

## Gluconate Test HiVeg™ Medium

MV483

Gluconate Test HiVeg Medium is used to check the ability of bacteria to oxidize gluconates to alpha keto-gluconate.

**Composition\*\* :**

Ingredients	Grams/Litre
HiVeg peptone	1.50
Yeast extract	1.00
Dipotassium hydrogen phosphate	1.00
Potassium gluconate	40.00

Final pH (at 25°C) 7.0 ± 0.2

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

**Directions :**

Suspend 43.5 grams in 1000 ml distilled water. Heat if necessary to dissolve the medium completely. Distribute in 10 ml quantities in screw capped bottles and sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes.

**For Gluconate test :**

After cultivating test organism for 48 hours (to be studied for detection of gluconate oxidizing ability) transfer 1 ml aliquots in clean sterile test tube. Add 1 ml of Benedict's qualitative reagent (R003). Mix well and place in boiling water bath (100°C) for 10 minutes.

**Principle and Interpretation :**

Gluconate Test HiVeg Medium is a modification of Gluconate Test Medium prepared by replacing Peptic digest of animal tissue with HiVeg peptone. Therefore this medium is free of TSE/BSE risk. Besides HiVeg peptone, yeast extract in the medium also serves as supplier of nitrogen, vitamins and other essential growth nutrients. Dipotassium hydrogen phosphate helps in buffering the medium. Potassium salt of gluconate in media serves as a readily available sole carbon source for the organism to be tested for gluconate metabolism.

This medium is used to check the ability of an organism to oxidize gluconates to alpha keto-gluconate which subsequently accumulates in the medium(1). *Pseudomonas aeruginosa* is known to accumulate at least 50% of ketogluconate after 48 hrs of incubation (2). The basis of the Gluconate test is the change from gluconate, a non-reducing compound to alpha keto-gluconate which is a reducing compound (1, 3). The alpha keto-gluconate formed when tested with a suitable reagent like Benedict's reagent, reduces copper sulphate (blue colour) to an insoluble cuprous oxide which is precipitated out. A yellow to orange to orange red precipitate is formed. The colour depends on the amount of reducing substance accumulated. Greater the amount of alpha keto-gluconate in the medium more orange to orange – red colour develops. However colours ranging from slight green to deep orange indicates oxidation. If the medium remains blue or bluish

**Product Profile :**

Vegetable based (Code MV)®	Animal based (Code M)
<b>MV483</b> HiVeg peptone	<b>M483</b> Peptic digest of animal tissue

**Recommended for** : To check the ability of bacteria to oxidize gluconates to alpha keto-gluconate.

**Reconstitution** : 43.5 g/l

**Quantity on preparation (500g)**: 11.49 L

**pH (25°C)** : 7.0 ± 0.2

**Supplement** : None

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

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

green after addition of Benedict's solution, it is considered as negative reaction which means no reducing substance is present or formed and potassium gluconate from the media is not metabolized.

**Quality Control :****Appearance of Powder**

Cream to yellow coloured, may have slightly greenish tinge, homogeneous, free flowing powder.

**Colour and Clarity**

Light straw coloured, clear solution without any precipitate.

**Reaction**

Reaction of 4.35% w/v aqueous solution is pH 7.0 ± 0.2 at 25°C.

**Cultural Response**

Cultural characteristics observed after an incubation at 35-37°C for 18-48 hours.

Organisms (ATCC)	Inoculum	Growth	Gluconate Test
<i>Citrobacter freundii</i> (8090)	10 <sup>2</sup> -10 <sup>3</sup>	poor-good	-
<i>Escherichia coli</i> (25922)	10 <sup>2</sup> -10 <sup>3</sup>	poor-good	-
<i>Klebsiella pneumoniae</i> (23357)	10 <sup>2</sup> -10 <sup>3</sup>	luxuriant	+
<i>Pseudomonas aeruginosa</i> (10145)	10 <sup>2</sup> -10 <sup>3</sup>	luxuriant	+

Key : + = positive reaction, slight green to deep orange precipitate

- = negative reaction, no change in colour or blue

**References :**

- Collee, J.G.; Marmin, B.P., Fraser, A.G and Simmons A (eds) Mackie and McCartney, Practical Medical Microbiology, (1996) 14<sup>th</sup> ed., Churchill Livingstone, New York.
- Haynes, W.C. 1951, J. Gen. Microbiology; 5(5):939.
- MacFaddin, J.F. (2000) (ed.) Biochemical Tests for identification of Medical Bacteria, 3<sup>rd</sup> edition, Lippincott Williams and Wilkins, New York.