



Instructions for Use

CHOCOLATE AGAR WITH BACITRACIN

Cat. no. E11	Chocolate with Bacitracin, 15x100mm Plate, 18ml	10 plates/bag
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INTENDED USE

Hardy Diagnostics Chocolate Agar with Bacitracin is recommended for use in the isolation and cultivation of *Haemophilus* species from respiratory specimens containing mixed flora.

SUMMARY

Chocolate Agar with Bacitracin consists of GC Agar Base with added hemoglobin and KoEnzyme Enrichment. GC Agar Base contains proteose peptone which provides nitrogenous nutrients. When heated, hemoglobin releases hemin (X-factor), a factor required by fastidious organisms such as *Haemophilus* species. The phosphate buffer is added to maintain the pH of the medium. Corn starch aids in neutralizing any toxic fatty acids present in the medium. KoEnzyme Enrichment is a chemically defined enrichment that provides NAD (V-factor), amino acids, vitamins, dextrose, ferric ions, and coenzymes which promote growth. Bacitracin inhibits most strains of streptococci, staphylococci, *Neisseria* and *Micrococcus* species. Because *Haemophilus* spp. are resistant to the concentration of bacitracin employed in this medium, Chocolate Agar with Bacitracin allows for their increased recovery, due to the decreased competition for nutrients in the medium.

FORMULA

Ingredients per liter of deionized water:*

Chocolate Agar with Bacitracin:				
Proteose Peptone	15.0gm			
Sodium Chloride	5.0gm			
Dipotassium Phosphate	4.0gm			
Monopotassium Phosphate	1.0gm			
Corn Starch	1.0gm			
Hemoglobin, Bovine	10.0gm			
KoEnzyme Enrichment	10.0ml			
Bacitracin	20,000U			
Agar	10.0gm			
KoEnzyme Enrichments:				
Dextrose	10.0gm			

L-Cystine HCl	2.59gm
L-Glutamine	1.01gm
L-Cystine	0.11gm
Adenine	0.101gm
Nicotinic Adenine Dinucleotide	25.0mg
Cocarboxylase	10.0mg
Guanine Hydrochloride	3.0mg
Ferric Nitrate	2.0mg
P-Aminobenzoic Acid	1.3mg
Vitamin B ₁₂	1.0mg
Thiamine	0.3mg

Final pH 7.2 +/- 0.2 at 25°C.

STORAGE AND SHELF LIFE

Storage: Upon receipt store at 2-8°C. away from direct light. Media should not be used if there are any signs of deterioration (shrinking, cracking, or discoloration), contamination, or if the expiration date has passed. Product is light and temperature sensitive; protect from light, excessive heat, moisture, and freezing.

The expiration date on the product label applies to the product in its intact packaging when stored as directed. The product may be used and tested up to the expiration date on the product label and incubated for the recommended incubation times as stated below.

Refer to the document "Storage" for more information.

PRECAUTIONS

This product may contain components of animal origin. Certified knowledge of the origin and/or sanitary state of the animals does not guarantee the absence of transmissible pathogenic agents. Therefore, it is recommended that these products be treated as potentially infectious, and handle observing the usual Universal Precautions for blood. Do not ingest, inhale, or allow to come into contact with skin.

This product is for *in vitro* diagnostic use only. It is to be used only by adequately trained and qualified laboratory personnel. Observe approved biohazard precautions and aseptic techniques. All laboratory specimens should be considered infectious and handled according to "standard precautions." Refer to the document "Guidelines for Isolation Precautions" from the Centers for Disease Control and Prevention.

For additional information regarding specific precautions for the prevention of the transmission of all infectious agents from laboratory instruments and materials, and for recommendations for the management of exposure to infectious disease, refer to CLSI document M29: *Protection of Laboratory Workers from Occupationally Acquired Infections*.

Sterilize all biohazard waste before disposal.

Refer to the document "Precautions When Using Media" for more information.

PROCEDURE

^{*} Adjusted and/or supplemented as required to meet performance criteria.

Specimen Collection: Consult listed references for information on specimen collection. (1-3,5,7) Infectious material should be submitted directly to the laboratory without delay and protected from excessive heat and cold. If there is to be a delay in processing, the specimen should be inoculated onto an appropriate transport media and maintained at an appropriate temperature. (1-3,5,7)

Method of Use: Inoculate the medium and streak the specimen as soon as possible after it is received in the clinical laboratory. If the specimen is being cultured directly from a swab, roll the swab over a small portion of the agar surface, and streak for isolation. Incubate in 5-10% CO₂ at $35-37^{\circ}$ C. for 24 to 48 hours. Extend incubation if needed. Examine plate for typical colonial morphology and characteristics.

INTERPRETATION OF RESULTS

Consult listed references for the interpretation of growth of fastidious species. (1-3,5,7)

LIMITATIONS

It is recommended that biochemical, immunological, molecular, or mass spectrometry testing be performed on colonies from pure culture for complete identification of bacteria and/or fungi.

E. coli, some Neisseria species, strains of Candida spp., Klebsiella, Proteus, and Pseudomonas spp., as well as other bacteria may grow on this medium.

Refer to the document "Limitations of Procedures and Warranty" for more information.

MATERIALS REQUIRED BUT NOT PROVIDED

Standard microbiological supplies and equipment such as loops, other culture media, swabs, applicator sticks, incinerators, and incubators, etc., as well as serological and biochemical reagents, are not provided.

QUALITY CONTROL

Hardy Diagnostics tests each lot of commercially manufactured media using appropriate quality control microorganisms and quality specifications as outlined on the Certificate of Analysis (CofA) and the CLSI document M22-A3 *Quality Assurance for Commercially Prepared Microbiological Culture Media*. The following microorganisms are routinely used for testing at Hardy Diagnostics:

Test Organisms	Inoculation	Incubation			Results
Test Organisms	Method*	Time	Temperature	Atmosphere	Results
Haemophilus influenzae ATCC® 10211	A	24hr	35°C	CO ₂ **	Growth
Staphylococcus epidermidis ATCC® 12228	В	24hr	35°C	Aerobic	Inhibited

^{*} Refer to the document "Inoculation Procedures for Media OC" for more information.

USER QUALITY CONTROL

End users of commercially prepared culture media should perform QC testing in accordance with applicable government regulatory agencies, and in compliance with accreditation requirements. Hardy Diagnostics recommends end users check for signs of contamination and deterioration and, if dictated by laboratory quality control procedures or regulation, perform quality control testing to demonstrate growth or a positive reaction and to demonstrate inhibition or a negative reaction, if applicable. Hardy Diagnostics quality control testing is documented on the certificate of analysis (CofA) available from Hardy Diagnostics Certificate of Analysis website. Also refer to the document "Finished Product Quality Control Procedures," and the CLSI document M22-A3 Quality Assurance for Commercially Prepared

Microbiological Culture Media for more information on the appropriate QC procedures. See the references below.

** Atmosphere of incubation is enriched with 5-10% CO₂.

PHYSICAL APPEARANCE

Chocolate Agar with Bacitracin should appear opaque, and brown in color.



Haemophilus influenzae (ATCC $^{\circledR}$ 10211) colonies growing on Chocolate Agar with Bacitracin (Cat. no. E11). Incubated in CO₂ for 24 hours at 35°C.



Staphylococcus epidermidis (ATCC[®] 12228) growth inhibited on Chocolate Agar with Bacitracin (Cat. no. E11). Incubated aerobically for 24 hours at 35°C.

REFERENCES

- 1. Anderson, N.L., et al. *Cumitech 3B; Quality Systems in the Clinical Microbiology Laboratory*, Coordinating ed., A.S. Weissfeld. American Society for Microbiology, Washington, D.C.
- 2. Jorgensen., et al. Manual of Clinical Microbiology, American Society for Microbiology, Washington, D.C.
- 3. Tille, P., et al. Bailey and Scott's Diagnostic Microbiology, C.V. Mosby Company, St. Louis, MO.
- 4. Carpenter, C.M. and H.E. Morton. 1947. Proc. N.Y. State Assoc. Public Health Labs, 27:58-60.
- 5. Isenberg, H.D. *Clinical Microbiology Procedures Handbook*, Vol. I, II & III. American Society for Microbiology, Washington, D.C.
- 6. Johnston, et al. 1945. J. Ven. Dis. Inf.; 26:239.
- 7. Koneman, E.W., et al. *Color Atlas and Textbook of Diagnostic Microbiology*, J.B. Lippincott Company, Philadelphia, PA.
- 8. Maritn, J.E., et al. 1967. Public Health Rep.; 82:361.
- 9. McLeod, J.W., et al. 1927. Br. J. Exp. Pathol.; 8:25.
- 10. *Quality Assurance for Commercially Prepared Microbiological Culture Media*, M22-A2, Vol. 16, No.16. 1996. Clinical Laboratory Standards Institute (CLSI formerly NCCLS), Villanova, PA.

ATCC is a registered trademark of the American Type Culture Collection.

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