

## Lead Acetate Agar Plate

MP180

### Intended Use:

Recommended for detection of hydrogen sulphide producing enteric bacteria from clinical and non-clinical samples.

### Composition\*\*

Ingredients	g / L
Peptone	15.000
Proteose peptone	5.000
Dextrose (Glucose)	1.000
Lead acetate	0.200
Sodium thiosulphate	0.080
Agar	15.000
Final pH ( at 25°C)	6.6±0.2

\*\*Formula adjusted, standardized to suit performance parameters

### Directions

Either streak, inoculate or surface spread the test inoculum (50-100 CFU) aseptically on the plate.

### Principle And Interpretation

*Salmonella*, *Shigella*, *Yersinia* species and certain strains of *Escherichia coli* cause severe gastroenteritis and life-threatening systemic illness in human (1,2). Of these, *Salmonella* Typhi can be differentiated due to their ability to form hydrogen sulphide (3). Lead Acetate Agar is the modification of the original formulation of Spray (4). This medium was successfully used to study hydrogen sulphide production (4,5). Lead Acetate Agar can also be used to differentiate between *Salmonella* Paratyphi A and *Salmonella* Paratyphi B (6). The latter produces hydrogen sulphide, observed as browning of the medium, within 18-24 hours, whereas the former fails to produce hydrogen sulphide.

Peptone, proteose peptone and dextrose provide all the essential nutrients for the growth of bacteria. Bacteria capable of using sulphur from sodium thiosulphate in their metabolic activities produce hydrogen sulphide. Lead acetate acts as an indicator of hydrogen sulphide production observed as browning of the medium. Dextrose is the fermentable carbohydrate source. Production of gas from dextrose is indicated by the presence of bubbles in the butt.

### Type of specimen

Pure isolate from clinical and non-clinical specimen.

### Specimen Collection and Handling:

For clinical samples, follow appropriate techniques for sample collection and processing as per guidelines (7,8). After use, contaminated materials must be sterilized by autoclaving before discarding.

### Warning and Precautions

In Vitro Diagnostic use. For professional use only. Read the label before opening the container. Wear protective gloves/ protective clothing/eye protection/ face protection. Follow good microbiological lab practices while handling specimens and culture. Standard precautions as per established guidelines should be followed while handling specimens. Safety guidelines may be referred in individual safety data sheets.

### Limitations

1. Individual organisms differ in their growth requirement and may show variable growth patterns on the medium
2. Each lot of the medium has been tested for the organisms specified on the COA. It is recommended to users to validate the medium for any specific microorganism other than mentioned in the COA based on the user's unique requirement
3. It is recommended to store the plates at 24-30°C to avoid minimum condensation.

### Performance and Evaluation

Performance of the medium is expected when used as per the direction on the label within the expiry period when stored at recommended temperature.

## Quality Control

### Appearance

Sterile Lead Acetate Agar in 90 mm disposable plates with smooth surface and absence of black particles/cracks/bubbles.

### Colour of medium

Amber coloured medium.

### Quantity of medium

25 ml of medium in 90 mm disposable plates.

### pH

6.40-6.80

### Cultural Response

Cultural characteristics observed after an incubation at 35-37°C for 18-24 hours.

Organism	Inoculum (CFU)	Growth	Gas Production	H <sub>2</sub> S production
<i>Escherichia coli</i> ATCC 25922 (00013*)	50-100	luxuriant	positive reaction	negative reaction
# <i>Klebsiella aerogenes</i> ATCC 13048 (00175*)	50-100	luxuriant	positive reaction	negative reaction
<i>Salmonella</i> Paratyphi A ATCC 9150	50-100	luxuriant	negative reaction	negative reaction
<i>Salmonella</i> Paratyphi B ATCC 8759	50-100	luxuriant	negative reaction	positive reaction, browning of the medium
<i>Salmonella</i> Typhi ATCC 6539	50-100	luxuriant	variable reaction	positive reaction, browning of the medium
<i>Salmonella</i> Typhimurium ATCC 14028 (00031*)	50-100	luxuriant	negative reaction	positive reaction, browning of the medium
<i>Shigella dysenteriae</i> ATCC 13313	50-100	luxuriant	negative reaction	negative reaction
<i>Shigella flexneri</i> ATCC 12022 (00126*)	50-100	luxuriant	negative reaction	negative reaction

Key : (\*) Corresponding WDCM numbers.

(#) Formerly known as *Enterobacter aerogenes*

## Storage and Shelf Life

On receipt store between 20-30°C. Use before expiry date on the label. Product performance is best if used within stated expiry period.

## Disposal

User must ensure safe disposal by autoclaving and/or incineration of used or unusable preparations of this product. Follow established laboratory procedures in disposing of infectious materials and material that comes into contact with clinical sample must be decontaminated and disposed of in accordance with current laboratory techniques (7,8).

## Reference

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- Murray P. R., Baron J. H., Pfaller M. A., Jorgensen J. H. and Tenover F. C., (Ed.), 2003, Manual of Clinical Microbiology, 8th Ed., American Society for Microbiology, Washington, D.C.
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- Spray R. S., 1936, J. Bacteriol., 32:135.

5. Morrison L. E. and Tanner F. W., 1922, J. Bacteriol., 7:343.
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7. Isenberg, H.D. Clinical Microbiology Procedures Handbook 2nd Edition.
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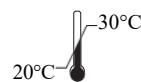
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HiMedia Laboratories Pvt. Limited,  
Plot No.C-40, Road No.21Y,  
MIDC, Wagle Industrial Area,  
Thane (W) -400604, MS, India



**IVD** *In vitro* diagnostic  
medical device



**Storage temperature**



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**CE Marking**



**Do not use if  
package is damaged**

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