Rappaport Vassiliadis Salmonella Enrichment Broth

Intended use

For selective enrichment of *Salmonella* species from pharmaceutical & clinical sample in accordance with harmonized methods of USP, EP, BP & JP.

Composition**

<table>
<thead>
<tr>
<th>Ingredients</th>
<th>Gms / Litre</th>
</tr>
</thead>
<tbody>
<tr>
<td>Soya peptone</td>
<td>4.500</td>
</tr>
<tr>
<td>Sodium chloride</td>
<td>8.000</td>
</tr>
<tr>
<td>Dipotassium hydrogen phosphate</td>
<td>0.400</td>
</tr>
<tr>
<td>Potassium dihydrogen phosphate</td>
<td>0.600</td>
</tr>
<tr>
<td>Magnesium chloride, hexahydrate</td>
<td>29.000</td>
</tr>
<tr>
<td>Malachite green</td>
<td>0.036</td>
</tr>
</tbody>
</table>

**Formula adjusted, standardized to suit performance parameters

Directions

Label the ready to use LQ104C bottle. Inoculate the sample and Incubate at specified temperature and time.

Principle And Interpretation

Rappaport Vassiliadis Salmonella Enrichment Medium is designed according to the revised formulation by Van Schothorst et al (10) and is recommended for the selective enrichment of Salmonellae from pharmaceutical products. This medium can also be used in direct enrichment of samples containing low inoculum. Present medium is a modification of the Rappaport Vassiliadis Enrichment Broth described by Van Schothorst and Renauld (11). It is prepared in accordance with the harmonized methodology of USP/EP/BP/JP/IP (9,1,2,5,3) has been found to be superior to other *Salmonella* selective medias. Addition of magnesium chloride to the medium was reported by Peterz et al (8). *Salmonella* species can be isolated from human faeces without pre-enrichment by using this medium.

*Salmonella* generally survive at little high osmotic pressure, grow at slightly low pH and are resistant to malachite green compared to other bacteria. These characteristics are exploited in this medium for selective enrichment of Salmonella. Magnesium chloride present in the medium raises the osmotic pressure. Natural sugars of soya peptone provide essential growth nutrients and enhance the growth of *Salmonella* (4). Phosphate buffers the medium to maintain constant pH. Sodium chloride maintains the osmotic balance. Malachite green inhibits many gram-positive bacteria, while selectively enriches *Salmonella*.

The relatively lower concentration of nutrition, also aids selective enrichment of *Salmonella*. This medium was reported to be superior to *Salmonella* selective medium like Tetrathionate Broth and Selenite enrichment broth and to Tetrathionate-Brilliant Green Broth for the detection of Salmonellae in milk samples. The enriched culture of Rappaport Vasiliadis Salmonella Enrichment Broth (MH1491) can be further subcultured and isolated on Xylose Lysine Deoxycholate Agar (MH031).

Type of specimen

Pharmaceutical samples; Clinical samples: faecal dilution

Specimen Collection and Handling

For pharmaceutical samples follow appropriate techniques for handling specimens as per established guidelines (9,1,2,5,3). For clinical samples follow appropriate techniques for handling specimens as per established guidelines (4,6).

After use, contaminated materials must be sterilized by autoclaving before discarding.

Please refer disclaimer Overleaf.
Warning and Precautions
In Vitro diagnostic Use only. Read the label before opening the container. Wear protective gloves/protective clothing/eye protection/ face protection. Follow good microbiological lab practices while handling specimens and culture. Standard precautions as per established guidelines should be followed while handling clinical specimens. Safety guidelines may be referred in individual safety data sheets.

Limitations
Due to variable nutritional requirements, some strains may show poor growth on this medium.

Performance and Evaluation
Performance of the medium is expected when used as per the direction on the label within the expiry period when stored at recommended temperature.

Quality Control
Appearance
Sterile clear Rappaport Vassiliadis Salmonella Enrichment Broth in bottles.

Colour
Bluish green coloured solution.

Quantity of medium
100 ml of medium in bottles.

Reaction
5.00- 5.40

Sterilization Method
Sterilized by autoclaving at 115 °C as per validated cycle

Sterility Assurance Level
Sterility assurance level of media was validated against *B. subtilis* Spore strips. The spore strips exposed at 115°C and unexposed strips were inoculated seperately in 100ml Soyabean Casein Digest Medium and incubated at 35°C for 7 days.

Exposed spore strips
No growth observed

Unexposed spore strips
Luxuriant growth observed

Sterility test
Passes release criteria

Cultural Response
Cultural characteristics observed after incubation at 30-35°C for 18-24 hours. Recovery is carried out using XLD Agar (M031).

Growth promoting properties
Clearly visible growth of microorganism comparable to that previously obtained with previously tested and approved lot of medium occurs at the specified temperature for not more than the shortest period of time specified inoculating <=100 cfu (at 30-35°C for <=18 hours).

Inhibitory properties
No growth of the test microorganism occurs for the specified temperature for not less than longest period of time specified inoculating >=100 cfu (at least 100 cfu) (at 30-35°C for >= 24 hours).

<table>
<thead>
<tr>
<th>Organism</th>
<th>Inoculum (CFU)</th>
<th>Growth</th>
<th>Observed Lot value (CFU)</th>
<th>Recovery</th>
<th>Colour of colony</th>
<th>Incubation temperature</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Growth promoting</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>Salmonella Typhimurium</em> ATCC 14028 (00031*)</td>
<td>50 -100</td>
<td>luxuriant</td>
<td>&gt;=35</td>
<td>&gt;=70 %</td>
<td>red with black centers</td>
<td>&lt;=18 hrs</td>
</tr>
<tr>
<td><em>Salmonella Abony NCTC 6017</em> (00029*)</td>
<td>50 -100</td>
<td>luxuriant</td>
<td>&gt;=35</td>
<td>&gt;=70 %</td>
<td>red with black centers</td>
<td>&lt;=18 hrs</td>
</tr>
<tr>
<td><strong>Inhibitory</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Staphylococcus aureus subsp. aureus ATCC 6538 (00032*)</td>
<td>&gt;=10³</td>
<td>inhibited</td>
<td>0</td>
<td>0 %</td>
<td></td>
<td>&gt;=24 hrs</td>
</tr>
</tbody>
</table>

Please refer disclaimer Overleaf.
### Additional Microbiological testing

<table>
<thead>
<tr>
<th>Organism</th>
<th>Concentration</th>
<th>Growth</th>
<th>pH</th>
<th>Color</th>
<th>Incubation Period</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Escherichia coli</em> ATCC 25922</td>
<td>50 - 100</td>
<td>none-poor</td>
<td></td>
<td>yellow</td>
<td>18 - 24 hrs</td>
</tr>
<tr>
<td><em>Escherichia coli</em> ATCC 8739</td>
<td>50 - 100</td>
<td>none-poor</td>
<td></td>
<td>yellow</td>
<td>18 - 24 hrs</td>
</tr>
<tr>
<td><em>Salmonella Enteritidis</em> ATCC 13076</td>
<td>50 - 100</td>
<td>luxuriant</td>
<td></td>
<td>red with black centre</td>
<td>18 - 24 hrs</td>
</tr>
<tr>
<td><em>Salmonella Paratyphi B</em> ATCC 8759</td>
<td>50 - 100</td>
<td>luxuriant</td>
<td></td>
<td>red with black centre</td>
<td>18 - 24 hrs</td>
</tr>
<tr>
<td><em>Staphylococcus aureus subsp. aureus</em> ATCC 25923</td>
<td>&gt;=10³</td>
<td>inhibited</td>
<td></td>
<td>&gt;=24 hrs</td>
<td></td>
</tr>
<tr>
<td><em>Pseudomonas aeruginosa</em> ATCC 9027</td>
<td>&gt;=10³</td>
<td>inhibited</td>
<td></td>
<td>&gt;=24 hrs</td>
<td></td>
</tr>
<tr>
<td><em>Pseudomonas aeruginosa</em> ATCC 27853</td>
<td>&gt;=10³</td>
<td>inhibited</td>
<td></td>
<td>&gt;=24 hrs</td>
<td></td>
</tr>
<tr>
<td><em>Enterococcus faecalis</em> ATCC 29212</td>
<td>&gt;=10³</td>
<td>inhibited</td>
<td></td>
<td>&gt;=24 hrs</td>
<td></td>
</tr>
<tr>
<td><em>E. coli + S. Typhimurium</em> (mixed culture)</td>
<td>50 - 100</td>
<td>luxuriant</td>
<td></td>
<td>red with black centre</td>
<td>18 - 72 hrs</td>
</tr>
</tbody>
</table>

Key: (*) Corresponding WDCM numbers.

### Storage and Shelf Life

Store between 15-25°C. Use before expiry date on the label. Product performance is best if used within stated expiry period.

### Disposal

User must ensure safe disposal by autoclaving and/or incineration of used or unusable preparations of this product. Follow established laboratory procedures in disposing of infectious materials and material that comes into contact with clinical sample must be decontaminated and disposed of in accordance with current laboratory techniques (4, 6).

### Reference

3. Indian Pharmacopoeia, 2018, Govt. of India, Ministry of Health and Family Welfare, New Delhi
In vitro diagnostic medical device

CE Marking

Storage temperature

Do not use if package is damaged

HiMedia Laboratories Pvt. Limited,
B/4-6, MIDC, Dindori, Nashik MH

www.himedialabs.com

CE Partner 4U, Esdoornlaan 13, 3951 DB Maarn The Netherlands,
www.cepartner4u.eu

Disclaimer:
User must ensure suitability of the product(s) in their application prior to use. Products conform solely to the information contained in this and other related HiMedia™ publications. The information contained in this publication is based on our research and development work and is to the best of our knowledge true and accurate. HiMedia™ Laboratories 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 or therapeutic use but for laboratory, diagnostic, research or further manufacturing use only, unless otherwise specified. 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.