

Dehydrated Culture Media Bases / Media Supplements

Technical Information

RS MiVeg Medium Base

Product Code : VM1576

Application:- RS (Rimler-Shotts) MiVeg Medium is used for the selective isolation, cultivation and presumptive identification of *Aeromonas hydrophila*.

Composition			
Ingredients	Gms / Litre		
Yeast extract	3.0		
Maltose	3.5		
L-Cysteine hydrochloride	0.3		
L-Lysine hydrochloride	5.0		
L-Ornithine hydrochloride	6.5		
Sodium thiosulphate	6.8		
Ferric ammonium citrate	0.8		
Synthetic detergent No. III	1.0		
Sodium chloride	5.0		
Bromo thymol blue	0.03		
Agar	13.5		
Final pH (at 25°C)	7.0 ± 0.2		
** Formula adjusted, standardized to suit perfo	rmance parameters.		

Principle & Interpretation

RS (Rimler-Shotts) MiVeg Medium Base is prepared by adding synthetic detergent No. III instead of sodium deoxycholate which makes the medium free of BSE/TSE risks. RS (Rimler-Shotts) MiVeg Medium Base is the modification of RS Medium which was developed by Rimler and Shotts used for the rapid isolation and identification of *Aeromonas hydrophila* (1). RS medium shows differentiation of characteristics based upon the biochemical reactions. Maltose fermenting organisms produces yellow colonies and organisms that gives H₂S production along with Maltose fermentation give rise to black centered yellow colonies on plates. The third colonial type give greenish-yellow to green colonies indicating lysine or ornithine decarboxylation or both. The colonies which are greenish-yellow to green with black center indicate the decarboxylation of both amino acids plus H₂S production.

The balance of ingredients provide a nutrient base and chemophysical stability to the medium. Synthetic detergent No. III and Novobiocin inhibits gram-positive organisms and *Vibrio* species. Bromo thymol blue is the indicator system for maltose fermentation. Incubation should be done at 37°C and the results should be seen after 20 hours but not after 24 hours, as plates observed after 26 hours demonstrate a slow reversion of the fermentation.

Methodology

Suspend 45.43 grams of powder media in 990 ml distilled water. Mix thoroughly and heat to boiling to dissolve the medium completely. DO NOTAUTOCLAVE. Cool to 45°-50°C and aseptically add Novobiocin Supplement (MS2096). Mix well before pouring into sterile petriplates.

Quality Control

Physical Appearance

Light green coloured, homogeneous, free flowing powder.





Bases / Media Supplements

Gelling

Firm, comparable with 1.35% Agar gel.

Colour and Clarity of prepared medium

Dark green coloured, clear gel forms in petri plates.

Reaction

Reaction of 4.54% w/v aqueous solution is pH 7.0 \pm 0.2 at 25°C.

pH Range

6.8-7.2

Cultural Response/Characteristics

Cultural characteristics observed after an incubation at 35-37°C for 24 hours on addition of Novobiocin supplement (MS2096). Organisms (ATCC) Inoculum (CELI) Pocovory Maltosa formantation Lyning/ цс

	moculum (CPO)	Recovery	Matose rementation	Ornithine*	F125
Aeromonas hydrophila (7966)	102-103	>50%	+	_	_
Citrobacter freundii (8090)	102-103	>50%	-	V	+
Escherichia coli (25922)	10 ² -10 ³	>50%	-	V	_
Proteus vulgaris (13315)	10 ² -10 ³	>50%	+	_	+
Salmonella serotype Typhi (6539)	102-103	>50%	+	-	-
Key : Maltose fermentation :	+ = yellow cole + = shades of	oured colonies	, to		

Lysine and/or Ornithine :	+ = shades of greenish yellow to
*decarboxylation	yellow coloured colonies
Hydrogen Sulphide (H ₂ S) production :	+ = black centered colonies

- = Negative reaction

V = Variable

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: 2-8⁰ in sealable plastic bags for 2-5 day.

Further Reading

. Shotts E. B. Jr. and Rimler R., 1973, Appl, Microbiol., 26(4):550.

Disclaimer:

- User must ensure suitability of the product(s) in their application prior to use. •
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