

Medium used for the isolation and cultivation of lipolytic bacteria.

#### CONTENTS (Liter)

Peptone	10.0 g
Sodium Chloride	5.0 g
Calcium Chloride	0.1 g
Agar	25.0 g
Final pH = 7.0 $\pm$ 0.2 at 25°C	

#### PROCEDURE

Suspend 40.1 G of powder in 990 mL of distilled or deionized water. Heat to boiling until completely dissolved. Add 10 mL of Tween 80 supplement (MB-T1861). Sterilize by autoclave at 121°C for 15 minutes. Cool to 45 - 50°C in water bath. Mix well. Pour into petri dishes.

#### INTERPRETATION

Sierra Lipolytic Agar is a medium used for the isolation and cultivation of lipolytic bacteria. Peptone provides nitrogen, carbon, amino acids and minerals. Sodium chloride maintains the osmotic balance. Lipolytic organisms split off the calcium chloride produces opaque zones around the colonies. Agar is the solidifying agent. Being soluble in water, Tween 80 is an ideal substrate for showing lipolytic activity.

#### TECHNIC

Inoculate the specimen using a sterile loop to the medium. Incubate at 35  $\pm$  2°C for 48 - 72 hours. Refer appropriate references for recommended test procedure.

#### • QUALITY CONTROL FOR USE

<u>Dehydrated medium</u> Appearance: free-flowing, homogeneous Color: light beige <u>Prepared medium</u> Appearance: slightly opalescent Color: light amber Incubation conditions:  $35 \pm 2^{\circ}C / 48 - 72$  hours

Microorganism	ATCC	Growth	Halo
Pseudomonas aeruginosa	27853	good	+
Staphylococcus aureus	25923	good	+
Candida albicans	10231	good	+
Escherichia coli	25922	good	-

# STORE

## REFERENCES

- 1. MacFaddin. (1985). Media for isolation-cultivation-identification identification-maintenance of medical bacteria, p. 695-699, vol. 1. Williams & Wilkins ,Baltimore, MD
- 2. Murray, Baron, Pfaller, Tenover, Yolken (ed.) (1995). .Manual of clinical microbiology, 6th ed. American Society for Microbiology, Washington, D.C.

### PACKAGE

