



# Technical Data

## Antibiotic HiVeg™ Assay Medium No. 11 (Neomycin, Erythromycin HiVeg™ Assay Agar) (Erythromycin Seed HiVeg™ Agar)

MV004

### Intended use

Recommended for microbiological assay of antibiotics.

### Composition\*\*

Ingredients	Gms / Litre
HiVeg™ peptone	6.000
HiVeg™ hydrolysate	4.000
Yeast extract	3.000
HiVeg™ extract	1.500
Dextrose	1.000
Agar	15.000
Final pH ( at 25°C)	8.3±0.2

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

### Directions

Suspend 30.5 grams in 1000 ml purified/distilled water. Heat to boiling to dissolve the medium completely. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Cool to 45-50°C. Mix well and pour into Petri plates or dispense as desired.

*Advice: Erythromycin, Chlortetracycline, Dihydrostreptomycin sulphate, Erythromycin estolate, Framycetin, Gentamicin, Gentamicin sulphate, Kanamycin sulphate, Kanamycin monosulphate, Kanamycin acid sulphate, Netilmicin sulphate, Netilmicin, Neomycin, Paromomycin, Sisomycin, Spiramycin, Streptomycin sulphate*

Other Tests :

Cup plate method is carried out using *B. pumilis* / kanamycin and *M. flavus* / erythromycin

1) Dilution : 16 mg Kanamycin in 10 ml distilled water

Stock : 1:10 dilution of above solution

concentration	stock (ml)	Distilled water (ml)	zone of inhibition
5	0.25	4.75	15 mm
20	1.00	4.00	20 mm
100	5.00	-	25 mm

2) Dilution : 9 mg Erythromycin in 10 ml distilled water

Stock : 1:10 dilution of above solution

Concentration	stock (ml)	Distilled water (ml)	zone of inhibition
5	0.25	4.75	22 mm
10	0.50	4.50	32 mm
100	5.00	-	41 mm

### Principle And Interpretation

Antibiotic Assay media are used in the performance of antibiotic assays. Grove and Randall have elucidated those antibiotic assays and media in their comprehensive treatise on antibiotic assays (1). Schmidt and Moyer have reported the use of antibiotic assay medium for the liquid formulation used in the performance of antibiotic assay (4). These media are recommended by USP (6) and FDA (5). Antibiotic HiVeg™ Assay Medium No. 11 is prepared by completely replacing animal based peptone with vegetable peptones to avoid BSE/TSE risks associate with animal peptones

Nutrients and growth factors are supplied by the ingredients like HiVeg™ peptone, HiVeg™ hydrolysate, yeast extract and HiVeg™ extract. Dextrose provides the carbon and energy source. Agar provides excellent medium for antibiotic diffusion

and gives well-defined zones of inhibition. Higher pH provides the optimal conditions for activity of antibiotic and also supports the growth of the test organisms. Freshly prepared plates should be used for antibiotic assays. Test organisms are inoculated in sterile seed agar pre-cooled to 40-45°C and spread evenly over the surface of solidified base agar. All conditions in the microbiological assay must be controlled carefully.

## Type of specimen

Pharmaceutical preparations

## Specimen Collection and Handling

For pharmaceutical samples follow appropriate techniques for handling specimens as per established guidelines (6). After use, contaminated materials must be sterilized by autoclaving before discarding.

## Warning and Precautions :

Read the label before opening the container. Wear protective gloves/protective clothing/eye protection/ face protection. Follow good microbiological lab practices while handling specimens and culture. Standard precautions as per established guidelines should be followed while handling specimens. Safety guidelines may be referred in individual safety data sheets

## Limitations :

1. Freshly prepared plates must be used or it may result in erroneous results.

## Performance and Evaluation

Performance of the medium is expected when used as per the direction on the label within the expiry period when stored at recommended temperature.

## Quality Control

### Appearance

Cream to yellow homogeneous free flowing powder

### Gelling

Firm, comparable with 1.5% Agar gel

### Colour and Clarity of prepared medium

Light yellow coloured, clear to slightly opalescent gel forms in Petri plates

### Reaction

Reaction of 3.05% w/v aqueous solution at 25°C. pH : 8.3±0.2

### pH

8.10-8.50

### Cultural Response

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

Organism	Inoculum (CFU)	Growth	Recovery	Antibiotics assayed
<i>Micrococcus luteus</i> ATCC 9341	50-100	luxuriant	≥70%	Erythromycin While assaying Tylosin, Tylosin, tartarate, Vancomycin hydrochloride, adjust the pH to 8.0±0.2
<i>Staphylococcus aureus</i> ATCC 6538P (00195*)	50-100	luxuriant	≥70%	Kanamycin monosulphate, Kanamycin acid sulphate, Netilmicin sulphate
<i>Staphylococcus epidermidis</i> ATCC 12228 (00036*)	50-100	luxuriant	≥70%	Gentamicin, Neomycin, Netilmicin, Paromomycin, Sisomycin
<i>Bacillus pumilis</i> ATCC 14884	50-100	luxuriant	≥70%	Chlortetracycline, Framycetin, Kanamycin sulphate
<i>Bacillus subtilis</i> subsp. <i>spizizenii</i> ATCC 6633 (00003*)	50-100	luxuriant	≥70%	Dihydrostreptomycin sulphate, Erythromycin estolate, Kanamycin monosulphate, Kanamycin acid

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<i>Bacillus subtilis</i> NCTC 8236	50-100	luxuriant	>=70%	Dihydrostreptomycin sulphate, Streptomycin sulphate
<i>Bacillus subtilis</i> NCTC 8241	50-100	luxuriant	>=70%	Erythromycin estolate, Gentamicin sulphate

\*- Corresponding WDCM Numbers

### Storage and Shelf Life

Store between 10-30°C in a tightly closed container and use freshly prepared medium . Use before expiry date on the label. On opening, product should be properly stored dry, after tightly capping the bottle in order to prevent lump formation due to the hygroscopic nature of the product. Improper storage of the product may lead to lump formation. Store in dry ventilated area protected from extremes of temperature and sources of ignition Seal the container tightly after use.

Product performance is best if used within stated expiry period.

### Disposal

User must ensure safe disposal by autoclaving and/or incineration of used or unusable preparations of this product. Follow established laboratory procedures in disposing of infectious materials and material that comes into contact with sample must be decontaminated and disposed of in accordance with current laboratory techniques (2,3).

### Reference

1. Grove and Randall, 1955, Assay Methods of Antibiotics Medical Encyclopedia, Inc, New York.
2. Isenberg, H.D. Clinical Microbiology Procedures Handbook 2<sup>nd</sup> Edition.
3. Jorgensen, J.H., Pfaller, M.A., Carroll, K.C., Funke, G., Landry, M.L., Richter, S.S and Warnock., D.W. (2015) Manual of Clinical Microbiology, 11th Edition. Vol. 1.
4. Schmidt and Moyer, 1944; J. Bact, 47:199.
5. Tests and Methods of Assay of Antibiotics and Antibiotic containing Drugs, FDA, CFR, 1983. Title 21, part 436, Subpart D, Washington, D.C. U.S Government printing office, paragraphs 436, 100-436, 106 pg 242-259 (April 1).
6. United States Pharmacopoeia 2018, US Pharmacopoeial Convention Inc, Rockville, MD

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### Disclaimer :

User must ensure suitability of the product(s) in their application prior to use. Products conform solely to the information contained in this and other related HiMedia™ publications. The information contained in this publication is based on our research and development work and is to the best of our knowledge true and accurate. HiMedia™ Laboratories Pvt Ltd reserves the right to make changes to specifications and information related to the products at any time. Products are not intended for human or animal or therapeutic use but for laboratory, diagnostic, research or further manufacturing use only, unless otherwise specified. Statements contained herein should not be considered as a warranty of any kind, expressed or implied, and no liability is accepted for infringement of any patents.