

GLP EFFICACY SUMMARY: RESIDUAL DISINFECTANT

CLAIM	TEST METHOD	MICROBE	TEST RESULTS
Residual Disinfectant for up to 24 hours	U.S. EPA's Interim Method (Modified EPA 01-1A)	Pseudomonas aeruginosa (ATCC #15442)	EPA Acceptance Criteria: >= 5 log reduction in =< 10 minutes Result: P. aeruginosa: >6-log ₁₀ reduction (>99.9999%) in 10 minutes
		Staphylococcus aureus (ATCC #6538)	EPA Acceptance Criteria: >= 5 log reduction in =< 10 minutes Result: >5-log ₁₀ reduction (>99.999%) in 10 minutes
		Human Coronavirus 229E (ATCC # VR740)	EPA Acceptance Criteria: >= 3 log reduction in =< 10 minutes Result: >4-log ₁₀ reduction (>99.99%) in 10 minutes

GLP EFFICACY SUMMARY: LIQUID DISINFECTANT

CLAIM	TEST METHOD	MICROBE	TEST RESULTS
Hospital/Healthcare Disinfectant	ASTM E2197	Pseudomonas aeruginosa (ATCC #15442) Staphylococcus aureus (ATCC #15442)	Acceptance Criteria: >5-log ₁₀ reduction Result: P. aeruginosa: >6-log ₁₀ reduction (>99.9999%) in 3 minutes S. aureus: >6-log ₁₀ reduction (>99.9999%) in 3 minutes
Broad-spectrum Virucide	ASTM E1053	Human Adenovirus type 5 (ATCC #VR-5)	Acceptance Criteria: >4-log ₁₀ reduction Result: >5-log ₁₀ reduction (>99.999%) in 10 minutes
Human Coronavirus 229E as Virus Surrogate for SARS-CoV-2	ASTM E2197	Human Coronavirus 229E (ATCC # VR740)	Acceptance Criteria: >4-log ₁₀ reduction Result: >5-log ₁₀ reduction (>99.999%) in 2 minutes