Soyabean Casein Digest Medium with 0.5% Soya Lecithin and 0.1% Polysorbate 80

Intended Use
Soyabean Casein Digest Medium with 0.5% Soya Lecithin and 0.1% Polysorbate 80 is used for determining efficiency of sanitization of containers, equipment surfaces, water miscible cosmetics etc.

Summary
Soyabean Casein Digest Medium with 0.5% Soya Lecithin and 0.1% Polysorbate 80 is used for the detection and enumeration of microorganisms on surfaces of sanitary importance, water miscible cosmetics, products containing antimicrobials or preservatives.

Principle
Pancreatic digest of casein and papaic digest of soyabean meal provide nitrogenous compounds and other nutrients essential for microbial replication. Soya Lecithin and polysorbate 80 (Tween 80) are neutralizers reported to inactivate residual disinfectants from where the sample is collected. Lecithin neutralizes quaternary ammonium compounds and polysorbate 80 neutralizes phenolic disinfectants, hexachlorophene, formalin and with lecithin ethanol. Dextrose serves as the carbohydrate source and dipotassium phosphate buffers the medium. Sodium chloride maintains the osmotic balance of the medium.

Formula*

Ingredients	**g/L**
Pancreatic Digest of Casein	17.0
Papaic Digest of Soyabean Meal	3.0
Sodium Chloride	5.0
Dextrose	2.5
Dipotassium Hydrogen Phosphate	2.5
Soya Lecithin	5.0
Polysorbate 80	1.0
Final pH (at 25°C)	7.3 ± 0.2

*Adjusted to suit performance parameters.

Storage and Stability
Store dehydrated medium below 30°C in tightly closed container and the prepared medium at 2°C-8°C. Avoid freezing and overheating. Use before expiry date on the label. Once opened keep powdered medium closed to avoid hydration.

Type of Specimen
Pharmaceutical samples

Specimen Collection and Handling
Ensure that all samples are properly labelled. Follow appropriate techniques for handling samples as per established guidelines. Some samples may require special handling, such as immediate refrigeration or protection from light, follow the standard procedure. The samples must be stored and tested within the permissible time duration. After use, contaminated materials must be sterilized by autoclaving before discarding.

Directions
1. Suspend 36.00 g of the powder in 1000 mL purified / distilled water.
2. Mix thoroughly.
3. Heat if necessary, to dissolve the powder completely.
4. Sterilize by autoclaving at 118°C-121°C respectively (12 to 15 psi) for 15 minutes as per validated cycle.

Quality Control

Dehydrated Appearance: Light yellow coloured, homogeneous, free flowing to moist powder, with a tendency to clump.

Prepared Appearance: Light yellow coloured, clear to slightly opalescent solution without any precipitation.

Growth Promotion Test: Growth promotion is carried out in accordance with the harmonized method of USP/EP/JP/IP and growth is observed after an incubation at 30°C-35°C for 18-24 hours for bacterial and ≤ 2 days for fungal.

Growth Promoting Properties: The test results observed are within the specified temperature and shortest period of time, inoculating ≤ 100 cfu (at 30°C-35°C for 18 hours for bacterial and 24 hours for fungal).

Growth Promoting Organism (ATCC)	**Growth**
Escherichia coli (8739)	Good
Escherichia coli (25922)	Good
Staphylococcus aureus subsp. *aureus* (6538)	Good
Staphylococcus aureus subsp. *aureus* (25923)	Good
Pseudomonas aeruginosa (9027)	Good
Pseudomonas aeruginosa Strain Boston 41501 (27853)	Good
Streptococcus pyogenes Strain Bruno (19615)	Good
Bacillus spizizenii (6633)	Good
Candida albicans 3147 (10231)	Good
Aspergillus brasiliensis WLRI 034(120) (16404)	Good

Note: Inoculum cfu for Good growth is 10 - 100.

Performance and Evaluation

Performance of the product is dependent on following parameters as per product label claim:

1. Directions
2. Storage
3. Expiry

Warranty

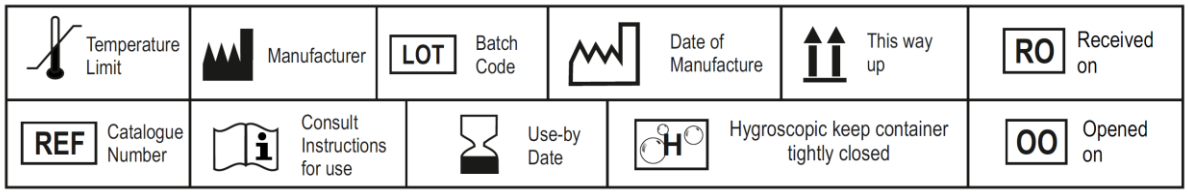
This product is designed to perform as described on the label and package insert. The manufacturer disclaims any implied warranty of use and sale for any other purpose.

Reference

1. Hall and Hartnett, 1964, Public Hlth. Rep., 79:1021.
2. Richardson (Ed).,1985, Standard Methods for Examination of Dairy Products, 15th ed.,APHA,Washington, D.C.
3. MacFaddin J.F., 1985, Media for Isolation-Cultivation-Identification-Maintenance of Medical Bacteria, Vol. I, Williams and Wilkins, Baltimore.
4. Brummer, 1976, Appl. Environ. Microbiol., 32:80.
5. Data on file: Microxpress®, A Division of Tulip Diagnostics (P) Ltd.

Product Presentation:

Cat No.	Product description	Pack Size
201191910100	Dehydrated Culture Media	100 g
201191910500	Dehydrated Culture Media	500 g



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Disclaimer

Information provided is based on our inhouse technical data on file, it is recommended that user should validate at his end for suitable use of the product.