

Reference : 01-211 Product :

XYLOSE LYSINE DEOXYCHOLATE AGAR (XLD Agar) (Eur. Pharm.)

Specification

Solid medium for the isolation of enteropathogenic species, especially *Salmonella* and *Shigella* according to Pharmacopeial Harmonised Method and ISO Standard.

Formula * in g/L			
Xylose	3.50	Sodium deoxycholate	2.50
L-Lysine	5.00	Sodium thiosulfate	6.80
Lactose	7.50	Ammonium ferric citrate	0.80
Sucrose	7.50	Agar	15.00
Sodium chloride	5.00		
Yeast extract	3.00	Final pH 7,4 ±0,2 at 25 °C	
Phenol red	0.08		

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

Directions

Suspend 56,68 g of powder in 1 L of distilled water. Heat with constant stirring until boiling (90-100 °C). Pour immediately into plates. Do not autoclave and avoid remelting.

Description

Xylose Lysine Deoxycholate Agar is a selective differential medium, suitable for the detection of pathogenic enterobacteria, especially *Shigella*. Gram positive microbiota are inhibited by the low amount of deoxycholate, whilst *Shigella* grows.

Xylose, lactose or sucrose fermentation produces the acidification of the medium, and this is seen by the indicator turning yellow, surrounding the colonies. This colour disappears after 24 hours, so observations must be carried out between 18 and 24 hours.

Hydrogen sulfide production from thiosulfate is easily detected because colonies become darker, due to the ferric sulfide precipitate. Lysine decarboxylation to cadaverine may also be observed in the medium, since it produces alkalinization and consequently the indicator turns to red.

All these reactions allow a good differentiation of *Shigella*. *Edwardsiella* and *Proteus inconstans* are the only enterobacteria other than *Shigella* which do not ferment xylose and therefore show negative fermentation reaction. *Salmonella* ferment xylose, but it is consumed quickly and alkalinization of the medium due to lysine decarboxylation, may mask the reaction. *Salmonella* colonies become darker due to ferrous sulfide precipitates, which is also a common property with *Edwardsiella*.

Other types of enterobacteria do not suffer this phenomenon, since acid accumulation due to lactose and sucrose fermentation is so high that it avoids pH reversion by decarboxylation and even ferrous sulfide precipitate in the first 24 hours.

In the quality control, typical colonial appearances on XLD medium after 18-48 hours of incubation at 30-35°C are described.

Note: in ready to use media (plates), After 24-48h at refrigeration temperature, slight precipitates may appear on the surface. This does not affect the oerformance of the medium.

Quality control

Incubation temperature: 30-35 °C

Incubation time: 18-48 h

Inoculum: Practical range 10 - 100 CFU (productivity)/ 10²-10⁴CFU (selectivity) according to Eur. Pharm. harm. Spiral

Plate Method. Microorganism	Growth	Remarks
Staphylococcus aureus ATCC® 6538	Inhibited	Selectivity
Salmonella abony NCTC [®] 6017	Productivity > 0.50	Colonies & cult. medium red / Black center (H ₂ S +)
Salmonella typhimurium ATCC [®] 14028	Productivity > 0.50	Colonies & cult. medium red / Black center (H $_2$ S +)

Scharlau Microbiology - Technical data sheet



Product : XYLOSE LYSINE DEOXYCHOLATE AGAR (XLD Agar) (Eur. Pharm.)

References

- ATLAS, R.M., L.C. PARK (1993) Handbook of Microbiological Mediafor the examination of Food. CRC Press Inc.Boca Ratón.
- · DOWNES, F.P. & K. ITO (2001) Compendium of Methods for the Microbiological Examination of Foods. 4th ed. APHA. Washington DC. USA.
- EUROPEAN PHARMACOPOEIA 10.0 (2020) 10th ed. § 2.6.13. Microbiological examination of non-sterile products: Test for specified microorganisms. Harmonised Method. EDQM. Council of Europe. Strasbourg.
- · HORWITZ, W. (2000). Official Methods of Analysis of the AOAC Internacional. 17th ed. Gaithersburg Md. USA.
- \cdot ICMSF (1978) Microorganisms in Foods 1. University of Toronto Press.

Reference : 01-211

- · PASCUAL ANDERSON, Mª R. (1992) Microbiología Alimentaria. Díaz de Santos, S.A. Madrid.
- TAYLOR, W.J. (1965) Isolation of Shigella. I. Xylose Lysine Agars: New media for isolation of enteric pathogens. Am. J. Clin. Path 44:471-475.
- · US FDA (Food and Drug Adminstrations). (1998) Bacteriological Analytical Manual. 8th ed. Revision A. AOAC International. Gaithersburg, Md. USA.
- · USP 33 NF 28 (2011) <62> Microbiological examination of non-sterile products: Test for specified microorganisms. Harmonised Method. USP Corp. Inc. Rockville. MD. USA.

Storage

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