

CM 20715 – THIOGLYCOLLATE MEDIUM, FLUID (FLUID THIOGLYCOLLATE MEDIUM) (as per USP)

INTENDED USE

For sterility testing of biological sand for cultivation of aerobes, anaerobes and microaerophiles.

PRODUCT SUMMARY AND EXPLANATION

Brewer formulated Fluid Thioglycollate Medium for rapid cultivation of aerobes as well as anaerobes including microaerophiles by adding a reducing agent and small amount of agar. The USP, BP, EP and AOAC have recommended the media for sterility testing of antibiotics, biologicals and foods and for determining the phenol coefficient and sporicidal effect of disinfectants. However, it is intended for the examination of clear liquid or water-soluble materials. Fluid Thioglycollate Medium is also routinely used to check the sterility of stored blood in blood banks

COMPOSITION

Ingredients	Gms / Ltr
Tryptone	15.000
Yeast extract	5.000
Dextrose (Glucose)	5.500
Sodium chloride	2.500
L-Cystine	0.500
Sodium thioglycollate	0.500
Resazurin sodium	0.001
Agar	0.750

PRINCIPLE

Dextrose, tryptone, yeast extract, L-cystine provide the growth factors necessary for bacterial multiplication. L-cystine and sodium thioglycollate allows Clostridium to grow in this medium even under aerobic conditions. Also the small amount of agar used in the medium favors the growth of aerobes as well as anaerobes in the medium, even if sodium thioglycollate is deleted from the medium. Sodium thioglycollate act as a reducing agent and neutralizes the toxic effects of mercurial preservatives and peroxides formed in the medium, thereby promoting anaerobiosis, and making the medium suitable to test materials containing heavy metals. Any increase in the oxygen content is indicated by a colour change of redox indicator, resazurin to red. The small amount of agar helps in maintaining low redox potential for stabilizing the medium.

INSTRUCTION FOR USE

Dissolve 29.75 grams in 1000 ml purified/distilled water.

Heat to boiling to dissolve the medium completely.

Sterilize by autoclaving at 15 psi pressure (121°C) for 15 minutes.

Cool to 25°C and store in a cool dark place preferably below 25°C.

Note: If more than the upper one-third of the medium has acquired a pink-purple colour, the medium may be restored once by heating in a water bath or in free flowing steam until the pink-purple colour disappears.

QUALITY CONTROL SPECIFICATIONS



Appearance of Powder : Cream to yellow homogeneous free flowing powder.
 Appearance of prepared medium : Light straw coloured, clear to slightly opalescent solution with upper 10% or less medium pink-purple on standing.
 pH (at 25°C) : 7.1±0.2

INTERPRETATION

Cultural characteristics observed after incubation.

Microorganism	ATCC	Inoculum (CFU/ml)	Growth	Recovery	Incubation Temperature	Incubation Period
Clostridium sporogenes	19404	50 -100	Luxuriant	>=70%	30-35°C	Not more than 3 Days
Clostridium sporogenes	11437	50 -100	Luxuriant	>=70%	30-35°C	Not more than 3 Days
Clostridium perfringens	13124	50 -100	Luxuriant	>=70%	30-35°C	Not more than 3 Days
Bacteroides fragilis	23745	50 -100	Luxuriant	>=70%	30-35°C	Not more than 3 Days
Bacteroides vulgatus	8482	50 -100	Luxuriant	>=70%	30-35°C	Not more than 3 Days
Staphylococcus aureus subsp. aureus	25923	50 -100	Luxuriant	>=70%	30-35°C	Not more than 3 Days
Staphylococcus aureus subsp. aureus	6538	50 -100	Luxuriant	>=70%	30-35°C	Not more than 3 Days
Pseudomonas aeruginosa	27853	50 -100	Luxuriant	>=70%	30-35°C	Not more than 3 Days
Pseudomonas aeruginosa	9027	50 -100	Luxuriant	>=70%	30-35°C	Not more than 3 Days
Streptococcus pneumoniae	6305	50 -100	Luxuriant	>=70%	30-35°C	Not more than 3 Days
Escherichia coli	8739	50 -100	Luxuriant	>=70%	30-35°C	Not more than 3 Days
Salmonella Typhimurium	14028	50 -100	Luxuriant	>=70%	30-35°C	Not more than 3 Days



Bacillus subtilis subsp. spizizenii	6633	50 -100	Luxuriant	>=70%	30-35°C	Not more than 3 Days
--	------	---------	-----------	-------	---------	----------------------------

PACKAGING:

In pack size of 100 gm and 500 gm bottles.

STORAGE

Dehydrated powder, hygroscopic in nature, store in a dry place, in tightly-sealed containers between 25-30°C and protect from direct sunlight. Under optimal conditions, the medium has a shelf life of 4 years. When the container is opened for the first time, note the time and date on the label space provided on the container. After the desired amount of medium has been taken out replace the cap tightly to protect from hydration.

Product Deterioration: Do not use if they show evidence of microbial contamination, discoloration, drying or any other signs of deterioration.

DISPOSAL

After use, prepared plates, specimen/sample containers and other contaminated materials must be sterilized before discarding.

REFERENCES

1. Brewer, 1940, J. Am. Med. Assoc., 115:598.
2. British Pharmacopoeia, 2020, The Stationery office British Pharmacopoeia
3. European Pharmacopoeia, 2020, European Dept. for the quality of Medicines.
4. Federal Register, 1992, Fed. Regist., 21:640.
5. Isenberg, H.D. Clinical Microbiology Procedures Handbook 2nd Edition.
6. Jorgensen, J.H., Pfaller, M.A., Carroll, K.C., Funke, G., Landry, M.L., Richter, S.S and Warnock., D.W. (2015) Manual of Clinical Microbiology, 11th Edition. Vol. 1.
7. MacFaddin J.F., 1985, Media for Isolation-Cultivation-Identification-Maintenance of Medical Bacteria, Vol. 1, Williams and Wilkins, Baltimore.
8. Marshall, Gunnison and Luxen, 1940, Proc. Soc. Exp. Biol. Med., 43:672.
9. Nungester, Hood and Warren, 1943, Proc. Soc. Exp. Biol. Med., 52:287.
10. Portwood, 1944, J. Bact., 48:255.
11. Quastel and Stephenson, 1926, J.Biochem., 20
12. The United States Pharmacopoeia, 2019, The United States Pharmacopoeial Convention, Rockville, MD. 13. Williams H., (Ed.), 2005, Official Methods of Analysis of the Association of Official Analytical Chemists, 19th Ed., AOAC, Washington, D.C



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

*For LabUse Only

