

# **Technical Information**

### **NIH Agar**

### Product Code: DM 1194

**Application:** NIH Agar is used for sterility testing and for the cultivation and maintenance of isolates from sterility testing of biological products.

Composition\*\*

| Composition                                      |                |  |
|--|----------------|--|
| Ingredients                                      | Gms / Litre    |  |
| Casein enzymic hydrolysate                       | 15.000         |  |
| Yeast extract                                    | 5.000          |  |
| Dextrose   | 5.500          |  |
| Sodium chloride                                  | 2.500          |  |
| L-Cystine  | 0.050          |  |
| Agar   | 15.000         |  |
| Final pH ( at 25°C)                              | 7.1±0.2        |  |
| **Formula adjusted standardized to suit performa | nce narameters |  |

# Principle & Interpretation

This medium can be used for sterility testing and also for cultivating the isolates from biological products tested for sterility. This medium is also recommended by the National Institute of Health (NIH) for sterility testing of biological products of turbid nature <sup>(3)</sup>. NIH Medium has a similar composition as Fluid Thioglycollate Medium, except sodium thioglycollate has resazurin and agar concentration is more in NIH Medium than in Fluid Thioglycollate Medium NIH Agar is formulated according to the agar medium specified by USPHS sterility test <sup>(1)</sup>.

NIH medium is a nutritious medium containing nutrients like casein enzymic hydrolysate, yeast extract and the amino acid L-cystine. It contains the fermentable carbohydrate dextrose and sodium chloride for maintaining osmotic equilibrium. NIH Medium is devoid of sodium thioglycollate. U.S. Pharmacopoeia (2) has recommended using this medium with sodium thioglycollate (0.05%) or thioglycollic acid (0.03%) for the sterility testing of biological products containing mercurial preservatives, since sodium thioglycollate neutralizes the bacteriostatic effect of mercuric compounds (4,5).

## Methodology

Suspend 43.05 grams of powder media in 1000 ml distilled water. Shake well and heat to dissolve the medium completely. Dispense into test tubes or flasks as desired. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes.

## **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

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

#### Reaction

Reaction of 4.3% w/v aqueous solution at 25°C. pH: 7.1±0.2

pH Range 6.90-7.30

### Cultural Response/ characteristices

DM 1194: Cultural characteristics observed after an incubation at 35-37°C for 18-24 hours and with addition of sodium thioglycollate.





| Organism                          | Inoculum (CFU) | Growth         | Recovery |
|-----------------------------------|----------------|----------------|----------|
| Escherichia coli ATCC 25922       | 50-100         | good-luxuriant | >=70%    |
| Streptococcus mitis ATCC 9895     | 50-100         | good-luxuriant | >=70%    |
| Staphylococcus aureus ATCC 25923  | 50-100         | good-luxuriant | >=50%    |
| Streptococcus pyogenes ATCC 19615 | 50-100         | good-luxuriant | >=70%    |
| Bacillus subtilis ATCC 6633       | 50-100         | good-luxuriant | >=50%    |
| Bacteroides vulgatus ATCC 8482    | 50-100         | good-luxuriant | >=50%    |
| Candida albi cans ATCC 10231      | 50-100         | good-luxuriant | >=50%    |
| Micrococcus luteus ATCC 9341      | 50-100         | good-luxuriant | >=50%    |
| Clostridium sporogenes ATCC 11437 | 50-100         | good-luxuriant | >=50%    |

### 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. USPHS Reg., 73, 730: Federal Register, 1970, Vol. 35, No. 0171, p. 13:930.
- 2. The United States Pharmacopoeia, 2006, USP29/NF24. The United States Pharmacopoeial Convention, Rockville, MD.
- 3. MacFaddin J. F., 1985, Media for Isolation-Cultivation-Identification-Maintenance of Medical Bacteria, Vol. 1, Williams and Wilkins, Baltimore
- 4. Nungester, Hood and Warren, 1943, Proc. Soc. Exp. Biol. Med. 52: 287
- 5. Portwood, 1944, J. Bacteriol., 48: 255.

### Disclaimer:

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