# **CORNMEAL AGAR w/ 1% DEXTROSE**

### **INTENDED USE**

Remel Cornmeal Agar w/ 1% Dextrose is a solid medium recommended for use in qualitative procedures for the cultivation of fungi and the chromogenesis of dermatophytes.

### SUMMARY AND EXPLANATION

In 1970, Hazen and Reed used Cornmeal Agar to isolate pathogenic fungi.<sup>1</sup> Cornmeal Agar w/ 1% Dextrose is recommended to demonstrate chromogenesis of some species of *Trichophyton.*<sup>2,3</sup> Identification of *Trichophyton* to the species level is necessary to allow correct treatment of diseases, such as tinea capitis, where drug dosage levels or treatment times may require species-specific adjustment.<sup>4</sup>

#### PRINCIPLE

Cornmeal Agar w/ 1% Dextrose consists of cornmeal extract and agar. The cornmeal extract provides the essential nutrients needed to support the growth of fungi. Agar is the solidifying agent. Dextrose is added to differentiate *Trichophyton rubrum*, which produces a red pigment on Cornmeal Agar w/ 1% Dextrose, from *Trichophyton mentagrophytes* which produces no pigment.

# **REAGENTS (CLASSICAL FORMULAE)\***

Cornmeal Extract	g g	Agar	g าI

pH 6.0 +/- 0.2 @ 25°C

\*Adjusted as required to meet performance standards.

### **PROCEDURE<sup>5</sup>**

- 1. Inoculate Cornmeal Agar w/ 1% Dextrose from a pure culture of the test isolate growing on primary isolation agar other than Cornmeal Agar.
- 2. Using a sterile inoculating needle or sterile teasing needle, inoculate a small fragment of the fungus colony onto the agar plate.
- 3. Incubate in ambient at 25-30°C for up to 14 days.
- 4. Examine at regular intervals for growth and pigmentation.

Agar Deep: Melt the agar deep in a boiling water bath and cool to 45-50°C. Mix and dispense into a sterile petri dish and proceed with the instructions above.

### **QUALITY CONTROL**

All lot numbers of Cornmeal Agar w/ 1% Dextrose have been tested using the following quality control organisms and have been found to be acceptable. Testing of control organisms should be performed in accordance with established laboratory quality control procedures. If aberrant quality control results are noted, patient results should not be reported.

#### CONTROL

Trichophyton rubrum ATCC<sup>®</sup> 38484 Trichophyton mentagrophytes ATCC<sup>®</sup> 9533

# INCUBATION

Ambient, up to 14 days @ 25-30°C Ambient, up to 14 days @ 25-30°C **RESULTS** Growth with red pigment

Growth with no pigment

# LIMITATIONS

1. Cornmeal agar with dextrose is not recommended for the production of chlamydospores by Candida albicans.<sup>2</sup>

#### BIBLIOGRAPHY

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- 2. MacFaddin, J.F. 1985. Media for Isolation-Cultivation-Identification-Maintenance of Medical Bacteria. Williams and Wilkins, Baltimore, MD.
- 3. Larone, D.H. 2011. Medically Important Fungi, A Guide to Identification. 5th ed. ASM Press, Washington, D.C.
- 4. Versalovic, J., K.C. Carroll, G. Funke, J.H. Jorgensen, M.L. Landry, and D.W. Warnock. 2011. Manual of Clinical Microbiology. 10<sup>th</sup> ed. ASM Press, Washington, D.C.
- 5. Conant, N.F., D.T. Smith, R.D. Baker, and J.L. Callaway. 1971. Manual of Clinical Mycology. W.B. Saunders Co., Philadelphia, PA.

Refer to the front of Remel *Technical Manual of Microbiological Media* for **General Information** regarding precautions, product storage and deterioration, specimen collection, storage and transportation, materials required, quality control, and limitations.

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