



## Technical Data Sheet

### Acetate Agar

**CLS-DCM-ACA-01**

#### Principle

Acetate agar is formulated by Whittenbury (1965) used for cultivation of *Leuconostoc* and *Pediococcus* species. The media is consisting of peptone, meat extract, yeast extract, dextrose, polysorbate 80, sodium acetate and agar. Peptone yeast extract and meat extract provide nitrogen, amino acids and essential elements required for the growth of organisms. Dextrose is the source of energy and sodium acetate is carbon source. Polysorbate 80 maintains the surface tension of the medium.

**Use:** Recommended for the isolation and cultivation of *Leuconostoc* and *Pediococcus* species.

#### Contents\*

##### Ingredients

	Gram/Litre
Peptone	5.000
Meat extract	5.000
Yeast extract	5.000
Dextrose	10.000
Polysorbate 80	0.500
Sodium acetate	16.400
Agar	20.000
pH at 25 °C	5.4±0.2

\* Formula adjusted for optimum performance and parameters

**Directions:** Dissolve 62.00 grams in 1000 ml distilled water, boil to dissolve the medium completely and sterilize by autoclaving at 15 lbs. pressure (121°C) for 15 min, cool it to 42-45 °C and mix well and pour into sterile petri plates. Ensure complete solidification and inoculate test sample aseptically.

#### Specimens' types analyzed

Brewery and food samples etc.

#### Precautions to be taken

These microbial media are intended for the in-vitro use only. All the handling, experiments, storage, and discarding should be performed with the help of skilled and knowledgeable technicians and as per the established guidelines. The material should be disposed only after proper sterilization by autoclaving. Please go through the MSDS of the media to avoid any accidents or in emergency.

#### Performance and Evaluation

The expected performance of the medium is liable to use as per the direction on the label when stored at optimum conditions and within expiry date.



## Quality Control

<b>Appearance</b>	Beige colored free flowing, homogeneous powder
<b>Reaction of 6.20% solution</b>	5.4 $\pm$ 0.2 at 25 °C
<b>pH</b>	5.20 – 5.60
<b>Gelling</b>	Firm comparable with 2% agar gel
<b>Color and clarity of ready medium</b>	Amber colored opalescent gel
<b>Growth Promotion properties</b>	Best at $\leq$ 100 CFU at 25-30 °C for 18-72 h
<b>Negative control</b>	Performed using sterile distilled water

## Different Microbial Response

<b>Organism</b>	<b>ATCC</b>	<b>Inoculum</b>	<b>Growth</b>	<b>Incubation</b>
<i>Leuconostoc mesenteroides</i>	Lab isolate	50-100	Luxurious	25-30 °C, 18 – 48 hours

## Storage and Shelf Life

Hygroscopic; keep container tightly closed. Store in cool dry place.

**Disposal:** To avoid the contamination or propagation of any hazardous microbes the used, unusable or modified preparation of this product must be disposed after autoclaving after completion of task.

## Reference

1. Atlas, R. M. (2005). *Handbook of media for environmental microbiology*. CRC press.
2. Jorgensen, J.H., Pfaller, M.A., Carroll, K.C., Funke, G., Landry, M.L., Richter, S.S and Warnock., D.W. (2015), *Manual of Clinical Microbiology*, 11<sup>th</sup> Edition. Vol. 1.
3. Whittenbury, R. (1965). A study of some *Pediococci* and their relationship to *Aerococcus viridans* and the enterococci. J. Gen. Microbiol. 40:97-106.