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Effect of Different Chemical Treatments for Breaking Seed Dormancy in Sesbania spp.

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> Fast growing N_2 fixing *Sesbania* spp. has shown great promise both as high quality fodder and green manure. In the present study seed dormancy and germination were

> evaluated for four Sesbania spp. viz., Sesbania sesban, S. grandiflora, S. aculeata and

S. rosrtata grown in different parts of the India. Sesbania grandiflora seeds were

highly dormant as compare to the seeds of other Sesbania spp. Amongst the physical treatments, soaking in distilled water (24 hr) showed maximum seed germination in all

plant species. By using chemical treatments (sulfuric acid, ethanol and potassium

ABSTRACT

Keywords

Germination, Seed dormancy, Sesbania spp, Soaking, Temperature

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Introduction

The genus Sesbania contains about 70 species prevalent over tropical and subtropical regions, including perennial, annual, herbaceous shrubs and trees. Sesbania spp. are native to Asian countries such as India, Malaysia, Indonesia and Philippins where it is generally growing on the dikes between rice paddies along roadsides and in backyard vegetable gardens. In and around India, many species of Sesbania are widespread with abundance of S. speciosa, S. grandiflora, S. sesban (L.), S. bispinosa (Jacq.), S. cannabina

germination.

nitrate), immersion in concentrated sulfuric acid (20 min.) followed by ethanol resulted the maximum seed germination in S. grandiflora while in Sesbania sesban, S. aculeata and S. rostrata immersion in ethanol (100% for 5 min.) showed highest seed (Retz.), S. aculeata and S. rostrata (Bhat, 2003). Most of the Sesbania spp. are described as soft, semi or slightly woody with

1-4 m height. These grow quickly and rapidly accumulate nitrogen (N₂) rich biomass suitable for soil fertility replenishment and also provide fuel wood, fodder and mulch.

During living stage of plant the seedling phase is a critical point, helpful in investigation the natural and developmental story of higher plants (Duke and Polhill, 1981). To sustain extensive periods of unfavourable environment, the seed is well set

mainly in a dry state. Without germinating, seeds can survive for an extended period more over as stored in dehydrated environment otherwise once covered under the soil. Seed enters a dormant position to optimize germination over time. Due to dormancy in seeds of Sesbania spp., seed reservoir in soil helps endurance and perpetuation in different conditions. Dormancy in seeds is forced by single or more of the kernel, endosperm and protective tissues that be related to double genesis and stand for two ploidy zone. Via avoidance of gas swap over, water uptake, light dispersion and escape of determent from the embryo, thick seed coat inhibits seed germination (Taylorson and Hendricks, 1977). To defeat state that are harsh for seedling, seed dormancy allows seeds to be established and is therefore significant for agriculture and plant ecology. Dormancy prevents pre-harvest germination as well. Failure of a viable seed, to sprout in positive water, temperature situation (e.g. and aeration) leads to seed dormancy (Bewley, 1997; Finch-Savage and Leubner-Metzger, 2006). To recognize the types of the key controller(s) that inhibit(s) the beginning of germination (dormancy) that initiate the germination practice and their communal interface are question in dormancy and germination study.

Germination is most important and sensitive stage in the life process of a seed bearing plant. Germination behaviour of Sesbania seeds could be improved either via diminishing the latency phase otherwise improving the eminence of seed batch. Seed component tissues that have been used to infer the involvement in dormancy obligation can be arrange into somatic, molecular and genetic groups on the basis of experimental approaches. To use embryos otherwise physically erase the protective tissues testa and pericarp housing caryopses in grass breed are somatic approaches used alternatively. To

fastest and most uniform germination of Chemical treatment also enhances the germination rate in different seeds. Moreover, research in laboratory, greenhouse and field conditions conducted by Oliveira et al., (2012) indicated that the success of local seed germination in the field has a low frequency. In array to recognize the probable function of different ecological aspect for efficient farming of Sesbania spp. it is extremely significant to follow seed germination necessities. Inspite of the potential uses of Sesbania spp. in agriculture and other field, accessible scarce data is agrobotanical character together with seed

deduce the occurrence of embryo or coat-

imposed dormancy the naked caryopses or

isolated embryos were germinated on a

special medium, as succeed in oat, barley,

Arabidopsis, wheat and rice (Takahashi,

1963; Morris et al., 1989; Foley, 1992; Wang

et al., 1995; Lee et al., 2010). Botumile et al.,

(2020) studied the highest mean germination

percentages for S. galpinii (92-100%) were

observed for seeds treated with sulphuric acid

(15, 30, 45 and 60 minutes). For V. robusta,

mechanical scarification resulted in the

highest mean germination percentage (96%),

and the boiling water treatments, except for

the treatment with 24 hours of cooling down,

resulted in the lowest mean germination

percentage. The two study species possess seed coat imposed dormancies that require

pre-sowing seed treatments. Based on the

results, the best treatments to release

dormancy in both S. galpinii and V. robusta

scarification, as these yielded the highest,

and

mechanical

on

their

acid

sulphuric

are

seeds.

dormancy, germination and seedling properties. Hence, in the current research seed germination behavior in four Sesbania spp. growing in different parts of India was evaluated by using different physical and chemical treatments.

Materials and Methods

Seeds of four *Sesbania* spp. *viz.*, *Sesbania sesban*, *S. grandiflora*, *S. aculeata* and *Sesbania rostrata* were collected during January, 2016 from Tamil Nadu Agricultural University, Coimbatore, Tamil Nadu. They were individually thrashed on paper and seeds in good physical shape were chosen by removing the trash and broken seeds (Plate 1). After washing with distilled water, constant moisture of seeds were obtained by putting in sunlight for 2-5 days and placed in airtight containers for further study.

Around 30-40 seeds from each Sesbania spp. without any previous treatment were rinsed in distilled water (DW). After that seeds were soaked in 25 ml DW flask up to 24 hr and placed consistently on a layer of moistened cotton bed in glass petri plates and incubated at 28±2°C including 12 hr photoperiod up to 7 days in growth chamber for germination. Growth chamber was maintained for relative humidity and temperature. Germination of seeds was considered on the basis of coming out of radical not less than 2 mm from the seed coat. Using randomized design the germination behavior of Sesbania seeds against different treatments was assessed regularly. Temperature treatment of seeds was done by preheated in petri plates up to 30-60 min. in an oven at 40, 50 and 60°C. Sulfuric acid treatment of Sesbania spp. seeds was performed with concentrated sulfuric acid (98%) for various time interval (5, 10, 15 and 20 min), then seeds were made acid free by rinsing (8-10 times) in DW to make sure whole elimination of acid. Ethanol treatment of seeds was performed by sopping wet in ethyl alcohol (70 and 100%) for 5 min, followed by 8-10 times rinsing in DW. For potassium nitrate treatments seeds soaking with various concentration (0.2, 0.3, 0.4, 0.5 and 0.6%) of 99% potassium nitrate were performed for 24 hr followed by (8-10 times)

rinsing in DW. Seeds with different therapy were permitted to grow on 1% agar medium plates. Percentage of cumulative germination (CG) of seeds was estimated on a daily basis up to 7 days (Bewley and Black, 1994):

 $CG(\%) = (\sum n \div N) \times 100$

(Where, n is the number of seeds germinated at each day and N is the total number of seeds sown). In control and physical scarification methods seed germination and seedling dimensions were calculated.

Results and Discussion

Despite the dormancy of Sesbania seeds, considerable germination was observed due to soaking in the distilled water. The maximum germination of seeds was observed on treatment with temperature at 50° C in four Sesbania spp. In India throughout summer. the seeds of Sesbania spp. in soil seem to be affected as a result of high temperature resultant in enhanced germination on the commencement of monsoon. Scarification has an imperative function among the diverse protocol used to reduce dormancy in legume seeds (Hartmann et al., 1997). Usually performed scarification processes consist of rasping with heat and acid treatment. Chemicals those are not growth regulators enhance metabolic could action and encourage germination. Seeds scarification with concentrated sulfuric acid (20 min), ethanol (100% for 5 min) and salts (0.4% for 5 min) increased seed germination and seedling dimension in four Sesbania spp. seeds. Enhanced germination of Sesbania grandiflora seeds were observed with concentrated sulphuric acid up to 20 min treatment, which was more than control, soaking, ethanol, potassium nitrate and temperature treatments (Table 1 and Plate 2). In S. grandiflora seeds to get better germination, acid-treatment increase

permeability of seed coat particularly of dormant. Seeds treatment with ethanol (100% for 5 min) resulted excellent germination and sprout development in *Sesbania aculeata*, *S. sesban* and *S. rostrata* seeds. However, seed

germination using treatment with potassium nitrate (0.4%) was higher in all four *Sesbania* spp. studied than other concentrations (0.2, 0.3, 0.5 and 0.6%) (Table 1).

Table.1 Percentage germination of untreated and treated seeds of Sesbania spp.

Species	Treatme	ent	Germination (%)
Sesbania sesban	Control	Untreated	17
	Soaking	Distilled water	32
	Temperature (⁰ C)	40	18
		50	27
		60	25
	Concentrated	5	44
	Sulfuric acid (min.)	10	50
		15	53
		20	55
	Ethanol (%)	70	60
		100	76
	Potassium nitrate (%)	0.2	47
		0.3	49
		0.4	53
		0.5	48
		0.6	45
Sesbania grandiflora	Control	Untreated	12
	Soaking	Distilled water	30
	Temperature (⁰ C)	40	20
		50	29
		60	26
	Concentrated	5	20
	Sulfuric acid (min.)	10	40
		15	55
		20	70
	Ethanol (%)	70	35
		100	47
	Potassium nitrate (%)	0.2	25
		0.3	27
		0.4	32
		0.5	26
		0.6	22
Sesbania aculeata	Control	Untreated	30
	Soaking	Distilled water	52
	Temperature (⁰ C)	40	33
		50	40
		60	37
	Concentrated	5	60

	Sulfuric acid (min.)	10	68
	× /	15	72
		20	80
	Ethanol (%)	70	75
	~ /	100	82
	Potassium nitrate (%)	0.2	51
	, , ,	0.3	57
		0.4	63
		0.5	59
		0.6	53
Sesbania rostrata	Control	Untreated	38
	Soaking	Distilled water	52
	Temperature (⁰ C)	40	30
		50	43
		60	38
	Concentrated	5	40
	Sulfuric acid (min.)	10	55
		15	68
		20	73
	Ethanol (%)	70	62
		100	70
	Potassium nitrate (%)	0.2	50
		0.3	52
		0.4	60
		0.5	56
		0.6	51

Plate.1 Dry seeds of *Sesbania sesban* (a) *Sesbania grandiflora* (b) *Sesbania aculeata* and (c) of *Sesbania rostrata* (d)



Plate.2 Germination behavior of *Sesbania sesban* treated with ethanol (70%) (a) *Sesbania grandiflora* using concentrated sulphuric acid (20 min) (b) *Sesbania aculeata* with potassium nitrate (0.4 %) and (c) of *Sesbania rostrata* at 50° C temperature (d) on 1% agar medium plates



Although potassium nitrate is normally cause seed germination but their function is not clear. Seeds of S. sesban and S. aculeata are less dormant than S. grandiflora followed by S. rostrata. Similarly Pollard et al., (2011) reported that pea-shaped seeds (e.g. S. formosa, S. tripettii, S. drummondii, S. grandiflora and S. speciosa) germinates superior as compare to cylinder-shaped seeds (e.g. S. cannabina, S. sesban, S. aculeata, S. javanica, S.egyptica, S. rostrata and S. exaltata). In Sesbania rostrata and S. grandiflora seed dormancy could be due to the seed coat rigidity. The temperature is very influential on seed germination. Shonjani reported that environmental (2002)conditions, for instance temperature, pH, low oxygen concentration, salinity, sediment accumulation and flooding influence seed germination in swampland. Ambika et al., (2014) reported the effect of seed size on seedling vigour and seed yields. Overall, meager soaking in distilled water, seeds of all plants resulted considerable germination. The highest seed germination was also achieved in

grandiflora and S. S. rostrata with concentrated sulfuric acid proceed by soaking in ethanol. Seeds of S. sesban and S. aculeata showed the highest germination on treatment with ethanol. Shreelalitha et al., (2015) concluded that seed germination rate of Sesbania could be enhanced by soaking the seeds in distilled water for 24 hours or in a solution of GA3 (250 mg/L) for 24 hours. Chanda et al., (2017) reported that seed size, both length and/or width, highly correlated with root and seedling length, base diameter, vigour index and biomass yield of Sesbania spp. Anshari et al., (2018) also reported germination time of 13 local plant species in two hydroseeding mulches containing diverse tackifier concentration, along with determined the optimal hydroseeding mulch media composition. Eight out of 13 species could be germinated: S. timorense, S. indicus, C. cajan, C. pallida, T. arundinaceae, S. sesban, S. grandiflora and T. purpurea. The fluid M1 medium was best for seeds germination of S. grandiflora (35%) and S. sesban (50%), while the thicker M2 medium was favorable for

seeds germination of S. timorense (50%) and S. sesban (67%) in 13 DAS. Carruggio et al., (2020) investigated morpho-anatomical traits and dormancy-breaking requirements in two taxa of the genus Bituminaria: the widespread B. bituminosa and the point endemic B. basaltica germination tests after mechanical, thermal, and chemical scarification treatments for seed dormancy breaking. Differences in morpho-anatomical traits were found, with B. basaltica having a thicker palisade cell layer and B. bituminosa showing larger pods. All of the scarification treatments proved to be able to break physical dormancy, with mechanical and chemical scarification being the most effective methods in both species.

In conclusion the results from the present study show that the barrier to germination of seeds of Sesbania grandiflora, S. sesban, S.aculeata and S.rostrata is the hard seed coat, which prevents water uptake. Hence, before the seeds can germinate, they require mechanisms, which overcome this barrier. The results also indicated that the barrier can be removed through different scarification pre-sowing treatments. Sesbania sesban, S. aculeata and S. rostrata immersion in ethanol (100% for 5 min.) showed highest seed germination. The future characterization of additional Sesbania genetic resources from other soil resources will have remarkable importance.

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