

## L.D. Esculin HiVeg™ Agar

MV743

L.D. Esculin HiVeg Agar is used for identification of anaerobic bacteria especially *Bacteroides* species on the basis of esculin hydrolysis.

**Composition \*\* :**

Ingredients	Grams/Litre
HiVeg hydrolysate	5.00
Yeast extract	5.00
Sodium chloride	2.50
L-Tryptophan	0.20
Vitamin K1	0.01
L-Cystine	0.40
Ferric pyrophosphate	0.01
Esculin	1.00
Ferric citrate	0.50
Agar	20.0

Final pH (at 25°C ) 7.4 ± 0.2

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

**Directions :**

Suspend 34.62 grams in 1000 ml distilled water. Heat to boiling to dissolve the medium completely. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes.

**Principle and Interpretation :**

L.D. Esculin HiVeg Agar is prepared by using HiVeg hydrolysate and Ferric pyrophosphate instead of Casein enzymic hydrolysate and Hemin respectively making the medium free from BSE/TSE risks. L.D. Esculin HiVeg Agar is the modification of L.D. (Lombard-Dowell) Esculin Agar developed by Lombard-Dowell (1) for identification of anaerobes especially *Bacteroides* species on the basis of esculin hydrolysis test.

HiVeg hydrolysate and yeast extract provide the nitrogenous nutrients to the organisms. L-Tryptophan and L-Cysteine serve as the amino acid source. Esculin is hydrolyzed by the organisms to form esculetin and dextrose. Esculetin reacts with the iron salt of ferric citrate to produce a dark brown to black complex. Also L-Cysteine is the sulphur containing amino acid and hence H<sub>2</sub>S production which in combination with ferric citrate gives black colouration to the colonies. Vitamin K1 and ferric pyrophosphate are the additional growth factors. Black colour of H<sub>2</sub>S positive colonies is rapidly lost after exposure to air, hence observe the plates in anaerobic glove box or immediately upon air exposure (2).

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

Yellow coloured, may have slightly greenish tinge, homogeneous, free flowing powder.

**Gelling**

Firm, comparable with 2.0% Agar gel.

**Colour and Clarity**

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

**Reaction**

Reaction of 3.46% w/v aqueous solution is pH 7.4 ± 0.2 at 25°C.

**Product Profile :**

Vegetable based (Code MV)Ⓢ	Animal based (Code M)
<b>MV743</b> HiVeg hydrolysate Ferric pyrophosphate	<b>M743</b> Casein enzymic hydrolysate Hemin

**Recommended for** : Identification of anaerobic bacteria especially *Bacteroides* species on the basis of esculin hydrolysis.

**Reconstitution** : 34.62 g/l

**Quantity on preparation (500g)** : 14.44 L

**(100g)** : 2.88 L

**pH (25°C)** : 7.4 ± 0.2

**Supplement** : None

**Sterilization** : 121°C / 15 minutes.

**Storage** : Dry Medium - Below 30°C, Prepared Medium 2 - 8°C.

**Cultural Response**

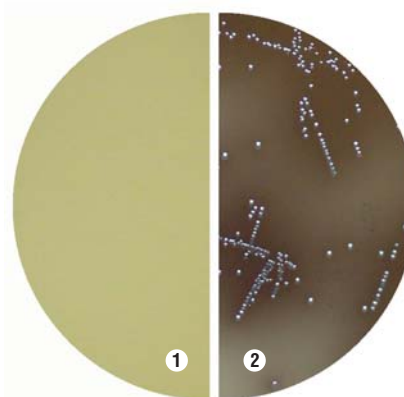
Cultural characteristics observed after an incubation at 35-37°C for 24-48 hours, under anaerobic condition.

Organisms (ATCC)	Inoculum (CFU)	Growth	Esculin hydrolysis	H <sub>2</sub> S production	Catalase
<i>Bacteroides asaccharolyticus</i>	10 <sup>2</sup> -10 <sup>3</sup>	luxuriant	-	-	-
<i>Bacteroides fragilis</i> (25285)	10 <sup>2</sup> -10 <sup>3</sup>	luxuriant	+	-	+
<i>Fusobacterium mortiferum</i>	10 <sup>2</sup> -10 <sup>3</sup>	luxuriant	+	+	-

Key : Esculin hydrolysis + = brown-black precipitate around colonies.  
H<sub>2</sub>S production + = blackening of colonies.

**References :**

- Dowell V., and Lombard G., 1977, U.S. DHEW, Centre for Disease Control (CDC), Atlanta.
- MacFaddin J.F., 1985 (ed), Media For Isolation-Cultivation-Identification-Maintenance of Medical Bacteria, Vol.1<sup>st</sup>, Williams and Wilkins, Baltimore.



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- Control
- Bacteroides fragilis*