

**Polysorbate 80 HiVeg™ Agar (Twin Pack)****MV1307**

Polysorbate 80 HiVeg Agar is recommended for the cultivation of variety of microorganisms.

**Composition \*\* :**

Ingredients	Grams/Litre
Part A:	
HiVeg peptone	10.00
Agar	15.0
Part B:	
Polysorbate 80	10.0

Final pH (at 25°C ) 7.2 ± 0.2

\*\* Formula adjusted, standardized to suit performance parameters.

**Directions :**

Suspend 25.0 grams of Part A in 990 ml distilled water. Heat to boiling to dissolve the medium completely. Add 10 ml of Part B. Mix well and sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes.

**Principle and Interpretation :**

Polysorbate 80 HiVeg Agar is prepared by using HiVeg peptone in place of peptic digest of animal tissue which is free from BSE/TSE risks. This medium is the modification of Polysorbate 80 Agar which is recommended for the cultivation of variety of microorganisms. It is a nutritional medium containing neutralizing agent. The medium contains HiVeg peptone which provide the necessary nutrients for the growth of the organisms. Polysorbate 80 provides fatty acids for the metabolism of the organisms and neutralizes phenolic disinfectants, hexachlorophene and formalin (1).

**Quality Control :****Appearance of powder**

Part A : Yellow coloured, may have slightly greenish tinge, homogeneous, free flowing powder

Part B : Colourless, clear viscous liquid

**Product Profile :**

Vegetable based (Code MV)Ⓞ	Animal based (Code M)
<b>MV1307</b> HiVeg peptone	<b>M1307</b> Peptic digest of animal tissue

<b>Recommended for</b>	: Cultivation of variety of microorganisms.
<b>Reconstitution</b>	: 25.0 g/lt of Part A+10 ml/lt of Part B
<b>Quantity on preparation (500g)</b>	: 20.0 L (Part A)
<b>pH (25°C)</b>	: 7.2 ± 0.2
<b>Supplement</b>	: None
<b>Sterilization</b>	: 121°C / 15 minutes.
<b>Storage</b>	: Dry Medium - Below 30°C, Prepared Medium 2 - 8°C.

**Gelling**

Firm, comparable with 1.5% Agar gel.

**Colour and Clarity**

Yellow coloured, clear to slightly opalescent gel forms in petriplates.

**Reaction**

Reaction of the medium (2.5% w/v part A + 1.0% v/v part B) is pH 7.2 ± 0.2 at 25°C

**Cultural Response**

Cultural characteristics observed after an incubation at 35-37°C for 18-24 hours.

Organisms (ATCC)	Inoculum (CFU)	Growth	Recovery
<i>Bacillus subtilis</i> (6633)	10 <sup>2</sup> -10 <sup>3</sup>	luxuriant	>70%
<i>Candida albicans</i> (10231)	10 <sup>2</sup> -10 <sup>3</sup>	luxuriant	>70%
<i>Escherichia coli</i> (25922)	10 <sup>2</sup> -10 <sup>3</sup>	luxuriant	>70%
<i>Staphylococcus aureus</i> (25923)	10 <sup>2</sup> -10 <sup>3</sup>	luxuriant	>70%

**References :**

1. Favero (chm.) 1967, Microbiological sampling of surfaces, Biological Contamination Control Committee, American Asso. For Contamination Control.