

# TM 1380 - M-BCG YEAST AND MOLD AGAR

## **INTENDED USE**

For the detection of fungi in routine analysis of beverages using membrane filter techinque.

### PRODUCT SUMMARY AND EXPLANATION

The microbiology of beverages will vary greatly depending upon the method of processing and the means of preservation. High microbial populations often indicate poor quality in raw material, unsanitary equipments or opportunity for growth in the food at some stage in the process. Heat processed beverages will be free of aciduric microorganism but may yield low numbers of viable spore forming bacteria when cultured on non-selective media. Bacteria cannot grow in the high acid environment and therefore direct microscopic count for yeast, bacteria or moulds may provide a clue to the conditions of sanitization during processing. Heat resistant spores may be present in low numbers. Because of their slow growth and poor competitive ability, yeast and moulds often manifest themselves on or in foods in which the environment is less favourable for bacterial growth.

M-BCG (Bromocresol Green) Yeast and Mould Agar is used for the detection of fungi in routine analysis of beverages using membrane filter technique.

### **COMPOSITION**

Ingredients	Gms / Ltr	
Yeast extract	9.000	
Dextrose	50.000	
Biopeptone	10.000	
Magnesium sulphate	2.100	
Potassium phosphate	2.000	
Diastase	0.050	
Thiamine hydrochloride	0.050	
Bromocresol green	0.026	
Agar	15.000	

### **PRINCIPLE**

The medium is highly nutritious for the growth of yeasts and moulds. Biopeptone and yeast extract provide nitrogenous compounds and vitamin B complex. Thiamine is also a B vitamin in the medium. Dextrose acts as the energy source. Diastase is a mixture of amylolytic enzymes. Bromocresol green is the pH indicator, which is green at acidic pH (pH 4.0) while blue at pH 5.6. Potassium phosphate helps in maintaining buffering action in the medium. The low pH inhibits bacterial growth. The membrane filter is directly placed on the agar surface of M-BCG Yeast and Mould Agar and incubated at 30-35°C for 48 hours.

## **INSTRUCTION FOR USE**

- Dissolve 8.82 grams in 100 ml distilled water.
- Heat to boiling to dissolve the medium completely.
- Dispense and sterilize by autoclaving at 12 15 psi pressure (118 121°C) for 10 minutes.

# **QUALITY CONTROL SPECIFICATIONS**















Appearance of Powder: Cream to yellow homogeneous free flowing powderAppearance of prepared medium: Green coloured opalescent gel forms in Petri plates

**pH (at 25°C)** : 4.6±0.2

### INTERPRETATION

Cultural characteristics observed after an incubation.

Microorganism	ATCC	Inoculum (CFU/ml)	Growth	Recovery	Incubation Temperature	Incubation Period
Aspergillus niger	16404	50-100	Good-luxuriant	>=50 %	25 - 30°C	48 - 72 Hours
Candida albicans	10231	50-100	Good-luxuriant	>=50 %	25 - 30°C	48 - 72 Hours
Saccharomyces cerevisiae	9763	50-100	Good-luxuriant	>=50 %	25 - 30°C	48 - 72 Hours

# **PACKAGING:**

In pack size of 500 gm bottles.

## **STORAGE**

Dehydrated powder, hygroscopic in nature, store in a dry place, in tightly-sealed containers between 25-30°C and protect from direct sunlight. Under optimal conditions, the medium has a shelf life of 4 years. When the container is opened for the first time, note the time and date on the label space provided on the container. After the desired amount of medium has been taken out replace the cap tightly to protect from hydration.

**Product Deterioration:** Do not use if they show evidence of microbial contamination, discoloration, drying or any other signs of deterioration.

# **DISPOSAL**

After use, prepared plates, specimen/sample containers and other contaminated materials must be sterilized before discarding.

# **REFERENCES**

1. MacFaddin J.F., 1985, Media for Isolation - Cultivation - Identification - Maintenance of Medical Bacteria, Vol. I, Williams and Wilkins, Baltimore.















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**NOTE:** Please consult the Material Safety Data Sheet for information regarding hazards and safe handling Practices. \*For Lab Use Only Revision: 08 Nov., 2019







