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Research Article

In vitro Seeds Germination and Seedling Growth of Begonia Malabarica Lam. (Begoniaceae) a Source for Anthocyanin

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Abstract

Begonia malabarica Lam. is an ornamental species. It occupies a prominent place in indoor landscaping programmes. These plants are pharmacologically useful due to their anti inflammatory, anti hepatotoxic, antioxidant and antiseptic activities, which have been attributed by anthocyanin. Generally, B. malabarica is propagated through seeds. The propagation of this species from seeds is particularly challenging because of dormancy of seeds and also lose their viability within a short span of time. In addition, propagation from seeds often leads in a high variability among the progeny. In this juncture, the aim of the present study was in vitro seed germination by breaking dormancy and determination of the optimal parameters for germination to propagate this plant on a large scale. Seeds were standardized with different surface sterilization protocols and inoculated on different basal media without growth regulators. Various culture media, its strengths, the type and concentration of carbon source were investigated. Responses to germination varied with the culture conditions also. Seeds in half strength MS medium germinated quickly and expressed remarkable germination rate. Similarly, the in vitro growth performances of plantlets varied with the basal medium composition. The optimal growth performances of plants were displayed on half strength MS basal medium. Further, 3% sucrose in the culture medium was more optimal for maximum growth of plantlets. B. malabarica is a source of anthocyanin. So, future studies are planned to induce callus, cell line culture from these in vitro germinated seedlings to extract, purify and fractionate the pigments.

Keywords: Begonia malabarica; In vitro; Germination media

Introduction

Indoor landscaping plants are produced worldwide for their aesthetic importance. Creation of novel variation in terms of qualitative characters like leaf nature, color of the flower, its fragrance, longevity, shape and architectural feature are the major economic objectives for the horticulture industries. Further, production of superior quality saplings through genetic modifications requires basic information regarding in vitro seed culture protocols [1]. In some ornamental genotypes shows non flowering or low production of seeds or often associated with irregular germination. Diseases also often accumulate in the saplings resulting in infected plants and low yields. Challenges with the vegetative multiplications include high perishability, handling and transportation labour costs and inconvenient weight and bulk of the samples [2].

Further, immature embryo, difficulty to obtain samples in all seasons of the year, poor germination rate, failure in seedling emergence and growth were the other hurdles associated with traditional cultivation techniques. Seed size has a special role in crop production. Small seed size indicates a low protein synthesizing ability, which is normally attributed to less availability of substrate, energy, active enzymes and machinery for protein synthesis [3].

Mass multiplication of ornamental species is labour oriented, and for that reason, in vitro multiplication of plants is an efficient system and used by many commercial nurseries and institutes globally for rapid plant propagation, germ plasm conservation, pathogen elimination, genetic manipulations, and for phytochemical production [4].

Begonia of Begoniaceae comprises approximately 1000 species and many of these are ideal indoor ornamentals and medicinal herbals. Begonia is herbaceous species with attractive foliage and can be utilized for horticultural purposes. Many begonias respond best to warm, moist conditions. Further, they prefer well drained
soil rich in compost or organic matter and shade [5]. Generally the seeds are very small with poor germination rate and time. The B. malabarica is commonly used by the locals for curing many skin borne disorders and also as an antioxidant herbal tonic. Ramesh et al. [6] reported the claim of the usefulness of the plant in respiratory tract infections and also suggests its use in diarrhoea and skin diseases caused by pathogenic bacteria. The plants are reported for its rich anthocyanin content. In this scenario, the objectives of the present study were to design a protocol for mass multiplication of the plants via the in vitro seeds culture. This includes initiation of aseptic cultures, surface sterilization, inoculation, effect of different media, MS medium strength, effect of carbohydrate source and germination % and growth parameters.

Materials and Methods

Plant Material and Initiation of Aseptic Cultures

The fresh healthy plants of Begonia malabarica were collected from the plant grown on wild habitat from Wayanad hills and Idukki district, Kerala and the voucher specimen was deposited in the herbarium of University College, Trivandrum (UCB 1207). Generally, due to the succulent and pubescent nature of Begonia, micropropagation seems to be laborious due to high rate of contamination. Thus, direct in vitro seed germination is an alternative and B. malabriica produces hundreds of dormant seeds in the capsule. So an attempt was made during the initial phase of the study using excised leaves, petioles, rhizomes and capsules (with seeds) as explants for in vitro micropropagation. These explants were thoroughly washed in running tap water for 1 h and treated with 5% teepol (v/v) for 20 min with continuous agitation. Subsequently, the teepol content from the explants were removed by repeated washing with double distilled sterile water.

Surface Sterilization and Inoculation

Various surface sterilization protocols were adapted using sterilants such as mercuric chloride 0.01%-0.1% for 3-8 min, sodium hypochlorite - 5%-15% for 5-10 min or calcium chloride 10-15% for 10 - 20 min for all the excised explants. Meanwhile, in the case of fully matured seeds that are about to burst out from the capsule were directly treated with mercuric chloride (0.1%) for 3-5 min. Invariably, Tween 20, the poly ethoxylated sorbitan ester, an emulsifying, wetting, dispersant, hydrophilic and solubilizing agent was also added with all the treatments.

The disinfected explants were rinsed with sterile distilled water for several times and were dissected into small pieces of explants like excised leaves, petioles, rhizomes and capsules (with seeds). Subsequently, the leaves and rhizome measuring approximately 1cm2 and petiole about 1cm long were cultured on different culture media such as MS medium [7], B5 [8], SH [9], MC [10], Chu N6 [11] and Whites medium with various combinations of hormones. The media were supplemented with 1-3% sucrose and gelled by 0.3%-0.7% agar. PH was adjusted to 5.8 ± 0.02 prior to autoclaving at a pressure of 1210 C for 20 min. The cultures were incubated at 25 ± 20C with 12/12. 8/16, 16/8 hours light/dark photoperiod through white fluorescent tubes (1000 lux).

Seed Sources and Surface Sterilization

Subsequently, the mature capsules of B. malabarica were collected from the natural habitat. The capsules were stored in air tight containers at room temperature. Under a laminar airflow cabinet, the capsules or dissected seeds from the capsules were initially surface-sterilized with 70% (v/v) ethanol (60 sec.). Subsequently, the capsules were treated with different sterilants such as 2.5, 5, 7.5, 10% (w/v) calcium hypochlorite solution (5-30 min), 0.01-0.1% mercuric chloride for 3-8 min, sodium hypochlorite - 5%-15% for 5-10 min or calcium chloride 10%-15% for 10 - 20 min and finally rinsed thoroughly with three times sterile distilled water. The treated capsules or seeds were soaked overnight in 100 mL beaker containing 50 mL of sterile distilled water (SDW). After soaking overnight, the water was discarded, the seeds were rinsed 3-4 times with SDW, and the seeds were used for in vitro germination trials.

Germination % and Growth Parameters

Germination was calculated as the appearance of 2 mm radial length and was considered as physiological state of germination. Germination was monitored regularly for until no further germination was recorded and the mean germination time (mgt) was calculated using the formula given below [12]

\[ \text{mgt} = \frac{\Sigma (nd)}{\Sigma n} \]

where \( n \) is the number of seeds newly germinated at each day at 25°C; \( d \) represents days from the beginning of the germination test; \( \Sigma n \) is the total number of seeds germinated at the termination of the experiment [13]. Four weeks after sowing, seedling height, epicotyl and primary root length, and the number of leaves, branches, and secondary roots were recorded. The dry weights of aerial and root systems were obtained by drying them in an oven at 65°C for 72 h until a constant dry weight.

Effect of Different Basal Media

The capsules or seeds were cultured on six basal media such as MS [7], B5 [8], SH [9], MC [10] and Chu N6 [11] (CHU) and Whites medium. Different types of sugars were added to these basal media without growth regulators. The pH of the culture media was adjusted to 5.8 before adding 0.6% (w/v) agar. Culture medium without any fortification of nutrients was considered as control.

Effect of MS Medium Strength

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To optimize germination of surface sterilized seeds and the in vitro development of plantlets, the seeds were inoculated on three strengths of MS medium, namely, full strength MS salts, half strength MS salts (½ MS), and quarter strength MS salts (¼ MS).

**Effect of Carbohydrate Sources**

The best culture medium from the previous experiment was selected for testing different sources of carbohydrate. Thus, in this germination medium contain different doses of sugars such as 1-3% of sucrose or glucose or fructose have been added to analyze their effectiveness in promoting the germination and subsequent growth of the seedling.

**Statistical Analysis of Data**

The whole experiment was completed out in a completely randomized design. 50 seeds were used for each of the following experiments. Germination days, % of germination, plant height, number of leaves, root length and biomass variables were analyzed against the media such as MS, B5, SH, MC, Chu N6, Whites and control. Subsequently, the plant height, number of leaves, root length and biomass variables were analyzed with different strength MS media such as full strength, half strength, and quarter strength MS salts. Half strength MS medium was supplemented with different carbon sources like glucose, fructose, sucrose vs plant height, number of leaves, root length and biomass variables were evaluated. Finally, the half strength MS medium against different concentrations of sucrose (1, 2 and 3%) were also analyzed in terms of plant height, number of leaves, root length and biomass.

Each experiment was repeated for six times. The data were subjected to normality test using descriptive data analysis. It was found that skewness for the above parameters were nearly zero and kurtosis was almost 3. Test also for the chi-square in terms for goodness of fit was analyzed. p - value > 0.05 was considered as probability level of significance. All these were carried using statistical software program Statistical version 7.

**Results and Discussion**

**Initiation of Aseptic Cultures, Surface Sterilization and Inoculation**

Standardization to develop aseptic in vitro cultures as per the pre-treatment protocols with mercuric chloride or calcium chloride or sodium hypochlorite with the selected explants resulted in high rate of contamination. This may be due to the abundance of epidermal trichomes and succulent nature. Thus, a protocol was designed to generate aseptic cultures of *B. malabarica* through in vitro seed germination.

**Seed Sources and Surface Sterilization**

Generally, in vitro responses in plants varied greatly based on the medium, explants, hormonal combinations and culture maintenance conditions. HgCl₂ sterulant yielded the optimal results with seeds when compared to other sterilants. Direct pre-treatment of capsule with 0.1% HgCl₂ for 5 min was effective when compared to direct treatment of seeds (i.e., it leads to dormancy, retarded growth with chlorosis with the germinated seedlings) (Figure 1).

**Figure 1:** Directly pretreated seeds with 0.1% mercuric chloride showing nitrification and seed dormancy.

While the seeds from the pre-treated excised capsule with HgCl₂ 0.1% for 5 min inoculated on various medium showed optimal seed germination with negligible rate of contamination. Sen et al. [14] evaluated the effect of sterilizing substances like ethanol, mercuric chloride, flugal, nystatin, and sodium hypochlorite on germination and development of explants of Achyranthes aspera. The percentage of contamination and germination, as well as seed color, growth pattern, and shoot let development varied depending on the sterilants.

**Effect of Different Basal Media**

After incubation on the culture media, seeds became swollen quickly and germination occurred within the four weeks of culture (Figure 2a & b). The mean germination time (mgt) and the mean germination % of the seed on different culture media after a 6-week incubation period are tabulated. Highest germination % and mean germination days was shown by MS medium i.e., 90 and 20 respectively. Mean growth % was not significantly influenced by the different culture media tested.

**Fig: 2a**

**Fig: 2b**
The media used in the present analysis varied from one another in their chemical composition. The remarkable feature of the MS inorganic salts is their high level of nitrate, potassium, and ammonium in comparison to other media formulations. MS medium is highly enriched with macro- and microelements and the inorganic salts in this medium were enough to support the maximum growth of the plant. The concentration and the quality of nitrogen in MS medium may be the reality of prolific growth obtained with plant derived seed types incubated on this medium. Indeed, nitrogen is supplied to medium in inorganic form as NO3- anion or the NH4+ cation.

Kone et al. [15] reported that the ammoniated form of nitrogen was more appropriate than the nitrate form yielded the fastest growth of Bambara groundnut seedlings at 50-100 mg-L-1 NH4NO3. Further, Chen and Chang [16] noticed that an optimal concentration of organic and inorganic nitrogen components can induce the growth of explants. In addition, better plant growth from embryonic axis was observed on MS medium compared to other basal media in Juglans regia [17].

Effect of MS Medium Strengths on the Growth of Seedlings Developed from the Seeds

Among the different strengths of MS basal medium employed, a remarkable difference was recorded for growth parameters when comparing the plantlets derived from seeds (Table 2). Full and ¼ strengths of MS medium, plantlet height, root length, and the biomass were similar. Significant difference was seen between the half strength MS basal medium with others in terms of leaf number, length of root and plantlets biomass derived from seeds. But a significant reduction in plantlet height was observed on 1/4 MS (Table 3). Thus, the result suggests that a low profile of macro- and micronutrients is not effective for plantlet growth. Half strength of MS produce optimal results for Begonia seedling development suggesting that an adjustment can be carried from the full composition of MS basal medium without any significant reduction in plantlet growth (Fig. 2c). Thus, half MS was selected as the culture medium for the subsequent assessment of carbohydrate sources.

Table 1: Mean germination time and germination % of B. malabarica seeds incubation on different basal media.

<table>
<thead>
<tr>
<th>Media culture</th>
<th>Plantlets height (cm)</th>
<th>Number of leaves</th>
<th>Root length (cm)</th>
<th>Biomass (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>3.76 ± 0.08</td>
<td>2.40 ± 0.24</td>
<td>2.07 ± 0.04</td>
<td>0.12 ± 0.001</td>
</tr>
<tr>
<td>MS</td>
<td>5.6 ± 0.84</td>
<td>8.76 ± 0.52</td>
<td>3.5 ± 0.43</td>
<td>0.51 ± 0.01</td>
</tr>
<tr>
<td>B5</td>
<td>4.10 ± 1.39</td>
<td>6.14 ± 1.57</td>
<td>2.65 ± 3.44</td>
<td>0.25 ± 0.03</td>
</tr>
<tr>
<td>McCown</td>
<td>3.99 ± 0.77</td>
<td>5.25 ± 0.98</td>
<td>2.98 ± 4.59</td>
<td>0.25 ± 0.02</td>
</tr>
<tr>
<td>SH</td>
<td>3.08 ± 0.60</td>
<td>4.82 ± 0.59</td>
<td>2.68 ± 3.29</td>
<td>0.22 ± 0.004</td>
</tr>
<tr>
<td>CHU</td>
<td>4.20 ± 1.48</td>
<td>4.43 ± 0.39</td>
<td>2.81 ± 3.12</td>
<td>0.21 ± 0.02</td>
</tr>
<tr>
<td>Whites</td>
<td>5.08 ± 0.60</td>
<td>4.55 ± 0.39</td>
<td>2.81 ± 3.12</td>
<td>0.21 ± 0.02</td>
</tr>
</tbody>
</table>

Table 2: Growth performance of B. malabarica in vitro plantlets derived from different basal media after four weeks of cultivation.

<table>
<thead>
<tr>
<th>Media culture</th>
<th>Plantlets height (cm)</th>
<th>Number of leaves</th>
<th>Root length (cm)</th>
<th>Biomass (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Full MS</td>
<td>4.4 ± 0.15</td>
<td>5.38 ± 0.52</td>
<td>2.99 ± 0.54</td>
<td>0.51 ± 0.05</td>
</tr>
<tr>
<td>Half MS</td>
<td>5.6 ± 0.84</td>
<td>8.76 ± 0.52</td>
<td>3.87 ± 0.43</td>
<td>0.67 ± 0.01</td>
</tr>
<tr>
<td>¼ MS</td>
<td>2.9 ± 0.04</td>
<td>6.11 ± 0.44</td>
<td>2.16 ± 0.1</td>
<td>0.38 ± 0.03</td>
</tr>
</tbody>
</table>

Table 3: Growth performance of B. malabarica in vitro plantlets derived from different strengths of Half MS basal medium after four weeks of culture.
Effect of Carbohydrate Sources

Plants growing under in vitro culture conditions are semi autotrophic and leaves formed during in vitro growth may never attain photosynthetic competence fully [18]. Moreover, plantlets growing under in vitro conditions have limited accessibility to CO₂ inside the culture vessel [19]. Therefore, sugar was supplemented as carbon source to maintain an optimal usage of carbon source for in vitro multiplication and growth of plant cell, tissue, and organs or whole plantlets. Continuous supply of carbohydrates to plants cultured in vitro is essential because the photosynthetic activity of in vitro grown tissues is usually low. These compounds are also act as osmolytes in the culture media. Gibson [20] revealed that sugars have potential role on the physiology, growth, and differentiation of cells. Therefore, the optimal carbon source needs to be considered. Thus, different sugars such as glucose, fructose, and sucrose at 3% (w/v) were incorporated into the 1/2 MS basal medium. After six weeks of growth, the plants were analyzed and presented in Table 4.

Table 4: Growth performance of B. malabarica in vitro plantlets growth on 1/2 MS containing glucose, sucrose, and fructose.

<table>
<thead>
<tr>
<th>Media culture</th>
<th>Plantlets height (cm)</th>
<th>Number of leaves</th>
<th>Root length (cm)</th>
<th>Biomass (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Glucose</td>
<td>5.30 ± 0.21</td>
<td>4.90 ± 0.43</td>
<td>5.0 ± 0.18</td>
<td>0.03 ± 0.01</td>
</tr>
<tr>
<td>Sucrose</td>
<td>6.4 ± 0.33</td>
<td>10.6 ± 0.09</td>
<td>5.4 ± 1.49</td>
<td>0.025 ± 0.01</td>
</tr>
<tr>
<td>Fructose</td>
<td>5.94 ± 0.57</td>
<td>5.35 ± 0.18</td>
<td>4.4 ± 0.65a</td>
<td>0.04 ± 0.004</td>
</tr>
</tbody>
</table>

Moreover, among the different concentrations of sucrose tested, a non significant difference was recorded for the leaf number and the root length. From these results, 3% sucrose concentration in the basal medium seems to be effective for normal plant growth. Sucrose is the most widely used carbon source in most of the plant in vitro cultures, as it is the major sugar form translocated via the phloem.

As a carbon source, sucrose supports growth of plant cells in culture. A sucrose concentration of 1-5% is generally used for in vitro tissue culture, since it is also synthesized naturally by the tissue. For tissue culture, researchers commonly use 3% sucrose in the medium as per recommendation. Besides serving as energy source, it also provides the carbon precursors for synthesis of structural and functional components.

Hoque et al. [23] analyzed seed dormancy status evaluated using different media with or without sugar (3%) in rice cultivars. Jhora cultivar showed strong seed dormancy than the wild and cultivated rice genotypes.

Fruit plants were widely analyzed for their economic importance and are cultivated mainly for fruit juice. To obtain a continuous source of material for screening of secondary metabolites, zygotic embryo culture was attempted among 62 Passiflora species, starting from seeds mainly collected in the wild. 29 species produced calli, which had different growth rates. Plants were successfully regenerated from calli of 13 different species [24]. Similarly, Hossain et al. [25] designed an efficient protocols for in vitro seed germination, neo-formation of secondary protocorms from primary protocorms and multiple shoot buds and protocorm-like body induction from pseudo-stem segments of in vitro-raised seedlings of Cymbidium giganteum using four nutrient media, such as MS, Phytamix, Mitra, and Knudson ‘C’. Hassanein and Aozo [26] reported seed drying decreased both the percentage of seed germination and the number of seedling per seed. Germination of seeds was better on MS medium supplemented with 0.5 mg benzyl amino purine than in soil. Bae and Yoon [27] reported that soaking of seeds in GA3 solution remarkably promoted germination up to 60%, but the control (0 mg/L) was not effective (> 5%). Toma and Rsheed [28] analyzed in vitro propagation through seed

Effect of Different Concentrations of Sucrose

Development of in vitro plantlets was further investigated to study the effect of different concentrations of sucrose, 1, 2, 3, 4, 5 and 6% (w/v), on plant growth. The results obtained after six weeks of culture were recorded in Table 5. Increasing the sugar concentration from 1 to 3% has visually stronger influence on plant height and biomass production. But, above 3% concentration of sucrose, no significant difference was noticed (Figs. 2c & d).
culture and regeneration of *Asparagus densiflorus* L. through callus cultures derived from hypocotyls.

**Hardening of in vitro culture raised plants**

Seed germinated and regenerated plantlets that are well developed were transplanted on sand medium and showed remarkable acclimatization to the environment (95% survivability) (Figure 3). The acclimatized plantlets were successfully transferred in the field with 93% survivability. The field transferred plants showed absence of variability with respect to the *in vivo* generated plants.

![Figure 3: Hardening of tissue culture raised plants.](image)

**Conclusion**

The overall objective of this investigation was to define the optimal conditions for *in vitro* seed germination and plant growth of *Begonia malabarica*. The major results showed that the composition of the germination medium influence the germination capacity of seeds used in *B. malabarica*. The best seedling growth was observed on the seeds on half MS medium containing 3% sucrose. This established protocol would provide sufficient materials as source of explants for initiating different types of *in vitro* callus and cell culturing in the species.

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**References**


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