

CLEANING LABORATORY EVALUATION SUMMARY

SCL #: 2023
 DateRun: 02/23/2024
 Experimenters: Alicia McCarthy, Aditi Patel, Namrata Chauhan, Serena Burkinshaw
 ClientType: Lab
 ProjectNumber: Project #12
 Substrates: Stainless Steel
 PartType: Coupon
 Contaminants: MS2 Bacteriophage
 Cleaning Methods: Pour Plate
 Analytical Methods: Organism count
 Purpose: To evaluate the efficacy of Thymol, glycolic acid, and plant based cleaners with inactivating MS2 on a hard surface without agitation.

Experimental Procedure: Pour Plate Method - MS2 Bacteriophage
 Six hours prior to the run, E.coli 15597 was subcultured into three milliliters of tryptic soy broth (TSB) screw-cap tubes and incubated at 37°C (98.6°F). 27 screw-cap tubes filled with 10ml of 0.5X tryptic soy agar (TSA) were autoclaved. The biosafety cabinet (BSC) was sprayed with 70% v/v isopropyl alcohol using a paper towel before spraying any items going into the BSC. Once autoclaving was complete, the TSA tubes were placed into a 45°C (113°F) D.I. water bath inside the biosafety cabinet (BSC). The four glass Petri dishes were marked using a black sharpie to designate the positive (P+), negative (N-), Test 1 (T1), and Test 2 (T2). Ten microliters of the organism were pipetted onto the P+, T1, and T2 stainless steel coupons and air-dried for 15 minutes. A motorized pipette with 10ml tips was used to pipet 15 ml of Dey-Engley (D/E) neutralizing broth into four separate 50ml conical tubes labeled P+, N-, T1, and T2. Once the MS2 bacteriophage dried on the coupons, the P+ coupon was placed into the conical tube. The N-, T1, and T2 were pipetted with 1000µl of the cleaning solution onto each coupon for 30 seconds before immediately placing them in the conical tube with an autoclaved forceps. The conical tubes were then placed on the shaker for 10 minutes. During this time, using the 1000ml pipette, 900ml of 1x phosphate-buffered saline (PBS) was pipetted into nine autoclaved dilution tubes, and serial dilutions were made for P+, T1, and T2 up to 10⁻⁴ using 100µl of the shaken D/E broth. Once the six-hour sub-time was complete, the E. coli 15597 subculture was removed from the incubator for use. For each variable (N-, P+, T1, and T2), 100µl of the stock and serial dilutions of MS2 bacteriophage, and 100µl of the E.coli 15597 subculture were combined into an empty dilution tube. A screwcap tube of 0.5X TSA was removed from the water bath, wiped with a paper towel to remove moisture, and poured into the dilution tube. The mixture was immediately poured into a sterile polystyrene petri dish; swirled to cover the entire plate surface, and then air-dried before covering. Dried Petri dishes were placed into a clean labeled zip lock bag that was partially closed and incubated at 37°C overnight. Plates were counted the following day based on the clear lysis zones in the bacterial lawn of growth (1 plate forming unit) to calculate log reduction and percent removal.

Results:

Product	Log Reduction	% Reduction
Seventh Generation Disinfecting Multisurface Cleaner	4.0012	99.9762
Mrs. Meyers Clean Day Multisurface Cleaner	0.0245	5.4055
PineSol Multisurface Cleaner	0.5688	64.2883
Everspring All Purpose Cleaner	0.8465	83.2228

Summary:

Substrates:		Stainless Steel				
Contaminants:		MS2 Bacteriophage				
Company Name:	Product Name:	Conc.:	Efficiency:	Effective:	Observations:	
Clorox Company	Pine Sol	100%	64.29	<input type="checkbox"/>		
Seventh Generation	Disinfecting Multisurface Cleaner Lemongrass Citrus	100%	99.98	<input checked="" type="checkbox"/>		
Mrs Myers Clean Day	Mrs Meyers Lemon Multisurface Cleaner	100	5.40	<input type="checkbox"/>		
Guy & O'Neill, Inc.	Everspring Citrus & Basil All Purpose Cleaner	100%	83.22	<input type="checkbox"/>		

Conclusion:

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Seventh Generation Disinfecting Multisurface cleaner had the highest log reduction of 4.0012 and % reduction of 99.9762 (claims to kill 99.9% of bacteria and viruses). Mrs. Meyers, PineSol, and Everspring had a log reduction below 1.000 and % reduction below 90%.