# Product Name : Azide Maltose Agar KF

Selective medium for faecal streptococci detection and enumeration in waters and foodstuffs.

## FORMULA (G/L)

Peptospecial	10.0
Yeast Extract	10.0
Sodium Chloride	5.0
Sodium Glycerophosphate	10.0
Maltose	20.0
Lactose	1.0
Sodium Azide	0.4
Brom Cresol Purple	0.015
Agar	20.0
Final pH = 7.2 $\pm$ 0.2 at 25°C.	

### DIRECTIONS

Suspend 76.42 G of powder in 990 mL of distilled or deionized water. Heat to boiling for approximately 5 minutes and gently agitate until completely dissolved. Do not sterilize. Cool to 45-50°C. Aseptically add 2 vials of TTC 1% supplement (MB-T1867). Mix well. Dispense in petri dishes.

### TTC 1% supplement

1 Vial contents (each vial is sufficient for 500mL of medium) Triphenyltetrazolium Chloride......0.05 G

## EXPLANATION

Azide Maltose Agar KF is a selective medium utilized for isolating and enumerating the faecal *streptococci* in waters and foods by the technique of membrane filtration or by poured plate method. Sodium azide inhibits growth of Gram-negative bacteria, of streptococci not belonging to group D and of lactic bacteria (*Leuconostoc mesenteroides, Lactobacillus lactis, Lactobacillus acidophilus*). Faecal *streptococci* reduce TTC and appear with pink-red colonies.

## TECHNIC

#### **Pour Plate Technique**

- 1. Prepare appropriate dilutions of test material.
- 2. Place the selected volume of sample in a Petri dish.
- 3. Pour 15 mL of the prepared medium at 45-50°C into each plate.
- 4. Thoroughly mix the medium and sample to uniformly disperse the organisms.
- 5. Allow the agar to solidify.
- 6. Incubate plates in the inverted position at 36  $\pm$  1°C for 48 hours.

#### **Membrane Filtration Method**

- 1. Distribute 4-5 mL of medium into 55 mm dishes.
- 2. Filter samples through a sterile membrane to give 20-200 colonies on the membrane surface.
- 3. Transfer the membrane to the agar, invert the plates and incubate at 36  $\pm$  1°C for 48 hours.

4. Count all red or pink colonies, if the case with the aid of a low power (10 to 15 magnifications) binocular wide field dissecting microscope.

5. Calculate the number of faecal streptococci and report as faecal streptococci per 100 mL.



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## QUALITY CONTROL

<u>Dehydrated medium</u> Appearance: free-flowing, homogeneous. Color: light-greenish beige. <u>Prepared medium</u> Appearance: very slightly to slightly opalescent. Color: light purple. Incubation conditions:  $36 \pm 1^{\circ}$ C / 24 hours.

Microorganism	ATCC	Growth	Characteristics
Escherichia coli	25922	partially or completely inhibited	
Streptococcus faecalis	19433	good	red centers
Streptococcus faecalis	29212	good	red centers
Enterobacter aerogenes	13048	partially or completely inhibited	

## PERFORMANCE AND LIMITATIONS

Many strains of S. bovis and S. equinus are inhibited by azide. Overheating may lower pH, causing the decrease of productivity by medium.

## PRECAUTIONS

Azide Maltose Agar KF contains sodium azide. This substance is harmful by inhalation and if swallowed, irritating to eyes, respiratory system and skin. Consult safety data sheet for further details. Sodium azide reacts with many metals, especially copper, to produce explosive metal azides.

### STORAGE

The powder is very hygroscopic: store the powder at room temperature, in a dry environment, in its original container tightly closed and use it before the expiry date on the label or until signs of deterioration or contamination are evident. Store prepared medium at 2-6°C.

### REFERENCES

1. APHA (1985). Compendium of Method for the Microbiological Examination of Foods.

2. Donnelly, C.W., R.E. Bracket. 1992. Compendium of methods for the microbiological examination of foods, 3 rd ed. American Public Health Association.

## PACKAGING

Cat. No : MB-A1154 Azide Maltose Agar KF

500 G

