

# **Technical Data**

## Loeffler Medium Base

## M537

Loeffler Medium Base with added horse serum is used for the cultivation of *Corynebacterium diphtheriae* from clinical specimens and in pure cultures, detection of chromogenesis, proteolysis and the production of ascospores.

Composition**		
Ingredients	Gms / Litre	
Peptone, special	2.500	
Beef extract	2.500	
Sodium chloride	1.250	
Dextrose	2.500	
Final pH ( at 25°C)	7.3±0.2	
**Formula adjusted, standardized to suit performance parameters		

### **Directions**

Suspend 8.75 grams in 250 ml distilled water. Dissolve the medium completely and sterilize by autoclaving at 10 lbs pressure (115°C) for 20 minutes. Cool to 50-55°C and aseptically add 750 ml of sterile horse serum (RM1239). Mix well and aseptically dispense into sterile tubes. Sterilize the medium by inspissation at 80-85°C for 2 hours in free flowing steam for atleast 3 consecutive days.

## **Principle And Interpretation**

Loeffler Medium was originally devised by Loeffler (1) and was further modified by Perry and Petran (2) and Buck (3). Loeffler medium enhances primary and secondary isolation and cultivation of fastidious pathogenic microorganisms especially from nose and throat. It also restores virulence and other identifying properties (microscopic and colonial) after they have been lost due to prolonged incubation or repeated subculturing.

The high serum content helps in determining proteolytic activity of organisms. It is also used for demonstration of pigmentation and ascospores. Peptic digest of animal tissue, beef extract provide essential growth nutrients. Dextrose is the source of fermentable carbohydrate and energy.

Rub the swabs directly over the surface of medium and after incubation; prepare the smears from surface of slope. For proteolysis testing, inoculate slant and prior to incubation, flood the slant with Brewer Thioglycollate Medium (M019). Incubation should be carried out for 3-4 days or much longer for appearance of proteolysis. Loeffler Medium should be used in parallel with Serum Tellurite Agar for selective isolation of Corynebacteria (4).

## **Quality Control**

#### Appearance

Cream to yellow homogeneous free flowing powder

Colour and Clarity of prepared medium

Basal: Light amber coloured clear to slightly opalescent solution. ; After addition of horse serum: Off-white coloured opaque solution

#### Reaction

Reaction of 3.52% w/v aqueous solution of base at 25°C. pH :  $7.3\pm0.2$ 

pН

## 7.10-7.50

#### **Cultural Response**

M537: Cultural characteristics observed with added 750ml horse serum, after an incubation at 35-37°C for 24-48 hours.

Organism	Inoculum	Growth
	(CFU)	
Cultural Response		

Corynebacterium	50-100	fair-good
diphtheriae ATCC 11913 Pseudomonas aeruginosa ATCC 10145	50-100	good (green colonies with
Staphylococcus aureus ATCC 25923	50-100	proteolysis) good (yellow to gold colonies)

#### **Storage and Shelf Life**

Store below 30°C in tightly closed container and the prepared medium at 2-8°C. Use before expiry date on the label. .

#### Reference

1.Loeffler F., 1887, Zentralb. Bakteriol. Parasitenkd., 2:102.

2.Perry and Petran, 1939, J. Lab. Clin. Med., 25:71.

3.Buck, 1949, J. Lab. Clin. Med., 34:582.

4.MacFaddin J. F., 1985, Media for Isolation-Cultivation-Maintenance of Medical Bacteria, Vol. I, Williams and Wilkins, Baltimore.

Revision : 1 / 2011

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