

Reference: 01-278

Scharlau Microbiology - Technical data sheet

Product:

TRYPTOSE SULFITE CYCLOSERINE AGAR (TSC AGAR)



Specification

Solid medium for the isolation and differentiation of Clostridium perfringens, according to ISO standards and other regulations.

Formula * in g/L

Enzymatic digest of casein (Tryptone) 15.0	0
Soy peptone5.0	0
Yeast extract5.0	0
Sodium disulphite1.0	0
Ammonium iron(III) citrate1.0	0
Agar18.0	0

Final pH 7.6 ±0.2 at 25 °C

Directions

Suspend 45 g of powder in 1L of distilled water and let soak minutes. Bring to a boil and distribute volumes of 250 ml or 100 ml in suitable containers. Sterilize the autoclave for 10 minutes at 121 ° C. Cool to 60 ° C and add 1 vial of Selective Supplement D-Cycloserine (Ref. 06-116LYO1 or 06-743LYO1) to each portion of media. Mix well and distribute on plates. If you wish yolk, while adding the antibiotic, sterile egg yolk (Ref. 06-016) at 80 ml/ L.

Note: Fluorogenic Supplement MUP may be used to identify Clostridium perfringens. (Ref. 06-744LYO1).

Description

The medium is a modification of the classical TSN Agar in which the traditional antibiotics, polymyxin and neomycin have been replaced by cycloserine. Cycloserine has been found more selective for Clostridium perfringens, and reduces the production of diffuse blackening. Clostridium perfringens is more resistant to cycloserine than to sulfadiazine, polymyxin and neomycin, hence reducing the dosage. The presence of sodium meta-bisulfite and ferric ammonium citrate allow three differential characteristics of this anaerobic species to be verified with just one assay. These characteristics are sulfite reduction, growth at 46°C and cycloserine resistance.

Cycloserine does not tolerate temperatures above 100 °C and its stability in a solution is variable. Therefore, it is advisable to prepare the exact number of plates that are going to be used.

A solution of cycloserine in phosphate buffer at pH 8,0 may be prepared (Di potassium phosphate 16.73 g/L and monopotassium phosphate 0.52 g/L) and if it is maintained refrigerated, can be used for approx. 5 days. This product, store at (-20±5) ° C can be use within 4 weeks of preparation. If stored frozen at (-20 ± 5) ° C could extend the expiration to 4 weeks or 12 months if stored at (-70 ± 10) ° C.

This lyophilized product, has a much higher expiration indicated on the manufacturer's label.

Necessary supplements

D-Cycloserine Selective Supplement (Ref. 06-743LYO1).

Vial contents:

Necessary amount for 100 mL of complete medium.

D-Cycloserine 40.00 mg

Distilled water (Solvent)

D-Cycloserine Selective Supplement (Ref. 06-116LYO1)

Vial contents:

Necessary amount for 250 mL of complete medium.

D-Cycloserine 100,00 mg

Distilled water (Solvent)

Revision date: 31/03/2021

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



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Technique

The standard procedure recommends surface inoculation of the samples or their dilutions, and once absorbed, to pour a second layer as a seal for anaerobiosis (TSC Agar or bacteriológical agar). After incubation à 44-46 $^{\circ}$ C for 24 \pm 3 h, proceed to enumerate the black colonies that appear in the plate.

Proceed according to standards or standardized methods.

Quality control

Incubation temperature: $44 \,^{\circ}\text{C} \pm 1.0$ Incubation time: $21 \pm 3 \,\text{h}$

Inoculum: Practical range 100±20 CFU. min. 50 CFU (productivity)/ 10⁴ -10□ CFU (selectivity), according to ISO 11133:2014/Amd 1:2018.

Microorganism

Clostridium perfringens ATCC® 10543 Clostridium perfringens ATCC® 13124 Bacillus subtilis ATCC® 6633

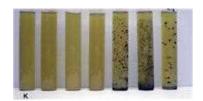
Growth

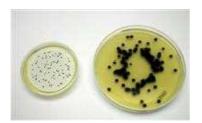
Productivity > 0.50 Productivity > 0.50 Inhibited

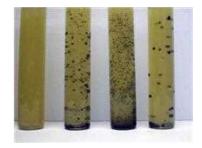
Remarks

Black colonies (Anaerobiosis) Black colonies (Anaerobiosis)

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Growth

Clostridium perfringens ATCC 13124

Clostridium perfringens ATCC 10543 Clostridium perfringens ATCC 13124

References

- · ATLAS, R.M., LC. PARKS (1993) Handbook of Microbiological Media. CRC Press, Inc. London.
- DIN Standard 10165. Referenz Verfahren fur Bestimmung von Clostridium perfringens. Fleisch und Fleischerzeugnissen.
- · DOWNES, F.P. & K. ITO (2001) Compendium of Methods for the Microbiological Examination of Foods. 4th ed. American Public Health Association. Washington.
- · DIRECTIVA 2015/1787/UE de la Comisión por la que se modifica la Directiva 98/ 83/CE relativa a la calidad de las aguas destinadas al consumo humano (DO L260 de 7.10.2015 pg 6 y ss)
- · FDA (Food and Drug Adminstrations) (1998) Bacteriological Analytical Manual. 8th ed. Revision A. AOAC International Inc. Gaithersburg. MD.
- · ISO 7937 (2004) Microbiology of Food and Animal Feeding Stuffs. Horizontal Method for Enumeration of C. perfringens. Colony-count technique.
- · ISO Norma 6461-2 (1986) Water Quality.- Detection and enumeration of the spores of sulfite-reducing anaerobes (Clostridia).- Part 2: Method by Membrane Filtration.
- . ISO 11133:2014/ Adm 1:2018. Microbiology of food, animal feed and water. Preparation, production, storage and performance testing of culture media.
- · ISO 14189 (2013 Water quality. Enumeration of Clostridium perfringens Method using membrane filtration
- · SMITH, L.D. (1981) Clostridial Anaerobic Infections, in Diagnostic Procedures for Bacterial Mycotic and Parasitic Infections. 6th ed. APHA. Washington.

Storage

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

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