



# **Specimen Preservative Medium Base (SP Hajna)**

**M300** 

Specimen Preservative Medium (SP Hajna) is used for collection, transportation and preservation of stool specimens or rectal swabs for the isolation of members of *Enterobacteriaceae*.

## **Composition\*\***

Ingredients	Gms / Litre	
Yeast extract	1.000	
Ammonium phosphate	4.000	
Monopotassium phosphate	2.000	
Sodium chloride	5.000	
Sodium citrate	5.000	
Magnesium sulphate	0.400	
Sodium deoxycholate	0.500	
Final pH ( at 25°C)	$7.0\pm0.2$	
**Formula adjusted, standardized to suit performance parameters		

Directions

Suspend 17.9 grams in 700 ml distilled water. Heat if necessary to dissolve the medium completely. Add 300 ml of neutral glycerol. Mix well and dispense as desired. Sterilize by autoclaving at 115°C for 15 minutes.

# **Principle And Interpretation**

Specimen Preservative Medium (SP Hajna) is designed for transport and preservation of clinical specimen. Transport media were primarily developed by Moffet et al (1) and Stuart et al (2) for carrying *gonococcal* specimens. Transport media are chemically defined, semisolid, non-nutritive, phosphate buffered media that provide a reduced environment. These media are formulated to maintain the viability and/ or infectivity of the microorganisms without significant growth during the period between collection and culture of the specimen. This medium is suitable for preserving gram-negative rods such as *Salmonella*, *Shigella* and *Klebsiella*. In comparative studies using Specimen Preservation Medium Base and glycerin preservation solution for different *Salmonella*, *Shigella* and *Klebsiella* strains, it was observed that preservation of organisms were twice more effective in Specimen Preservation Medium Base.

Specimen Preservative Medium is made inhibitory for gram-positive organisms by sodium deoxycholate and sodium citrate. Sodium, magnesium, potassium salts control permeability of bacterial cells. Sodium chloride helps in maintaining osmotic balance in medium. Yeasts extract acts as the source of nitrogen, vitamins and growth factors.

# **Quality Control**

Appearance Cream to yellow homogeneous free flowing powder Colour and Clarity of prepared medium Light amber coloured clear solution Reaction Reaction of 1.79% w/v aqueous solution at 25°C. pH : 7.0±0.2 pН 6.80-7.20 **Cultural Response** M300: Cultural characteristics observed after an incubation at 35-37°C for 18-24 hours. Organism Inoculum **Recovery** (on (CFU) Soyabean **Casein Digest** Agar

#### **Cultural Response**

Enterobacter aerogenes ATCC 13048	50-100	good-excellent, no increase in number
Escherichia coli ATCC 25922	50-100	good-excellent, no increase in number
Klebsiella pneumoniae ATCC 13883	50-100	good-excellent, no increase in number
Shigella flexneri ATCC 12022	50-100	good-excellent, no increase in number
Salmonella Typhimurium ATCC 14028	50-100	good-excellent, no increase in number

## **Storage and Shelf Life**

Store below 30°C in tightly closed container and the prepared medium at 2 - 8°C. Use before expiry date on the label.

### Reference

1. Moffett M., Young D. and Stuart R. D., 1948, Brit, Med. J., 2:241.

2. Stuart S. D., Toshach S. R. and Patsula T. M., 1954, Can. J. Public Health, 45:73.

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HiMedia Laboratories Pvt. Ltd. A-516, Swastik Disha Business Park, Via Vadhani Ind. Est., LBS Marg, Mumbai-400086, India. Customer care No.: 022-6147 1919 Email: techhelp@himedialabs.com

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