# **PRODUCT DATA SHEET**



# TM 411 – CHOLERA MEDIUM BASE

### **INTENDED USE**

For selective isolation of Vibrio species from samples contaminated with Enterobacteriaceae.

# **PRODUCT SUMMARY AND EXPLANATION**

*Vibrio cholerae* is the etiological agent of cholera in humans in which the disease is caused not by tissue invasion of microorganisms but through the production of toxins that interrupt normal intra-intestinal exchanges of water and electrolytes. *Vibrios* grow readily on most isolation media. Adding sodium chloride to the medium enhances growth of all species. Cholera Medium Base is a selective medium used for the isolation of *Vibrio* species from specimens contaminated with enteric bacteria. It is based on the formulation described by Felsenfeld and Watanabe for the isolation of *V. cholerae* and similar *Vibrios* from specimens contaminated with Enterobacteriaceae.

# **COMPOSITION**

Ingredients	Gms / Ltr	
Peptone	10.000	
Beef extract	10.000	
Sucrose	10.000	
Sodium lauryl sulphate	0.100	
Sodium chloride	20.000	
Sodium carbonate	5.000	
Agar	10.000	

#### PRINCIPLE

Peptone and beef extract provide nitrogenous nutrients whereas sucrose serves as the fermentable carbohydrate source for the metabolism of *Vibrios*. Sodium lauryl sulphate inhibits many contaminating organisms. Potassium tellurite also inhibits many gram-positive and gram-negative bacteria except *Vibrios*. Sodium chloride maintains osmotic equilibrium.

#### **INSTRUCTION FOR USE**

- Dissolve 65.1 grams in 1000 ml purified / distilled water.
- Heat to boiling to dissolve the medium completely. DO NOT AUTOCLAVE.
- Cool to 70°C and add 2 ml of sterile 1% Potassium Tellurite Solution and 5 ml of sterile defibrinated blood.
- Maintain at 70°C for a few minutes. Cool to 45-50°C. Mix well and pour into sterile Petri plates.

# QUALITY CONTROL SPECIFICATIONS

Appearance of Powder	: Cream to yellow homogeneous free flowing powder.
Appearance of prepared medium	: Basal medium: Yellow coloured clear to slightly opalescent gel. After Addition of blood & Tellurite and on heating : Brownish red coloured opaque gel forms in Petri plates.
pH (at 25°C)	: 8.5±0.2

#### **INTERPRETATION**

Cultural characteristics observed after incubation with added 1% Potassium Tellurite Solution and sterile defibrinated blood.

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Microorganism	ATCC	lnoculum (CFU/ml)	Growth	Recovery	Incubation Temperature	Incubation Period
Bacillus subtilis	6633	>=10 <sup>3</sup>	Inhibited	0%	35-37°C	18-24 Hours
Escherichia coli	25922	>=10 <sup>3</sup>	Inhibited	0%	35-37°C	18-24 Hours
Proteus mirabilis	25933	>=10 <sup>3</sup>	Inhibited	0%	35-37°C	18-24 Hours
Pseudomonas aeruginosa	27853	>=10 <sup>3</sup>	Inhibited	0%	35-37°C	18-24 Hours
Vibrio cholerae	15748	50-100	Luxuriant	>=70%	35-37°C	18-24 Hours
Vibrio parahaemolyticus	17802	50-100	Luxuriant	>=70%	35-37°C	18-24 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.

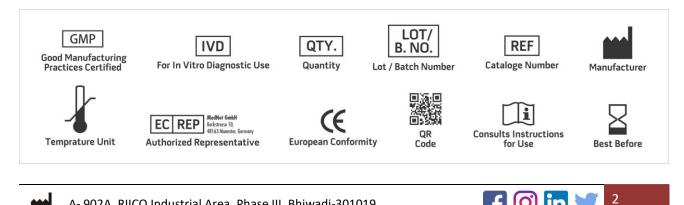
# DISPOSAL

After use, prepared plates, specimen/sample containers and other contaminated materials must be sterilized before discarding.

#### REFERENCES

1. Felsenfeld O. and Watanabe Y., 1958, U.S. Armed Forces Med. J., 9 (7): 975.

2. Isenberg, H. Clinical Microbiology Procedures Handbook. 2nd Edition.



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

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