



Abbott Analytical



Consulting Scientists to the Disinfectant Industry

Certificate of Analysis

Sample(s) : One sample of Remuvit

Received from: Klenitise Ltd. Unit 59, Vale Business Park, Llandow,
Cowbridge, CF71 7PF

Date received: 23 November 2009 **Date tested:** 27 November 2009

Certificate no: 09L.148SB-KR.ENZ **Certificate date:** 4 December 2009

Sample ref: 9L/148 **Page:** 1 of 2

Analysis required: EN 13697, Chemical disinfectants and antiseptics -
Quantitative non-porous surface test for the evaluation of
bactericidal and/or fungicidal activity of chemical
disinfectants used in food, industrial, domestic and
institutional areas - Test method and requirements without
mechanical action (phase 2, step 2)

Product stored at: Room temperature

Active substance: Not declared

Test conditions: 'Clean'

Interfering substance: 0.3g/l bovine albumin +
1g/l tryptone

Product test concentration: 10.0% v/v

Product diluent used during test: Sterile hard water 300mg/l CaCO₃

Contact time: 5 minutes

Test temperature: 20°C ± 0.5°C

Neutralising solution: 30g/l polysorbate 80, 3g/l lecithin,
1g/l histidine, 1g/l cysteine

Incubation temperature: 37°C ± 1°C

**Identification of bacterial strain(s)
used:**

<i>Pseudomonas aeruginosa</i>	ATCC 15442
<i>Escherichia coli</i>	NCTC 10418
<i>Staphylococcus aureus</i>	NCTC 10788
<i>Enterococcus hirae</i>	ATCC 8043

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Page: 2 of 2

Test results:

Test Organism	<i>Pseudomonas aeruginosa</i>		<i>Escherichia coli</i>		<i>Staphylococcus aureus</i>		<i>Enterococcus hirae</i>	
Validation Suspension	10 ⁻¹	Vc1 654 Vc2 638	Vc1 314 Vc2 426	Vc1 536 Vc2 447	Vc1 411 Vc2 285			
		Nv0 6.46 x10 ³	Nv0 3.70 x10 ³	Nv0 4.92 x10 ³	Nv0 3.48 x10 ³			
Experimental Control	10 ⁰	Vc1 566 Vc2 592	Vc1 350 Vc2 308	Vc1 440 Vc2 412	Vc1 222 Vc2 306			
		A 5.79 x10 ²	A 3.29 x10 ²	A 4.26 x10 ²	A 2.64 x10 ²			
Neutraliser Control	10 ⁰	Vc1 588 Vc2 542	Vc1 372 Vc2 214	Vc1 378 Vc2 400	Vc1 344 Vc2 258			
		B 5.65 x10 ²	B 2.93 x10 ²	B 3.89 x10 ²	B 3.01 x10 ²			
Method Validation	10 ⁰	Vc1 582 Vc2 514	Vc1 300 Vc2 336	Vc1 404 Vc2 362	Vc1 277 Vc2 260			
		C 5.48 x10 ²	C 3.18 x10 ²	C 3.83 x10 ²	C 2.69 x10 ²			
Surface Inoculum	10 ⁻⁵	Vc1 536 Vc2 468	Vc1 392 Vc2 428	Vc1 356 Vc2 394	Vc1 344 Vc2 272			
	10 ⁻⁶	Vc1 54 Vc2 46	Vc1 46 Vc2 53	Vc1 44 Vc2 56	Vc1 31 Vc2 46			
		N 5.01 x10 ⁷	N 4.53 x10 ⁷	N 4.38 x10 ⁷	N 3.47 x10 ⁷			
Results	10 ⁻¹	Vc 0	Vc 0	Vc 0	Vc 0			
		Na <1.00 x10 ¹ R >5.01 x10 ⁶	Na <1.00 x10 ¹ R >4.53 x10 ⁶	Na <1.00 x10 ¹ R >4.38 x10 ⁶	Na <1.00 x10 ¹ R >3.47 x10 ⁶			
Log ₁₀ Reduction		> 6.70	> 6.66	> 6.64	> 6.54			

Vc = Viable count

Nv = cfu/ml in the validation suspension

N = cfu/ml in the surface inoculum

Na = cfu/ml in the test mixture

R = Reduction in viability

Requirements & Conclusion:

To pass EN 13697 a log₁₀ reduction of at least 4 is required.

This batch of Remuvit, when diluted to 10.0% v/v, passes the requirements of EN 13697 for bactericidal activity in 5 minutes at 20°C under 'clean' conditions against the reference organisms detailed.

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