

TECHNICAL DATA SHEET

BISMUTH SULFITE AGAR

DETECTION OF *SALMONELLA* TYPHI AND OTHER *SALMONELLAE*

1 INTENDED USE

Bismuth Sulfite (modified Wilson Blair agar) Agar is a selective medium used to isolate *Salmonella* Typhi and other salmonellas in pathological products of animal origin, water, dairy and other food products. Bismuth Sulfite agar can be used in the normalized methods for *Salmonella* detection as the second isolation media.

The typical composition corresponds to that defined in the standards NF EN ISO 6579, NF EN ISO 6785 and NF EN ISO 19250.

2 HISTORY

In 1926, Wilson and Blair combined bismuth and sodium sulfite in a medium destined to isolate *Salmonella* of the typhi and paratyphi groups. In 1956, Hajna and Damon described a modified formula which was recommended by the United States Pharmacopoeia.

3 PRINCIPLES

The concentrations of brilliant green and bismuth sulfite inhibit accompanying Gram-positive flora and most enterobacteria, except for *Salmonella* and several *Shigella*.

Using the sulfur compounds in the medium, *Salmonella* releases hydrogen sulfide which produces a metallic precipitate in the presence of ferrous sulfate, giving the colonies a black or sometimes green color.

It is particularly recommended to first enrich using Tetrathionate, Selenite or Rappaport-Vassiliadis Broths and to simultaneously inoculate onto other less selective media : MacConkey, XLD or Hektoen Enteric Agars, for example.

Because of its elevated inhibitory power, this medium enables a highly contaminated inoculum to be used.

4 TYPICAL COMPOSITION

The composition can be adjusted in order to obtain optimal performance.

For 1 liter of media :

- Tryptone	5,00 g
- Peptic digest of meat	5,00 g
- Meat extract	5,00 g
- Glucose	5,00 g
- Disodium phosphate	4,00 g
- Ferrous sulfate	0,30 g
- Bismuth ammonium citrate.....	1,85 g
- Sodium sulfite	6,15 g
- Brilliant green	25,0 mg
- Bacteriological agar.....	14,70 g

pH of ready-to-use media at 25 °C : 7,6 ± 0,2.

5 PREPARATION

- Dissolve 47,0 g of dehydrated media (BK004) in 1 liter of distilled or demineralized water.
- Slowly bring to boiling, stirring with constant agitation until complete dissolution.
- Do not autoclave.
- Cool and maintain the media in a molten state at 44-47 °C.
- Homogenize well in order to disperse the precipitate.
- Pour into sterile Petri plates.
- Let solidify on a cold, flat surface.
- Dry in an incubator with the covers partially removed.

✓ **Reconstitution :**
47,0 g/L

✓ **Sterilization :**
Bring to boil

Note :

Immediately after its preparation, the medium has optimal selectivity which gradually decreases with time. This is why it is not recommended to store the ready-to-use medium more than 4 days at 2-8°C.

6 INSTRUCTIONS FOR USE

- Inoculate by streaking the medium with the enrichment media used.
- Incubate at 37 ± 1 °C for 24 and 48 hours.

✓ **Inoculation :**
On surface

✓ **Incubation :**
24 h and 48 h at 37 °C

7 RESULTS

Characteristic colonies of *Salmonella* Typhi appear black, flat, dry colonies surrounded by a black-brown zone with metallic reflections.

Green to brownish colonies without dark zones are characteristic of *Salmonella* Enteritidis, *Salmonella* Gallinarium, *Salmonella* Choleraesuis, and *Salmonella* Paratyphi.

Coliforms, *Proteus* and *Shigella* are strongly inhibited but may sometimes yield small greenish or brownish colonies.

See ANNEX 1 : PHOTO SUPPORT.

8 QUALITY CONTROL

Dehydrated media : greenish-beige powder, free-flowing and homogeneous.

Prepared media in plates : green-beige agar.

Typical culture response after 48 hours of incubation at 37 °C:

Microorganisms	Growth	Characteristics
<i>Salmonella</i> Typhimurium WDCM 00031	Good	Black colonies with metallic reflection
<i>Salmonella</i> Enteritidis WDCM 00030	Good	Greenish brown colonies
<i>Escherichia coli</i> WDCM 00012	Partially inhibited	Green colonies
<i>Shigella sonnei</i> WDCM 00127	Partially inhibited	Green colonies
<i>Enterococcus faecalis</i> WDCM 00087	Inhibited	-
<i>Staphylococcus aureus</i> WDCM 00034	Inhibited	-

9 STORAGE / SHELF LIFE

Dehydrated base media : 2-30 °C.

The expiration date is indicated on the label.

Prepared media in plates (*) : 4 days at 2-8 °C, shielded from light.

(*) Benchmark value determined under standard preparation conditions, following manufacturer's instructions.

10 PACKAGING

Dehydrated media :

500 g bottle BK004HA

11 BIBLIOGRAPHY

Wilson, J.W., and Blair, E.M. 1931. Further experience of the Bismuth Sulphite Media in the isolation of *Bacillus typhosus* and *Bacillus paratyphosus* B from faeces, sewage and water. J. Hyg., 31: 138.

Wilson, J.W. 1938. Isolation of *Bact. typhosum* by means Bismuth Sulphite Medium in water and milk born epidemics. J.Hyg., 38: 507-519.

NF EN ISO 19250. Juin 2013. Qualité de l'eau. Recherche de *Salmonella* spp.

NF EN ISO 6579. Décembre 2002. Microbiologie des aliments - Méthode horizontale pour la recherche des *Salmonella* spp.

NF EN ISO 6785. Avril 2008. Lait et produits laitiers – Recherche de *Salmonella* spp.

12 ADDITIONAL INFORMATION

The information provided on the labels take precedence over the formulations or instructions described in this document and are susceptible to modification at any time, without warning.

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Bismuth Sulphite Agar (modified Wilson Blair)

Detection of *Salmonella* Typhi and other *Salmonellae*

Results :

Growth obtained after 24 hours of incubation at 37 °C (surface inoculation).

***Salmonella* Typhi**
Characteristic colony :
Black color with metallic reflects,
flat and dry, surrounded by a
brown-black halo.

