



## Potato Dextrose Agar

M096

Potato Dextrose agar is recommended for the isolation and enumeration of yeasts and moulds from dairy and other food products.

### Composition\*\*

Ingredients	Gms / Litre
Potatoes, infusion from	200.000
Dextrose	20.000
Agar	15.000
Final pH ( at 25°C)	5.6±0.2

\*\*Formula adjusted, standardized to suit performance parameters

### Directions

Suspend 39 grams in 1000 ml distilled water. Heat to boiling to dissolve the medium completely. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Mix well before dispensing. In specific work, when pH 3.5 is required, acidify the medium with sterile 10% tartaric acid / lactic acid. The amount of acid required for 100 ml of sterile, cooled medium is approximately 1 ml. Do not heat the medium after addition of the acid.

### Principle And Interpretation

Potato Dextrose Agar is recommended by APHA (1) and F.D.A. (2) for plate counts of yeasts and moulds in the examination of foods and dairy products (3). Potato Dextrose Agar is also used for stimulating sporulation, for maintaining stock cultures of certain dermatophytes and for differentiation of typical varieties of dermatophytes on the basis of pigment production (4). It is also recommended by USP (5), BP (6), EP (7) and JP (8) for growth of fungi.

Potato infusion and dextrose promote luxuriant fungal growth. Adjusting the pH of the medium by tartaric acid to 3.5, inhibits the bacterial growth. Heating the medium after acidification should be avoided as it may hydrolyse the agar which can render the agar unable to solidify.

### Quality Control

#### Appearance

Cream to yellow homogeneous free flowing powder

#### Gelling

Firm, comparable with 1.5% Agar gel

#### Colour and Clarity of prepared medium

Light amber coloured clear to slightly opalescent gel forms in Petri plates

#### Reaction

Reaction of 3.9% w/v aqueous solution at 25°C (after sterilization).pH:-5.6±0.2

#### pH

5.40-5.80

#### Cultural Response

M096: Cultural characteristics observed after incubation at 20-25 °C for 2-5 days. Recovery rate is considered as 100% for fungus growth on Sabouraud Dextrose Agar.

Organism	Inoculum (CFU)	Growth	Observed Lot value (CFU)	Recovery	Incubation temperature	Incubation period
<i>Candida albicans</i> ATCC 10231	50 -100	luxuriant	35 -100	≥70 %	20 -25 °C	2 -3 d

* <i>Aspergillus brasiliensis</i> ATCC 16404	50 -100	luxuriant	25 -100	>=70 %	20 -25 °C	5 -7 d
<i>Aspergillus fumigatus</i> ATCC 9197	50 -100	luxuriant	25 -100	>=70 %	20 -25 °C	5 -7 d
<i>Saccharomyces cerevisiae</i> ATCC 9763	50 -100	luxuriant	35 -100	>=70 %	30 -35 °C	2 -5 d
<i>Rhodotorula mucilaginosa</i> DSM 70403		luxuriant			20 -25 °C	3 -5 d
<i>Geotrichum candidum</i> DSM 1240		good- luxuriant			25 -30 °C	3 -5 d
<i>Penicillium commune</i> ATCC 10248		fair -good			25 -30 °C	3 -5 d
<i>Trichophyton ajelloi</i> ATCC 28454		fair-good			25 -30 °C	3 -7 d
<i>Fusarium solani</i> ATCC 36031		luxuriant			20 -25 °C	3 -5 d

\*Key:-Formerly known as *Aspergillus niger*

### Storage and Shelf Life

Store below 30°C in tightly closed container and the prepared medium at 2 - 8°C. Use before expiry date on the label.

### Reference

1. Downes F. P. and Ito K., (Eds.), 2001, Compendium of Methods for the Microbiological Examination of Foods, 4th Ed., APHA, Washington, D.C.
2. FDA Bacteriological Analytical Manual, 2005, 18th Ed., AOAC, Washington, DC.
3. Wehr H. M. and Frank J. H., 2004, Standard Methods for the Microbiological Examination of Dairy Products, 17th Ed., APHA Inc., Washington, D.C.
4. MacFaddin J. F., 1985, Media for the Isolation-Cultivation-Identification-Maintenance of Medical Bacteria, Vol.1, Williams and Wilkins, Baltimore
5. The United States Pharmacopoeia, 2016, The United States Pharmacopoeial Convention. Rockville, MD.
6. British Pharmacopoeia, 2016, The Stationery office British Pharmacopoeia
7. European Pharmacopoeia, 2014, European Dept. for the quality of Medicines.
8. Japanese Pharmacopoeia, 2008.

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