



Technical Information

Fern Micropropagation Medium

w/ Vitamins, NAA and Kinetin; w/o Sucrose and Agar

Product Code: PT1087

rroduct code: F11087		
Composition**		
Ingredients	mg/Litre	
Potassium nitrate	1900.00	
Ammonium nitrate	1650.00	
Calcium chloride.2H₂O	440.00	
Magnesium sulphate	180.69	
Potassium phosphate monobasic	170.00	
Sodium phosphate monobasic	221.71	
Manganese sulphate.H₂O	16.90	
Boric acid	6.20	
Potassium iodide	0.83	
Molybdic acid (sodium salt).2H₂O	0.25	
Zinc sulphate.7H₂O	8.60	
Copper sulphate.5H₂O	0.025	
Cobalt chloride.6H₂O	0.025	
Ferrous sulphate.7H₂O	27.80	
EDTA disodium salt.2H₂O	37.30	
myo - Inositol	100.00	
Thiamine hydrochloride	0.40	
α- Naphthalene acetic acid	0.10	
Kinetin	2.00	
Total	4.76 gm/litre	

Principle And Interpretation

Fern Micropropagation medium has been specially formulated for the in vitro micropropagation of ferns. Ammonium nitrate and potassium nitrate serves as sources of nitrate. NAA and kinetin serves as plant growth regulators. Medium does not contain sucrose and agar; hence these components have to be added to the medium before use.





Product Specification

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Directions

Suspend 4.76 grams of dehydrated medium# in 600ml of distilled water and rinse media vial with small quantity of distilled water to remove traces of powder. Apply constant gentle stirring to the solution till the powder dissolves completely. Add desired heat stable supplements prior to autoclaving. Adjust the medium to the desired pH using 1N HCI/NaOH.

Make up the final volume to 1000ml with distilled water. Sterilize the medium by autoclaving at 15 lbs or 121ºC for 15 minutes. Cool the autoclaved medium to 45ºC before adding the filter sterilized heat labile supplements. Dispense the desired amount of medium aseptically in sterile culture vessels.

Weight after vacuum drying to remove all water

Quality Control

Appearance: White to off-white, homogeneous, free flowing powder.

Solubility : 4.63 gm/litre soluble in distilled water.

Colour and Clarity : Colourless to light yellow, clear solution.

pH at 25°C : 3.9 ±0.5 of 0.463% w/v dehydrated powder.

Cultural Response:

Cultural condition:

 $\begin{array}{lll} \cdot \mbox{ Incubation period} & : 5 \mbox{ weeks} \\ \cdot \mbox{ Relative humidity} & : 60\% \pm 2\% \\ \cdot \mbox{ Temperature} & : 22^{\circ}\mbox{C} \pm 2^{\circ}\mbox{C} \\ \cdot \mbox{ Photoperiod (D:N) in hours} & : 16:8 \\ \end{array}$

Cell Line	Types Of Culture	Results	
Musa species	Shoot culture	No structural deformity observed	
		No necrotic tissues,	
		Actively growing shoots,	
		No toxicity to shoots	
Daucus species	Callus culture	No necrotic tissues,	
		Actively growing callus,	
		No toxicity to callus	

[The medium is prepared as per direction. The growth promoting activity of this plant tissue culture medium is evaluated using two plant species viz. Artemisia species and Daucus species through three passages. Plant growth hormones (e.g. 2,4-D, NAA, Kinetin and 6-BAP) are added in suitable combinations and concentrations.]

Storage and Shelf Life

Dehydrated macroelements powder is extremely hygroscopic and should be protected from atmospheric moisture. If possible, the entire content of each bottle should be used immediately after opening or else the unused portion should be stored in a desiccator and refrigerated at 2-8°C. Use before the expiry date.





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Further Reading

1. Murashige T. & Miller L.R., In Vitro, (1976), 12, 797 - 813

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.
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