

CM 22433 - SOYBEAN CASEIN DIGEST MEDIUM W/ LTHTh

INTENDED USE

For determining the efficiency of sanitization of containers, equipments, surfaces, water miscible cosmetics.

PRODUCT SUMMARY AND EXPLANATION

Soyabean Casein Digest Medium w/ LTHTh is used for the detection and enumeration of microorganisms for products of sanitary importance, water miscible cosmetics, Products containing antimicrobials or preservatives. The formulation of the basic medium (Soybean-Casein Digest Broth) is prepared according to the recommendations of the current European, Japanese and United States Pharmacopoeia.

COMPOSITION

Ingredients	Gms / Ltr
Casein enzymic hydrolysate	15.000
Papaic digest of soyabean meal	5.000
Sodium chloride	5.000
Polysorbate 80 (Tween 80)	5.000
Lecithin	0.700
Histidine	0.500
Sodium thiosulphate	0.500

PRINCIPLE

Casein enzymic hydrolysate and papaic digest of soyabean meal provide nitrogenous compounds and other nutrients essential for microbial replication. Lecithin, polysorbate 80 (Tween 80) and thiosulphate act as neutralizing agents reported to neutralize the activity of antimicrobial agents. Lecithin and polysorbate 80 neutralizes quaternary ammonium compounds and parahydroxy benzoates. Sodium thiosulphate neutralizes mercurial, halogens, aldehydes etc. Histidine acts as a reducing agent.

INSTRUCTION FOR USE

Inoculate the sample and incubate at specified temperature and time.

QUALITY CONTROL SPECIFICATIONS

Appearance of prepared medium	:	Light yellow to yellow coloured, clear solution
Quantity of Medium	:	10 ml of medium in tubes.
pH (at 25°C)	:	7.3 ± 0.2
Sterility Check	:	Passes release criteria

INTERPRETATION

Cultural characteristics observed after incubation.



Microorganism	ATCC	Inoculum (CFU/ml)	Growth	Growth with Disinfectant*	Incubation Temperature	Incubation Time
<i>Escherichia coli</i>	25922	50 - 100	Luxuriant	Good-Luxuriant	35 ± 2°C	18 - 24 Hours
<i>Escherichia coli</i>	8739	50 - 100	Luxuriant	Good-Luxuriant	35 ± 2°C	18 - 24 Hours
<i>Staphylococcus aureus</i>	25923	50 – 100	Luxuriant	Good-Luxuriant	35 ± 2°C	18 - 24 Hours
<i>Staphylococcus aureus</i>	6538	50 – 100	Luxuriant	Good-Luxuriant	35 ± 2°C	18 - 24 Hours
<i>Pseudomonas aeruginosa</i>	27853	50 – 100	Luxuriant	Good-Luxuriant	35 ± 2°C	18 - 24 Hours
<i>Bacillus subtilis</i>	6633	50 - 100	Luxuriant	Good-Luxuriant	35 ± 2°C	18 - 24 Hours
<i>Salmonella typhimurium</i>	14028	50 - 100	Luxuriant	Good-Luxuriant	35 ± 2°C	18 - 24 Hours
<i>Candida albicans</i>	10231	50 - 100	Luxuriant	Good-Luxuriant	20-25°C	≤ 5 days

* depends on concentration of quarternary ammonium compounds

PACKAGING:

Pack of 25 Ready-To-Use Liquid Medium tubes containing 10 ml in each tube.

Pack of 50 Ready-To-Use Liquid Medium tubes containing 10 ml in each tube.

STORAGE

Onreceipt, store tubes in the dark at 10-25 °C. Avoid freezing and overheating. Do not open until ready to use. Minimize exposure to light. Tubed media stored as labeled until just prior to use may be inoculated up to the expiration date and incubated for the recommended incubation times. Allow the medium to warm to room temperature before inoculation.

DISPOSAL

Usermustensure safe disposal by autoclaving and/or incineration of used or unusable preparations of this product. Follow established laboratory procedures in disposing of infectious materials and material that comes into contact with clinical sample must be decontaminated and disposed of in accordance with current laboratory techniques.

REFERENCES

1. Brown, M.R.W. (1966): Turbidimetric method for the rapid evaluation of antimicrobial agents. Inactivation of preservatives by nonionic agent. – J. Soc. Cosm. Chem., 17; 185-195.
2. European Pharmacopoeia 8.0 (2014): 2.6.1. Sterility; 2.6.12. Microbial examination of non-sterile products (total viable aerobic count);
3. Japanese Pharmacopoeia 16th edition (2011): 4.05 Microbial Limit Test; 4.06 Sterility Test.
4. Russel, A.D., Ahonkhai, I. and Rogers, D.T. (1979) Microbiological applications of the inactivation of antibiotics and other antimicrobial agents. Journal of Applied Bacteriology 46:207-245.
5. United States Pharmacopoeia 38 NF 33 (2015): Sterility Tests; Microbiological Examination of Non-Sterile Products: Microbial Enumeration Tests.



Quantity



Lot / Batch Number



Temperature Unit



Manufacturer



Best Before



Certification of Good Manufacturing Practices



Catalogue No.



Authorized Representative



European Conformity



Consults Instructions for use :



QR Code



For In Vitro Diagnostic Use

NOTE: Please consult the Material Safety Data Sheet for information regarding hazards and safe handling Practices.



***For Lab Use Only**

