

### Specification

Liquid medium used for the enrichment of *Salmonella* and *Shigella* from clinical specimens and other products according to ISO standards.

### Presentation

20 Tubes  
Tube 16 x 113 mm  
with: 10 ± 0,2 ml

#### Packaging Details

16x113 mm glass tubes, ink labelled, metal-Non  
injectable cap. - 20 tubes per box

#### Shelf Life

9 months

#### Storage

8-25°C

### Composition

Composition (g/l):

Tryptone.....5.0  
Lactose.....4.0  
Disodium hydrogenphosphate.....10.0  
Sodium selenite.....4.0  
L-Cystine.....0.01

### Description /Technique

It is essentially an enrichment medium for *Salmonella* found in food or pathological materials, such as faeces or urine. It is used as an initial step prior to isolation on selective media such as SS Agar or Hektoen Agar.

Essentially, it is an enrichment medium for *Salmonella* coming from food or pathologic materials, such as faeces or urine, in a previous step to isolation in selective media plates, such as Agar SS or Hektoen Agar.

For normal assays it is advisable an incubation at 37°C for a period never superior to 18 hours, since within that period a good nutrition of coliforms and an enhancement of pathogens is reached, but after 24 hours that effect seems to disappear and the growth of companion flora may hide salmonella.

Red precipitate apparition before inoculation means the medium has been overheated, in which case the selective properties are worse. Presence of copious sample residuum may also inactivate the selective power of the medium, overall if sample is faeces and egg powder. In this case, it is better to make a dilution 1:10 and let it settle to separate the biggest particles, then inoculate Selenite cystine broth with an aliquot portion of it, maintaining the proportion 1:10 between sample and medium.

It has been demonstrated that when it is desired to isolate *Salmonella* from faeces, results are better if enrichment medium incubation is performed at 43°C. This procedure only seems to fail with *Salmonella typhi*.

When starting material is urine, the best procedure is to use Selenite cystine broth in double concentration, and to inoculate it in an equal volume of urine. Anyway, subculture must always be done after 6 hours of incubation and before 24 hours. Most authors recommend the simultaneous use of another enrichment broth, such as Tetrathionate broth.

### Quality control

#### Physical/Chemical control

Color : Clear and yellowish      pH: 7 ± 0.2 at 25°C

#### Microbiological control

Inoculate: Practical range 100 ± 20 CFU; Min. 50 CFU (Productivity)/ 10<sup>4</sup>-10<sup>6</sup> (Selectivity).

Aerobiosis. Incubation at 37 ± 1°C, reading after 24 ± 3 h

#### Microorganism

*Escherichia coli* ATCC® 25922, WDCM 00013  
*Salmonella enterica* ATCC® 13076, WDCM 00030  
*Salmonella typhimurium* ATCC® 14028, WDCM 00031  
Mezcla : *Salmonella* (14028) + *E. coli* (8739)

#### Growth

Partial Inhibition - Pink  
Good - Colourless colonies  
Good - Colourless colonies  
Conform

#### Sterility Control

Incubation 48 hours at 30-35°C and 48 hours at 20-25°C: NO GROWTH  
Check at 7 days after incubation in same conditions

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