

## Certificate of Analysis

Certificate No: BMDX/COA/P/416/2026

**Product Name:** Mueller Hinton II Agar  
**Product Code:** FP90M2001  
**Lot No:** 260427001  
**Expiry date:** 03/08/2026  
**Date of Analysis:** 27/04/2026 – 30/04/2026  
**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)
<b><i>Escherichia coli</i> ATCC® 25922</b>	Gentamicin 10 µg	19-26	25
	Amikacin 30 µg	19-26	24
	Ampicillin 10 µg	15-22	19
	Ceftazidime 30 µg	25-32	27
	Tigecycline 15 µg	20-27	24
<b><i>Pseudomonas aeruginosa</i> ATCC® 27853</b>	Ceftazidime 30 µg	22-29	22
	Amikacin 30 µg	20-26	25
	Cefepime 30 µg	25-31	26
	Imipenem 10 µg	20-28	24
	Ciprofloxacin 5 µg	25-33	26
	Gentamicin 10 µg	17-23	21
	Colistin	0.25-2	2.0
<b><i>Enterococcus faecalis</i> ATCC® 29212</b>	Vancomycin 5 µg	10-16	14
	Trimethoprim-sulfamethoxazole 1.25/23.75 µg	≥ 20	26
	Nitrofurantoin 300 µg	18-24	22
	Penicillin	1-4	2
<b><i>Staphylococcus aureus</i> ATCC® 25923</b>	Penicillin 10 units	26-37	30
	Gentamicin 10 µg	19-27	25
	Erythromycin 15 µg	22-30	24
	Clindamycin 2 µg	24-30	24
	Trimethoprim-sulfamethoxazole 1.25/23.75 µg	24-32	26
<b><i>Staphylococcus aureus</i> ATCC® 29213</b>	Vancomycin	0.5-2	0.75
<b>Uninoculated plate</b>	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:**

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QC Executive  
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**Date:** 30/04/2026

