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Influence of Pre-Sowing Seed Treatments on Germination Pattern of *Senna auriculata*, (L.) Roxb.; (Family-Fabaceae)

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Abstract: Seed dormancy is a state in which seeds are prevented from germinating even under environmental conditions normally favourable for germination such as adequate water supply, suitable temperature and the normal composition of the atmosphere. *Senna auriculata*, (L.) Roxb.; is a much branched shrub with brown bark and closely pubescent branchlets. Dormancy breaking is of economic importance. Under laboratory conditions and in agriculture certain means of rendering the seed coat permeable have been adopted. The mechanism of artificial dormancy breaking and the natural process leading to the same effect are frequently similar. In nature seed dormancy is broken automatically due to development of growth hormones to counter growth inhibitors, leaching of germination inhibitors, maturation and after-ripening of embryo. Artificial breaking of seed dormancy includes stratification, exposure to light, scarification, alternating temperatures, hormone treatment and chemicals, pressure etc. This study aims at discovering efficient method of breaking seed dormancy in *Senna auriculata*, (L.) Roxb.; These seeds were subjected to hot water treatment and H_2SO_4 scarification for 30, 45 and 60 minutes. H_2SO_4 scarification was found to be efficient treatment in breaking the seed dormancy.

Keywords: Seed dormancy, H_2SO_4 scarification, Germination index, Germination percentage, Mean germination time.

I. INTRODUCTION

The seeds of angiosperms are essentially simple in structure. It is enclosed in a covering called the seed coat; usually store some food along with. It is the product of fertilization. The seeds insure that the gene pool will continue to the next generation. Plants have many ways to disperse and spread the population through their seeds. Seeds are fairly resistant to extreme external conditions, provided they are in a state of desiccation. As a result seeds can retain their ability to germinate, or viability, for considerable periods. Seed has a pivotal role in human and animal nutrition and life. Many of the seeds, after distribution of mother plants or harvest do not germinate in optimal conditions due to a period of dormancy. For the seed, to germinate it must be placed in environmental conditions favourable to this process. Among the conditions required are an adequate supply of water, a suitable temperature and composition of the gases in the atmosphere, as well as light for certain seeds. The requirement for these conditions varies according to the species and variety and is determined both by the conditions which prevailed during seed formation and even more by hereditary factors, (Mayer and Poljakoff -Mayber, 1982).

Hard seed coat is found in *Senna auriculata* (L.) Roxb.; To overcome the problem associated with germination, seeds need to be subjected to a specific treatment for breaking dormancy, increasing of per cent and acceleration of uniform seed germination (Frett, 1987). The seed dormancy is considered to have importance in various aspects, like perennation, dispersal, germination under favourable conditions, storage etc. Seed dormancy facilitates their storage and transport to the areas of deficiency. However its presence causes delayed and sporadic germination which is undesirable. Lack of seed germination at certain times and suitable conditions are a big problem for seed researchers, botanists, and farmers. The present study aims to find out, suitable methods for breaking seed dormancy. It is essential for achieving a uniform cultivation and proper weed control.

II. MATERIALS AND METHODS

A. Study Area (Plate-2)

Tamil Nadu is the eleventh largest state in India and covers an area of 130,058 square kilometers. It is heavily dependent on monsoon rains. Nirmala College for Women is located in Coimbatore, Tamil Nadu. Coimbatore district is a district in the Kongu Nadu region of the state of Tamil Nadu. It is located on the banks of the Noyyal River and surrounded by the Western Ghats. The climate of the state ranges from dry sub-humid to semi-arid. The soil types commonly found are loamy soil, clayey soil and calcareous black cotton soil.

Plate 1: Location Map



=N

Plate 2: Study Area



B. Sample Collection

For the present study the seeds of, *Senna auriculata* (L.) Roxb.; were collected from Sowripalayam Pirivu, Coimbatore. The seeds were obtained from the collected pods and kept in dry place till treatments.

Systematic position

Kingdom : Plantae

Sub Division : Spermatophyta

Division : Magnoliophyta

Class : Magnoliopsida

Sub Class : Rosidae

Order : Fabales

Family : Fabaceae

Genus : *Senna*

Species : *S. auriculata* (L.) Roxb.;

C. Plant Description

- 1) *Senna auriculata*, (L.) Roxb.; is a native of India. It is commonly called Avaram tree. The plant is wild in dry regions of Madhya Pradesh, Tamil Nadu and Rajasthan. It is cultivated in other parts of India. It is a much branched shrub with brown bark and closely pubescent branchlets. The leaves are alternate, stipulate, paripinnate compound. Leaflets are very shortly stalked, oval oblong, obtuse. Its flowers are irregular, bisexual, bright yellow and large, the pedicels are glabrous. The racemes are few-flowered, short; erect, to form terminal inflorescence. The fruit is a short legume, 12-20 seeds per fruit are carried each in its separate cavity.
- 2) *Uses*: The pod husk contains nonacosane, chrysophanol, emodin and rubiadin. The roots are used in skin diseases and asthma and flowers are used in diabetes, urinary disorders and nocturnal emissions. Its Bark is used as astringent. The leaves and flowers possess anti-diabetic activity.

D. Pre-Sowing Treatments

Dormancy breaking treatments were imposed with a control. The seeds were subjected for hot water scarification and H_2SO_4 scarification.

T0 – Control

T1 – Hot water treatment for 30 min

T2 – Hot water treatment for 45 min

T3 – Hot water treatment for 60 min

T4 –Scarification with H_2SO_4 for 30 min

T5 –Scarification with H_2SO_4 for 45 min

T6 –Scarification with H_2SO_4 for 60 min

12 seeds of the plant are subjected for six dormancy breaking treatments each as indicated above. For hot water soaking treatments, the seeds were soaked in hot water (80°C) for different durations. In H_2SO_4 scarification treatments, the seeds were scarified with concentrated sulphuric acid for different durations. Immediately after scarification, the seeds were washed with distilled water

thoroughly. The seeds were taken in clean petri plates after different treatments. After imposing the treatments, the seeds were sown in fertile soil taken in small pots for germination. Each pot is labelled corresponding to the treatment imposed for the sown seeds. Proper irrigation was provided to ensure adequate water availability for seeds. Germination of each of the lots was noted for the next 20 days. A comparison of calculated data is done among the seeds to derive a conclusion on the effect of seed dormancy breaking treatments.

The following parameters were found:

1) **Germination Percentage (%)** (International Seed Testing Association (ISTA), 1999): Twelve seeds of each of the plant subjected for six dormancy treatments each were sown in soil. After the test period of forty five days the normal seedlings were counted and the mean value was expressed as percent. $GP = (\text{Final no. of seeds germinated in a seed lot} \div \text{Total no. of seeds sown}) \times 100$

2) **Mean Germination Time (MGT)**: $MGT = \sum (f \cdot x) / \sum f$

Where, f = Seeds germinated on day x

3) **Germination Index (GI)**: $GI = (20 \times n_1) + (19 \times n_2) + \dots + (1 \times n_{20})$

Where, n_1, n_2, \dots, n_{20} = No. of germinated seeds on the first, second and subsequent days until the 20th day; 20, 19 . . . and 1 are weights given to the number of germinated seeds on the first, second and subsequent days, respectively.

Plate: 3 - H₂SO₄ Scarification of *Senna auriculata* (L.) Roxb.; (Sample- 1)



III. RESULTS AND DISCUSSION

Table: 1- Determination of Germination Percentage (GP), Germination Index (GI) and Mean Germination Time (MGT) of *Senna auriculata* (L.) Roxb.;

Day	Control	Hot Water Treatment			H ₂ SO ₄ Scarification		
		30 Minutes	45 Minutes	60 Minutes	30 Minutes	45 Minutes	60 Minutes
1	0	0	0	0	0	0	0
2	0	0	0	0	0	0	0
3	0	0	0	0	0	0	0
4	0	0	0	0	1	1	0
5	0	0	0	0	1	1	0
6	0	0	0	0	2	2	1
7	0	1	0	1	1	0	0
8	0	0	1	2	2	0	0
9	0	2	0	1	0	0	0
10	0	0	0	0	0	0	0
11	1	0	0	0	0	0	0
12	2	0	0	0	0	0	0
13	0	0	0	0	0	0	0
14	1	0	0	0	0	0	0
15	0	0	0	0	0	0	0
16	0	0	0	0	0	0	0
17	0	0	0	0	0	0	0
18	0	0	0	0	0	0	0
19	0	0	0	0	0	0	0
20	0	0	0	0	0	0	0
GP	33.33%	25%	8.33%	33.33%	50%	25%	8.33%
GI	35	38	13	52	103	63	15
MGT	12.25	8.3	8	8	6.28	5.25	6

Table- 1 Shows that the germination percentage (GP) is higher in seeds of *Senna auriculata* (L.) Roxb.; subjected to H₂SO₄ scarification for 30 minutes, that is 50%. 25 percentage of germination was found in seeds subjected to hot water treatment for 30 minutes and H₂SO₄ scarification for 45 minutes. Low germination percentage was found in seeds subjected to hot water treatment for 45 minutes and H₂SO₄ scarification for 60 minutes. The germination index (GI) is greater in seeds subjected to H₂SO₄ scarification for 30 minutes and lowest in that treated with hot water for 45 minutes. The mean germination time (MGT) is higher for untreated seeds followed by that treated with hot water for 30 minutes. It is low for seeds subjected to H₂SO₄ scarification for 45 minutes. The length of time elapsed between the first seed to germinate and the last, the variation in germination speed and the timing that the majority of seeds germinate all have impacts on diverse cultural operations like fertilizing, harvesting and field maturity of crops (Roberts,1981, Washitani and Saeki, 1986, Kader and Jutzi, 2001). 'High' (the time at which the majority of seeds germinate) and 'low' (the time at which the minority of seeds germinate) (Kader *et al.*, 1998) germination events are also important indicators of seed vigor and stress resistance (Kader and Jutzi, 2002). The higher the GP value, the greater the germination of a seed population (Scott *et al.*, 1984). In the context of the parameters tested in the investigation comparing seed germination calculations and the associated interpretation, it appears that the GI is the most accurate among various parameters (Khader, 2005). For seeds of *Senna auriculata* (L.) Roxb.; the germination percentage (GP) is higher in those subjected to H₂SO₄ scarification for 30 minutes, that is 50%. This seed lot is having greater germination. The mean germination time (MGT) is low for those subjected to H₂SO₄ scarification for 45 minutes. These seeds germinated faster when compared to others. When the Germination Index of seeds of *Senna auriculata* (L.) Roxb.; are observed, H₂SO₄ scarification for 30 minutes is found to be effective in enhancing the percentage and rate of germination. The Germination Percentage and Germination Index is found to be greater in seed lots subjected to H₂SO₄ scarification. The Mean Germination Time is low in the seeds subjected to H₂SO₄ scarification, which is desirable. *Senna auriculata* (L.) Roxb.; shows higher values of MGT for control. Hence the control germinate at a low speed.

IV. CONCLUSION

The seed germination depends on both internal and external factors. Greater Germination Percentage and thereby greater germination is found in the seed lots of all the three samples subjected to H₂SO₄ scarification. The seed lots of *Senna auriculata* (L.) Roxb.; subjected to H₂SO₄ scarification for 30 minutes shows greater germination. A higher percentage and rate of germination is found in those seeds with higher values of germination index. The seeds subjected to H₂SO₄ scarification for 30 minutes are found as the lots with higher percentage and rate of germination. Also faster germination is found in those seed lots treated with H₂SO₄ for 45 minutes. These are the seed lots with low MGT values. Observing the parameters Germination Percentage, Germination Index and Mean Germination Time of seeds of *Senna auriculata* (L.) Roxb.; H₂SO₄ scarification is found to be effective in enhancing the overall germination pattern. Over hot water treatment and control, H₂SO₄ scarification provides better results in shrub. The given treatments gave better results than the untreated seeds (control). Hence it is preferable to subject the dormant seeds to pre-sowing treatments for fast and better germination. The higher seed germination due to H₂SO₄ scarification might be due to the weakening of seed coat by distributing and dissolving the lignins and pectins present on epidermal layer of the seed coat, which render them impermeable to water and oxygen. It is inconvenient for seed researchers, botanists, and farmers if seeds do not germinate at certain times and suitable conditions. Subjecting the dormant seeds for pre-sowing treatments are useful in germinating seeds when needed. A wide range of seeds overcomes seed dormancy after pre-sowing treatments like hot water treatment and H₂SO₄ scarification. H₂SO₄ scarification is most preferred for the seeds of *Senna auriculata* (L.) Roxb.; over the control.

REFERENCES

- [1] Agrawal P.K., Dadlani M., (1995). Techniques in Seed Science and Technology. 2nd Edition, South Asian Publishers, New Delhi. ISBN; 10: 0412063018.
- [2] Bench A.R., Fenner, M., and Edwards, P., (1991). Changes in germinability, ABA content and ABA embryonic sensitivity in developing seeds of Sorghum bicolor (L.) Moench induced by water stress during grain filling. New Phytologist, pp-118, 339-347.
- [3] Frett, J.J., (1987). Seed germination of *Cycas revoluta*. Journal of Environmental Horticulture. ISSN; 0738-2898. Vol-5. Issue-3. pp-105-6.
- [4] Kader, M., and Jutzi, S., (2002). Time-course changes in high temperature stress and water deficit during the first three days after sowing in hydro-primed seed: germinative behaviour in sorghum. Journal of Agriculture and Rural Development in the Tropics and Subtropics, ISSN; 16129830. Vol-103, pp-157-168.
- [5] Kumar, R.N., S. Chakraborty., N.J.I. Kumar., (2011). Methods to break seed dormancy of (*Burm.f.Nees*): an important medicinal herb of tropical Asia. Asian Journal of Experimental Biological Sciences, ISSN; 0975-5845. Vol-2. Edition- 1. pp- 143-146.
- [6] Mayer, A.M., and A. Poljakoff- Mayber., (1982). The germination of seeds, British Library Cataloguing in Publication Data, 3rd Ed. ISBN: 0 080288537.
- [7] Qu, X.X., Huang, Z.Y., Baskin, J.M., and Baskin, C.C., (2008). Effect of temperature, light and salinity on seed germination and radicle growth of the geographically widespread halophyte shrub *Halocnemum strobilaceum*. Annals of Botany. ISSN; 0305-7364. Vol -101. Issue -2. pp- 293.
- [8] Roberts, E.H., (1981). The interaction of environmental factors controlling loss of dormancy in seeds. Annals of Applied Biology. ISSN; 0003-4746. Vol-98, pp-552-555.
- [9] Washitani, I., and Saeki, T., (1986). Germination responses of *Pinus densiflora* seeds to temperature, light and interrupted imbibition. Journal of Experimental Botany. ISSN; 0022-0957. pp- 37, 1376-1387.



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