

Test Report: EN 1276:2019 Chemical disinfectants and antiseptics – Quantitative suspension test for the evaluation of bactericidal activity of chemical disinfectants and antiseptics used in food, industrial, domestic and institutional areas — Test method and requirements (phase 2, step 1)

Test Laboratory**BluTest Laboratories Ltd**

5 Robroyston Oval, Nova Business Park, Glasgow G33 1AP

Identification of sample

Name of the product

Batch number

Client

Client address

Project code

Date of delivery

Storage conditions

Active substances

Surface Disinfectant Concentrate

K5093

Fluid science Limited

Unit 5 Pride Point, Ashcroft Road, Knowsley Ind. Park,
Liverpool L33 7TW

BT-GEN-03

16 April 2020

Ambient

L-lactic acid

Test Method and Neutralisation

Method

Filtration neutralisation

Experimental Conditions

Period of analysis

13 May 2020 to 22 May 2020

Product diluent used

Sterile synthetic hard water

Product test concentrations

10.0% v/v; 5.0% v/v; 2.5% v/v

Appearance product dilutions

No changes noted

Appearance in test mixture

No changes noted

Contact time

t = 5 mins ± 10 s

Test temperature

20°C ± 1°C

Interfering substance

3.0g/l bovine albumin

Temperature of incubation

37°C ± 1°C

Identification of strains

Staphylococcus aureus ATCC 6538*Enterococcus hirae* ATCC 10541*Pseudomonas aeruginosa* ATCC 15442*Escherichia coli* ATCC 10536

EN 1276:2009 Results

Results for the efficacy of Surface Disinfectant Concentrate from Fluid Science Ltd under Dirty conditions									
Test organisms	Validation test				Bacterial Test Suspension (N)	Test procedure at concentration % (V/V)			
	Bacterial Suspension (Nv)	Experimental Conditions (A)	Filtration Control (B)	Filtration Test Control (C)		2.5%	5.0%	10.0%	
<i>Pseudomonas aeruginosa</i>	Vc: 80 ; 94	Vc: 62 ; 64	Vc: 51 ; 60	Vc: 58 ; 79	10 ⁻⁶ : >330 ; >330 10 ⁻⁷ : 40 ; 36 N: 3.80E+08	Vc: 30 ; 6 Na: 1.80E+02 R: >10(5)	0 ; 0 <1.40E+02 >10(5)	0 ; 0 <1.40E+02 >10(5)	
ATCC 15442	Nv: 8.70E+02	A: 6.30E+01	B: 5.55E+01	C: 6.85E+01	Q: n/a				
Validation	30 ≤ Nv ₀ ≤ 160 ? yes	A ≥ 0.5 x Nv ₀ ? yes	B ≥ 0.5 x Nv ₀ ? yes	C ≥ 0.5 x Nv ₀ ? yes	7.17 ≤ log N ₀ ≤ 7.70 ? yes	Test is valid			
<i>Escherichia coli</i>	Vc: 80 ; 73	Vc: 79 ; 82	Vc: 69 ; 80	Vc: 84 ; 89	10 ⁻⁶ : >330 ; >330 10 ⁻⁷ : 29 ; 39 N: 3.40E+08	Vc: 0 ; 0 Na: <1.40E+02 R: >10(5)	0 ; 0 <1.40E+02 >10(5)	0 ; 0 <1.40E+02 >10(5)	
ATCC 10536	Nv: 7.65E+02	A: 8.05E+01	B: 7.45E+01	C: 8.65E+01	Q: n/a				
Validation	30 ≤ Nv ₀ ≤ 160 ? yes	A ≥ 0.5 x Nv ₀ ? yes	B ≥ 0.5 x Nv ₀ ? yes	C ≥ 0.5 x Nv ₀ ? yes	7.17 ≤ log N ₀ ≤ 7.70 ? yes	Test is valid			
<i>Staphylococcus aureus</i>	Vc: 62 ; 51	Vc: 49 ; 51	Vc: 55 ; 38	Vc: 51 ; 52	10 ⁻⁶ : >330 ; >330 10 ⁻⁷ : 26 ; 17 N: 2.15E+08	Vc: 3 ; 5 Na: <1.40E+02 R: >10(5)	0 ; 0 <1.40E+02 >10(5)	0 ; 0 <1.40E+02 >10(5)	
ATCC 6538	Nv: 5.65E+02	A: 5.00E+01	B: 4.65E+01	C: 5.15E+01	Q: n/a				
Validation	30 ≤ Nv ₀ ≤ 160 ? yes	A ≥ 0.5 x Nv ₀ ? yes	B ≥ 0.5 x Nv ₀ ? yes	C ≥ 0.5 x Nv ₀ ? yes	7.17 ≤ log N ₀ ≤ 7.70 ? yes	Test is valid			
<i>Enterococcus hirae</i>	Vc: 65 ; 68	Vc: 43 ; 84	Vc: 47 ; 49	Vc: 53 ; 52	10 ⁻⁶ : >330 ; >330 10 ⁻⁷ : 22 ; 26 N: 2.40E+08	Vc: 9 ; 1 Na: <1.40E+02 R: >10(5)	1 ; 0 <1.40E+02 >10(5)	0 ; 0 <1.40E+02 >10(5)	
ATCC 10541	Nv: 6.65E+02	A: 6.35E+01	B: 4.80E+01	C: 5.25E+01	Q: n/a				
Validation	30 ≤ Nv ₀ ≤ 160 ? yes	A ≥ 0.5 x Nv ₀ ? yes	B ≥ 0.5 x Nv ₀ ? yes	C ≥ 0.5 x Nv ₀ ? yes	7.17 ≤ log N ₀ ≤ 7.70 ? yes	Test is valid			

Please note: the upper limit for counting bacterial plates is 330 cfu. Enter as >330. For filter membranes, this is 165 cfu. Enter as >165.

Definitions: Vc = viable count; N = number of cfu/ml in the bacterial test suspension; Q = quotient of control of weighted mean counts; Nv = number of cfu/ml in the bacterial validation suspension; A = number of cfu/ml in the experimental conditions validation; B = number of cfu/ml in the filtration control; C = number of cfu/ml in the membrane-filtration neutralisation method validation; Na = number of cfu/ml in test mixture after contact time; R = reduction in viability (Log10)

Conclusion

According to EN 1276:2019, **Surface Disinfectant Concentrate POSSESSES BACTERICIDAL** activity at a concentration of **2.5% v/v** as tested after **5 MINUTES** at **20°C** under **DIRTY** conditions (3.0 g/l bovine albumin) against *Pseudomonas aeruginosa* ATCC 15442, *Escherichia coli* ATCC 10536, *Staphylococcus aureus* ATCC 6538 and *Enterococcus hirae* ATCC 8043.

Authorised signatory



Dr Chris Woodall, Director
BluTest Laboratories Ltd
Glasgow, UK.
Date: 27 MAY 2020



4597

Expanded Uncertainty of Measurement $U = \pm 0.0044$ logs

DISCLAIMER

The results in this test report only pertain to the sample supplied.

BluTest (BT) has performed the testing detailed in this report using reasonable skill and care and has used reasonable endeavours to carry out the testing in accordance with an EN 1276 protocol. All forecasts, recommendations and results contained in this report are submitted in good faith. However, other than as expressly set out in this report, no warranty is given (i) in relation to the testing or the use(s) to which any results or deliverables produced in the course of the testing are or may be put by the Client or their fitness or suitability for any particular purpose or under any special conditions notwithstanding that any such purpose or conditions may have been made known to BT or (ii) that the intended results or deliverables from the testing can be achieved or (iii) that the Client can freely make use of the results or the deliverables without infringing any third party intellectual property rights and the Client will be deemed to have satisfied itself in this regard. BT shall have no liability (which is hereby excluded to the fullest extent permissible by law) in respect of any loss, liability or damage, including without limitation any indirect and/or consequential loss such as loss of profit or loss of business, market or goodwill, that the Client may suffer directly or indirectly as a result of or in connection with: (i) the performance of the testing; (ii) the use of any materials, samples or other information provided by the Client for use in the testing; and (iii) the Client's reliance upon or use of any results or deliverables provided as part of the testing.