

## Technical Information

### Brilliant Green Agar with Phosphates

**Product Code: DM 19711**

**Application:** - Brilliant Green Agar with Phosphates is recommended for detection and enumeration of Salmonellae.

#### Composition\*\*

Ingredients	Gms / Litre
Peptic digest of animal tissue	10.000
Meat extract	5.000
Yeast extract	3.000
Lactose	10.000
Sucrose	10.000
Disodium phosphate	1.000
Monosodium phosphate	0.600
Phenol red	0.090
Brilliant green	0.005
Agar	15.000
Final pH (at 25°C)	6.9±0.2

\*\*Formula adjusted, standardized to suit performance parameters

#### Principle & Interpretation

*Salmonella* species is responsible for many types of infections, ranging from mild self-limiting gastroenteritis to life threatening typhoid fever. The most common type of *Salmonella* disease is self-limiting gastroenteritis with fever lasting less than 2 days and diarrhoea less than 7 days.<sup>(8)</sup> This medium contains peptic digest of animal tissue, meat extract and yeast extract as sources of carbon, nitrogen, vitamins, amino acids and essential nutrients. The two sugars, namely lactose and sucrose act as energy sources. Fermentation of lactose and / or sucrose in the medium results in the formation of acidic pH which is detected by phenol red indicator. This medium is formulated according to the recommendation of Rijks Institute Voorde Volksgezondheid (National Institute for Public Health), Utrecht<sup>(1, 2)</sup>. It is widely accepted and used in ISO Standards also<sup>(3, 4, 5)</sup> because of its improved performance with respect to recovery of even smaller numbers of *Salmonella* species, and inhibition of *Escherichia coli*, *Proteus* and *Pseudomonas* species<sup>(6)</sup>. The medium can be further supplemented with sulphacetamide (1 g/l) and sodium mandelate (0.25 g/l) to inhibit contaminating microorganisms when the sample is suspected to be containing large number of competing organisms along with *Salmonella* species<sup>(7)</sup>.

#### Methodology

Suspend 54.69 grams of powder media in 1000 ml distilled water. Shake well & heat with occasional agitation and bring just to the boil to dissolve the medium completely. DO NOT AUTOCLAVE. For maximum recovery, Sulpha Supplement (MS2068) may be aseptically added. Cool to 50°C. Mix well and pour into sterile Petri plates.

#### Quality Control

##### Physical Appearance

Light yellow to pink coloured homogeneous free flowing powder.

##### Gelling

Firm, comparable with 1.5% Agar gel.

##### Colour and Clarity of prepared medium

Greenish brown coloured clear to slightly opalescent gel forms in Petri plates.

**Reaction:** Reaction of 5.46 % w/v aqueous solution at 25°C. pH : 6.9±0.2

**pH Range** 6.70-7.10

**Cultural Response/ characteristics**

**DM 1971I:** Cultural characteristics observed after an incubation at 35-37°C for 18-24 hours.

Organism	Growth	Colour of colony
<i>Escherichia coli</i> ATCC 25922	Inhibited	
<i>Proteus vulgaris</i> ATCC 13315	none-poor	Red
<i>Pseudomonas aeruginosa</i> ATCC 10145	none-poor	Red
<i>Salmonella Enteritidis</i> ATCC 13076	Luxuriant	bright red
<i>Salmonella Typhimurium</i> ATCC 14028	luxuriant	bright red

## Storage and Shelf Life

**Dried Media:** Store below 30°C in tightly closed container and use before expiry date as mentioned on the label.

**Prepared Media:** 2-8° in sealable plastic bags for 2-5 days.

## Further Reading

1. Edel W. and Kampelmacher E.H., 1969, Bull. W.H.O., 41:297.
2. Edel W. and Kampelmacher E.H., 1969, Bull. W.H.O., 39:487.
3. Anon, 1975, International Organization for Standardization, Meat and Meat products. Ref. Method, ISO: 3 565.
4. Anon, 1981, International Organization for Standardization, Microbiology Ref. Methods, ISO: 6579.
5. Anon, 1985, International Organization for Standardization, Milk and Milk Products, Ref. Method, ISO: 6785.
6. Read R. B. and Reyes A.L., 1968, Appl. Microbiol., 16:746.
7. Watson U.C. and Walker A.P., 1978, J. Appl. Bact., 45:195.
8. Murray P.R., Baron J. H., Pfaller M.A., Jorgensen J.H. and Tenover F.C., (Ed.), 2003, Manual of Clinical Microbiology, 8th Ed., American Society for Microbiology, Washington, D.C.

## Disclaimer :

- User must ensure suitability of the product(s) in their application prior to use.
- The product conforms 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.
- **Central Drug House Pvt. Ltd.** reserves the right to make changes to specifications and information related to the products at any time.
- Products are not intended for human or animal diagnostic or therapeutic use but for laboratory, research or further manufacturing of diagnostic reagents extra.
- Statements contained herein should not be considered as a warranty of any kind, expressed or implied, and no liability is accepted for infringement of any patents.
- Do not use the products if it fails to meet specifications for identity and performance parameters.