

CM 20381 – CASEIN HYDROLYSATE AGAR W/ 2.5% AGAR

INTENDED USE

For large scale cultivation of *Vibrio cholerae* for cholera vaccine production.

PRODUCT SUMMARY AND EXPLANATION

CaseinHydrolysateAgarw/2.5%is themodification of medium recommended by APHA and is a highly selective medium, recommended particularly for the production of cholera vaccine by *Vibrio* species.

COMPOSITION

Ingredients	Gms / Ltr
Tryptone	5.000
Beef infusion from	150.000
Peptone	5.000
Yeast autolysate	1.500
Sodium phosphate	2.500
Sodium chloride	5.000
Agar	25.000

PRINCIPLE

IthasTryptone, beef infusion B from, and Peptone which serves as a rich source of nitrogen and carbon. Yeast autolysate provides necessary growth factors and vitamin supplement required for metabolism of wide number of bacteria. Sodium phosphate helps buffering of media whereas sodium chloride balances the osmotic equilibrium.

INSTRUCTION FOR USE

- Dissolve 45.5gramsin 1000 ml purified / distilled water containing 22 ml glycerol.
- 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.
- Mix well and pour into sterile Petri plates.

QUALITY CONTROL SPECIFICATIONS

- Appearance of Powder : Yellow coloured homogeneous free flowing powder.
- Appearance of prepared medium : Light yellow coloured clear to slightly opalescent gel forms in petri plates.
- pH (at 25°C) : 7.8±0.2

INTERPRETATION

Culturalcharacteristics observed after incubation.

Microorganism	ATCC	Inoculum (CFU/ml)	Growth	Recovery	Incubation Temperature	Incubation Period



Vibrio cholerae	15748	50-100	Luxuriant	>=70%	35-37°C	18-24 Hours
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PACKAGING:

Inpacksizeof500 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

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

REFERENCES

1. VanderzantCand Splittstoesser D (Eds) 1992. Compendium of Methods for the Microbiological Examination of Foods, 3rd ed, APHA, Washington, DC.

 GMP Good Manufacturing Practices Certified	 IVD For In Vitro Diagnostic Use	 QTY. Quantity	 LOT/ B. NO. Lot / Batch Number	 REF Cataloge Number	 Manufacturer
 Temperature Unit	 EC REP Authorized Representative <small>MedNet GmbH Birkstrasse 10, 48143 Münster, Germany</small>	 European Conformity	 QR Code	 Consults Instructions for Use	 Best Before

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

*For LabUse Only