Germination of Pollen Grains of *Spathodia companulata* P. Beauv in *in vitro* and *in vivo* like Condition with the Influence of a Promoter Medium

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Abstract  *In vitro*, germination is the most commonly used technique in pollen physiology. This provides a simple experiment method to study the pollen germination and pollen tube growth, in the endogenous as well as exogenous conditions. *In vivo* conditions (endogenous) can also apply to the study of pollen system. In the present investigation, the pollen germination of *Spathodia companulata* P. Beauv was studied in the different parts of the carpel viz., style, stigma, ovary, style-stigma, style-ovary and whole gynoecium. *Spathodia companulata* P. Beauv is planted for reforestation schemes, for soil conservation. In India, it is seen that there is a rare formation of fruit and seeds in this plant. Thus for the study of the growth of the plant for cultivation purposes, the growth of the pollen tube was studied which could be one of the reason for the less fruit and seed formation. Pollen tubes were found to be stimulated by those components present in the different selected parts of the carpel.

Keywords  *In vitro, in vivo* like Condition, Pollen Germination, Pollen System Female Reproductive Parts (FRP)

1. Introduction

*Spathodea campanulata* is an African plant introduced into South America and other tropical and subtropical areas for ornamental purposes. The flowers bloom with great profusion, and the trees can be seen from great distances. It is not browsed by domestic animals and although a popular decorative tree for avenues. It has shallow roots and a tendency for branches to break off in a storm. The unopened flower buds contain a sweet, watery liquid that is considered to be tonic [1]. The seeds, flowers and roots are used as medicine in most of the African countries. The tree is planted for soil improvement, reforestation, erosion control and land rehabilitation.

Before attempting any breeding experiment, testing the germinability of pollen from different sources is necessary. Methods for testing pollen grain viability usually employ either the flurochromatic reaction test or germinability test under near optimum germination conditions. *In vitro* pollen germination studies not only help in understanding different aspects of pollen biology but are also crucial for assisted breeding programmes, in which regular monitoring of germinability of pollen samples that are stored, is required [2]. BK (Brewbaker and Kwack) medium is the most frequently used and has been found to be suitable for over number plant species. However, flowering plant species do not always respond satisfactorily to BK medium and thus adjustment in the concentration of the medium component is required sometimes [3]. Therefore, pollen germination media recommended so far range from simple sugar/boric acid
media [4] to complex ones containing polyethylene glycol (PEG) formulated by Zhang and Croes (1982) and Tandon et al. (1999) [5, 6].

The aim of this study provides the details of formulation of an optimized pollen germination medium as an in vitro study with the association of the conditions of in vivo provided to study by observing the effect of different parts of carpels on the pollen tube growth in Spathodea companulata.

2. Materials and Method

2.1. Collection of Plant Material

The present study was conducted on S. companulata P. Beauv collected from RTMNU, Nagpur, Campus premises (Figure 1). The identification was confirmed by the flora of Maharashtra Vol. I and II [7].

Classification

Kingdom: Plantae
Subkingdom: Tracheobionta
Superdivision: Spermatophyta
Division: Magnoliophyta
Class: Magnoliopsida
Subclass: Asterideae
Family: Bignoniaceae
Genus: Spathodia
Species: companulata
P. Beauv

2.2. In vitro and in vivo like Conditions for Pollen Germination and Tube Growth

To standardize the germination medium, for pollen germination and pollen tube development, the different extracts of female reproductive part (gynoecium) viz., stigma, style, ovary, stigma-style, style-ovary and whole gynoecium were raised in various concentration of Brewbacker and Kwack (BK) medium (10%, 20% and 30%) which contain Sucrose- 25%, Boric acid: 0.01% (10mg/100ml), Calcium nitrate: 0.03% (30mg/100ml), Magnesium sulphate: 0.02% (20mg/100ml), Potassium nitrate: 0.01% (10mg/100ml). Pollen grains were dusted in each grade of BK medium and incubated at room temperature for 1 - 4 hours. After incubation one drop of this solution was placed on a clean glass slide and it was stained with 1% aceto-orcein and glass cover was placed over it. It was then observed after few minutes under microscope. Pollen germination percentage and tube growth were evaluated. The germination response of pollen was expressed as percentage. The length of pollen tube was measured using a calibrated ocular micrometer.

3. Results

The percentage of pollen germination and pollen tube growth was assessed in the extract of different Female Reproductive parts (FRP) respectively. An attempt made to test the effect of various concentrations of FRP extract was able to enhance the germination response and showed highest pollen tube length. Initially no germination response was achieved up to 30 min of incubation in BK medium supplemented with the extract of all female reproductive parts (FRP). However, the pollen grains responded to germination after the incubation period was extended up to 1 - 4 hours at room temperature with the same composition. Ovary extract gave the best response to germination while maximum tube growth was noticed in the extract of whole gynoecium (Table 1 and Figure 2, 3). This shows that pollen germination and tube growth regulated by all the components present in the whole gynoecium. Overall 20% and 30% BK concentration shows maximum germination in one and other selected parts of gynoecium.

Figure 1. Spathodea companulata: A. Inflorescence with young buds and open flowers. B. Open flower showing anthers and stigma. C. Pistil with terminal style (St), bilobed and closed stigma (s) and the nectariferous disc at the base of ovary (ov)
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**Table 1.** Pollen Tube Length (in µm) in Different Female Reproductive Parts

|          | Style       | Ovary      | Stigma-Style | Style-Ovary | Whole Gynoecium | Stigma       |
|----------|-------------|------------|--------------|-------------|-----------------|--------------|
| 10%      | 17.73 ± 3.37| 9.33 ± 0.93| 20.53 ± 0.93| 27.13 ± 3.35| 60.67 ± 5.19    | 11.5 ± 0.3   |
| 20%      | 21.47 ± 6.53| 7.47 ± 0.93| 17.73 ± 1.87| 38.27 ± 6.73| 59.73 ± 4.94    | 12.61 ± 0.67 |
| 30%      | 26.13 ± 6.12| 6.53 ± 0.94| 16.8 ± 1.61 | 29.87 ± 10.4| 56.93 ± 6.12    | 13.71 ± 0.44 |

Values are Mean (Pollen tube Length in µm) ± SEM (n = 3)

Values are Mean ± SEM (n = 3)

**Figure 2.** Effect of Female Reproductive parts (FRP) on pollen germination

**Figure 3.** A-Germinated pollen grain in Gynoecium; B-Measurement of Pollen tube with ocular

### 4. Discussion

The pollen grains of *Spathodea companulata* failed to respond well *in vitro*. Thus it became essential to optimize a germination medium suitable in *in vivo* for this species. The sugar in the medium acts as an osmotic regulant which regulates the diffusion rate of water from the medium into pollen grains. The failure of pollen to germinate and its bursting may indicate lack of sugars within pollen grains and critical dependence on external supply [8].

Pollen grains are believed to be deficient in boron, which is normally compensated by high levels of boron present in stigma and style. Boron combines with sugar to form a sugar-borate complex which facilitates translocation of sugar molecules [9]. Its exogenous application, as observed in many trees, significantly influences both pollen germination and tube growth [10, 11, 12].
In both *in vitro* and *in vivo* conditions, optimal pollen germination and continued tube growth require a high level of calcium (Ca) (300-5000 ppm). Calcium is involved in cationic balance and is essential for tube elongation [3, 13, 14].

In the present study a noteworthy increase in the germination rate of *Spathodea campanulata* pollen in Female Reproductive parts (FRP) extract supplemented medium indicated its positive influence. The pollen grains of *Spathodea campanulata* lack those substances (like Boron) which were endogenously present in the Female Reproductive part (FRP) of flower. Thus the application of the BK medium with the extract improved the response. Thus our study demonstrates that the pollen grains of *Spathodea campanulata* might be deficient in those mineral components which required for the pollen grain germination and tube growth. But under *in vivo* conditions the same nutrients are perhaps supplied by the stigmatic exudate for attaining successful fertilization and seed-set. Also BK medium increases the rate of pollen tube development giving *in vitro* condition.

5. Conclusions

The growth of pollen tube is stimulated by those components which are only present in the whole gynoecium. The pollen tube requires the natural substances for elongation and which are possibly present in *in vivo* condition but there are some limitations in the natural conditions causes less fruit and seed formation. BK medium as a promoter influence the growth of pollen tube more rapidly. Thus a simple medium may be developed to suit the germination requirement of pollen.

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