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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)

Identification of the test laboratory: Abbott Analytical Ltd

Unit 2, Hickmans Road, Birkenhead, CH41 1JH, United Kingdom

Identification of the client: Coventry Chemicals Ltd

Woodhams Road, Coventry, CV3 4FX, United Kingdom

Identification of the sample: 21B/114

Name of the product: Antibac Concentrate

Batch number/reference and expiry date (if available):

AQSB1 P177/2

Date of delivery: 25 February 2021

Storage conditions: Room temperature in darkness

Product diluent recommended by

the manufacturer for use:

Not disclosed

Active substance(s) and their

concentrations (s) (optional):

Not disclosed

Appearance of the product: Clear colourless liquid

Notes:

- 1) The test results in this report relate only to the sample(s) tested.
- 2) This test report may not be reproduced except in full, adapted, altered or used to create a derivative work, without written approval from Abbott Analytical Ltd.

Abbott Analytical Ltd Unit 2, Hickmans Road, Birkenhead, CH41 1JH, United Kingdom







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Test method and its validation:

Method: Dilution-neutralisation

Neutraliser: 100.0 g/l Polysorbate 80 + 30.0 g/l Lecithin +

30.0 g/l Tryptone Soya Broth + 5.0 g/l Sodium thiosulphate +

1.0 g/l L-histidine (Neutraliser B)

Neutraliser validation: Validated in accordance with EN 1276:2019 (5.5.2)

Experimental conditions:

Period of analysis: 13 April 2021 to 15 April 2021

Product test concentration(s): 10% v/v

Diluent used for product test

solution(s):

Hard water

Contact time(s): $5 \min \pm 10 \text{ s}$

Test temperature(s): $20^{\circ}C \pm 1^{\circ}C$

Interfering substance: 3.0 g/l bovine albumin (dirty conditions)

Temperature of incubation: $36^{\circ}\text{C} \pm 1^{\circ}\text{C}$

Identification of the bacterial Pseudomonas aeruginosa (NCIMB 10421)

strain(s) used: Escherichia coli (NCTC 10418)

Staphylococcus aureus (NCTC 10788) Enterococcus hirae (NCIMB 8192)

Deviations: None

Remarks:

1) All test conditions are as requested by the client, irrespective of whether these are in accordance with EN 1276:2019 (5.4.2) or EN 1276:2019 (5.5.1.1).





& Murden



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Requirements:

The product shall demonstrate at least a 5 decimal log (lg) reduction against every test organism.

Conclusion:

According to EN 1276:2019, this sample of Antibac Concentrate possesses bactericidal activity against all of the referenced strains of *Pseudomonas aeruginosa*, *Escherichia coli*, *Staphylococcus aureus* and *Enterococcus hirae*, when tested at a concentration of 10% with a contact time of 5 minutes at 20°C under dirty conditions.

Report prepared by:

Signed:

Name: Tony Watson

Position: General Manager

Date: 16 April 2021

Approved by:

Signed:

Name:

Kirsty Murden

Position: Laboratory Manager

Date: 16 April 2021







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Results: EN 1276:2019

Test organism: Pseudomonas aeruginosa (NCIMB 10421)

Date of test: 13 April 2021 Test temperature: 20°C ± 1°C

Interfering substance: 3.0 g/l bovine albumin

Dilution-neutralisation method: Pour plate Number of plates: 1/ml Neutraliser: B Incubation temperature: $36^{\circ}C \pm 1^{\circ}C$

Validation and controls:

Validation	n suspensio	on (Nv _o)	Experime	ntal condit	ions	Neutralis	er or filtrat	tion	Method v	Method validation (C)		
				control (A)			control (B)			Product conc.: 10%		
Vc1	48	<u></u> =	Vc1	50	<u></u> =	Vc1	51	<u></u> =	Vc1	53	<u> </u>	
Vc2	52	50	Vc2	55	52.5	Vc2	51	51	Vc2	50	51.5	
$30 \le \overline{\mu}$ of $Nv_0 \le 160$?			$\overline{\mu}$ of $A \ge 0.5 \times \overline{\mu}$ of Nv_0 ?			$\overline{\mu}$ of $B \ge 0.5 \times \overline{\mu}$ of Nv_0 ?			$\overline{\mu}$ of $C \ge 0.5 \times \overline{\mu}$ of Nv_0 ?			
⊠ yes □ no			⊠ yes	□no		⊠ yes □ no			⊠ yes □ no			

Test suspension (N and N_o):

Ν	Vc1	Vc2	$\pi \text{ wm} = 2.58 \times 10^8 \text{ ;}$	lg N =	8.41
10 ⁻⁶	248	264	$N_0 = N / 10$; $\lg N_0 =$	7.41	
10 ⁻⁷	29	26	$7.17 \le \lg N_0 \le 7.70$?	⊠ yes	□no

Conc. of the	Contact	Vc1	Vc2	Na	lg Na	lg R
product	time			(π x 10)		(lg N _o - lg Na)
10%	5 min	0	0	<140	<2.15	>5.26







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Results: EN 1276:2019

Test organism: Escherichia coli (NCTC 10418) Date of test: 13 April 2021 Test temperature: $20^{\circ}\text{C} \pm 1^{\circ}\text{C}$

Interfering substance: 3.0 g/l bovine albumin

Dilution-neutralisation method: Pour plate Number of plates: 1/ml Neutraliser: B Incubation temperature: $36^{\circ}C \pm 1^{\circ}C$

Validation and controls:

Validation	n suspensio	on (Nv _o)	Experime	ntal condit	tions	Neutralis	Neutraliser or filtration			Method validation (<i>C</i>)		
				control (A)			control (B)			Product conc.: 10%		
Vc1	50	<u></u> =	Vc1	47	<u></u> =	Vc1	48	<u> </u>	Vc1	47	<u> </u>	
Vc2	46	48	Vc2	49	48	Vc2	51	49.5	Vc2	49	48	
$30 \le \overline{\mu}$ of $Nv_0 \le 160$?			$\overline{\mu}$ of $A \ge 0.5 \times \overline{\mu}$ of Nv_0 ?			$\overline{\varkappa}$ of $B \ge 0.5 \times \overline{\varkappa}$ of Nv_0 ?			$\overline{\mu}$ of $C \ge 0.5 \times \overline{\mu}$ of Nv_0 ?			
⊠ yes □ no		⊠ yes □ no			⊠ yes □ no			⊠ yes □ no				

Test suspension (N and N_0):

	Ν	Vc1	Vc2	$\overline{\mu}$ wm = 2.25 x 10 ⁸ ;	lg N =	8.35
Ī	10 ⁻⁶	232	216	$N_0 = N/10$; $\lg N_0 =$	7.35	
	10 ⁻⁷	26	22	$7.17 \le \lg N_0 \le 7.70$?	⊠ yes	□no

Conc. of the	Contact	Vc1	Vc2	Na	lg Na	lg R
product	time			($\overline{\mu}$ x 10)		(lg N _o - lg Na)
10%	5 min	0	0	<140	<2.15	>5.20







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Results: EN 1276:2019

Test organism: Staphylococcus aureus (NCTC 10788)

Date of test: 13 April 2021 Test temperature: $20^{\circ}\text{C} \pm 1^{\circ}\text{C}$

Interfering substance: 3.0 g/l bovine albumin

Dilution-neutralisation method: Pour plate Number of plates: 1/ml Neutraliser: B Incubation temperature: $36^{\circ}C \pm 1^{\circ}C$

Validation and controls:

Validatio	n suspensio	on (Nv _o)	Experime	ntal condit	tions	Neutralis	er or filtrat	tion	Method v	Method validation (C)		
				control (A)			control (B)			Product conc.: 10%		
Vc1	50	<u></u> =	Vc1	55	<u></u> <u>n</u> =	Vc1	50	<u></u> =	Vc1	54	<u> </u>	
Vc2	53	51.5	Vc2	51	53	Vc2	53	51.5	Vc2	51	52.5	
$30 \le \overline{\varkappa}$ of $Nv_0 \le 160$?			$\overline{\mu}$ of $A \ge 0.5 \times \overline{\mu}$ of Nv_0 ?			$\overline{\mu}$ of $B \ge 0.5 \times \overline{\mu}$ of Nv_0 ?			$\overline{\mu}$ of $C \ge 0.5 \times \overline{\mu}$ of Nv_0 ?			
⊠ yes □ no			⊠ yes	□no		⊠ yes □ no			⊠ yes □ no			

Test suspension (N and N_0):

	Ν	Vc1	Vc2	$\overline{\mu}$ wm = 2.53 x 10 ⁸ ;	lg N =	8.40
ſ	10 ⁻⁶	256	240	$N_0 = N/10$; $\lg N_0 =$	7.40	
ſ	10 ⁻⁷	28	32	$7.17 \le \lg N_0 \le 7.70$?	⊠ yes	□no

Conc. of the	Contact	Vc1	Vc2	Na	lg Na	lg R
product	time			($\overline{\mu}$ x 10)		(lg N _o - lg Na)
10%	5 min	0	0	<140	<2.15	>5.25







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Results: EN 1276:2019

Test organism: Enterococcus hirae (NCIMB 8192)

Date of test: 13 April 2021 Test temperature: $20^{\circ}\text{C} \pm 1^{\circ}\text{C}$

Interfering substance: 3.0 g/l bovine albumin

Dilution-neutralisation method: Pour plate Number of plates: 1/ml Neutraliser: B Incubation temperature: $36^{\circ}C \pm 1^{\circ}C$

Validation and controls:

Validatio	n suspensio	on (Nv _o)	Experime	ntal condit	tions	Neutralis	Neutraliser or filtration			Method validation (C)		
				control (A)			control (B)			Product conc.: 10%		
Vc1	36	<u></u> =	Vc1	35	<u></u> =	Vc1	41	<u> </u>	Vc1	39	<u> </u>	
Vc2	40	38	Vc2	39	37	Vc2	37	39	Vc2	35	37	
$30 \le \overline{\mu}$ of $Nv_0 \le 160$?			$\overline{\mu}$ of $A \ge 0.5 \times \overline{\mu}$ of Nv_0 ?			$\overline{\varkappa}$ of $B \ge 0.5 \times \overline{\varkappa}$ of Nv_0 ?			$\overline{\mu}$ of $C \ge 0.5 \times \overline{\mu}$ of Nv_0 ?			
⊠ yes □ no			⊠ yes	□no		⊠ yes □ no			⊠ yes □ no			

Test suspension (N and N_o):

Ν	Vc1	Vc2	$\overline{\mu}$ wm = 2.16 x 10 ⁸ ;	lg N =	8.33
10 ⁻⁶	224	208	$N_0 = N/10$; $\lg N_0 =$	7.33	
10 ⁻⁷	21	22	$7.17 \le \lg N_0 \le 7.70$?	⊠ yes	□no

Conc. of the	Contact	Vc1	Vc2	Na	lg Na	lg R
product	time			($\overline{\mu}$ x 10)		(lg N _o - lg Na)
10%	5 min	0	0	<140	<2.15	>5.18







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Explanations:

Vc count per ml (one plate or more)

 $\overline{\mu}$ average of Vc1 and Vc2 (1 + 2 duplicate)

 $\overline{\mu}$ wm weighted mean of $\overline{\mu}$

N number of cells per ml in the test suspension

 N_0 number of cells in the test mixture at the beginning of the contact time ($N_0 = N / 10$)

Na number of survivors per ml in the test mixture at the end of the contact time (before neutralisation or

filtration)

R reduction ($\lg R = \lg N_0 - \lg Na$)

Nv number of cells per ml in the validation suspension

 Nv_0 number of cells in the validation mixtures at the beginning of the contact time ($Nv_0 = Nv / 10$)

A number of survivors per ml in the experimental conditions control mixture

B number of survivors per ml in the neutraliser or filtration control mixture

C number of survivors per ml in the method validation mixture

All test results have an associated uncertainty of measurement, details of which are available on request.