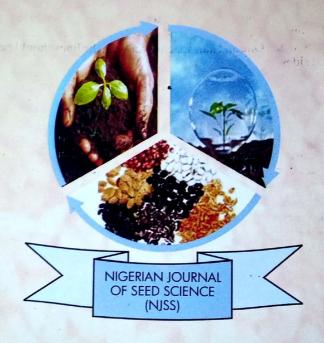


A Scientific Journal Serving Seed Scientists and Technologists

Volume 2, 2018









Association of Seed Scientists' of Nigeria (ASSN)

NIGERIAN JOURNAL OF SEED SCIENCE

Scope of the Journal: Nigerian Journal of Seed Science (NJSS) is a peer review journal published by The Association of Seed Scientists' of Nigeria (ASSN). The journal is published regularly and is available as paper copies. The scope of the journal covers all basic and applied research in all aspects of seed science including seed biology, physiology, production, processing, testing, certification, storage, pathology etc. Short communications from seed experts and technologists may also be published as Seed technical notes if such notes depict new laboratory techniques, anatomical and pathological observations of seed and seedling development etc. The journal may also accept review articles in all areas of seed science and technology relating to agriculture and seed system development.

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Manuscript format:

Each manuscript should be arranged in the following order: Title, Author(s), Affiliations, Abstract, Text (separated into introduction, methodology, results (including tables and figures), discussion and conclusion), Acknowledgments and References.

Title: The title should be concise and informative, containing key words which describe the subject matter. It should not exceed 15 words.

Author/s's name: The name of the author(s) should be written starting with the surname followed by initials and placed below the title. Numbers in superscript are placed at the end of last initial to indicate affiliation of each author. Also an asterisk (*) is placed after the name of the corresponding author.

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Germination and seedling growth of golden shower (Cassia fistula L) as influenced by method of breaking hard seed coat dormancy and dipping duration

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Abstract

Laboratory and field experiments were conducted at the Federal University of Agriculture, Abeokuta on Latitude 7o 14' N and longitude 3o 21' E to determine the best method of breaking hard seed coat dormancy and duration of dipping for rapid and complete germination and growth of golden shower seedlings. Seeds of golden shower (Cassia fistula) soaked in H2SO4 (95%); Gibberelic acid (-GA3) (0.5 g/L); hot water (100 0C) for 5, 10 and 15 minutes and untreated (control) were arranged in a Completely Randomized Design with three replications. Data collected on germination and seedling growth parameters were subjected to Analysis of Variance and treatment means separated using Least Significant Difference (LSD) at 5% level of probability. In the laboratory experiment, seeds soaked in GA3 had highest germination percentage (100%), followed by those soaked in hot water (87%), H2SO4 (81%) and untreated seeds did not germinate (0%) at 19 days after sowing. Seeds soaked for 5, 10 and 15 minutes had germination percentages of 100, 87 and 81 %, respectively. On the field, seeds treated with H2SO4 had significantly (P ≤ 0.05) higher seedling emergence (100% and 79%), followed by seeds treated with hot water (87% and 53%) while seeds treated with GA3 and untreated had zero germination. Soaking of seeds for 5 and 10 minutes resulted in highest (87 and 68%) and least (58 and 53%) seedling emergence, respectively for the two field trials. Dry matter of seedlings from seeds soaked in either H2SO4 or hot water was not different, but seeds soaked for 5 minutes had seedlings with higher values. Therefore, soaking of seeds of Cassia fistula in hot water for 5 minutes is the best practicable and economical procedure for nurserymen to induce rapid germination and seedling growth.

Keywords: Seed scarification, Seed dormancy, Chemical treatment, Seed soaking

Introduction

Cassia fistula Linn, commonly called golden shower belongs to the genius Cassia subfamily Caesalpiniacaea and family fabaceae (Sartorelli, 2009). It is a native of Southern Asia, Indian, and Malaysia but now widely grown in tropical areas as flowering ornamental plant. Golden shower is primarily grown for its beautiful bright yellow flowers, shade for relaxation and has many medicinal values (Bhakta, 1999; Grupta et al., 2000; Duraipardiyan

and Idnacimuthu, 2007). It could also be used to conserve soil, prevent erosion, improve soil fertility, as wind breaker and source of industrial raw materials in tanning and dye stuff production (Perry and Hay, 1982). However, presence of hard seed coat hinders quick germination of seeds of Cassia fistula, thereby affecting attainment of uniform establishment of seedlings in the nursery. Rapid germination and proper management of the seedlings in the nursery is very crucial for the uniform

production of seedlings for transplanting. The inability of viable seeds to germinate in the presence of all necessary external environmental factors that are required for seed germination poses a serious bottleneck to commercial production of seedlings of ornamental plants. scarification with hot water, acid treatment, hormonal treatment and heat among others have been previously reported to break hard seed coat dormancy in order to ensure quick and proper germination, seedling emergence and growth in many plants (Kim et al., 2017; Ahmed et al., 2006 and Fagbayide and Fawusi, 1994).

Cassia fistula seeds exhibit hard seed coat dormancy which prevents water imbibition, exchange of oxygen and activation of embryo for germination, resulting in poor germination rate (Ramamoorthy, 2005). However, information on the best seed treatment methods and duration of the treatment to overcome seed dormancy in Cassia fistula is scanty. Hence this study seeks to investigate the best dipping duration in H2S04, GA3, and hot water to attain optimum and even germination, seedling emergence, and growth in Cassia fistula.

Materials and Methods

The experiments were carried out in 2014 and 2015 at the laboratory and the nursery of the Department of Horticulture, Federal University of Agriculture, Abeokuta, Ogun state, Nigeria, located on Latitude 70 14' N and longitude 30 21' E. The annual rainfall was 1200 mm and the vegetation was mainly secondary forest.

In both experiments, treatments consisted of scarification methods (H2SO4 (95%)) GA3 (0.5g/l), hot water (100 °C) and untreated seeds as control) and dipping durations (5, 10 and 15 minutes) factorially combined in a Completely Randomized Design (CRD) in five replications and there were twenty five seeds each per replicate. In the laboratory, Petri-dishes lined with 7 g of cotton wool were arranged on a flat surface and treated seeds were placed in each Petri-dish with 25 seeds per Petri-dish at 1cm depth for observation. On the field, 14 kg of the air dried soil was weighed into each perforated planting pot to prevent water logging. Then 25 treated seeds were sown per pot at 3 cm depth at 5 x 5 cm spacing and each planting pot separated by 75 x 45

Mature brown pods of *C. fistula* were collected from the fully grown trees used for avenue planting at Federal University of Agriculture, Abeokuta. The pods were broken to release the seeds and seeds were washed with distilled water and the clean seeds were air dried and tested for germination. Seed germination was determined by placing seeds in Petridishes under laboratory conditions and seeds were sown in planting pots after treatment in the nursery.

Three sets of 100 seeds of Cassia fistula were soaked in 3 different glass beakers containing equal volume (20 ml) each of H2SO4, hot water and GA3 at dipping durations of 5, 10 and 15 minutes, respectively. As the duration of dipping lapsed, the acid solution was poured away and seeds were thoroughly rinsed with

distilled water to remove remnant of H2SO4, blotted dry before sowing. The control was direct germination of untreated seeds.

Though the experiment was rain fed, watering was done occasionally and weed control was done manually within the pots as required.

Data collected on the days to first emergence, days to 50% germination, percentage germination and dry weight (g) of leaves, stem and root were subjected to analysis of variance (ANOVA) using GenStat discovery and means were separated using Least Significant Difference (LSD) at 5% level of probability.

Results

Statistical analysis of the result showed that effect of method of breaking hard seed coat dormancy was significant (p< 0.05) on germination of golden shower across sampling period (Figure 1). It took seeds treated with GA3 and hot water four days (4) to start germinating followed by those treated with H2SO4 that took seven days (7) while none of the untreated seeds germinated 16 days after sowing (DAS). Similarly, seeds treated with GA3 attained 50% germination within 9 days, followed by those treated with H2SO4 in 12 days and hot water in 14 days. Seeds of golden shower dipped in GA3 had significantly (p< 0.05) highest germination percentage (100%), followed by those dipped in hot water (87%), then those dipped in H2SO4 (81%) and none of the untreated seeds germinated at 16 DAS (Figure 1). However, total germination percentage of seeds treated with H2SO4 or hot water was statistically similar.

Dipping duration significantly (p< 0.05) influenced germination of golden shower (Figure 2). Seeds treated for 5 minutes attained 50% germination in 12 days and completed germination (100%) in 14 days, while seeds dipped for 10 minutes attained 50% germination in 13 days and maximum germination of 85% in 15 days. Seeds dipped for 15 minutes attained 50% germination in 14 days and maximum germination of 77% in 16 days. Effect of dipping duration on earliness to germination and 50% germination was not significant (p<0.05).

Also, the interactive effect of method of breaking hard seed coat dormancy and dipping duration on germination percentage of golden shower showed seeds were significant (p< 0.05) indicating that they were dependent (Table 1). With seeds dipped in H2SO4, highest germination percentage (81%) was recorded with 5 minutes dipping while those treated for 10 minutes had the least (68%) at 16 DAS. Conversely, seeds treated with GA3, for 5, 10 or 15 minutes had 100% germination. However, seeds dipped in hot water for 5 and 10 minutes had the highest of 87% and lower value of 82% (Table 1).



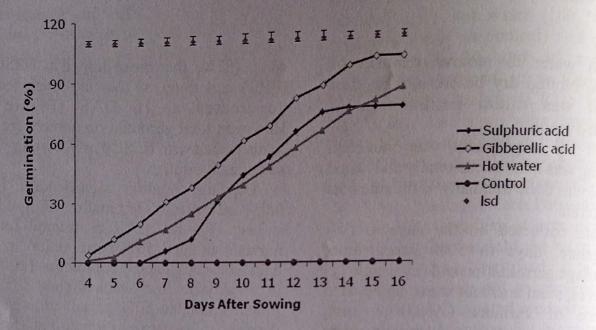


Figure 1: Effects of method of breaking hard seed coat dormancy on seed germination (%) of Cassia fistula in the laboratory

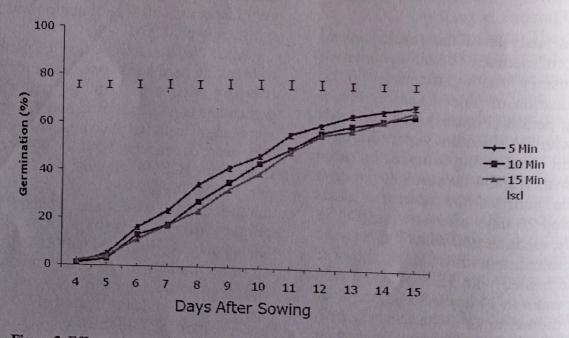


Figure 2: Effects of dipping duration on seed germination (%) Cassia fistula in the laboratory

Table 1: Effects of method of breaking hard seed coat dormancy and dipping duration on germination (%) of golden shower

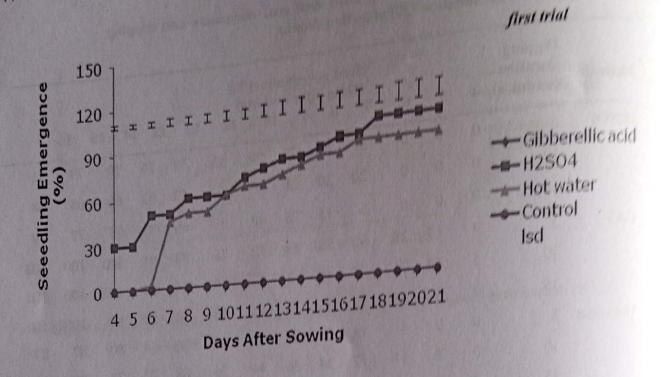
Dipping Seed germination(%) duration (Days after soaking)														
Methods	(minutes)	4	5	6	7	8	9	10	11	12	13	14 1	5 16	
H2SO4	5	0	0	0	6	17	43	54	62	78	79	79	79	81
	10	0	0	0	20	20	21	29	39	48	62	68	68	68
	15	0	0	0	6	15	28	48	59	69	71	75	77	79
GA3	5	4	17	28	42	50	59	72	73	84	91	100	100	100
	10	5	14	21	33	40	51	66	76	84	89	100	100	100
	15	4	6	10	19	25	36	45	56	74	80	88	100	100
Hot water	5	0	0	8	14	24	33	39	47	56	64	80	87	87
	10	0	3	11	17	24	36	44	56	65	72	78	82	82
	15	4	6	14	20	24	29	35	42	49	59	63	69	85
Control	0	0	0	0	0	0	0	0	0	0	0	0	0	0
LSD (P=0	0.05)	3.0	3.1	4.1	4.8	5.0	4.8	5.7	5.7	5.7	5.7	4.2	3.6	3.8

Effect of method of breaking hard seed coat dormancy on seedling emergence of golden shower was significant (p< 0.05) in both trials (Figure 3). Seeds dipped in H₂SO₄ and hot water attained 50% emergence in 9 and 12 days, respectively. However, seeds dipped in H₂SO₄ reached the highest of 100% in 19 days and those dipped in hot water reached highest of 80% in 21 days in the first and second trials, respectively. Seeds treated with GA₃ and the untreated seeds did not emerge at 21 DAS.

Dipping duration significantly (p < 0.05) affected seedling emergence of golden shower in the first and second trials

(Figure 4). Seeds of golden shower treated for 5 minutes attained 50% seedling emergence in 9 days and had highest of 100% and 80% seedling emergence, respectively in 19 days in the first and second trials, followed by those dipped for 15 and 10 minutes that attained 50% seedling emergence in 12 days and had highest of 86% and 79%, and 77% and 74% in the first and second trials, respectively.

However, the interactive effect of method of breaking hard seed coat dormancy and dipping duration on seedling emergence of golden shower seeds was significant (p< 0.05) in the first



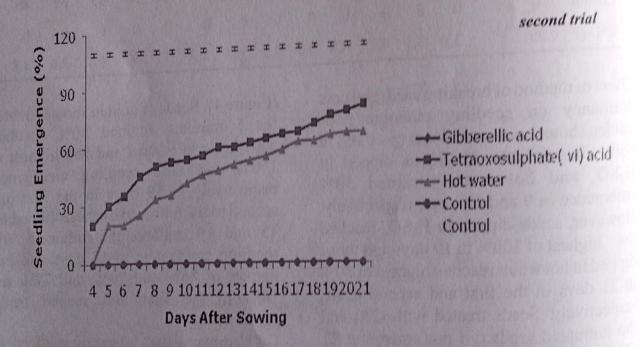
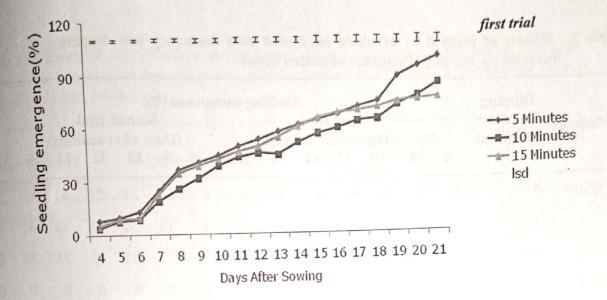


Figure 3: Effects of method of breaking hard seed coat dormancy on seedling emergence of golden shower.



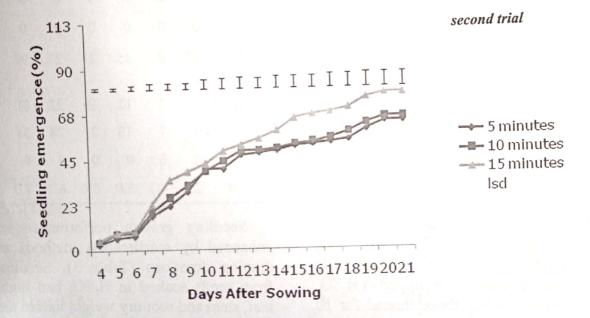


Figure 4: Effects of dipping duration on seedling emergence of golden shower.

Table 2: Effects of method of breaking hard seed coat dormancy and dipping duration on seedling emergence of golden shower

Methods	Dipping duration	Seedling emergence (%) First trial (Days after soaking) (Days after soaking)													
Methods	(minutes)	6	8	(Day 10	s afte 12	r soa 14	king) 16	18	6	8	10	12	14	16	18
H2SO4	5	25	35	36	44	66	74	79	8	18	28	37	41	61	70
	10	17	24	25	35	50	59	66	4	13	20	27	30	47	53
	15	19	28	34	41	57	64	70	7	15	23	36	38	53	61
GA3	5	0	0	0	0	0	0	0	0	0	0	0	0	0	0
	10	0	0	0	0	0	0	0	0	0	0	0	0	0	0
	15	0	0	0	0	0	0	0	0	0	0	0	0	.0	0
Hot water	5	12	17	20	30	42	47	53	4	8	15	23	27	39	44
	10	9	14	18	25	35	38	42	2	7	12	19	22	31	37
	15	10	15	18	27	36	40	44	4	7	13	21	24	34	38
Control	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
LSD (P=	0.05)	4.8	5.4	5.7	6.9	8.4	8.8	8.9	2.7	4.2	5.0	6.0	6.7	7.9	8.5

and second trials (Table 2). Highest seedling emergence of 79% and 70%, respectively in the first and second trials were recorded from seeds dipped in H₂SO₄ for 5 minutes while those treated for 10 minutes had the least of 66% and 53%, respectively at 18 DAS. Seeds dipped in hot water for 5 minutes had highest seedling emergence of 53% and 44% followed by those dipped for 15 minutes with 44% and 38%, while seeds dipped for 10 minutes had the lowest of 42 and 37% in the first and second trials, respectively. Seeds treated with gibberellic acid and the untreated (control) seeds did not emerge.

Seedling growth performance was enhanced by scarification methods and dipping durations (Table 3). Seedlings from seeds soaked in H₂SO₄ had higher leaf, stem and root dry weight values than those soaked in hot water. Soaking of seeds for 5 minutes resulted in superior seedlings growth with higher leaf and stem dry weight than other treatments, but was not different in terms of dry root weight. The interactive effects show that increase in soaking duration in either H₂SO₄ or hot water resulted in lower leaf, stem and root dry weight except for dry root weight of seedlings from seeds soaked in hot water.

Table 3: Effects of scarification methods and dipping duration interactions on leaf, stem and root dry weight of golden shower at 18 weeks after sowing (mean of both trials)

	were all and the first and the first	Dry	weight (g)	
Scarification Methods (SM)	Dipping Duration (min) (DD)	Leaf	stem	root
Sulphuric acid	Adular of no beautiful	2.7	1.3	3.2
enia han ta	10	2.2	1.1	2.6
	15 0 414	2.0	0.9	2.6
Gibberellic acid	5	0.0	0.0	0.0
	10	0.0	0.0	0.0
ap was bee ato	spinsor 115	0.0	0.0	0.0
Hot water	standard 5	1.5	0.7	0.5
	10	1.3	0.6	0.6
	15	1.2	0.4	0.6
Control	0	0.0	0.0	0.0
LSD(p=0.05) SMXDD		0.25	0.16	0.77
LSD (p=0.05) SM		0.14	0.09	0.44
LSD (p= 0.05) DD		0.12	0.08	ns

should be given to concentration and duration of soaking in chemical acids to avoid possible damage of embryo which may result in deformed seedlings.

Soaking in hot water is economical and eliminates problem of scorching and Previous death of emerged seedlings. works (Seiler, 1994; Fagbayide and Fawusi, 1994; Rehman and Park, 2000 and Chuanren et al., 2004) emphasize pretreating seeds of legumes with hard seed coat at a specified dipping duration for faster and complete germination. Chuanren et al., (2004) described seeds of Cassia fistula as similar to those of many other hard coated legumes and that scarification could enhance its germination ability. Past studies have been inconclusive or at best requesting further investigation on appropriate duration of soaking for chemical treatment of seeds with hard seed coat. This study established soaking period of 5 minutes as adequate for rapid germination of Cassia fistula seeds.

Germination success of golden shower seeds soaked in GA₃ in the laboratory was unique compared with poor observation on the field. This might be due to prevailing environmental factors that may inhibit germination. Application of growth regulators such as GA₃ has a positive effect on germination and could double the germination rate of dormant seeds with hard seed coat (Seiler, 1994; Borghetti et al. (2002)). However, Li et al., (2005) suggested that gibberellins synthesis and perception are often affected by light and temperature among other numerous

environmental signals that can influence release from seed dormancy. Better seedling performance of Cassia fistula seeds treated with H₂SO₄ and hot water suggests positive relationship between rapid germination and superior seedling establishment.

Conclusion

Based on the result from the experiment, it could be concluded that, highest germination was recorded in seeds treated with GA₃ acid for 5 minutes in the laboratory while optimum seedling emergence occurred in seeds treated with H₂SO₄ on the field. However, soaking seeds of Cassia fistula in hot water for 5 minutes is the most practicable and economical procedure for faster germination and uniform growth of seedlings in the nursery.

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