AsureQuality Kaitiaki Kai

MICROBIOLOGICAL MEDIA QUALITY PERFORMANCE REPORT # 26105

Test Media: *E. coli* / Coliform Count Plate

Brand: Neogen Petrifilm™ **Batch No. / Expiry Date:** 418326088C/2027-09-27

Reference Media: Standard Plate Count (SPC) Agar

Brand: Oxoid **Batch No. / Expiry Date:** 6288928 / Jul 2030

Date Tested: 11 Aug 2026 **Project No:** 26-223377-1A

BATCH PERFORMANCE CERTIFICATE

Qualitative Analysis

Test Organism	Observed Growth
Staphylococcus aureus NZRM 87	No Growth
Escherichia coli NZRM 916	Positive (*E. coli*)
Enterococcus faecalis NZRM 1106	No Growth
Pseudomonas aeruginosa NZRM 918	Negative
Enterobacter cloacae NZRM 2375	Positive (*Coliform*)
Blank	No Growth

Positive: Typical growth as described by method.

Negative: Growth not described as typical.

No Growth: No visible colonies observed.

Method: Direct inoculation i.e. 1ml of a diluted overnight culture was used to inoculate the test media.

Blank: 5 plates inoculated with 1ml sterile diluent.

Incubation Conditions: Aerobic 35°C/24 hours

Quantitative Analysis

-	Test Organism	
-	*Escherichia coli* NZRM 916	-
Ave. No. Colonies Neogen Petrifilm	66.4	-
Ave. No. Colonies SPC	60.4	-
% Difference	9.9%	-

Quantitative analysis compares the recovery rate of the test media to a reference media.

Calculation: The percentage difference =

$$\frac{[\text{Ave. No. of Test Media Colonies} \times 100]}{\text{Ave. No. of Reference Media Colonies}} - 100$$

Method: In House

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 Neogen as a report interpretation guideline.

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

Authorised by: Michele

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Scientific Analyst

Report Date: 18 Aug 2026

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