



Product Specification

cdhfinechemical.com

Technical Information

Eriksson (ER) Medium With Vitamins Without Sucrose and Agar

Product Code: PT1033

Application: Eriksson (ER) Medium has been developed by Eriksson in 1965 for the in vitro cell suspension culture of *Haplopappus* gracilis, family Asteraceae/Compositae. The formulation is a nutrient blend of inorganic salts, vitamins and amino acid.

Eriksson (ER) Medium provides all the essential macroelements and microelements. Potassium nitrate and ammonium nitrate serve as sources of nitrogen that initiates morphogenesis. This mixture of cation and anion also helps to maintain pH of the medium. Potassium dihydrogen phosphate provides phosphate and stimulates cell growth. Microelements like Boron, Manganese, Molybdenum, Copper, lron and Zinc play vital role in plant metabolism and improve growth of cells. Thiamine, pyridoxine, nicotinic acid act as enzymatic cofactors in universal pathways including glycolysis and TCA cycle along with primary and secondary metabolism in the plants. Glycine serves as a source of amino acid.

The product is plant tissue culture tested but it is the sole responsibility of the user to ensure the suitability of the medium for individual species.

Composition**		
Ingredients	mg/Litre	
MACROELEMENTS		1
Ammonium nitrate	1200.000	
Calcium chloride	332.200	
Magnesium sulphate	180.690	
Potassium nitrate	1900.000	
Potassium phosphate monobasic	340.000	
MICROELEMENTS		
Boric acid	0.630	
Copper sulphate pentahydrate	0.003	
Cobalt chloride hexahydrate	0.003	
EDTA disodium salt dehydrate	37.300	
Ferrous sulphate heptahydrate	27.800	
Manganese sulphate monohydrate	1.689	
Molybdic acid (sodium salt)	0.021	
Potassium Iodide	0.830	
Zinc sulphate heptahydrate	9.145	
VITAMINS		
Nicotinic acid (free acid)	0.500	
Pyridoxine HCl	0.500	
Thiamine hydrochloride	0.500	
AMINO ACID		
Glycine	2.000	
Total	4.0 gms/litre	

Material required but not provided

- Autoclaved distilled water
- Sucrose (PCT1607)
- Plant growth regulators
- Gelling agents like Agar (PCT1901) or CleriGel (PCT1903)
- 1N NaOH/HCl





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Quality Control

Appearance

White to off-white, homogenous, free flowing powder

Solubility

4.0 gms/litre soluble in distilled water

Colour and Clarity

Colourless to light yellow, clear solution

pH at 25ºC

3.30 – 4.30

Plant Tissue Culture Test

The growth promoting properties of medium is assessed by providing plant cultures with relative humidity of about 60%±2%, temperature 22°C±2°C and photoperiod of about 16:8. The plant species showed actively growing callus and shoots with no structural, necrotic and toxic deformity.

Directions

- Reconstitute medium by adding required quantity of powder in two-third of total volume with constant, gentle stirring till the medium gets completely dissolved.
- Add heat stable supplements prior to autoclaving.
- Make up the final volume with distilled water.
- Adjust the pH of the medium to 5.75 ± 0.5 using 1N NaOH/HCl.
- Add gelling agent and heat the medium to boiling till complete dissolution of gelling agent.
- Sterilize the medium by autoclaving at 15 lbs and 121°Cfor 15 min.
- Cool the autoclaved medium to about 45°C before adding heat labile supplements.
- Aseptically dispense the desired amount of medium under a laminar airflow unit in sterile culture vessels

Storage and Shelf Life

- The plant tissue culture medium powder is extremely hygroscopic and must be stored at 2-8°C in air tight containers.
- Preferably, entire content of each package should be used immediately after opening.
- Use before the expiry date.

Disclaimer

- User must ensure suitability of the product(s) in their application prior to use.
- The product conforms solely to the technical information provided in this booklet and to the best of knowledge research and development work carried at **CDH** is true and accurate.
- Central Drug House 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 diagnostic or therapeutic use but for laboratory, research or further manufacturing of diagnostic reagents extra.
- 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.
- Do not use the products if it fails to meet specifications for identity and performance parameters.

Precautions

- Ensure appropriate pH of the medium before addition of gelling agent as acidic pH will lead to decreased gelation resulting in semi solid flowing gel while alkaline pH will lead to formation of hardened gel.
- Use of Distilled water/Tissue culture grade water is recommended for media preparation as tap water or lower grade water may lead to salt precipitation and improper gelation.
- Avoid preparation of concentrated solutions, as it will lead to precipitation of salts.