

Test Report

Number: GZHH00358761

Applicant: Orcagel Ltd, Blackhouse Circle, Blackhouse Industrial Estate, Peterhead, AB42 1BN

Date: Apr 22, 2020

Sample Description:

One (1) style of submitted sample said to be :

Item Name : **Orcagel Hand Sanitiser Gel – 70% alcohol.**
 Client's reference information : Handsafe Hand Sanitiser Gel – 70% Alcohol.
 Item Quantity : 5pcs.
 Date Sample Received : Apr 08, 2020

Tests conducted:

As requested by the applicant, refer to attached page(s) for details.

Conclusion:

<u>Tested Sample</u>	<u>Standard</u>	<u>Result</u>
Tested component(s) of submitted sample(s)	BS 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)	Pass

Intertek GM Testing Service Zhuhai Co. Ltd.

Sarah Xu

Sarah Xu
Asst. Manager
Healthcare and Beauty Products



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1 Quantitative suspension test for evaluation of bactericidal activity of chemical disinfectants and antiseptics

With reference to BS EN 1276:2019

Dilution recommended for use:	No dilution
Product test concentration:	80% v/v
Active ingredient in product:	Alcohol
Appearance:	Blue transparent gel with bubbles
Contact time:	1 minute ± 10s
Test temperature:	20°C
Interfering substance:	0.3g/L bovine albumin (clean condition)
Inhibition method:	Dilution-neutralization
Neutralizing solution:	D/E neutralizing broth
Incubation:	(37 ± 1)° C, 48 hours
Agar medium:	Trypticase Soy Agar Escherichia coli K12 (NCTC 10538) Pseudomonas aeruginosa (ATCC 15442)
Test culture:	Staphylococcus aureus (ATCC 6538) Enterococcus hirae (ATCC 10541)



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Controls & validation:

<u>Test microorganism</u>	<u>Validation suspension (cfu/ml) N_v</u> N _{v0} =1/10N _v Criteria: 300 ≤ N_v ≤ 1600	<u>Experimental conditions control (cfu/ml)</u> A Criteria: A ≥ 0.5 N_{v0}	<u>Neutralizer control (cfu/ml)</u> B Criteria: B ≥ 0.5 N_{v0}	<u>Method validation (cfu/ml)</u> C Criteria: C ≥ 0.5 N_{v0}	<u>Validity</u>
Escherichia coli K12 (NCTC 10538)	550	54	50	46	Valid
Pseudomonas aeruginosa (ATCC 15442)	1000	100	98	96	Valid
Staphylococcus aureus (ATCC 6538)	1500	170	150	140	Valid
Enterococcus hirae (ATCC 10541)	1300	130	140	130	Valid



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Result:

Test microorganism	Initial suspension(N) (cfu/ml) $N_0=1/10N$ Criteria: $1.5 \times 10^8 \leq N \leq 5.0 \times 10^8$	Final count (cfu/ml) N_a	R(Log ₁₀ Reduction) =Log N_0 -Log N_a Criteria: $R \geq 5.0$	%Reduction Criteria: ≥ 99.999	Assessment
Escherichia coli K12 (NCTC 10538)	2.3×10^8	<140	>5.0	>99.999	Meet
Pseudomonas aeruginosa (ATCC 15442)	3.5×10^8	<140	>5.0	>99.999	Meet
Staphylococcus aureus (ATCC 6538)	4.4×10^8	<140	>5.0	>99.999	Meet
Enterococcus hirae (ATCC 10541)	2.0×10^8	<140	>5.0	>99.999	Meet

Remark:

- N = Test suspension, Number of cells per ml in bacterial suspensions.
- N_0 = ($N_0=1/10N$), 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 at the end of the contact-time.
- N_v = Validation suspension, Number of cells per ml in bacterial suspensions.
- N_{v0} = ($N_{v0} = 1/10N_v$), Number of cells per ml in the test mixtures at the beginning of the contact time (time 0).
- A,B,C = Represent the different control test mixtures, A(experimental conditions control), B(Neutralizer control), C(Method validation)



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Criteria: According to BS EN 1276:2019, in order to satisfy the requirement of bactericidal efficacy of chemical disinfectants and antiseptics, the product shall demonstrate at least a 5 decimal logarithm (lg) reduction (3 lg for handwashes) of the specified test organisms under the obligatory sample contact time, test temperature, and the simulated clean conditions according to its practical applications when the product is tested at its intended use dilution.

Sample received condition: sample in closed original package.

Date sample received: Apr 08, 2020

Testing period: Apr 08, 2020 to Apr 21, 2020

End of report

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