

Product Insert

TRYPTIC SOY BLOOD AGAR (TSBA)

Products

AS-542 Tryptic Soy Blood Agar (TSBA)

4 plate / pkg

Intended Use

Tryptic Soy Blood Agar (TSBA) is an enriched nonselective medium used for the isolation and cultivation of many fastidious and non-fastidious microorganisms found in clinical specimens. This medium is also suitable for use in the CAMP test for the presumptive identification of group B *Streptococci*.

Summary

TSBA is an enriched nonselective medium used for the isolation and cultivation of many microorganisms. The nutritive base consists of casein and soy peptone, which has been supplemented with sheep blood for the observation of hemolytic reactions, like the typical double zone β -hemolytic reactions of *Clostridium perfringens*. The media is supplemented with hemin and vitamin K₁ for the enhanced recovery of some fastidious anaerobes. This media is prepared, dispensed, and packaged under oxygen-free conditions to prevent the formation of oxidized products prior to use.

Formulation*

| | | |
|--|-------|----|
| Pancreatic Digest of Casein | 15.00 | g |
| Soy Peptone | 5.00 | g |
| Sodium Chloride | 5.00 | g |
| Hemin (0.1% solution) | 5.00 | mL |
| Vitamin K ₁ (1.0% solution) | 1.00 | mL |
| Agar | 15.00 | g |
| Sheep Blood | 45.50 | mL |
| DI Water | 1.00 | L |

Final pH: 7.1 \pm 0.2 at 25° CFinal weight: 16.0 g \pm 1.6 g

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

Precautions

For *IN VITRO DIAGNOSTIC USE* only. Utilize approved biohazard precautions and aseptic technique when using this product. This product is for use only by properly-trained and qualified personnel. Sterilize all biohazard waste prior to disposal.

Storage and Shelf Life

Storage: Upon receipt, store at room temperature in original package until used. Avoid overheating or freezing. Do not use media if there are signs of deterioration (shrinking, cracking, or discoloration due to oxidation of media) or contamination. The expiration date applies to the product in its original packaging and stored as directed. Do not use product past the expiration date shown on the label.

Shelf Life: 90 days from date of manufacture.

Procedure

Specimen Collection: Protect specimens for anaerobic culture from oxygen during collection, transportation, and processing. Consult appropriate references for detailed instructions concerning collection and transportation of anaerobes.

Methods for Use: TSBA should be inoculated directly with clinical specimen or from a broth that has been inoculated from a clinical specimen. Streak plates with inoculum to obtain isolated colonies and immediately place into an anaerobic atmosphere, incubating at 35-37°C for 18-48 hours. Extended periods of incubation may be required to recover some anaerobes. Detailed instructions for processing anaerobic cultures can be found in the listed references.

Materials Required, But Not Provided

Standard microbiological supplies and equipment such as loops, saline blanks, slides, staining supplies, microscope, incinerator / autoclave, incubators, anaerobic chamber / anaerobic jars, disinfectant, other culture media, and serological / biochemical reagents.

Interpretations of Results

If used properly, TSBA will support good growth or many fastidious microorganisms found in clinical infections. In addition, this media will support the typical double zone of β -hemolysis around colonies of *Clostridium perfringens*.

Limitations

TSBA agar will not provide complete information for identification of bacterial isolates. Additional test procedures and media are required for complete identification. In some cases, TSBA agar may be overgrown with swarming *Proteus* spp. or *Clostridium* spp. It is recommended that selective media such as Brucella Laked Blood Agar with Kanamycin and Vancomycin (LKV, catalog #: AS-112) and/or Brucella Blood Agar with Phenylethyl Alcohol (PEA, catalog #: AS-113) also be inoculated from clinical specimens to prevent such overgrowth and thus provide isolated colonies. Consult reference materials for additional information.

Quality Control

The following organisms are routinely used for quality control testing at Anaerobe Systems.

| Organism Tested | ATCC # | Results | Time | Special Reaction |
|---|----------------|---------|----------------------------|-----------------------------------|
| Bacteroides fragilis* | 25285 | Growth | 24 hrs | |
| Prevotella melaninogenica* | 25845 | Growth | 48 hrs | |
| Fusobacterium nucleatum* | 25586 | Growth | 24 – 48 hrs | |
| Clostridium perfringens* | 13124 | Growth | 24 hrs | Double Zone of β -hemolysis |
| Peptostreptococcus anaerobius* | 27337 | Growth | 24 – 48 hrs | |
| Staphylococcus aureus or Enterococcus faecalis | 25923 29212 | Growth | 24 hrs | |
| Escherichia coli | 25922 | Growth | 24 hrs | |
| Proteus mirabilis | 12453 | Growth | 24 hrs | |
| Propionibacterium acnes or Clostridium difficile | 6919 9689 | Growth | 24 – 48 hrs 24 – 48 hrs | |

* Organisms specified by CLSI for Quality Control testing of Anaerobic Blood Agars.

User Quality Control: The final determination to the extent and quantity of user laboratory quality control must be determined by the end user.

If sterility testing is to be performed on this product, a representative sample of the lot(s) should be incubated anaerobically and aerobically for 48 – 96 hours.

If the nutritive capacity of this media is to be tested for performance, it is recommended that the following ATCC organisms be evaluated for growth.

| Organism | ATCC # | Expected Results | Special Reactions |
|----------------|--------|------------------|-----------------------------------|
| B. fragilis | 25285 | 24 hrs | |
| E. faecalis | 29212 | 24 hrs | |
| C. perfringens | 13124 | 24 hrs | Double zone of β -hemolysis |
| P. anaerobius | 27337 | 24 hrs | |

Physical Appearance: TSBA should appear opaque burgundy red in color.

References

1. Dowell, V. R., Jr., G. L. Lombard, F. S. Thompson and A. Y. Armfield. 1977. *Media for the Isolation, Characterization and Identification of Obligately Anaerobic Bacteria*. USDHHS, CDC. Atlanta, GA 30333.
2. Engelkirk, P. G., Duben-Engelkirk, J. and Dowell, V. R. 1992. *Principles and Practices of Clinical Anaerobic Bacteriology*. Star Publishing Co., Belmont, CA 94002.
3. Holdeman, L. V., F. P. Cato and W. E. C. Moore. 1987. *Anaerobe Laboratory Manual*. Virginia Polytechnic Institute and State University. Blacksburg, VA 24061
4. Jousimeis-Somer, H. R., Summanen, P., Citron, D. M., Baron, E. J., Wexler, H. M. and S. M. Finegold. 2002. *Wadsworth – KYL Anaerobic Bacteriology Manual*. Star Publishing Co., Belmont, CA 94002.
5. CLSI. *Quality Control for Commercially Prepared Microbiological Culture Media; Approved Standard- Third Edition*. (2004). CLSI document M22-A3. CLSI, 940 West Valley Road, Suite 1400, Wayne, PA 19087-1898.

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