

TMH 113 - CETRIMIDE AGAR (as per USP/EP/JP/BP/IP)

INTENDED USE

For selective isolation of *Pseudomonas aeruginosa*.

PRODUCT SUMMARY AND EXPLANATION

Cetrimide Agar was described by King et al. This media formulation is in accordance with the harmonized method of USP/EP/BP/JP/IP. It is used as a selective medium for the isolation of *Pseudomonas aeruginosa* from pharmaceutical products. This medium is also recommended for microbial limit testing of non-sterile products. Lowbury first reported the use of cetrimide as an agent for selective isolation of *Pseudomonas* spp. This medium is also used for determining the ability of an organism to produce fluorescein and pyocyanin.

COMPOSITION

Ingredients	Gms / Ltr
Pancreatic digest of Gelatin	20.000
Agar	13.600
Dipotassium sulphate	10.000
Magnesium chloride	1.400
Cetrimide	0.300

PRINCIPLE

Cetrimide (N-acetyl-N,N,N-trimethylammonium bromide) is incorporated in the medium to inhibit bacteria other than *Pseudomonas aeruginosa*. This compound a cationic detergent acts as a quaternary ammonium compound, which causes nitrogen and phosphorus to be released from bacterial cells other than *Pseudomonas aeruginosa*. Magnesium chloride and potassium sulphate incorporated in the medium enhances the production of pigment pyocyanin, which is a blue-green pigment, diffusing into the medium. This improves detection of *Pseudomonas* spp. on this medium. Presence of magnesium ions can also neutralize the EDTA, if present in the sample. Pancreatic digest of gelatin provides the essential nutrients for growth of *Pseudomonas* spp., while glycerol serves as slow and continuous carbon source for the growing cells.

INSTRUCTION FOR USE

- Dissolve 45.30 grams in 1000 ml distilled water containing 10 ml glycerol.
- Gently heat to boiling with swirling to dissolve the medium completely.
- Sterilize by autoclaving at psi (121°C) for 15 minutes.
- Cool to 45-50 °C.
- Mix well and pour into sterile petri plates.

QUALITY CONTROL SPECIFICATIONS

Appearance of Dehydrated powder	:	Cream to yellow colour, homogeneous free flowing powder
Appearance of Prepared medium	:	Light amber colour, opalescent gel
pH (at 25°C)	:	7.2±0.2

INTERPRETATION

Culture characteristics observed after incubation. Recovery rate is considered as 100% for bacteria growth on Soyabean Casein Digest Agar.



Microorganism	ATCC	Inoculum (CFU/ml)	Growth	Pigmentation	Recovery	Incubation Temperature	Incubation Period
<i>Pseudomonas aeruginosa</i>	27853	50-100	Good-Luxuriant	green color fluorescence	≥50%	30 - 35°C.	18-72 Hours
<i>Pseudomonas aeruginosa</i>	9027	50-100	Good-Luxuriant	green color fluorescence	≥50%	30 - 35°C.	18-72 Hours
<i>Staphylococcus aureus</i>	25923	≥1000	Inhibited		0%	30 - 35°C.	> 72 Hours
<i>Staphylococcus aureus</i>	6538	≥1000	Inhibited		0%	30 - 35°C.	> 72 Hours
<i>Escherichia coli</i>	25922	≥1000	Inhibited		0%	30 - 35°C.	> 72 Hours
<i>Escherichia coli</i>	8739	≥1000	Inhibited		0%	30 - 35°C.	> 72 Hours
<i>Salmonella typhimurium</i>	14028	≥1000	Inhibited		0%	30 - 35°C.	> 72 Hours

PACKAGING

In 100 & 500 gm packaging size.

STORAGE

Dehydrated powder, hygroscopic in nature, store in a dry place, in tightly-sealed containers below 25°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 powder show evidence of microbial contamination, discoloration, drying, or other signs of deterioration.

DISPOSAL

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

REFERENCES

1. King, Ward and Raney, 1954, J. Lab. Clin. Med., 44:301.
2. British Pharmacopoeia, 2016 The Stationery Office British Pharmacopoeia.
3. European Pharmacopoeia, 2017, European Dept. for the quality of Medicines.
4. Japanese Pharmacopoeia, 2016.
5. The United States Pharmacopoeia, 2019, The United States Pharmacopoeial Convention. Rockville, MD.
6. Indian Pharmacopoeia, 2018, Govt. of India, the controller of Publication, Delhi, India.
7. Lowbury E J L., 1951, J.Clin.Path., 4:66.

 GMP Good Manufacturing Practices Certified	 Best Before	 QTY. Quantity	 REF Catalogue Number	 Manufacturer
 Temperature Unit	 LOT/ B. NO. Lot / Batch Number	 Consults Instructions for Use	 QR Code	

NOTE: Please consult the Material Safety Data Sheet for information regarding hazards and safe handling Practices.

***For professional use only.**

Revision: 10th July, 2020

