

Dehydrated Culture Media Bases / Media Supplements

# **Technical Information**

## **Malt Extract Broth Base**

### Product Code: DM 1255

Application: - Malt Extract Broth Base is recommended for the detection, isolation and enumeration of yeasts and moulds.

Composition**	
Ingredients	Gms / Litre
Malt extract	17.000
Mycological peptone	3.000
Mycological peptone Final pH (25°C)	5.4±0.2

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

#### Principle & Interpretation

The laboratory diagnosis of fungal infection mainly depends on direct verses indirect methods. Considerable importance should be given to direct microscopy in addition to isolation of the organisms. The use of malt and malt extracts for the propagation of yeasts and moulds is quite common. Reddish <sup>(1)</sup> described a culture medium prepared from malt extract that was a satisfactory substitute for wort. Malt Extract Medium is similar to the formula of Galloway and Burgess <sup>(2)</sup> used for the detection, isolation and enumeration of yeasts and moulds. Malt Extract Broth is recommended for the examination of yeasts and moulds in the U.S. Food and Drug Administrations Bacteriological Analytical Manual <sup>(2)</sup>. For mycological counts preforably more acidic medium is prepared which can suppress bacterial growth.

Malt extract provides an acidic environment and nutrients favourable for growth and metabolism of yeasts and moulds. Mycological peptone rapidly gives a luxuriant growth with typical morphology and pigmentation. For mycological count, it is advisable to adjust the reaction of medium more acidic with addition of 10% lactic acid. In order to suppress bacterial growth antibiotics may be added as sterile solutions to the molten medium immediately before dispensing into sterile tubes <sup>(3)</sup>

Malt Extract Broth Base has been widely used in the maintenance, isolation and identification of fungi and it is also proposed in several pharmacopeias as a medium for the control of sterility in pharmaceutical products, though it is mostly used for comparative morphological studies.

### Methodology

Suspend 20 grams of powder media in 1000 ml distilled water and soak for 15 minutes. Shake well & heat to dissolve the medium completely. Sterilize by autoclaving at 10 lbs pressure (115°C) for 10 minutes. Mix well before dispensing. Avoid overheating. If desired, to adjust acidic pH use 10% Lactic Acid (MS2095).

## Quality Control

Physical Appearance Cream to yellow homogeneous free flowing powder

**Colour and Clarity of prepared medium** Amber coloured clear solution in tubes

Reaction

Reactionof 2.0% w/v aqueous solution at 25°C.pH:-5.4±0.2

**pH range** 5.20-5.60

Cultural Response/ characteristices DM 1255: Cultural characteristics observed after an incubation at 25-30°C for 48-72 hours.





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Organism	Inoculum (CFU)	Growth
*Aspergillus brasiliensis ATCC 16404	50-100	luxuriant
Candida albi cans ATCC 10231	50-100	luxuriant
Saccharomyces cerevisiae ATCC 9763	50-100	luxuriant
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\*Key: Formerly known as Aspergillus niger ATCC 16404

### Storage and Shelf Life

**Dried Media:** Store below 30°C in tightly closed container and use before expiry date as mentioned on the label. **Prepared Media**: 2-8° in sealable plastic bags for 2-5 days.

#### **Further Reading**

- 1. Reddish A., 1919, Abstr. Bacteriol., 3:6.
- 2. FDA Bacteriological Analytical Manual, 2005, 18th Ed., AOAC, Washington, DC.
- 3. Gallowey L. D. and Burgess R., 1952, Applied Mycology and Bacteriology, 3rd Ed., Leonard Hill, London, pg. 54 and 57.

#### **Disclaimer**:

- User must ensure suitability of the product(s) in their application prior to use.
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