

Certificate of Analysis

Certificate No: BMDX/COA/P/240/2025

Product Name: Mueller Hinton II Agar

Product Code: FP90M2001

Lot No: 251209001

Expiry date: 17/03/2026

Date of Analysis: 11/12/2025 – 12/12/2025

Parameter Requirement: Physical, Chemical and Microbiological Analysis

| Physical & Chemical Test | Specification | Results |
|--------------------------|---|---------|
| Color | Light to medium, yellow to tan, trace hazy to slightly hazy | Passed |
| pH | 7.30 ± 0.1 | Passed |
| Hardness | Semi-solid with a firm, smooth, & bounces surface | Passed |
| Pour Depth/Thickness | 4.00-4.50mm | Passed |

| Microbiological Test | Antimicrobial Agent | Acceptable range (mm or (µg/mL) | Zone Diameter (mm) or MIC (µg/mL) |
|---|---|---------------------------------|-----------------------------------|
| <i>Escherichia coli</i> ATCC® 25922 | Gentamicin 10 µg | 19-26 | 22 |
| | Amikacin 30 µg | 19-26 | 22 |
| | Ampicillin 10 µg | 15-22 | 16 |
| | Ceftazidime 30 µg | 25-32 | 28 |
| | Tigecycline 15 µg | 20-27 | 26 |
| <i>Pseudomonas aeruginosa</i> ATCC® 27853 | Ceftazidime 30 µg | 22-29 | 25 |
| | Amikacin 30 µg | 20-26 | 24 |
| | Cefepime 30 µg | 25-31 | 27 |
| | Imipenem 10 µg | 20-28 | 22 |
| | Ciprofloxacin 5 µg | 25-33 | 26 |
| | Gentamicin 10 µg | 17-23 | 21 |
| | Colistin | 0.25-2 | 2.0 |
| <i>Enterococcus faecalis</i> ATCC® 29212 | Trimethoprim-sulfamethoxazole 1.25/23.75 µg | ≥ 20 | 25 |
| | Nitrofurantoin 300 µg | 18-24 | 22 |
| | Penicillin | 1-4 | 3.0 |
| <i>Staphylococcus aureus</i> ATCC® 25923 | Penicillin 10 units | 26-37 | 30 |
| | Gentamicin 10 µg | 19-27 | 23 |
| | Erythromycin 15 µg | 22-30 | 24 |
| | Clindamycin 2 µg | 24-30 | 25 |
| | Trimethoprim-sulfamethoxazole 1.25/23.75 µg | 24-32 | 26 |
| <i>Staphylococcus aureus</i> ATCC® 29213 | Vancomycin | 0.5-2 | 0.75 |
| Uninoculated plate | No growth | - | - |

Test Statement:

1. Incubation at $35 \pm 2^{\circ}\text{C}$ for 24 hours in ambient air.
2. Antibiotic susceptibility tests are performed in accordance with, and meet the acceptance limits of, the current ISO/TS 16782. Performance is assessed using CLSI methodology.

Sample statement:

1. Culture media were prepared according to procedure SP-PD-BMDX-02 Production Process Control and SP-PD-BMDX-11 Product Specification.
2. Culture media were kept at temperature conditions $2-8^{\circ}\text{C}$.
3. QC Organisms were prepared and followed according to procedure SP-QC-BMDX-07 Quality Control Testing and SP-PD-BMDX-11 Product Specification.

Result:

The information given is believed to be correct, all results reported in this certificate relate only to the product in this certificate of analysis according to SP-PD-BMDX-11 Product Specification.

Sign by:

Fairuz Ayuni Binti Abdul Halim
QC Executive
BSc. in Microbiology with Honours
Universiti Putra Malaysia



Biomed MDX Sdn Bhd
8, Jalan IAN 3, Industri Angkasa Nuri,
76100 Durian Tunggal, Melaka, Malaysia

Date: 15/12/2025

