

# TM 439 – TELLURITE GLYCINE AGAR BASE

#### **INTENDED USE**

For quantitative detection of coagulase positive Staphylococci from foods and other sources.

# PRODUCT SUMMARY AND EXPLANATION

Bacteria in the genus Staphylococcus are pathogens of man and other mammals. Traditionally they were divided into two groups on the basis of their ability to clot blood plasma (the coagulase reaction). Coagulase-positive strains of Staphylococcus aureus form the most pathogenic staphylococci. The presence of staphylococci in a lesion might first be suspected after examination of a direct gram stain. However, small numbers of bacteria in blood preclude microscopic examination and require culturing first. Tellurite Glycine Agar was originally developed by Ludlam and modified by Zebovitz et al. It is used for the quantitative detection of coagulase-positive staphylococci from foods and other sources like skin, mucous membranes, air and soil etc. This medium supports better growth of coagulase-positive cocci even if present in small numbers.

Coagulase positive staphylococci produce black colonies within 24 hours after an incubation at 37°C. Generally other organisms produce no growth during this incubation period with the exception of an occasional coagulase-negative strain that may produce small grey colonies, not readily confused with black coagulase positive colony.

# **COMPOSITION**

Ingredients	Gms / Ltr		
Casein enzymic hydrolysate	10.000		
Yeast extract	5.000		
Mannitol	5.000		
Dipotassium phosphate	5.000		
Lithium chloride	5.000		
Glycine	10.000		
Agar	16.000		

#### **PRINCIPLE**

Casein enzymic hydrolysate and yeast extract provide nitrogenous compounds, vitamin B complex and other essential growth nutrients. Lithium chloride and potassium tellurite are the inhibitors of the coagulase negative staphylococci and a wide variety of other bacteria. Potassium tellurite also serves as a differential agent since coagulase-positive staphylococci reduce tellurite and form black colonies. Mannitol is a source of fermentable carbohydrate for coagulase positive staphylococci.

# **INSTRUCTION FOR USE**

- Dissolve 56 grams in 1000 ml distilled water.
- Heat to boiling to dissolve the medium completely.
- Sterilize by autoclaving at 15 psi pressure (121°C) for 15 minutes.
- Cool to 45-50°C and to each 100 ml of base add 2 ml of 1% Potassium Tellurite Solution. Mix well before pouring into sterile Petri plates.

Caution: Lithium chloride is harmful. Avoid bodily contact and inhalation of vapours. On contact with skin, wash with plenty of water immediately.

# **QUALITY CONTROL SPECIFICATIONS**













**Appearance of Powder** : Cream to yellow homogeneous free flowing powder.

: Light amber coloured clear to slightly opalescent gel forms in Petri plates Appearance of prepared medium

: 7.2±0.2 pH (at 25°C)

### INTERPRETATION

Cultural characteristics observed after incubation with added 1% Potassium Tellurite Solution.

Microorganism	ATCC	Inoculum (CFU/ml)	Growth	Recovery	Colour of colony	Incubation Temperature	Incubation Period
Escherichia coli	25922	>=10³	Inhibited	0%	-	35-37°C	24-48 Hours
Salmonella Typhimurium	14028	>=10³	Inhibited	0%	-	35-37°C	24-48 Hours
Staphylococcus aureus	25923	50-100	Good- luxuriant	>=50%	Black	35-37°C	24-48 Hours
Staphylococcus epidermidis	12228	50-100	Poor-fair	10-30%	Grey	35-37°C	24-48 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. Easmon C. S. F., Adlam C, 1983, Staphylococci and Staphylococcal infections. Vol. I and II, Academic Press, London.
- 2. Zebovitz E., Evans J. B. and Niven C. F., 1955, J. Bacteriol., 70:687.
- 3. MacFaddin J. F., 1985, Media for Isolation-Cultivation-Identification-Maintenance of Medical Bacteria, Vol. 1, Williams and Wilkins, Baltimore.



































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







