

A Comparison Between Manual and Automated Microbial Testing Methods

Organized by: Neutec group

Moderator: Dr. Brady Carter, Neutec Group, Senior Application Scientist

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Moderator Bio



Brady Carter
Neutec Group

Dr. Brady Carter is a Senior Application Scientist with Neutec Group. He specializes in Water Activity and Moisture Sorption applications.

Dr. Carter earned his Ph.D. and M.S Degree in Food Engineering and Crop Science from Washington State University and a B.A. Degree in Botany from Weber State University. Dr. Carter has 20 years of experience in research and development has been the instructor for water activity seminars in over 23 different countries and has provided on-site water activity training for companies around the world.

He has authored over 20 white papers on water activity, moisture sorption isotherms, and complete moisture analysis. He has participated in hundreds of extension presentations and has given talks at numerous scientific conferences.

He developed the shelf life simplified paradigm and hygrothermal time shelf life model and is the leading expert in applying water activity to shelf life prediction.

Presenter Bio



Blanca Ruiz
University of Georgia

Blanca Ruiz is a Research Coordinator at the University of Georgia in the Department of Food Science & Technology.

Prior to that, she was an Associate in research at Washington State University working with fresh produce safety.

She also got her master's degree at Washington State University focusing on environmental monitoring programs to control Listeria in apple packinghouses.



College of Agricultural &
Environmental Sciences
UNIVERSITY OF GEORGIA

Food Microbiology lab – Department of Food Science and Technology

Principal investigator: Dr. Faith Critzer

Research Coordinator: Blanca Ruiz-Llacsahuanga

Postdoctoral Researcher: Martha Sanchez

Graduate students: 1 PhD, 4 MS , 2 undergraduate students



Peru



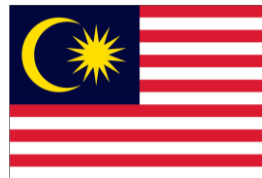
Colombia



Lebanon



India



Malaysia



USA



Food Microbiology lab

(Biosafety level II)

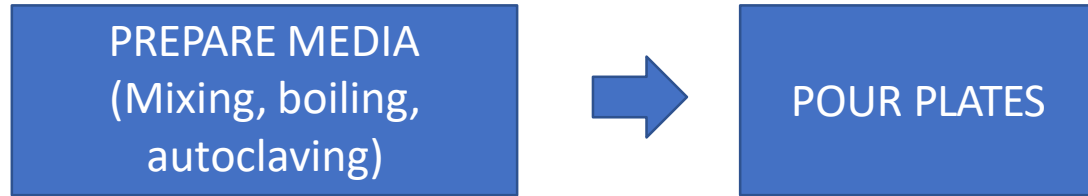
Microbiological and molecular analysis related to bacterial foodborne pathogens to improve fresh-produce safety:

- Ongoing projects:
 1. Evaluating the efficacy of commercially available sanitizers in reducing *Escherichia coli* and *Salmonella* in dump tank water used in packinghouses.
 2. Inactivation of foodborne pathogens on organic apples by application of antimicrobial waxes containing essential oils.
 3. Effective strategies to sanitize apple harvesting supplies against *Listeria monocytogenes*, *STEC* and *Salmonella*.

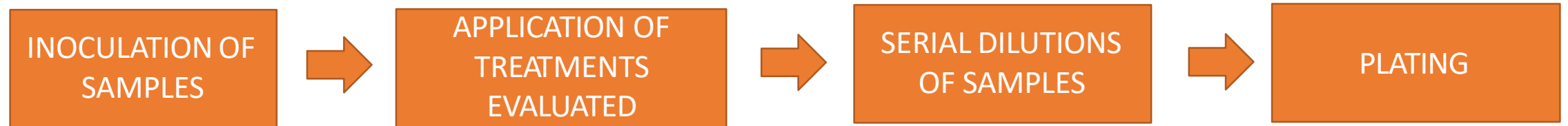


Microbial testing procedures overflow

Prepping for experiments:



Day of experiment:



24h, 48h after:



Types of automation implemented in our lab

1. Media preparation

- To prepare up to 9 L of media (around 500 plates at once)

2. Media filler

- Faster plate pouring
- Keeps consistency in the amount of media poured in each plate.

3. Spiral plating

- To reduce the amount of media plates and dilutions used.

4. Automated colony counter

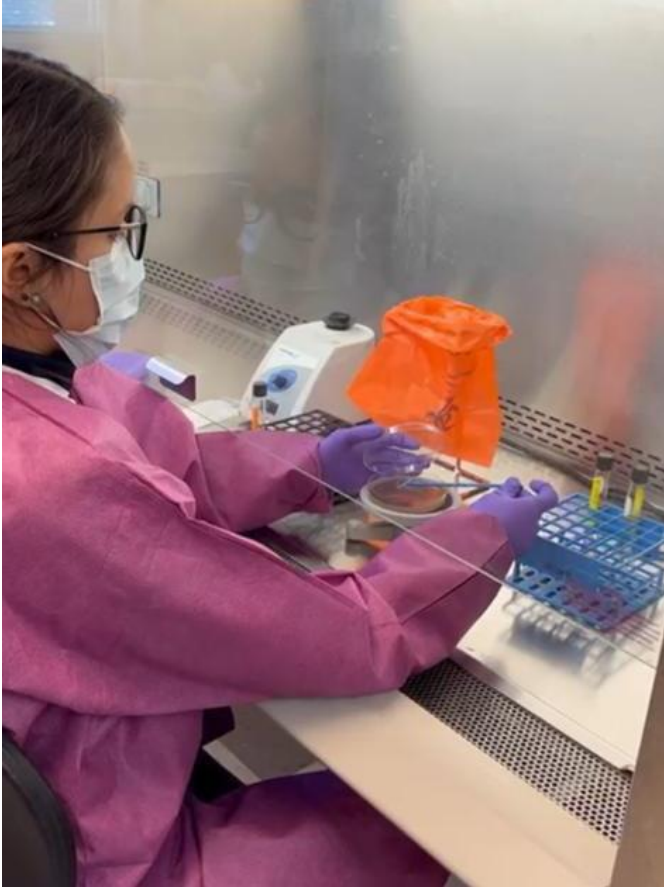
- Keep consistency , reduce labor hours



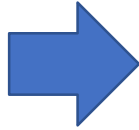
Media preparator and filler



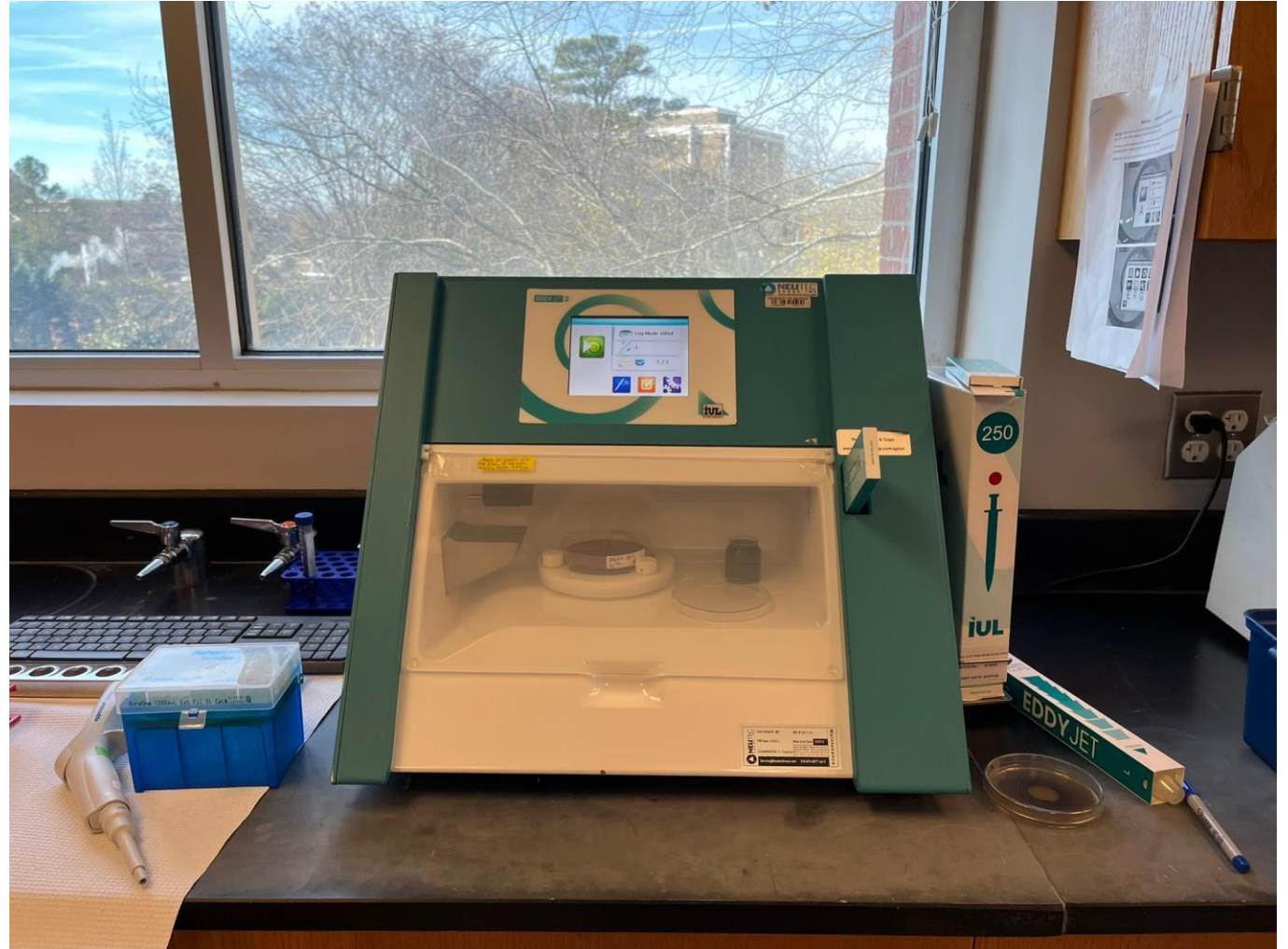
Spread plating



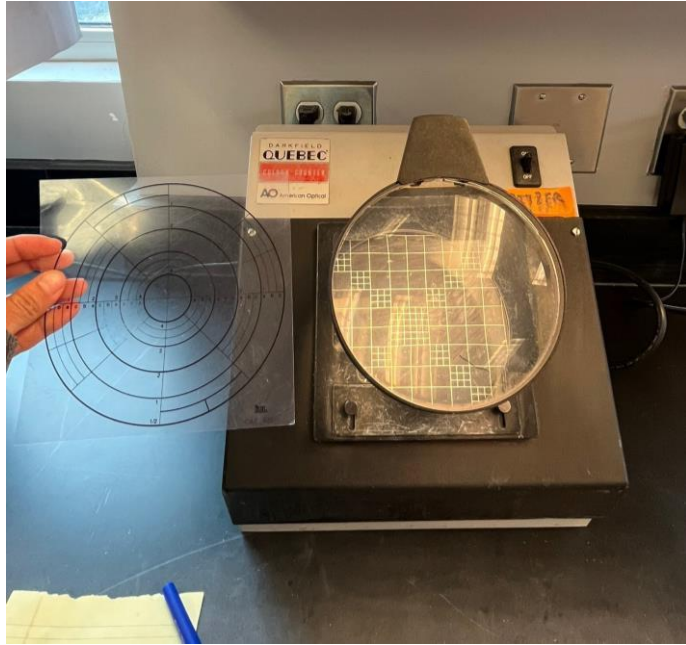
SPIRAL PLATER



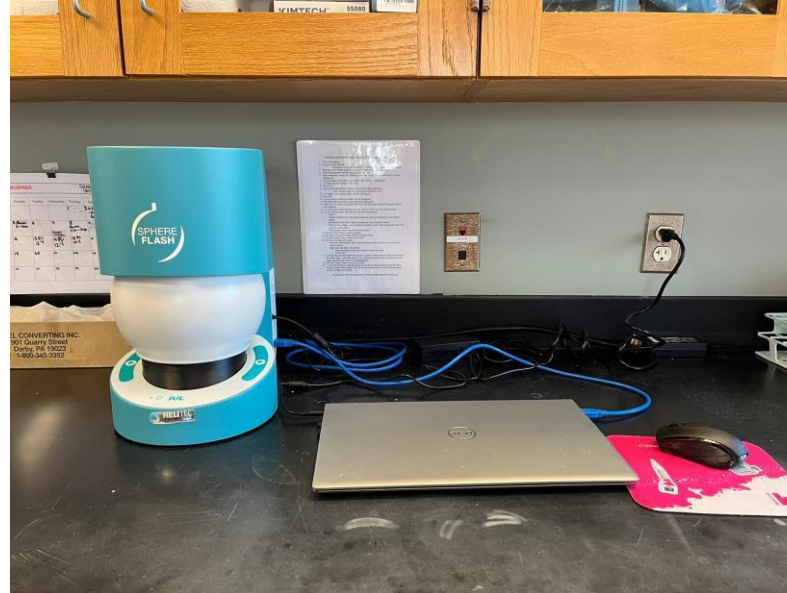
Spiral plating



Manual counting



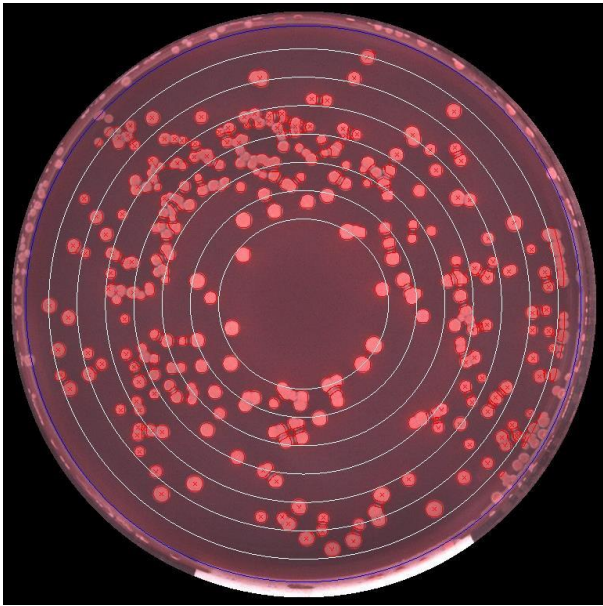
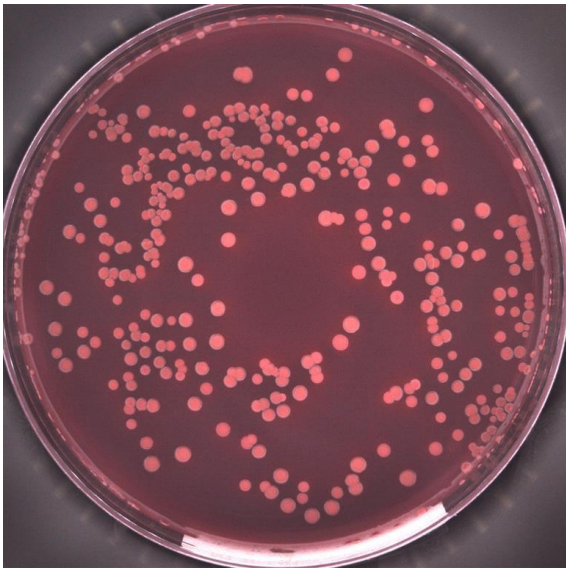
Automated colony counter



COLONY COUNTER

Example: Excel spreadsheet and plate captures.

Plate Id	Counted Colonies	Counted Volume	Concentration	Dilution Factor	Inoculated Volume	Inoculation mode
200 ppm –R1 plate 2	162	0.098400004	1646341.393	1000	0.0984	Log Mode 100ul
200 ppm R1 - plate 1	236	0.059999999	3933333.421	1000	0.0984	Log Mode 100ul
200 ppm R2 - plate 2	161	0.021119999	7623106.505	1000	0.0984	Log Mode 100ul
200 ppm R2- plate 1	269	0.036399998	7390110.343	1000	0.0984	Log Mode 100ul
200 ppm R3 - plate 2	249	0.036399998	6840659.76	1000	0.0984	Log Mode 100ul
200 ppm R3 - plate 1	175	0.021119999	8285985.332	1000	0.0984	Log Mode 100ul



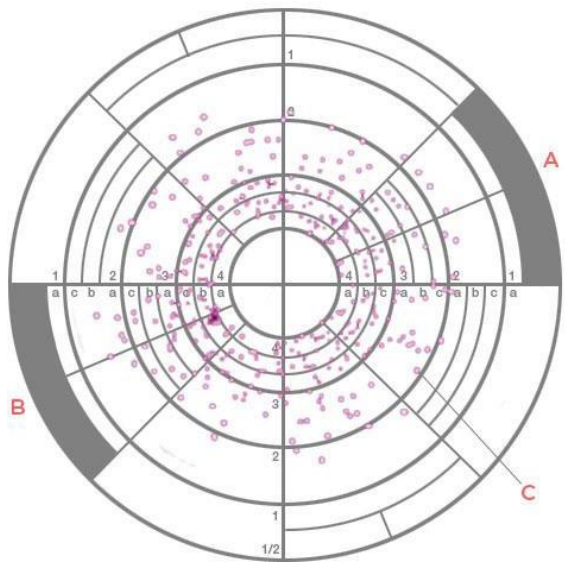
Challenge:

- Setting up methods for each microorganism and type of media.

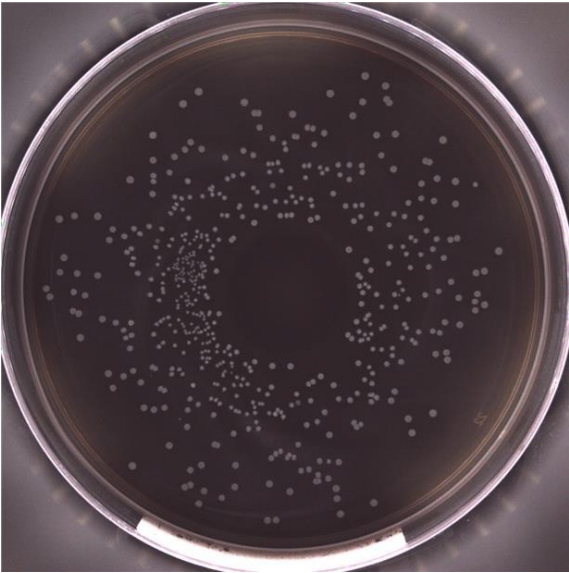
COLONY COUNTER - RESULTS

E. Coli – Macconkey agar plates

Manual counting (Grid and formulas)

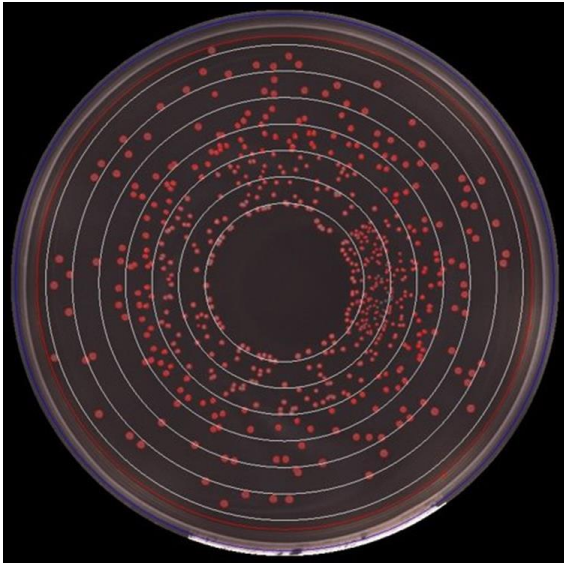


ID	Dilution	Section	N'	N''	N'+N''	V/4	Concentration	Log (CFU/ml)
R1 - plate 1	0.001	3cba	37	27	64	0.01932	3312629.4	6.56
R1 - plate 2	0.001	3cba	22	15	37	0.01932	1915113.872	6.35
R2 - plate 1	0.001	3cb	22	22	44	0.01932	2277432.712	6.41
R2 - plate 2	0.001	3cb	20	16	36	0.01932	1863354.037	6.34
R3 - plate 1	0.001	3cba	24	20	44	0.01932	2277432.712	6.41
R3 - plate 2	0.001	3cba	30	19	49	0.01932	2536231.884	6.45



Automated colony counter

ID	Automated colony counter Log (CFU/ml)
R1 - plate 1	6.88
R1 - plate 2	6.86
R2 - plate 1	6.83
R2 - plate 2	6.84
R3 - plate 1	6.82
R3 - plate 2	6.76



COLONY COUNTER - RESULTS

Listeria monocytogenes – MOX plates

Salmonella – CHROMagar plates



Salmonella Newport concentrations (Log CFU/ml)

ID	Colony counter	Manual counting
R1	9.128	9.117
R1	9.147	9.158
R2	8.833	8.799
R2	8.598	8.613
R3	8.484	8.415
R3	8.405	8.301

SPIRAL PLATER MODE: Log mode 100 μ L

COLONY COUNTER -
RESULTS

Take away message:

- Automation in our lab has shown significant great benefits reducing labor hours, keeping consistency during experiments, and less use of resources.
- SOPs for a correct use of the devices is always needed. Training of personnel using equipment is essential.
- Maintenance and calibration of equipment are essential.
- As anything in science challenges will still be present.



Questions

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