

# **Technical Information**

# **Pseudomonas Asparagine Broth**

#### Product Code: DM 2096

Application: - Pseudomonas Asparagine Broth is used for presumptive determination of *Pseudomonas aeruginosa* from recreational or natural water as per APHA.

Composition**			
Ingredients	Gms / Litre		
DL-Asparagine	3.000		
Dipotassium phosphate	1.000		
Magnesium sulphate	0.500		
Final pH ( at 25°C)	7.0±0.2		
**Formula adjusted, standardized to suit performa	nce parameters		

#### Principle & Interpretation

Recreational water in swimming pool is like a body of water holding in a structure. Microorganisms of concern in swimming pool are those causing infections of ear, skin and upper respiratory tract etc. *Pseudomonas aeruginosa* is one of those organisms which in responsible for a large percentage of swimming pool associated intection. Asparagine Medium is recommended for the microbiological analysis of water. This is an excellent enrichment medium for *P. aeruginosa*; It is also used in the multiple-tube technique for microbiological analysis of recreational waters. *from different sources* <sup>(1)</sup>. Pseudomonas Asparagine Broth medium is a relatively simple medium containing an amino acid DL-asparagine and two salts dipotassium phosphate and magnesium sulphate. Asparagine is the amino acid and carbon source while phosphate and sulphate provide the ions for the growth of *P. aeruginosa*. Dipotassium phosphate also helps in maintaining the buffering conditions of the medium. This medium is only a presumptive medium for *P. aeruginosa*, and further confirmatory tests are necessary for the identification. For five tubes multiple tube test, use 10 ml of single strength Aspargine Broth for inocula of 1 ml or less and 10 ml double strength Aspargine Broth for 10 ml inocula. For swimming pools, higher dilutions may be necessary. Incubate inoculated tubes at 35-37°C. After 24 hours and again after 48 hours of incubation examine tubes under long water ultraviolet light in a darkened room. Production of a green fluorescent pigment indicates a positive presumptive test. Confirmation is performed by subculturing a loop from each tube in Acetamide Medium (DM2148). Development of purple colour within 24-36 hours of incubation at 35-37°C is a positive confirmed test for *P. aeruginosa*.

#### Methodology

Suspend 4.5 grams of powder media in 1000 ml distilled water. Shake well & Gently boil to dissolve the medium completely. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Mix well and dispense as desired.





#### uality Control

Physical Appearance
White to cream homogeneous free flowing powder
Colour and Clarity of prepared medium
Colourless clear solution with slight precipitate.
Reaction
Reaction of 0.45% w/v aqueous solution at 25°C. pH : 7.0±0.2
pH range
6.80-7.20
Cultural Response/Characteristics
DM 2096: Cultural characteristics observed after an incubation at 35-37°C for 20 - 24 hours.

Organism	lnoculum (CFU)	Growth
Pseudomonas aeruginosa	50-100	luxuriant
ATCC 27853		

# Storage and Shelf Lifez

Dried Media: Store below 30°C in tightly closed container and use before expiry date as mentioned on the label. **Prepared Media:** 2-8<sup>0</sup> in sealable plastic bags for 2-5 days.

### Further Reading

1.1. Eaton A. D., Clesceri L. S. and Greenberg A W., (Eds.), 2005, Standard Methods for the Examination of Water and Wastewater, 21st Ed., APHA, Washington, D.C.

#### Disclaimer :

- User must ensure suitability of the product(s) in their application prior to use.
- The product conform solely to the technical information provided in this booklet and to the best of knowledge research and development work carried at CDH is true and accurate
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