

# **Technical Information**

### Rogosa Agar, Modified

Product Code: DM 2899

Application: - Used for the selective cultivation of Lactobacilli from food.

### Composition\*\*

Ingredients	Gms / Litre		
Tryptone	10.000		
Yeast extract	5.000		
Glucose	20.000		
Potassium dihydrogen orthophosphate	6.000		
Tween 80	1.000		
Triammonium citrate	2.000		
Sodium acetate	15.000		
Magnesium sulphate.7H₂O	0.575		
Manganese (II) sulphate.H₂O	0.110		
lron (II) sulphate.7H₂O	0.034		
Agar	15.000		
Final pH ( at 25°C)	6.2±0.1		
**Formula adjusted, standardized to suit performance parameters			

### Principle & Interpretation

Rogosa Agar is primarily a selective medium for the cultivation of *Lactobacillus* (1). High acetate concentration and low pH effectively suppress other bacteria, but also many strains of other lactic acid bacteria. The modification of the pH to 6.2 instead of 5.5 alters the selectivity of the medium for the whole group of lactic acid bacteria (2, 3).

Casein enzymic hydrolysate, yeast extract supply nitrogen compounds, sulphur, trace elements and vitamin B complex, essential for growth of Lactobacilli. Glucose acts as fermentable carbohydrate. Polysorbate 80 is the source of fatty acids. Ammonium citrate and sodium acetate inhibit moulds, Streptococci and many other organisms. Monopotassium phosphate buffers the medium. Magnesium sulphate, manganese sulphate and ferrous sulphate are sources of inorganic ions. Low pH of the medium and addition of acetic acid makes the medium selective for Lactobacilli, inhibiting other bacterial flora (4). It is recommended that the plates should be incubated at 30°C for 5 days or at 37°C for 3 days in an atmosphere of 95% hydrogen and 5% carbon dioxide (5). High acetate concentration and acidic pH suppress many strains of other lactic acid bacteria.

## Methodology

Suspend 74.40 grams of dehydrated powder media in 1000 ml distilled water. Mix thoroughly & heat to boiling to dissolve the medium completely. Add 1.32 ml glacial acetic acid and mix well. DO NOT AUTOCLAVE. Shake well before pouring into sterile Petri plates.

## **Quality Control**

#### Appearance

Cream to yellow homogeneous soft lumps which can be easily broken down to powder form.

#### Gelling

Firm, comparable with 1.5% Agar gel

#### Colour and Clarity

Light yellow coloured opalescent gel forms in Petri plates





#### Reaction

Reaction of 7.44% w/v aqueous solution with 0.132% v/v acetic acid at 25°C. pH: 6.2±0.1

#### pH Range

6.10-6.30

#### **Cultural Response**

DM2899: Cultural characteristics observed in presence of 5% Carbon dioxide ( $CO_2$ ) and 95%  $H_2$  after an incubation at 35-37°C for 40-48 hours.

#### **Cultural Response**

(	Organism	Inoculum (CFU)	Growth	Recovery
	Cultural Response actobacillus casei ATCC 9595	50-100	good - luxuriant	>=50%
L	actobacillus fermentum ATCC 9338	50-100	good - luxuriant	>=50%
L	actobacillus leichmanni ATCC 4797	50-100	good - luxuriant	>=50%
L	actobacillus plantarum ATCC 8014	50-100	good -luxuriant	>=50%
9	Staphylococcus aureus ATCC 25923	>=10³	inhibited	0%

## Storage and Shelf Life

**Dried Media:** Store below 30°C in tightly closed container and the prepared medium at 2 - 8°C. Use before expiry date on the label. **Prepared Media:** 2-8° in sealable plastic bags for 2-5 days.

### **Further Reading**

- 1. Rogosa,J.,Mitchell J.A. and Wiseman,R.F.(1951)A selective medium for the isolation and enumeration of oral and fecal lactobacilli.J.Bacteriol.62,132-133.
- 2. ISO(1984)Drafts reports.Enumeration of Lactobacteriaceae in meat and meat products.ISO/TC 34/SC 6/WG 15,no.3and no.5. International Organization for Standardization,Geneva.
- 3. Reuter, G. (1985) Elective and selective media for lactic acid bacteria . Int. J. Food Microbiol. 2,55-68.
- 4. MacFaddin J. F., 1985, Media for Isolation-Cultivation-Identification- Maintenance of Medical Bacteria, Vol. I, Williams and Wilkins, Baltimore. Md.
- 5. Sharpe M. L. (Ed.), 1960, Lab-Practice, 9(4): 223.

### Disclaimer :

- User must ensure suitability of the product(s) in their application prior to use.
- The product conform solely to the technical information provided in this booklet and to the best of knowledge research and development work carried at CDH is true and accurate
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