

Also known as

Xylose Lysine Tergitol ™ 4 Agar Base

Specification

Solid culture medium for the selective and differential isolation of Salmonella spp., (Except S.typhi and S.paratyphi).

Formula * in g/L	
Proteose peptone No 3	1,60
Yeast extract	3,00
Xylose	3,75
Lactose	7,50
Sucrose	7,50
L-Lysine HCI	5,00
Sodium chloride	5,00

Sodium thiosulphate	6,80
Ferric ammonium citrate	0,80
Phenol red	0,08
Agar	15,00

Final pH 7,4 ±0,2 at 25 °C

* Adjusted and /or supplemented as required to meet performance criteria

Directions

Suspend 56 g of powder in one litre of distilled water containing 4 mL of Tergitol®4 (or Niaproof @ 4). Heat gently, stirring constantly until boiling or totally dissolved. Do not overheat or autoclave. Cool to approx. 50°C and pour plates. The medium when ready for use is clear and red.

NOTE: Excessive or prolonged heating may cause precipitation, therefore re-melting solidified medium or maintenance of the medium at melting temperatures (> 45 ° C) for more than an hour is not recommended.

Description

This medium was developed in 1990 by Miller and Tate as a modification of XLD agar for the isolation of non-typhoid Salmonella species in samples from poultry farms.

The replacement of deoxycholate by the tensioactive anionic Tergitol ® 4 (tetradecyl sodium sulphate, sodium salt 7-ethyl -2-methyl-4-undecanol or Niaproof ® 4) provides for greater selectivity, inhibiting most strains of Proteus, Providencia Pseudomonas and accompanying microflora.

The differential diagnostic features of the culture medium is based on the simultaneous use of three indicator systems: The utilization of xylose, lactose and sucrose as indicated by phenol red, which also allows recognition of the decarboxylation of lysine and thirdly the production of H_2S from thiosulphate which is manifested by a black precipitate of iron sulphide. Later, in 1995, the same authors were able to enhance the latter feature by adding a small amount of peptone medium allowing faster growth of black colonies.

Technique

For enumeration, The dry surface of the culture medium may be inoculated by, spreading 0.1 mL of sample, but if a count is not required it is more desirable to carry out pre-enrichment in an appropriate broth and then streak the sample on the XLT4 agar plate to obtain isolated colonies. The inoculated plates are incubated at 35-37 ° C and growth is observed at 18-24 hours, with additional observations at 48 h if the results are negative.

After 24 hours of incubation typical colonies of Salmonella appear totally black or with a black core and coloured periphery, usually yellow or red. The non-H2S producing Salmonella spp colonies are normal and yellow-pink in colour. The other gram-negative bacteria are either inhibited or grow poorly, giving rise to small yellow, or red-pink colonies, but never black in colour.

Limitations:

This media is designed for the detection and isolation of Salmonella based on selectivity and typical colonial characteristics, especially the production of H2S. The existence of non-producing strains of H2S can cause the appearance of false negative colonies. Also, some strains of bacteria such as Citrobacter and Escherichia that grow well in this medium and produce yellow colonies can be found however they are easily differentiated from salmonella. It is recommended that presumptive colonies are confirmed by biochemical or immunological methods.

In certain circumstances, multicoloured crystals or precipitates with a metallic finish may appear on the surface of the prepared medium, however they do not interfere with the behavior of the medium.

The plates of medium can be used for up to three months after preparation if the prepared medium is stored, away from light, refrigerated and water loss prevented, although it is advisable to use them within fifteen days.



Reference : 01-708 Product : XLT4 AGAR BASE

Quality control

Incubation temperature: 36°C ±2,0

Incubation time: 24 ± 3h

Inoculum: Practical range 100 ± 20 CFU. Min. 50 CFU (Productivity) / 10⁴-10⁶ CFU (Selectivity) / ≥ 10³ CFU (specificity) according to ISO 11133:2014/Amd 1:2018 .. Spiral plate Methods.

Microorganism

Salmonella enteritidis ATCC[®] 13076 Salmonella typhimurium ATCC[®] 14028 Escherichia coli ATCC[®] 25922 Enterococcus faecalis ATCC[®] 29212

Growth

Productivity > 0.50 Productivity > 0.50 Poor to good Inhibited

Remarks

red colonies / Black center (H2S+) red colonies / Black center (H2S+) Yellow colonies







Salmonella typhimurium ATCC 14028

uninoculated plate

Salmonella enterica ATCC 13076

References

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Storage

For laboratory use only. Keep tightly closed, away from bright light, in a cool dry place (+4 °C to 30 °C).