

Applicant: SAVEWO LIMITED Number: HKGH02870816

266-270 TEXACO RD Date: Jul 14, 2022 TSUEN WAN NT

HK

Attn: ZEN DING

Sample and Information provided by customer

FreshQ Portable Electrolytic Sterilizer – Disinfectant 3g Salt/300ml water, Mode 2 (5 mins) Item Name

For and on behalf of: Intertek Testing Services HK Ltd.

Cindy I.K. Chan Vice President







HKGH02870816 Number:

Conclusion:

The submitted sample was tested under the following requirements requested by the applicant, subject to the information stated in the remark and attached page(s) for details:

<u>Requirement</u> BS EN 14476:2019 (with modification) See details enclosed

Decision Rule(s):
When a statement of conformity to a specification or standard is provided on test report, the decision rule shall be applied. For details, please refer to Intertek's "Decision Rule Document" and is available on Intertek's website. https://intertekhk.grd.by/decision-rule-doc. If decision rule already inhered in the requested specification or standard, Intertek's "Decision Rule Document" is not applicable and indication of "..."

was shown as above table.



Number: HKGH02870816

(1) Quantitative suspension test for evaluation of virucidal activity of chemical disinfectants and antiseptics

Reference standard: BS EN 14476:2019 (with modification)

Dilution recommended for use: No dilution

Product test concentration: 80% (v/v)

Active ingredient in product: Hypochlorite

Appearance: Colorless liquid

Contact time: 10 seconds

Test temperature: 23°C

Interfering substance: 0.3 g/L bovine albumin (clean condition)

Incubation: 37°C/5% CO₂

Viral culture: SARS-CoV-2 (hCoV-19/Czech Republic/NRL_6632_2/2020)

Concentration of inoculum: 10⁶ PFU/mL







Number: HKGH02870816

Result:

Test Organism	Control Titre PFU/mL	Sample Titre PFU/mL	Log ₁₀ Reduction Criteria: R ≥ 4.0	% Reduction Criteria: R ≥ 99.99	Assessment
SARS-CoV-2					
(hCoV-19/Czech Republic/NRL_6632_2/ 2020)	283,589	0	>5.0	>99.999	Satisfactory

Criteria: According to BS EN 14476, in order to satisfy the requirement of virucidal efficacy of chemical

disinfectants and antiseptics, the product shall demonstrate at least 4.0 log₁₀ reduction of the

specified test organism.

Date sample received: Apr 29, 2022

Testing period : Apr 29, 2022 to Jul 07, 2022

End of report

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Applicant: SAVEWO LIMITED

266-270 TEXACO RD

TSUEN WAN NT

HK

ZEN DING Attn:

Sample and Information provided by customer

Item Name

FreshQ Portable Electrolytic Sterilizer - Disinfectant

Number:

Date:

3g Salt/300ml water, Mode 2 (5 mins)

Conclusion:

The submitted sample was tested under the following requirements requested by the applicant, subject to the information stated in the remark and attached page(s) for details:

Result Antibacterial Activity of Chemical Disinfectants and Antiseptics BS EN 1276:2019

Satisfactory

HKGH0285411901

Apr 19, 2022

Decision Rule(s):
When a statement of conformity to a specification or standard is provided on test report, the decision rule shall be applied. For details, please refer to Intertek's "Decision Rule Document" and is available on Intertek's website. https://intertekhk.grd.by/decision-rule-doc. If decision rule already inhered in the requested specification or standard, Intertek's "Decision Rule Document" is not applicable and indication of "..." was shown as above table.

For and on behalf of: Intertek Testing Services HK Ltd.

Cindy I.K. Chan Vice President



Number: HKGH0285411901

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

Test Standard: BS EN 1276:2019.

Dilution recommended for use: No dilution

Product test concentration: 80% (v/v)

Active ingredient in product: Hypochlorite

Appearance: Colorless liquid

Contact time: 10 seconds

Test temperature: 20°C

Interfering substance: 0.3 g/L bovine albumin (clean condition)

Inhibition method: Dilution-neutralization

Neutralizing solution: D/E neutralizing broth - double strength (sodium thioglycollate 2.0 g/L,

sodium thiosulfate 12.0 g/L, sodium bisulfite 5.0 g/L, Polysorbate 80 10.0

g/L, lecithin 14.0 g/L)

Incubation: 37°C, 48 hours

Agar medium: Trypticase Soy Agar

Test culture: Escherichia coli (ATCC 10536)

Pseudomonas aeruginosa (ATCC 15442) Staphylococcus aureus (ATCC 6538) Enterococcus hirae (ATCC 10541)

Controls & validation:

Test microorganism	$\frac{\text{Validation suspension}}{(N_v)}$ Criteria: $300 \le N_v \le 1600$	Method validation Criteria: ≥ 0.05 N _v	Validity
Escherichia coli (ATCC 10536)	865	64	Valid
Pseudomonas aeruginosa (ATCC 15442)	1460	126	Valid
Staphylococcus aureus (ATCC 6538)	1280	122	Valid
Enterococcus hirae (ATCC 10541)	815	92	Valid







HKGH0285411901 Number:

Result:

Test microorganism	Initial suspension (N) No = (1/10N) Criteria: 1.5x10 ⁸ ≤ N ≤ 5x10 ⁸	(Na)	R (Log ₁₀ Reduction) = Log No - Log Na Criteria: R ≥ 5.0	% Reduction Criteria: R ≥ 99.999	Assessment
Escherichia coli (ATCC 10536)	3.4 x 10 ⁸	<140	>5.3	>99.999	Satisfactory
Pseudomonas aeruginosa (ATCC 15442)	5.0 x 10 ⁸	<140	>5.5	>99.999	Satisfactory
Staphylococcus aureus (ATCC 6538)	4.6 x 10 ⁸	<140	>5.5	>99.999	Satisfactory
Enterococcus hirae (ATCC 10541)	3.4 x 10 ⁸	<140	>5.3	>99.999	Satisfactory

Criteria:

According to EN 1276, in order to satisfy the requirement of bactericidal efficacy of chemical disinfectants and antiseptics (for general purpose disinfection), the product shall demonstrate at least 5.0 log₁₀ reduction 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.

Date sample received: Mar 25, 2022

Testing period: Mar 28, 2022 to Apr 06, 2022

End of report

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