

**POTATO DEXTROSE AGAR****TM 344**

For cultivation and enumeration of yeasts and molds

**Composition**

Ingredients	Gms/Ltr.
Dextrose	20.00
Agar	15.00
Potato infusion from	4.00

\* Dehydrated powder, hygroscopic in nature, store in a dry place, in tightly-sealed containers below 25°C and protect from direct Sunlight.

**[\*4.0gm of potato extract is equivalent to 200gm of infusion from potatoes]**

**Instructions for use**

Dissolve 39gms in 1000ml distilled water. Gently heat to boiling with gentle swirling and dissolve the medium completely. Sterilize by autoclaving at 15 psi (121°C) for 15 minutes. Cool at room temperature and dispense into sterile petri - plates.

**Appearance:** Light yellow colour, slightly opalescent

**pH (at 25°C):** 5.6 ± 0.2

**Principle**

**POTATO DEXTROSE AGAR** is used for the isolation and enumeration and culturing of yeast and molds from samples. It can also be used in the identification of fungi and yeasts in parallel with their cellular morphology or in methods of micro cultivation in slides. Dextrose and Potato infusion (BEEVER and BOLLARD 1970) promote the growth of yeasts and moulds while the low pH value partially inhibits the growth of the accompanying bacterial flora. Agar is a solidifying agent. If the medium is to be used for fungal counts and enumeration of yeasts and mold, the pH should be adjusted to approximately 3.5 to inhibit bacterial count. By the addition of sterilized 10% tartaric acid solution to 1000ml of sterilized medium to obtain a pH of 3.5. Fungi grow on this medium to develop typical morphology. This general-purpose medium can be supplemented with acid or antibiotics to inhibit bacterial growth.

**NOTE:** Do not re-heat the adjusted medium after adding the acid because the agar may hydrolyses and may not solidify.

**Interpretation**

Cultural characteristics observed after inoculating (10<sup>3</sup>CFU/ml), on incubation at 25°C for 3 - 5 days (fungi) and at 30- 35°C for 72 hours (yeast & molds).

Microorganisms	ATCC	Inoculum (CFU/ml)	Growth
* <i>Aspergillus brasiliensis</i>	16404	Point inoculation	Good
<i>Candida albicans</i>	10231	10 <sup>3</sup>	Good
<i>Sacchromyces cerevisiae</i>	9763	10 <sup>3</sup>	Good
<i>Trichophyton sp.</i>	9533	10 <sup>3</sup>	Good
<i>Trichophyton rubrum</i>	28188	10 <sup>3</sup>	Good

Note:\* - Formerly known as *Aspergillus niger*.



## References

1. Mac Faddin, J. F. Media for isolation-cultivation-identification-maintenance of medical bacteria, vol.1. Williams & Wilkins, Baltimore, MD. (1985).
2. Marshall, (ed.). Standard methods for the examination of dairy products, 16th ed. American Public Health Association, Washington, D.C. (1993).
3. Association of Official Analytical Chemists. Bacteriological analytical manual, 8th ed. AOAC International, Gaithersburg, MD. (1995).
4. American Public Health Association. Recommended Methods for the Microbiological Examination of Foods. APHA). New York. (1958).
5. Bacteriological analytical manual, 8th ed. AOAC International. Gaithersburg, MD. European Pharmacopoeia 6th edition. (2007).