PHOTOGRAPH
721657042	Sample in the bag	

TEST METHOD

BS EN 13727-2012+A2:2015

Chemical disinfectants and antiseptics - Quantitative suspension test for the evaluation of bactericidal activity in the medical area-Test method and requirements(phase 2, step 1)

REQUIREMENT(S)

Method: Neutralizer dilution method
Sample Pretreatment: Extrusion liquid
Test temperature: 20°C
Contact time: 15 min
Neutralizer : D/E Broth
Interfering substance: 0.3g/L bovine albumin= clean conditions
Appearance of the product test solutions: No precipitation, Transparent liquid

TEST ORGANISMS

Pseudomonas aeruginosa ATCC 15442



TESTING RESULTS

Test organism	Validity test		Test suspension N (CFU/mL)	Product test		Comment Reduction≥5
				Number of survivors (CFU/mL) Na	Reduction (lgR=lgN ₀ -lgNa)	
<i>Pseudomonas aeruginosa</i> ATCC 15442	Validation suspension (CFU/mL) N _{v0}	66	1.7×10 ⁸	<140	>5.08	Pass
	Experimental conditions control A (CFU/mL)	66				
	Neutralizer control B (CFU/mL)	78				
	Method validation C (CFU/mL)	82				

Note:

R = reduction (lg R = lg N₀ - lg N_a)

If Na<140 , lg R = > [lg N₀ - 2.15]

If Na > X*10 , lg R = < [lg N₀-lgX](X=upper limit for V_c)

Where:

N: N is the number of cells per ml in the test suspension.

N₀: (N₀= N/10), Number of cells per ml in the test mixtures at the beginning of the contact time (time 0)

N_a: Number of survivors per ml in the test mixtures

N_v: N_v is the number of cells per ml in the validation suspension.

N_{v0}: (N_{v0} = N_v/10), N_{v0} is the number of cells per ml in the mixtures A,B and C at the beginning of the contact time(time 0).

Conditions for passing the validity test: 30 ≤ N_{v0} ≤ 160; A ≥ 0.5 N_{v0}; B ≥ 0.5 N_{v0}; C ≥ 0.5 N_{v0}

This report is for internal use only such as internal scientific research ,education, quality control, product R&D.

This report replaces report 721657042-1-R1, 721657042-1-R1 is obsolete.

-END OF THE TEST REPORT-

