# MICROBIOLOGICAL MEDIA
# QUALITY PERFORMANCE REPORT # 24141

**Test Media:** Rapid Yeast and Mould Count Plate

**Brand:** 3M Petrifilm™ **Batch No. / Expiry Date:** 33Y6EX/2025-12-25

**Reference Media:** Yeast Dextrose Chloramphenicol (YDC) Agar

**Brand:** Millipore **Batch No. / Expiry Date:** VM988800 /Oct 2026

**Date Tested:** 30 Oct 2024 **Project No:** 24-290583-1A

## BATCH PERFORMANCE CERTIFICATE

### Qualitative Analysis

| Test Organism | Observed Growth |
|---|---|
| *Staphylococcus aureus* NZRM 87 | No Growth |
| *Escherichia coli* NZRM 916 | No Growth |
| *Saccharomyces cerevisiae* NZRM 2241 | Positive |
| *Pseudomonas aeruginosa* NZRM 918 | No Growth |
| *Aspergillus brasiliensis* NZRM 2578 | Positive |
| 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 25°C/2 days

### Quantitative Analysis

| - | Test Organism | |
|---|---|---|
| - | *Saccharomyces cerevisiae* NZRM 2241 | - |
| Ave. No. Colonies 3M Petrifilm | 79.2 | - |
| Ave. No. Colonies YDC | 81.2 | - |
| % Difference | -2.5% | - |

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 25°C/2 days

**Quantitative Interpretation:** The media performance is considered acceptable if the percentage difference between the test and non-selective reference media does not exceed ±10%.

**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:

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Scientific Analyst

Report Date: 06 Nov 2024

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