

MICROBIOLOGICAL MEDIA QUALITY PERFORMANCE REPORT # 24042

Test Media: E. coli / Coliform Count Plate

Brand: 3M Petrifilm™

Batch No. / Expiry Date: 418323334C/2025-05-30

Reference Media: Standard Plate Count (SPC) Agar

Brand: Oxoid

Batch No. / Expiry Date: 3573499 / 30 Nov 2027

Date Tested: 14 Mar 2024

Project No: 24-65542-1A

	BATCH PERFOR	MANCE CERT	FIFICATE		
Qualitative Analysis		Quantitative Analysis			
Test Organism	Observed Growth	-	- Test Organism		
Staphylococcus aureus NZRM 87	No Growth	-	Escherichia coli NZRM 916	-	
<i>Escherichia coli</i> NZRM 916	Positive (<i>E. coli</i>)	Ave. No. Colonies	104.2	-	
<i>Enterococcus faecalis</i> NZRM 1106	No Growth	3M Petrifilm			
Pseudomonas aeruginosa NZRM 918	Negative	Ave. No. Colonies	103.6	-	
Enterobacter cloacae NZRM 2375	Positive (Coliform)	SPC			
Blank	No Growth	% Difference	0.6%	-	
Positive: Typical growth as described by method. Negative: Growth not described as typical.		Quantitative analysis compares the recovery rate of the test media to a reference media.			
<u>No Growth:</u> No visible colonies observed. <u>Method:</u> Direct inoculation i.e. 1ml of a diluted overnight culture was used to inoculate the test media.		[Ave. No. of]	Calculation: The percentage difference = [<u>Ave. No. of Test Media Colonies x 100</u>] - 100 Ave. No. of Reference Media Colonies		
Blank: 5 plates inoculated with 1ml sterile diluent.		Method: In Ho	Method: In House		
Incubation Conditions: Aerobic 35°C/24 hours		Incubation Co	Incubation Conditions: Aerobic 35°C/24 hours		

Quantitative Interpretation: The media performance is considered acceptable if the percentage difference between the test and non-selective reference media does not exceed ±30%.

Qualitative Interpretation: The media performance is considered acceptable if both positive growth of the target organism(s) and negative or no growth of non-target organism(s) is expressed.

The above criteria are testing laboratory In-house limits, which have been endorsed by 3M as a report interpretation guideline.

Results apply to samples of the Batch as received by the laboratory.

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Authorised by:

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