

# **Technical Data**

## Thioglycollate Medium w/o Dextrose

**M190** 

Thioglycollate Medium without Dextrose is used for cultivation of aerobes, microaerophiles, anaerobes and for fermentation studies with various carbohydrates.

## Composition\*\*

Ingredients	Gms / Litre
Casein enzymic hydrolysate	20.000
Sodium chloride	2.500
Dipotassium phosphate	1.500
Sodium thioglycollate	0.600
L-Cystine	0.400
Sodium sulphite	0.200
Methylene blue	0.002
Agar	0.500
Final pH ( at 25°C)	7.2±0.2

<sup>\*\*</sup>Formula adjusted, standardized to suit performance parameters

#### **Directions**

Suspend 25.7 grams in 1000 ml distilled water. If the medium is to be used for fermentation studies or for diagnostic work adds 0.5 to 1% carbohydrate of choice. Heat to boiling to dissolve the medium completely. Dispense and sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Alternatively, sterile carbohydrate solutions may be added to the broth after sterilization. The prepared medium should be stored in the dark at room temperature.

Note: If more than the upper one-third has acquired a green colour, the medium may be restored once by heating in a waterbath or free flowing steam until the green colour disappears.

## **Principle And Interpretation**

Thioglycollate Medium without Dextrose is the modification of original Thioglycollate medium (1, 2) used for the fermentation study of anaerobes and for enhancement of sporulation. Omission of dextrose facilitates it to be used in fermentation studies with the addition of desired carbohydrate. Some Clostridia remain viable for a longer period and sporulate better in the absence of carbohydrate and thus this medium could be used for sporulations.

Casein enzymic hydrolysate, L-cystine and salts provide essential nutrients like nitrogenous compounds, carbon, sulphur, minerals and amino acids. The reducing action provided by sodium thioglycollate and sodium sulphite binds molecular oxygen, thereby maintaining a low Eh (3). A small amount of agar is added to retard the absorption of oxygen by reducing convection currents in the medium. Methylene blue is a redox indicator.

## **Quality Control**

## **Appearance**

Cream to yellow homogeneous free flowing powder

## Colour and Clarity of prepared medium

Light yellow coloured very slightly opalescent viscous solution with upper 10% or less medium green on standing

#### Reaction

Reaction of 2.57% w/v aqueous solution at 25°C. pH: 7.2±0.2

рH

7.00-7.40

#### **Cultural Response**

M190: Cultural characteristics observed after an incubation at 35-37°C for 48 hours(in an appropriate atmosphere).

Organism Inoculum Growth

(CFU)

HiMedia Laboratories Technical Data

#### **Cultural Response**

Bacillus subtilis ATCC 6633	3 50-100	good
Bacteroides vulgatus ATCC	50-100	fair
8482		
Candida albicans ATCC	50-100	good
10231		
Clostridium sporogenes	50-100	good-luxuriant
ATCC 11437		
Micrococcus luteus ATCC	50-100	good
10240		
Neisseria meningitidis ATC	C50-100	good
13090		
Streptococcus pyogenes	50-100	good-luxuriant
ATCC 19615		-

## **Storage and Shelf Life**

Store below 30°C and the prepared medium at 2-8°C. Use before expiry date on the label.

#### Reference

1.Brewer J. H., 1940, J. Am Med. Assoc., 115, 598.

2.Brewer J. H., 1940, J. Bacteriol., 39:10.

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

Revision: 02 / 2015

#### Disclaimer:

User must ensure suitability of the product(s) in their application prior to use. Products conform solely to the information contained in this and other related HiMedia<sup>™</sup> publications. The information contained in this publication is based on our research and development work and is to the best of our knowledge true and accurate. HiMedia<sup>™</sup> Laboratories Pvt Ltd reserves the right to make changes to specifications and information related to the products at any time. Products are not intended for human or animal or therapeutic use but for laboratory, diagnostic, research or further manufacturing use only, unless otherwise specified. Statements contained herein should not be considered as a warranty of any kind, expressed or implied, and no liability is accepted for infringement of any patents.