

Schenk & Hildebrandt Medium w/ Vitamins, Sucrose and Agar

PT138

Composition :

| Ingredients | milligrams/litre |
|---|------------------|
| Potassium nitrate | 2500.00 |
| Ammonium phosphate monobasic | 300.00 |
| Calcium chloride.2H ₂ O | 200.00 |
| Magnesium sulphate | 195.34 |
| Manganese sulphate.H ₂ O | 10.00 |
| Boric acid | 5.00 |
| Potassium iodide | 1.00 |
| Molybdic acid (sodium salt).2H ₂ O | 0.10 |
| Zinc sulphate.7H ₂ O | 1.00 |
| Copper sulphate.5H ₂ O | 0.20 |
| Cobalt chloride.6H ₂ O | 0.10 |
| Ferrous sulphate.7H ₂ O | 15.00 |
| EDTA disodium salt.2H ₂ O | 20.00 |
| myo - Inositol | 1000.00 |
| Thiamine hydrochloride | 5.00 |
| Pyridoxine hydrochloride | 0.50 |
| Nicotinic acid (Free acid) | 5.00 |
| Sucrose | 10000.00 |
| Agar | 8000.00 |
| TOTAL gm/litre | 22.26 |

Directions :

Suspend 22.20 grams of dehydrated medium[#] in 600ml of distilled water and rinse media vial with small quantity of distilled water to remove traces of powder. Add desired heat stable supplements prior to autoclaving. Adjust the medium to the desired pH using 1N HCl/NaOH. Make up the final volume to 1000 ml with distilled water. Boil the medium to dissolve agar completely. Sterilize the medium by autoclaving at 15 lbs or 121°C for 15 minutes. Cool the autoclaved medium to 45°C 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

Principle and Interpretation :

Schenk & Hildebrandt medium has been specially formulated for plant cell, tissue and organ cultures. Potassium nitrate serves as the nitrate source. Sucrose serves as the source of carbohydrate. Agar is incorporated into the medium to provide firm base to the explants.

Quality Control :

| | |
|--------------------|---|
| Appearance | : White to off-white, homogeneous, free flowing powder. |
| Solubility | : 22.20 gm/litre soluble after boiling in distilled water. |
| Colour and Clarity | : Colourless to light yellow, hazy gel is formed on cooling. |
| pH at 25°C | : 4.1 ±0.5 (under observation) of 2.220% w/v dehydrated medium. |

Cultural Response :

Cultural condition :

| | |
|------------------------------|--------------|
| · Incubation period | : 5 weeks |
| · Relative humidity | : 60% ± 2% |
| · Temperature | : 22°C ± 2°C |
| · Photoperiod (D:N) in hours | : 16:8 |

| Cell Line | Type of Culture | Results |
|-----------------------|-----------------|---|
| <i>Musa</i> species | Shoot culture | No structural deformity observed No necrotic tissues, Actively growing shoots, No toxicity to shoots |
| <i>Daucus</i> 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 media 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.

Reference :

1. Schenk R.U. & Hildebrandt A.C., Can. J. Bot., (1972), 50, 199 - 204

Disclaimer :

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