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Effect of Seed Priming Using GA₃, KNO₃, ZnSO₄ on Germination Parameters of *Vigna unguiculata*

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ABSTRACT

Cowpeas (Vigna unguiculata), an important legume is a versatile crop, also commonly known as southern pea, black eye pea, crowder pea, lubia, niebe, coupea or Frijole. The establishment of this crop is poor due to poor germination. Seed priming technique helps to overcome this difficulty. There is only less research have been carried out to enhance seed germination in cowpea by seed priming technique. This research paper focuses to study on the effects of different priming agents like GA₃ at 20 ppm, 40 ppm and 60 ppm, KNO₃ at 1 %, 2%, 3%, ZnSO₄ at 1 %, 2%, 3% and hydro priming all in 3 hrs and 6 hrs soaking conditions. The untreated seeds serve as control. The result from the present study revealed that the germination parameters viz., germination value, germination index, imbibition, co-efficient of variation of germination, seed length, root length, seedling length, vigor index were at its maximum

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for the seeds primed with GA₃ at 60 ppm, KNO₃ at 2% and ZnSO₄ at 2%. Priming of seeds at lower concentration improves the seed germination parameters which in turn increases the yield.

Keywords Seed priming, Hydro priming, Osmo priming, Hormonal priming, Nutri priming, Nano priming.

INTRODUCTION

Cowpea also known as Vigna unguiculata belong to the family of Fabaceae. It was originated in Africa. It is known by the local names as Lobia in India, Caupi in Brazil, long bean in China. Humans use cowpea as food, fodder for cattles and as manure for agricultural purposes. Cowpea can be grown for its grain, pod as vegetable, leaves as fodder, hay, silage, mulching material, intercrop with many cereal crops, fixes atmospheric nitrogen and many more. The protein content is around 22.4%, carbohydrate 55-66%, iron levels varied from 2.0 to 2.4 mg/1 kg seeds, whereas calcium levels ranged from 9 to 36 mg/100 g (Gondwe et al. 2019). It also contains vitamins such as thiamine, riboflavin and niacin, aminocaids like lysine, leucine and phenylalanine. As the crop has many uses the production of crop is very low in developing countries due to the usage of seeds which are not stored in a proper way. The lack of knowledge about handling of seeds after harvesting, nutritional qualities of the seeds and the production practices are some of the reasons for low production. The seeds stored by the farmers does not achieve recommended seed standards to uses as a seed. Many researchers have

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confirmed this (Eskandari and Kazemi 2011, Kamara et al. 2019, Njonjo et al. 2019). Seed priming may improve germination parameters like germination index, seed vigour, reduce emergence time, imbibition, speed of germination. Seed priming is a process by which seeds are induced into a state of pre-germinative metabolism by controlled rehydration to increase germination rates and germination vigor (Paparella et al. 2015). In this method seeds are hydrated up to a point where pre germinative metabolic activities may continue but radicle emergence is prevented. Seed priming with different priming methods like hydropriming, osmo priming, hormonal priming, nutri priming, nano priming can be done to improve the germination, uniform seedling establishment as these are essential stage of a plant life. In the present study seed priming was done using three agents viz., GA₃, KNO₃ and ZnSO₄. GA₃ helps to overcome seed dormancy arising due to weak endosperm. Priming of sunflower seeds (Ulfat et al. 2017) and wheat seeds (Jafri et al. 2015) with GA₃ increased seed yield. Seed priming with KNO, incerases seed germination and seed priming with ZnSO, doubled the yield in maize. These three agents are easily available to farmers. Hydropriming has many advantages which has been shown in many crops viz., wheat (Harris et al. 2001), chickpea (Musa et al. 2001), maize (Ashraf and Rauf 2001), sunflower (Kaya et al. 2006) and barley (Abdulrahmani et al. 2007). But not many authors investigated the effects of seed priming in cowpea. Hence, this research was aimed to investigate the effects of priming using GA₂, KNO₂ and ZnSO₄ and hydropriming for 3 hrs and 6 hrs soaking time at three concentrations on germination parameters of this crop.

MATERIALS AND METHODS

The study was conducted using genetically pure seeds of cowpea cv VBN 3 obtained from Krishi Vigyan Kendra, Sandhiyur, Tamil Nadu. The experiment was conducted at the Department of Genetics and Plant Breeding, Faculty of Agriculture, Annamalai University, Annamalai Nagar (11°24 N latitude and 79°44 'E longitude with an altitude of +5.79 m above mean sea level) with four replications in a Factorial Completely Randomized Block Design (FCRD) in three sets. The bulk seeds were first dried to below

 Table 1. Priming agents, their concentrations, and duration of priming in cowpea cv VBN 3.

Priming	g agent	Concentration (Factor 1)	Duration of soa- king (h) (Factor 2)
	Hydropriming		
Set 1	GA ₃	20 ppm 40 ppm 60 ppm	3 and 6
	Control	_	_
	Hydropriming	_	
Set 2	KNO ₃	1% 2%	3 and 6
		3%	
	Control	_	_
Set 3	Hydropriming	_	
	ZnSO ₄	1%	
		2%	3 and 6
		3%	
	Control	_	_

9% moisture content, cleaned, and then graded with suitable sieves. The concentrations of the priming agents as factor 1 and factor 2 being the duration of soaking hours. Their treatment details are given in Table 1.

Seeds were treated with their respective solutions for a specified duration. Following treatment, the seeds were extracted from the solutions, rinsed with water two to three times, air-dried in shade until they regained their original moisture content, and evaluated for various seed quality parameters, such as germination index, root length, shoot length, dry weight, seedling length (cm), mean germination rate and time (Ranal et al. 2009), imbibition rate (Tian et al. 2014), water content, speed of germination, germination percentage, peak value germination value, (Czabator 1962) days to 50% germination, germination percentage and vigour index were recorded with hydroprimed seeds and control in each set. Germination test was performed using glassplate and petri-plate method and Germination (%) = No. of normal seedlings germinated × 100/ Total no. of seeds placed for germination. Speed of germination was calculated according to the equation of Ellis and Roberts (1981): MGT = $\sum (n \times d) / N$, where N = Number of seeds germinated on each day, D =Number of days from the beginning of the test, and N

Priming agen	Wat Soaki i	ter cont ng dura n h (d)	ent ation	Imbi Soakir i	tion rate ng durat n h (d)	e ion	Germ Soak	ination ting hrs	%	Mean ge ti Soa	rminati me king hrs	on	Mean g Soa	erminati aking hrs	on rate
	3	6	Mean	3	6	Mean	3	6	Mean	3	6	Mean	3	6	Mean
Hydro	13.74	13.96	13.85	1.164	1.258	1.211	84	80	82	4.593	4.338	4.045	0.218	0.21	0.214
GA ₃ 20 ppm	14.34	15.05	14.7	1.163	1.344	1.253	85	82	84	4.322	4.343	4.332	0.233	0.23	0.232
GA, 40 ppm	13.88	15.35	14.62	1.159	1.383	1.271	89	86	88	4.112	4.444	4.278	0.246	0.226	0.236
GA, 60 ppm	16.04	14.2	15.12	1.243	1.156	1.2	92	87	90	3.91	4.181	4.466	0.258	0.233	0.245
Mean	12.08	11.23	11.65	0.946	1.028	0.987	75	75	75	3.2	3.2	3.2	0.19	0.19	0.19
С							84	83	83	4.027	4.101	4.264	0.229	0.218	0.223
	D	Т	D*T	D	Т	D*T	D	Т	D*T	D	Т	D*T	D	Т	D*T
SEm	0.096	0.152	0.215	0.008	0.012	0.018	0.501	0.792	1.121	0.023	0.037	0.052	0.001	0	0.003
SEd	0.133	0.21	0.298	0.011	0.017	0.025	0.708	1.121	1.585	0.033	0.052	0.074	0.001	0.003	0.004
Cd(p = 0.05)	0.286	0.452	0.639	0.024	0.037	0.053	1.489	2.355	3.33	0.07	0.111	0.157	0.004	0.006	0.009

Table 2. Effect of seed priming using GA_3 on water content imbibition rate, germination (%) and mean germination time and mean germination rate of *Vigna unguiculata*.

= Total number of seeds germinated at the termination of the experiment . Root length and shoot length test was carried out by glassplate method. Vigor Index was also calculated by Abdul-Baki and Anderson (1973) as Vigor index-I = Germination (%) × Seedling length (cm). Vigor index -II= Germination (%) × Seedling dry weight (g). Data from the experiment underwent analysis using the 'F' test for significance, as outlined by Panse and Sukhatme (1985). In cases where necessary, percentage values were converted to angular (arc-sine) values before analysis. Critical differences (CD) were determined at a 5% probability level. Non-significant results from the 'F' test were

denoted by the abbreviation NS.

RESULTS AND DISCUSSION

Significant differences were observed in all the seed and seedling quality parameters in Cowpea VBN 3 due to the priming agents and priming duration. The interaction effect between priming treatments and durations was also significant for germination characters, seedling length, dry matter production and vigor index. Between the soaking durations of 3 and 6 hrs, three hours of soaking recorded the maximum physiological and morphological parameters of seed

Table 3. Effect of seed priming using GA, on mean daily germination, peak value, CV, CVG and germination index of Vigna unguiculata.

	Mean	Daily Ge tion %	ermina-	Р	eak Val	ue		(CV _t)			(CVG)		Germ	nination	Index
Priming agent	S	oaking h	irs	Se	oaking ł	nrs	S	oaking h	rs	Se	oaking l	nrs	S	oaking	hrs
	3	6	Mean	3	6	Mean	3	6	Mean	3	6	Mean	3	6	Mean
Hydro	10.5	10.167	10.333	12.85	12.45	12.65	29.403	30.261	29.832	21.95	21.5	21.73	5.61	5.32	5.47
GA ₃ 20 ppm	10.667	10.833	10.75	13.87	13.56	13.71	31.318	32.49	31.904	23.3	23.03	23.17	5.48	5.71	5.6
GA ₃ 40 ppm	11	9.345	10.173	14.82	13.75	14.29	32.78	30.883	31.831	24.58	22.56	23.57	5.97	5.71	5.84
GA ₃ 60 ppm	11.5	10.167	10.834	15.07	14.1	14.59	38.359	33.756	36.058	25.85	23.25	24.55	6.22	5.67	5.94
Mean	9.33	9.33	9.33	11.5	11.5	11.5	30.043	30.043	30.043	20.12	20.12	20.12	5.13	5.13	5.13
С	10.599	9.968	10.284	13.62	13.07	13.35	32.381	31.487	31.934	23.16	22.09	22.63	5.68	5.51	5.6
	D	Т	D*T	D	Т	D*T	D	Т	D*T	D	Т	D*T	D	Т	D*T
SEm	0.062	0.098	0.139	0.078	0.124	0.175	0.182	0.288	0.407	0.138	0.218	0.309	0.034	0.053	0.076
SEd	0.088	0.139	0.197	0.111	0.175	0.248	0.257	0.407	0.576	0.195	0.309	0.437	0.048	0.076	0.107
Cd (p = 0.05)	0.185	0.292	0.414	0.233	0.369	0.522	0.542	0.856	1.211	0.411	0.65	0.92	0.101	0.16	0.226

Priming agent	Shoo	t length	(cm)	Root	length	(cm)	Seedl	ing leng (cm)	th	Vigor in Soaking in	ndex I g durati h (d)	on	Vigor in Soaking in	ndex II g durati h (d)	on
	So	aking h	rs	So	aking h	rs	So	aking h	rs						
	3	6	Mean	3	6	Mean	3	6	Mean	3	6	Mean	3	6 N	Mean
Hydro	14.9	14	14.45	10.3	9.8	10.05	26.9	23.8	25.35	2260	1936	2098	73	44	58
GA ₃ 20 ppm	17.1	16.6	16.85	12	10.3	11.15	29.1	26.9	28	2483	2331	2407	65	36	50
GA ₃ 40 ppm	16.6	15.2	15.9	11.4	9.8	10.6	26.3	25	25.65	2314	2267	2291	48	59	53
GA, 60 ppm	18.2	16.6	17.4	13	9.5	11.25	31.2	26.1	28.65	2870	2332	2601	109	80	95
Mean	10.1	10.1	10.1	6.7	6.7	6.7	16.8	16.8	16.8	1254	1254	1254	25	25	25
С	15.38	14.5	14.94	10.68	9.22	9.95	26.06	23.72	24.89	2236	2024	2130	64	49	56
	D	Т	D*T	D	Т	D*T	D	Т	D*T	D	Т	D*T	D	Т	D*T
SEm	0.086	0.137	0.194	0.061	0.096	0.136	0.147	0.233	0.329	13.573	21.461	30.351	0.433	0.685	0.968
SEd	0.12	0.194	0.274	0.086	0.136	0.192	0.208	0.329	0.465	19.195	30.35	42.921	0.612	0.968	1.37
Cd (p = 0.05)	0.258	0.408	0.577	0.181	0.286	0.405	0.437	0.692	0.979	40.328	63.765	5 90.178	1.287	2.035	2.878

Table 4. Effect of seed priming using GA₃ on, shoot length, root length, seedling length, vigor index I and vigor index II of *Vigna unguiculata*.

and seedling quality parameters compared to control.

In the current research the seeds of VBN₃ were primed using GA₃ at 20, 40 and 60 ppm, KNO₃ and ZnSO₄ at three different concentrations viz., 1%, 2% and 3% along with hydropriming. Among all the treatments values for all the germination parameters were higher in the seeds soaked for three hours than for six hours. using 3 concentrations viz., 20, 40 and 60 ppm. The uptake of water by seeds is the first phase in seed germination which is called as imbibition. The principle of seed priming is based on seed imbibition (Ruttanaruangboworn *et al.* 2017). Water content and seed imbibition rate was higher in the seeds primed with GA₃ at 60 ppm concentration for 3 hrs (Table 2). Germination percentage was the highest in GA₃ 60 ppm concentration soaked for 3 hrs. The important impact of GA₃ is the enhancement of seed germination, which has been demonstrated in ornamental plants (Joshi *et al.* 2022). During germination, GA₃ is

Effect of GA₃

In the current research cowpea seeds were primed

Table 5. Effect of seed priming using KNO₃ on water content imbibition rate, germination (%) and mean germination time and mean germination rate of *Vigna unguiculata*.

Priming agent	Wate	er conte	nt	Imb Soaki	ition rat	e tion	Gern	nination	n% Me	ean gern	nination	time	Mean g	erminat	ion rate
i inning again	Douil	in h (d)		in	h (d)		Soa	king h	rs	Soa	king hrs		So	aking hi	rs
	3	6	Mean	3	6	Mean	3	6	Mean	3	6	Mean	3	6	Mean
Hydro	13.89	13.44	13.67	1.01	1.1	1.05	83	81	82	4.2	4.1	4.15	0.21	0.215	0.213
KNO, 1%	14.95	12.92	13.94	1.32	1.13	1.22	88	83	86	4.16	4.14	4.15	0.242	0.244	0.243
KNO ² 2%	15.44	13.81	14.63	1.54	1.07	1.3	89	87	88	4.83	4.32	4.58	0.287	0.241	0.264
KNO, 3%	12.01	14.09	13.05	0.96	1.18	1.07	87	80	84	4.25	4.44	4.35	0.221	0.222	0.221
Mean	11.26	10.85	11.06	0.97	0.89	0.93	79	79	79	3.6	3.6	3.6	0.19	0.19	0.19
С							85	82	84	4.21	4.12	4.16	0.23	0.222	0.226
	D	Т	D*T	D	Т	D*T	D	Т	D*T	D	Т	D*T	D	Т	D*T
SEm SEd Cd (p = 0.05)	0.094 0.13 0.279	0.148 0.206 0.442	0.21 0.291 0.625	0.008 0.011 0.025	0.013 0.018 0.039	0.019 0.026 0.056	0.465 0.657 1.382	0.735 1.04 2.185	1.04 1.471 3.09	0.026 0.036 0.077	0.041 0.058 0.122	0.058 0.082 0.173	0.001 0.002 0.004	$0.002 \\ 0.003 \\ 0.007$	$0.003 \\ 0.004 \\ 0.009$

	Me	an daily	y n %	Pea	k value		(C	V _t)		(0	CVG)	(Germina	tion inc	lex
Priming agent	Sc	aking h	rs	Soa	king hrs	5	Soa	king hr	s	Soa	king hr	5	Soak	ing hrs	
	3	6	Mean	3	6	Mean	3	6	Mean	3	6	Mean	3	6	Mean
Hydro	10.2	10.17	10.18	13.93	12.93	13.43	30.45	30.72	30.59	22.11	22.17	22.14	5.3	5.4	5.35
KNO, 1%	10.5	10.32	10.41	14.04	14.48	14.26	34.5	33.64	34.07	24.18	24.36	24.27	5.7	5.8	5.75
$KNO_3^2 2\%$	11.48	10.53	11.01	14.8	14.27	14.53	39.16	35.05	37.11	26.85	24.11	25.48	6.85	5.95	6.4
KNO ₃ 3%	10.15	10.23	10.19	12.7	12.53	12.62	31.25	32.34	31.79	24.12	22.17	23.14	5.35	5.17	5.26
Mean	9.13	9.13	9.13	11.06	11.06	11.06	26.25	26.25	26.25	18.92	18.92	18.92	4.25	4.25	4.25
С	10.29	10.08	10.18	13.31	13.05	13.18	32.32	31.6	31.96	23.24	22.35	22.79	5.49	5.31	5.4
	D	Т	D*T	D	Т	D*T	D	Т	D*T	D	Т	D*T	D	Т	D*T
SEm	0.065	0.104	0.147	0.081	0.129	0.182	0.206	0.326	0.462	0.135	0.213	0.302	0.035	0.056	0.08
SEd	0.093	0.147	0.208	0.115	0.182	0.258	0.292	0.462	0.653	0.191	0.302	0.427	0.05	0.08	0.113
Cd (p = 0.05)	0.195	0.309	0.437	0.242	0.384	0.543	0.614	0.971	1.373	0.401	0.635	0.898	0.106	0.168	0.238

Table 6. Effect of seed priming using KNO₃ on mean daily germination, peak value, CV, CVG and germination index of Vigna unguiculata.

releazed from the embryo hence stimulating mRNA and production of α -amylase (Blankenship 2021). α -Amylase helps in the hydrolysis of the starch during seed germination. It may also be responsible for the maintenance of requisite water potential, by providing solute sugars during the seed germination phase. The germination rate of 14–27 % was observed in *Leymus chinensis* when treated with GA₃ (Ma *et al.* 2018). Other germination parameters like mean germination time, mean germination rate, mean daily germination percentage, peak value, CV, CVG and germination index were also highest for the seeds primed with GA₃ at 60 ppm concentration for 3 hrs (Table 3). The decrease in germination time and increase in germination rate is observed in *Brassica napus* compared to control when treated with GA₃ (Zhu *et al.* 2021). The seed priming treatment increased the length of seedling than control. The maximum length of shoot, root and seedling was observed in the seeds primed with GA₃ at 60 ppm concentration for 3 hrs (Table 4). GA₃ treatment accelerates germination by secreting hydrolytic enzymes which helps in weakening the seed testa and speed up seed germination by extending the shoots and roots (Pradhan *et al.* 2022). The increased seed length after GA₃ treatment is the result of the activation of dormant embryos to perform cell

Table 7. Effect of seed priming using KNO# on, shoot length, root length, seedling length, vigor index I and vigor index II of Vigna unguiculata.

Priming agent	Shoo Si	t length oaking l	(cm) hrs	Root l Soa	ength (c aking hr	sm)	Seedlin So	ng lengt aking hi	h (cm) rs	Vigor Soakir	r index index ing durat	I ion	Vigo Soaki	r index ng dura in h (d)	II tion
	3	6	Mean	3	6	Mean	3	6	Mean	3	6	Mean	3	6	Mean
Hydro	14.7	14.5	14.6	8.2	7.5	7.9	22.9	22.5	22.7	1901	1823	1862	58	41	50
KNO ₃ 1%	16.6	15.7	16.2	10.3	9.8	10.1	26.9	23.8	25.4	2367	1975	2171	77	45	61
KNO ₃ 2%	16.5	15.5	16	11.1	10.4	10.8	27.6	25.9	26.8	2466	2245	2355	82	69	76
KNO ₃ 3%	16.1	14.2	15.2	9.8	8.6	9.2	25.9	22.8	24.4	2253	1915	2084	72	54	63
Mean	10.1	10.1	10.1	6.7	6.7	6.7	16.8	16.8	16.8	1327	1327	1327	27	27	27
C	14.8	14	14.4	9.2	8.6	8.9	24	22.4	23.2	2063	1857	1960	63	47	55
	D	Т	D*T	D	Т	D*T	D	Т	D*T	D	Т	D*T	D	Т	D*T
SEm	0.091	0.144	0.204	0.058	0.092	0.131	0.149	0.236	0.334	13.444	21.257	30.063	0.429	0.679	0.96057
SEd	0.129	0.204	0.289	0.082	0.131	0.185	0.211	0.334	0.472	19.012	30.062	42.514	0.607	0.96	1.3584
Cd (p = 0.05)	0.272	0.43	0.608	0.174	0.275	0.389	0.444	0.702	0.993	39.946	63.16	89.322	1.276	2.018	2.854

Priming agent	Wa Soak	iter cont ing dur	tent ation	Im Soak	bition rating dur	ate ation	Germ Soal	ination king hr	s [%]	Mean gei Soa	rminatio king hrs	on time	Mean g Soa	germinat king hrs	ion rate
	3	6 f	Mean	3	6	Mean	3	6	Mean	3	6	Mean	3	6	Mean
Hydro ZnSo 1%	12.85	13.1	12.98	0.9	1.1	1	81 85	79 85	80 85	4.92	4.75	4.84	0.25	0.26	0.255
$ZnSo_4 2\%$	15.91	13.25	14.58	1.32	1.15	1.235	89	83	86	3.45	4.51	3.98	0.230	0.220	0.235
$ZnSo_{4}^{4}3\%$	12.04	13.8	12.92	0.95	1.131	1.04	85	83	84	4.57	4.04	4.3	0.222	0.251	0.236
Mean	11.12	10.78	10.95	0.844	0.892	0.868	76	76	76	5.4	5.4	5.4	0.28	0.28	0.28
С							83	81	82	4.51	4.62	4.57	0.24	0.252	0.246
	D	Т	D*T	D	Т	D*T	D	Т	D*T	D	Т	D*T	D	Т	D*T
SEm SEd Cd (p = 0.05)	0.095 0.132 0.284	0.151 0.209 0.449	0.213 0.296 0.635	0.007 0.01 0.023	0.012 0.017 0.036	0.017 0.024 0.052	0.43 0.608 1.277	0.68 0.961 2.02	0.961 1.36 2.857	0.027 0.038 0.08	0.042 0.06 0.126	0.06 0.085 0.179	0.001 0.002 0.004	0.002 0.003 0.006	$0.003 \\ 0.004 \\ 0.009$

Table 8. Effect of seed priming using $ZnSO_4$ on water content imbibition rate, germination (%), mean germination time and mean germination rate of *Vigna unguiculata*.

division, elongation, and multiplication (Nedunchezhiyan *et al.* 2023). The proportion of viable seed in a sample is determined by seed vigour testing, which also evaluates the seed's capacity to develop healthy seedlings in unfavorable or subpar growth circumstances that are comparable to those that would arise in the field. Seed vigour test is essential for seeds stored under unfavourable conditions or seeds sourced from unauthorized sources. In the present study seed vigour index I and II was estimated. The results revealed that the seeds primed with GA₃ at 60 ppm for 3 hrs had the highest seed vigour. Okra seeds primed with GA₃ resulted in seed viability of 74.59% (Bereded Sheferie 2023). GA₃ treatment promotes cell division by promoting DNA replication, repair seed damage and increases ATP availability which all inturn increases seed viability (Junhaeng *et al.* 2015). A study done by Arun, *et al.* (2016) in cowpea shows that priming with GA₃ 100 ppm reduced the time to 50% germination and mean germination time and increased the germination percentage in low vigour seeds of cowpea.

Effect of KNO,

The role of potassium nitrate as seed priming agent to

Table 9. Standardization of bio priming using $ZnSO_4$ on mean daily germination, peak value, CV, CVG and germination index of *Vigna unguiculata*.

	Me	an daily	y %	Ре	eak valu	e	(0	CVt)		(C	CVG)		Germin	nation i	ndex
Priming agent	Soa	aking hi	ſS	So	oaking h	rs	Soal	king hrs		Soal	king hrs		Soa	aking h	rs
	3	6	Mean	3	6	Mean	3	6	Mean	3	6	Mean	3	6	Mean
Hvdro	8.45	8.74	8.6	12.34	12.93	12.64	32.78	27.88	30.33	21.3	20.03	20.67	5.19	5.21	5.2
ZnSo, 1%	10.83	9.24	10.04	13.77	11.6	12.68	36.36	29.1	32.73	23.78	22.83	23.31	5.91	5.31	5.61
$ZnSo_{4}^{4}2\%$	11	10.33	10.67	14.8	13.2	14	39.26	34.1	36.68	25.85	22.34	24.1	6.54	5.32	5.93
$ZnSo_{4}^{3}$ 3%	9.76	9.86	9.81	13.65	14.28	13.97	27.91	30.48	29.2	22.16	25.13	23.65	5.27	5.78	5.52
Mean	7.45	7.45	7.45	10.05	10.05	10.05	25.49	25.49	25.49	19.24	19.24	19.24	4.65	4.65	4.65
С	9.5	9.12	9.31	12.92	12.41	12.67	32.36	29.41	30.89	22.47	21.91	22.19	5.51	5.25	5.38
	D	Т	D*T	D	Т	D*T	D	Т	D*T	D	Т	D*T	D	Т	D*T
SEm	0.059	0.093	0.132	0.084	0.133	0.188	0.202	0.32	0.453	0.146	0.231	0.327	0.035	0.056	0.079
SEd	0.083	0.132	0.186	0.119	0.188	0.266	0.286	0.453	0.641	0.207	0.327	0.463	0.05	0.07	0.112
Cd (p = 0.05)	0.175	0.277	0.392	0.25	0.395	0.559	0.602	0.953	1.347	0.435	0.688	0.973	0.105	0.167	0.236

Priming agent	Shoot Soal	length (king hrs	cm)	Root l Soa	ength (c aking hr	m) s	Seedlin Soa	ng lengt aking hr	h (cm) rs	Vigor Soakin	index I g durati h (d)	on	Vig Soal	or index ting dura	II ation
	3	6	Mean	3	6	Mean	3	6	Mean	3	6	Mean	3	6	Mean
Hydro ZnSo_1%	15.5 16.1	14 14 8	14.8	8.5 9.7	8.1 8.8	8.3 9.3	24 25 8	22.1	23.05 24 7	1950 2205	1756	1853 2104	32 74	43 49	38 62
$ZnSo_4 2\%$	16.6	15.4	16	10.3	9.2	9.8	26.9	24.6	25.75	2406	2002	2220	108	72	90
$ZnSo_{4}^{4}3\%$	14.2	13.1	13.7	9.4	9.3	9.4	23.6	22.4	23	1995	1865	1930	66	46	56
Mean	10.1	10.1	10.1	6.7	6.7	6.7	16.8	16.8	16.8	1277	1277	1277	26	26	26
С	14.5	13.5	14	8.9	8.4	8.7	23.42	21.9	22.66	1967	1787	1877	61	47	54
	D	Т	D*T	D	Т	D*T	D	Т	D*T	D	Т	D*T	D	Т	D*T
SEm	0.089	0.141	0.199	0.054	0.086	0.122	0.144	0.227	0.32	12.817	20.265	28.66	0.522	0.825	1.168
SEd Cd (p = 0.05)	0.126 0.265	0.199 0.419	0.282 0.593	0.077 0.162	0.122 0.257	0.173 0.364	0.203 0.427	0.322 0.676	0.455 0.956	18.125 38.082	28.659 60.212	40.53 85.154	0.738 1.552	1.168 2.454	1.651 3.47

Table 10. Effect of seed priming using $ZnSO_4$ on, shoot length, root length, seedling length, vigor index I and vigor index II of *Vigna unguiculata*.

promote has been known for long time. In the current investigation KNO3 was used in three concentrations viz., 1%, 2%, 3% as a priming agent with no treatment as control. The imbibition rate of the seeds primed with 2% KNO, was the highest compared with the other concentrations studied (Table 5). The imbibition time was lower when the concentration was higher and soaking duration was long. This was in agreement with the results of study by Ruttanaruangboworn et al. (2017) in rice where imbibition rate was lower in higher concentration of KNO₂. The germination characters like germination percentage, mean germination time, mean germination rate, mean daily germination percentage and peak value of germination were highest when the seeds were primed with KNO₂ at 2% for 3 hrs. At 3% KNO₂ treatment the germination was the lowest (Table 6). Higher concentrations exert decreasing effects on seed germination by causing death of cells and ultimately result in loss of seed viability. These were against the findings of Ruttanaruangboworn et al. (2017) where the germination in rice was higher in the seeds primed with 1% KNO₃ than 2 % KNO₃ in rice. This suggests that there is toxic effect of KNO, due to ion accumulation in the embryo (Song et al. 2007). In accordance with our results, priming of Paspalum vaginatum seeds with KNO, had significant effects on germination. During the process of halo-priming with potassium nitrate (KNO₂), ions derived from potassium nitrate solutions gather within the seeds,

consequently lowering water potential and enhancing water absorption Shim et al. (2008). The availability of nitrate during imbibition provides additional sources for protein synthesis to enhance germination while seed priming (Basra et al. 2005). The coefficient of germination and germination index were highest for the treatment with KNO₂ 2 % for 3 hrs. 0.1% KNO₂ was regarded as the best treatment for parameters like CVG and germination index in Apple. Similar results was also observed in crops like Citrullus colocynthis (Saberi et al. 2011), Foeniculum vulgare, Cuscuta epithymum (Tavili et al. 2010). The decrease in ABA sensitivity of imbibed seeds may due to the effect of KNO₂ (Bethke et al. 2006). The highest shoot length, root length and seedling length was observed in the treatment of KNO₂ at 2%. Maximum seedling length was achieved in tomato seed primed with 0.75% KNO₂ (Moaaz Ali et al. 2020). Seedling vigour is the combined result of the emerged seeds under a wide range of biotic and abiotic stresses. Seedling vigour is a combined measure of seedling length, seedling fresh weight, and seedling dry weight (ISTA 2015). Seed vigor I and II were highest in the treatment of KNO₂ at 2% for 3 hrs. Similar results were found in corn when the seeds were primed done with 1% KNO₂ (Hadinezhad et al. 2014) (Table 7). Many studies revealed the effect of KNO, viz., increase in germination potential and seedling vigour in tomato at 50 mmol KNO₂ (Ebrahimi et al. 2014), increase in seedling vigor in wheat (Shu et al. 2016).

Priming agent	Germi Soal	nation val king hrs	ue	Priming agent	Germii Soa	nation val aking hrs	ue	Priming agent	Germi So	nation val aking hrs	ue
	3	6	Mean	-	3	6	Mean	-	3	6	Mean
Hydro GA ₃ 20 ppm GA ₃ 40 ppm GA ₃ 60 ppm C Mean	142.33 152.33 162.66 165.33 112.38 147.01	138.75 155.75 152.45 161.66 112.38 144.2	140.54 154.04 157.56 163.5 112.38 145.6	Hydro KNO ₃ 1% KNO ₃ 2% KNO ₃ 3% C Mean	129.31 154.53 156.93 147.3 112.38 140.09	129.73 155.43 154.93 131.33 112.38 136.76	129.52 154.98 155.93 139.32 112.38 138.43	Hydro $ZnSo_4 1\%$ $ZnSo_4 2\%$ $ZnSo_4 3\%$ C Mean	149.82 150.05 154.43 150.8 112.38 143.5	131.33 145.33 148.23 153.47 112.38 138.15	140.58 147.69 151.33 152.13 112.38 140.82
	D	Т	D*T		D	Т	D*T		D	Т	D*T
SEm SEd Cd (p=0.05)	0.88 1.245 2.616	1.392 1.968 4.136	1.969 2.784 5.85	SEm SEd Cd (p=0.05)	0.872 1.233 2.591	1.379 1.95 4.098	1.95 2.758 5.795	SEm SEd Cd (p=0.05)	0.906 1.282 2.693	1.433 2.027 4.258	2.027 2.866 6.022

Table 11. Effect of seed priming using GA₃, KNO₃, ZnSO₄ on germination value of *Vigna unguiculata*.

Effect of ZnSO₄

Zinc is plays an important role in germination as it is an important component of various enzymes (Shazia Anwer Bukhari et al. 2021). In the current investigation seeds of cowpea were primed with ZnSO₄ solution at 1, 2, 3% for 3 and 6 hrs. The water content and imbibition rate were the highest for the primed seeds 2% concentration for 3 hrs (Table 8). The germination parameters like germination percentage, mean germination time, mean germination rate, mean daily germination and peak value of germination were maximum for the seeds primed at 2% concentration for 3 hrs (Table 9). Seed priming increased the germination rate resulting which in turns aids in germination (Sajjan et al. 2017). This is due to the faster conversion of substances into energy by various physical and chemical process. Priming of seeds with $ZnSO_4$ either at 0.025 or 0.05% shown higher germination percentage and germination index than control (Raj et al. 2019). This result was against result of the present study in which germination index was the higher at the treatment ZnSO₄ at 2% than at 1%. Seed priming with 0.05 mM of ZnSO₄ recorded higher values for germination %, mean germination time (MGT), mean germination rate (MGR), coefficient of velocity of germination (CVG), germination rate index (GRI) (% / day), germination index (GI), shoot, root and seedling length in wheat (Singhal and Bose 2020). In the current investigation, seed priming with ZnSO₄ at 2% recorded the maximum values of growth parameters like shoot length, root length and seedling length, coefficient of velocity of germination and seed vigor. Priming the seeds with $ZnSo_4$ resulted in increased root and shoot length compared to unprimed seeds (Farooq *et al.* 2005) (Table 10).

Effect of hydropriming

The hydroprimed seeds performed better than unprimed seeds for each and every character studied. Many studies have reported that hydropriming of seeds improved the seed germination and seedling emergence of rice (Basra *et al.* 2005). Hydropriming of seeds of *Pisum sativum* for 12 hrs had a significant on its germination characters (Singh *et al.* 2017). (Table 11).

From the present study it can be clearly known that priming seeds with GA_3 at 60 ppm, KNO_3 at 2%, $ZnSO_4$ at 2% increases the seed germination parameters which in turn increases the yield.

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