

# THIOGLYCOLLATE MEDIUM w/ INDICATOR and DEXTROSE

## INTENDED USE

Remel Thioglycollate Medium w/ Indicator and Dextrose (Fluid Thioglycollate Medium) is a liquid medium recommended for use in qualitative procedures as a general purpose medium for the cultivation of aerobes and anaerobes.

## SUMMARY AND EXPLANATION

Thioglycollate Medium was first described by Brewer in 1940.<sup>1</sup> The medium contains a small amount of agar and a reducing substance to initiate the growth of anaerobes. Marshall et al. reported satisfactory cultivation of anaerobes in this medium in the presence of mercurial preservatives.<sup>2</sup> This observation was confirmed by Nungester et al., who demonstrated the neutralization of the bacteriostatic effect of mercurial compounds by sodium thioglycollate.<sup>3</sup>

## PRINCIPLE

Sodium thioglycollate is a reducing agent which removes molecular oxygen from the medium and prevents the accumulation of peroxides which may be lethal to some microorganisms. Sulfhydryl groups inactivate mercury and other heavy metals neutralizing the antibacterial effect of mercurial preservatives. A small amount of agar is added to impede diffusion of oxygen. Casein peptone and cystine supply nitrogen and carbon compounds necessary for the growth of aerobes and anaerobes. Sodium chloride is a source of essential electrolytes and maintains osmotic equilibrium. Resazurin is an oxidation-reduction indicator which turns pink when increased oxidation occurs.

## REAGENTS (CLASSICAL FORMULA)\*

Casein Peptone.....	15.0 g	L-Cystine.....	0.5 g
Dextrose .....	5.0 g	Sodium Thioglycollate.....	0.5 g
Yeast Extract.....	5.0 g	Resazurin.....	1.0 mg
Sodium Chloride.....	2.5 g	Agar.....	0.75 g
		Demineralized Water .....	1000.0 ml

pH 7.1 ± 0.2 @ 25°C

\*Adjusted as required to meet performance standards.

## PROCEDURE

- Liquid media for anaerobic incubation should be reduced prior to inoculation by one of the following methods:
  - Place tubes with caps loosened in an anaerobic environment for 18-24 hours prior to use.
  - Boil tubes with caps loosened and cool to room temperature before inoculation.
- Consult appropriate references for the recommended procedure for sample inoculation and cultivation.
- Incubate inoculated medium in ambient air at 33-37°C for 24 hours to 7 days or following established laboratory procedures.
- Turbidity or growth in Thioglycollate Medium must be confirmed by Gram stain and growth following subculture to appropriate medium.

## QUALITY CONTROL

All lot numbers of Thioglycollate Medium w/ Indicator and Dextrose have been tested using the following quality control organisms and have been found to be acceptable. This quality control testing meets or exceeds CLSI standards.<sup>4</sup> Testing of control organisms should be performed in accordance with established laboratory quality control procedures. If aberrant quality control results are noted, patient results should not be reported.

### CONTROL

\**Bacteroides fragilis* ATCC® 25285  
*Clostridium sporogenes* ATCC® 11437  
*Pseudomonas aeruginosa* ATCC® 9027  
*Staphylococcus aureus* ATCC® 6538  
\**Staphylococcus aureus* ATCC® 25923

\*CLSI recommended organism

### INCUBATION

Ambient, up to 48 h @ 33-37°C  
Ambient, 3 days @ 30-35°C  
Ambient, 3 days @ 30-35°C  
Ambient, 3 days @ 30-35°C  
Ambient, 18-24 h @ 33-37°C

### RESULTS

Growth  
Growth  
Growth  
Growth  
Growth

## LIMITATIONS

- A slight turbidity (haziness) may be present due to the small amount of agar in the medium.
- When the medium has been boiled, it is clear. Do not boil more than once; frequent boiling results in a development of toxic products.<sup>5</sup>

## BIBLIOGRAPHY

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- Nungester, W.J., M.N. Hood, and M.K. Warren. 1943. Proc. Soc. Exp. Biol. Med. 52:287-289.
- Clinical and Laboratory Standards Institute (CLSI). 2004. Quality Control for Commercially Prepared Microbiological Culture Media; Approved Standard, 3<sup>rd</sup> ed. M22-A3. CLSI, Wayne, PA.
- MacFaddin, J.F. 1985. Media for Isolation-Cultivation-Identification-Maintenance of Medical Bacteria. Vol. 1. Williams & Wilkins, Baltimore, MD.

Refer to the front of Remel *Technical Manual of Microbiological Media* for **General Information** regarding precautions, product storage and deterioration, specimen collection, storage and transportation, materials required, quality control, and limitations.

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