

Dehydrated Culture Media Bases / Media Supplements

# **Technical Information**

# Inhibitory Mold Agar, Ulrich

## Product Code: DM 1246

Application: - Inhibitory Mould Agar, Ulrich is used for selective isolation of pathogenic fungi.

Composition**		
Ingredients	Gms / Litre	
Casein enzymic hydrolysate	3.000	
Peptic digest of animal tissue	2.000	
/east extract	5.000	
Dextrose	5.000	
Starch, soluble	2.000	
Dextrin	1.000	
Sodium phosphate	2.000	
Ferrous sulphate	0.040	
Magnesium sulphate	0.800	
Sodium chloride	0.040	
Manganese sulphate	0.160	
Chloramphenicol	0.125	
Agar	15.000	
Final pH ( at 25°C)	6.7±0.2	
**Formula adjusted, standardized to suit pe	rformance	
parameters		

### Principle & Interpretation

Fungi with the potential to cause human diseases constitute a very small group and belong to the genera Aspergillus, Candida, Cryptococcus, Histoplasma and Pneumocystis. Based on their methods of reproduction members of pathogenic fungi group are classified into four taxonomic group's viz. Zygomycetes, Basidiomycetes, Ascomycetes and Deuteromycetes (Fungi Imperfecti)<sup>(2)</sup>. To confirm the existence and nature of infection by fungi and yeasts, Identification of the organisms is much more useful than demonstrating the humoral and cellular responses of the host<sup>(1)</sup>. Inhibitory Mould Agar formulated by Ulrich<sup>(3)</sup> is used as a general-purpose medium for the selective isolation and cultivation of pathogenic fungi.

Casein enzymic hydrolysate and peptic digest of animal tissue provide essential growth nutrients. Yeast extract is a rich source of vitamin B complex. Dextrose, starch and dextrin are energy sources for the metabolism of fungi. Sodium chloride and metallic salts provide essential ions and minerals. Chloramphenicol inhibits a wide variety of gram-positive and gram-negative bacteria. Potential contaminants of cosmetics and toiletries like *Pseudomonas aeruginosa* and *Serratia marcescens* are effectively inhibited by chloramphenicol. Sodium phosphates buffer the medium.

## Methodology

Suspend 36.17 grams of powder media in 1000 ml distilled water. Mix thoroughly and heat to dissolve the medium completely. Sterilize by autoclaving at 118 - 121°C for 15 minutes. Mix well and pour into sterile Petri plates.





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## **Quality Control**

#### Physical Appearance

Cream to yellow homogeneous free flowing powder

#### Gelling

Firm, comparable with 1.5% Agar gel

#### Colour and Clarity of prepared medium

Amber coloured, clear to slightly opalescent gel forms in Petri plates.

#### Reaction

Reaction of 3.62% w/v aqueous solution at 25°C. pH : 6.7±0.2

#### pH Range 6.50-6.90

#### Cultural Response/Characteristics

DM 1246: Cultural characteristics observed after an incubation at 25-30°C for upto 7 days ii) Bacterial cultures are incubated at 35-37°C.

Organism	Inoculum (CFU)	Growth	Recovery
Candida albi cans ATCC 10231	50-100	luxuriant	>=50%
Escherichia coli ATCC 25922	>=10 <sup>°</sup>	luxuriant	0%
Staphylococcus aureus ATCC 25923	>=10	luxuriant	0%
Trichop hyton mentagrophytes ATCC 9533	50-100	luxuriant	

### 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<sup>0</sup> in sealable plastic bags for 2-5 days.

### **Further Reading**

1.Cruikshank R., Marmion B. P., Duguid J. P., Swain R.H.A., (Eds.), Medical Microbiology, 12th Edition, Vol. II, Churchill Livingstone 2.Frey D., Oldfield R. J., Bridger R. C., A Colour Atlas of Pathogenic Fungi, Wolfe Medical Publications, London. 3.Ulrich J. A., 1956, Bact. Proc., S.A.B., M75: 87.

### **Disclaimer**:

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