

LYSINE IRON AGAR

Differential medium for Enterobacteria isolation.

TYPICAL FORMULA (g/l)	
Peptospecial	5.0
Glucose	1.0
L-Lysine Hydrochloride	10.0
Ferric Ammonium Citrate	0.5
Yeast Extract	3.0
Sodium Thiosulfate	0.04
Brom Cresol Purple	0.02
Agar	14.5
Final pH 6.7 ± 0.2 at 25 °C	

DESCRIPTION

LYSINE IRON AGAR is a medium used for differentiating microorganisms, especially *Salmonella* spp. on the basis of lysine decarboxylation/deamination and H_2S production.

PRINCIPLE

Peptospecial is the source of proteins. Glucose is the substrate for the fermentation. L-Lysine, Ferric Ammonium Citrate and Sodium Thiosulfate, are the specific substrates for the reactions of identification. Yeast extract is a source of amino acids and vitamins of group B. Brom Cresol Purple is the pH indicator. Agar is the solidifying agent.

PREPARATION

Completely dissolve the bottle content in water bath at 100°C. Cooling at 45-50°C and gently dispenseinto final tubes in aseptic condition. Allow the medium to solidify in a position that provides a short slant and a deep butt.

TECHNIQUE

Pick the center of a well-isolated colony from a fresh, pure culture with a needle and inoculate it by stabbing to the base of the butt and streaking the slant of the medium in the tube.

Cap the tube loosely to ensure aerobic conditions. Incubate at $36 \pm 1^{\circ}$ C for 18-24 hours.

Examine after 18-24 hours and 40-48 hours for growth and color change in the butt and the slant of the medium and blackening at the apex of the slant.

INTERPRETATION OF RESULTS

Lysine decarboxylase reaction results are: Positive: purple (alkaline) butt, purple slant. Negative: yellow (acid) butt, purple slant. Lysine deaminase reaction results are: Positive: red slant. Negative: purple slant. Hydrogen sulphide reaction: Positive: blackened medium at the apex of the slant.

STORAGE

10-25°C away from light, until the expiry date on the label or until signs of deterioration or contamination are evident.

WARNING AND PRECAUTIONS

The product is not classified as hazardous by current legislation and does not contain harmful substances in concentrations of \geq 1%. The product is designed for *In vitro* diagnostic use and must be used only by properly trained operators.

DISPOSAL OF WASTE

Disposal of waste must be carried out according to national and local regulations in force.

REFERENCES

- 1. Edwards, P.R., and M.A. Fife. 1961. Appl. Microbiol. 9 : 478.
- MacFadding, J.F. 1985. Media for isolation-cultivation-identificationmaintenance of medical bacteria, vol. 1. Williams & Wilkins, Baltimore, MD.



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PRODUCT SPECIFICATIONS

NAME

LYSINE IRON AGAR

PRESENTATION

Ready-to-use glass bottles containing 200 ml of medium.

STORAGE

10-25°C

DACKGING

Acitolito							
Code	Content	Packaging					
412040	6 bottles x 200 ml	6 bottles in cardboard box					

pH OF THE MEDIUM

6.7 ± 0.2

USE

LYSINE IRON AGAR is a medium used for differentiating microorganisms, especially Salmonella spp. on the basis of lysine decarboxylation/deamination and H₂S production.

TECHNIQUE

Refer to technical sheet of the product.

APPEARANCE OF THE MEDIUM

Very slightly opalescent purple medium without precipitate.

SHELFLIFE

1 year.

QUALITY CONTROL

Control of general characteristics, label and print. 1.

- 2. Sterility control:
 - 7 days at $25 \pm 1^{\circ}$ C, in aerobiosis. 7 days at $36 \pm 1^{\circ}$ C, in aerobiosis.
- Microbiological control: 3. Inoculum for productivity: 10-100 UFC/ml. Inoculum for selectivity: 104-105 UFC/ml. Inoculum for specificity: $\leq 10^4$ UFC/ml. Incubation conditions: 18-48 h at $36 \pm 1^{\circ}$ C.

Microorganism	ATCC	Butt	Slant	H₂S
Escherichia coli	25922	purple	purple	-
Salmonella typhimurium	14028	purple	purple	+
Klebsiella pneumoniae	13883	purple	purple	-
Citrobacter freudi	8090	yellow	purple	+
Proteus mirabilis	25933	yellow	red	-

TABLE OF SYMBOLS

LOT Batch code	IVD	<i>In vitro</i> Diagnostic Medical Device	•••	Manufacturer	\Box	Use by	Ţ	Fragile, handle with care
REF Catalogue number	Ł	Temperature limitation	\sum	Contains sufficient for <n> tests</n>		Caution, consult accompanying documents	\otimes	Do not reuse



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