

Reference: 5000

Technical Data Sheet

Product: TRYPTICASEIN SOY AGAR (T.S.A) (EUROPEAN CEIVD PHARM)

Specification

General purpose solid medium containing animal and plant peptone according to Pharmacopoeial Harmonised Method and ISO Standards.

Presentation

10 Prepared bottle	Packaging Details	Shelf Life	Storage
Bottle 125 ml	1 box with 10 bottles 125 ml	16 months	8-25°C
with: 100 ± 3 ml	Non injectable cap		

Composition

Composition (g/l):	
Peptone from casein	. 15.0
Soy peptone	. 5.00
Sodium chloride	. 5.00
Agar	. 15.0



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Description /Technique

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TSA is a widely used medium containing two peptones which support the growth of a wide variety of organisms, even that of very fastidious ones such as Neisseria, Listeria, Brucella, etc. It is frequently used for routine diagnostic purposes due to its reliability and its easily reproducible results.

The following list includes some of its most common applications:

1. Sensitivity testing, either by the Kirby-Bauer system or by following the WHO guidelines. Both systems recommend the use of the Mueller-Hinton Agar or verification purposes.

2. The medium provides, with added blood, perfectly defined haemolysis zones, while preventing the lysis of erythrocytes due to its sodium chloride content.

3. It can be used for the preparation of an exceptionally nutrient 'chocolate' agar, thanks to the richness of its peptones.

4. In a reducing environment or with a CO2 enriched atmosphere, it provides an excellent medium for the isolation of Brucella and Neisseria. It may be made selective by using additives.

5. Most streptococci grow in this medium though clear differences can be observed from one species to another.

6. Tryptic Soy Agar can be used as a selective medium for the count of urine samples although differentiation must be done on selective differential media.

7. Several tests for the differentiation and identification of staphylococci can be performed on this medium, provided suitable additives are used.

8. Yeast, particularly Candida species, can grow in this medium forming very characteristic colonies.

9. Chromogenic pseudomonas frequently produce pigmentation on TSA and are therefore easily recognized.

10. A vast bibliography documents its applications in the food industry.

11. It has been frequently used in the Health industry to produce antigens, toxins, etc...

12. Its simple and inhibitor-free composition makes it suitable for the detection of antimicrobial agents in food and other products.

13. A balanced and high nutrient value together with a lack of fermentable carbohydrates make this medium ideal for maintaining bacterial strains.

14. Classical media for microbiological examination of non-sterile products according to Pharmacopeial Harmonised Methods.

<u>Technique:</u>

Melt the medium contained in the bottles in a water bath or in a microwave oven, avoiding overhating before pouring into Petri dishes when cooled to room temperature.

Once solidified on a flat surface, Spread the plates by streaking methodology or by spiral method.

The inoculated plates are incubated at 30-35 ° C for 24-72 h (bacteria) and 3-5 days for fungi (yeast & molds). Examined daily. (Incubation times greater then those mentioned above or different incubation temperatures may be required dpending on the sample, on the specifications,... This medium can be inoculated directly or after enrichment broth).

After incubation, enumerate all the colonies that have appeared onto the surface of the agar.

Each laboratory must evaluate the results according to their specifications.

Calculate total microbial count per ml of sample by multiplying the average number of colonies per plate by the inverse dilution factor if streaked a diluted sample. Report results as Colony Forming Unit (CFU's) per ml or g along with incubation time and temperature.



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Growth

Quality control

Physical/Chemical control

Color : Straw-coloured yellow

pH: 7.3 ± 0.2 at 25°C

Microbiological control

Melt Medium - Prepare Plates - According to harmonized pharmacopoeial monographs, ISO standards and test methods Spiral Spreading: Practical range 50 - 100 CFU (Productivity).

Aerobiosis.Incubation at 30-35 °C. Read after 18-24h to 72 h for bacteria and 3-5 days for fungi.

Microorganism

Good (≥70 %) Escherichia coli ATCC[®] 8739, WDCM 00012 Good (≥70 %) Staphylococcus aureus ATCC[®] 6538, WDCM 00032 Good (≥70 %) Bacillus subtilis ATCC® 6633, WDCM 00003 Candida albicans ATCC[®] 10231, WDCM 00054 Good (≥70 %) Ps. aeruginosa ATCC[®] 9027, WDCM 00026 Good (≥70 %) Salmonella typhimurium ATCC[®] 14028, WDCM 00031 Good (≥70 %) Aspergillus brasiliensis ATCC[®] 16404, WDCM 00053 Good (≥70 %) L. monocytogenes ATCC[®] 13932, WDCM 00021 Good (≥70 %) **Sterility Control**

Incubation 48 hours at 30-35°C and 48 hours at 20-25°C: NO GROWTH Check at 7 days after incubation in same conditions

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