



# **Technical Information**

Nitsch Medium w/ Vitamins and Sucrose; w/o CaCl₂ and Agar

Product Code: PT1012

## Composition\*\*

Ingredients	mg/Litre
Potassium nitrate	950.00
Ammonium nitrate	720.00
Magnesium sulphate anhydrous	90.34
Potassium phosphate monobasic	68.00
Manganese sulphate.H₂O	18.94
Boric acid	10.00
Molybdic acid (sodium salt).2H₂O	0.25
Zinc sulphate.7H₂O	10.00
Copper sulphate.5H₂O	0.025
Ferrous sulphate.7H <sub>2</sub> O	27.85
EDTA disodium salt.2H₂O	37.25
myo - Inositol	100.00
Thiamine hydrochloride	0.50
Pyridoxine hydrochloride	0.50
Nicotinic acid (Free acid)	5.00
Folic acid	0.50
Biotin	0.05
Glycine (Free base)	2.00
Sucrose	20000.00
TOTAL	22.04 gm/litre

## **Principle And Interpretation**

Nitsch medium has been specially formulated for the in vitro culture of plant cell, tissue and organ culture. Ammonium nitrate and potassium nitrate serves as the sources of nitrate. Glycine serves as the source of amino acid. Sucrose serves as the carbohydrate source. Medium is devoid of calcium chloride and agar; hence these components have to be added to the medium prior to use.





# **Product Specification**

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### Directions

Suspend 22.04 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 : 22.04 gm/litre soluble in distilled water.

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

pH at 25°C : 3.8 ±0.5 of 2.204% w/v dehydrated medium.

#### **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	
<i>Musa</i> 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. Musa 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 plant tissue culture spowder 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. Nitsch J.P. & Nitsch C., Science, (1969), 163, 85 - 87

### 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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