

Technical Data

HiEncapTM YT Broth (HiEncapTM 2X YT Broth)

EC1251CCL

HiEncapTM YT Broth (HiEncapTM 2X YT Broth) is used for the cultivation of recombinant strains of *Escherichia coli*.

Composition**	
Ingredients	Gms / Litre
Casein enzymic hydrolysate	16.000
Yeast extract	10.000
Sodium chloride	5.000
Final pH (at 25°C)	7.0±0.2

**Formula adjusted, standardized to suit performance parameters

Directions

Each capsule contains 7.75 grams of medium. Suspend 1 capsule in 250 ml (4 capsules in 1000 ml)distilled or purified water. Heat to boiling to dissolve the medium completely. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes.

Principle And Interpretation

YT Broth is recommended for use in the cultivation of recombinant strains of *Escherichia coli* (1, 2, 4). It is also used in culture of *E. coli* strains for propagation of M 13 bacteriophages (2,3,4).

These media contain casein enzymic hydrolysate and yeast extract, which supply nitrogenous compounds, vitamin B complex and other essential nutrients and co-factors necessary for the luxuriant growth of recombinant *E. coli* and allows the bacteria to recover from the stress of transformation and grow well. Sodium chloride helps in maintaining isotonic conditions in the medium.

Quality Control

Appearance

Gelatin capsule containing light yellow to beige coloured granular media

Quantity

Each capsule contains 7.75 grams of medium sufficient for 250ml media.

Color and Clarity of Prepared Medium

Light amber coloured clear solution in tubes.

Reaction

Reaction of 3.1% w/v aqueous solution at 25°C. pH : 7.0±0.2

pН

6.80-7.20

Cultural Response

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

Cultural Response

Organism	Inoculum (CFU)	Growth
Cultural Response		
Escherichia coli ATCC	50-100	good-luxuriant
23724		
Escherichia coli ATCC	50-100	good-luxuriant
53868		

Storage and Shelf Life

Store below 30°C in tightly closed container and the prepared medium at 2-8°C. Use before expiry date on the label.

Reference

1. Miller H., 1987, Meth. Enzymol; 152, 145.

2. Ausubel F. M., Brent R., Kingston R. E., Moore B. D., Seidman J. G., Smith J. A. and Strohl K., 1994, Current Protocols in Molecular Biology, Vol. I, Current Protocols, New York, N.Y.

3. Davis L. G., Dibner M. D., Battey J. F., 1986, Basic Methods in Molecular Biology, Elsevier, New York, N.Y.

4. Sambrook J., Fritsch E. E. and Maniatis T., 1989, Molecular Cloning: A Laboratory Manual, 2nd Ed., Cold Spring Harbor Laboratory, Cold Spring Harbor, N.Y.

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HiMedia Laboratories Pvt. Ltd. A-516,Swastik Disha Business Park,Via Vadhani Ind. Est., LBS Marg, Mumbai-400086, India. Customer care No.: 022-6147 1919 Email: techhelp@himedialabs.com Website: www.himedialabs.com