



Chemical-Induced Dormancy Breaking of Freshly Harvested Potato Minitubers and Its Effect on Subsequent Growth and Yield

Zishan Gul ^{a++*} and Nayyar Iqbal ^a

^a Hazara Agriculture Research Station, Abbottabad, Pakistan.

Authors' contributions

This work was carried out in collaboration between both authors. Both authors read and approved the final manuscript.

Article Information

DOI: 10.9734/AJAAR/2023/v23i4474

Open Peer Review History:

This journal follows the Advanced Open Peer Review policy. Identity of the Reviewers, Editor(s) and additional Reviewers, peer review comments, different versions of the manuscript, comments of the editors, etc are available here: <https://www.sdiarticle5.com/review-history/109232>

Original Research Article

Received: 16/09/2023

Accepted: 22/11/2023

Published: 29/11/2023

ABSTRACT

Tuber dormancy is an important phenomenon in potato seed production cycle and its duration can positively affect the number of crops and crop performance during the seed production. The present study was conducted to evaluate the effect of different factors which include potato genotypes (Roko and SM kaghan varieties), chemical treatments i.e GA₃, thiourea, Thiourea+GA₃ and sugar under light and dark storage conditions. It was found that between varieties, Roko responded better than SM kaghan to various chemical treatments regarding number of days to dormancy breaking. It was found that the growth hormone GA₃ (gibberellic acid) in combination with thiourea can significantly increase sprouting percentage to 100% under dark conditions after 31 days of application. However, it was observed that after planting of the treated tubers the emergence percentage, and minituber yield did not show any significant effect of the three

⁺⁺ Senior Research Officer;

^{*}Corresponding author: Email: gul.zishan@gmail.com;

experimental treatments in comparison with control. It is therefore concluded that tuber treatment with thiourea+GA₃ and store in dark can decrease the dormancy duration without having any negative effect on plant physiological growth and tuber yield.

Keywords: Seed-potato; tuber treatment; thiourea; GA₃; sugar; sprouting%.

1. INTRODUCTION

Potato (*Solanum tuberosum* L.) is considered as an important food crop worldwide and a good source of energy, fats, proteins, minerals and vitamins [1,2]. Among the root and tuber crops, it ranks first followed by cassava, sweet potatoes and yams [3,4]. There are a number of constraints that negatively affect potato production in the world which include many biotic and abiotic stresses among which soil and tuber borne diseases and insect pests are major biotic problems while heat, salinity, cold/frost stress, nutrient deficiency, drought are the key abiotic factors [5] and among them lack of quality seed potato has vital significance [6]. In Pakistan, it is also an important vegetable and is grown on 0.19 million hectares, with an estimated annual production of 4.6 million tons, at an average yield of 24.2 tons per hectare [7]. Potato is grown round the year in Pakistan as autumn, spring and summer crop. The year round potato cropping in Pakistan shortens the interval between harvesting and planting consecutive crop and the growers usually adapted the autumn to autumn cycle by storing the seed of autumn crop of medium to long dormancy period genotypes for the coming autumn crop which declines the seed quality [8,9].

Dormancy is the final stage of tuber life serving to preserve tubers as organs of vegetative reproduction under unfavorable growth conditions. Endogenous hormones have been proposed to play a significant role in tuber dormancy regulation. Generally dormancy is divided into three categories. During deep dormancy (endodormancy) growth is stopped under the influence of internal physiological factors even under ideal conditions. The second category of dormancy i.e paradormancy is forced or induced dormancy in which growth is arrested by unfavorable external factors. The third stage of dormancy is ecodormancy during which meristematic activity is halted by external environmental factors. Immediately after harvest tubers are in the state of deep dormancy and can not sprout even under favorable environmental conditions [10].

After the termination of the deep dormancy tubers can sprout but they remain at rest under unfavorable external conditions (at temperature 0 to 4°C). In the beginning of sprouting, lateral buds are in the state of forced dormancy (depending on internal physiological factors) because they experience a correlative inhibition from the apical bud sprouting earlier [11,12].

The tuber dormancy duration and sprouting time have significant economic importance in seed production cycle. The tuber dormancy period depends on potato cultivar characteristics (genotype) and also on growing season or pre-harvest and post-harvest conditions such as temperature and light. Depending on the genotype tuber storage at high temperature (up to 30°C) and humidity (up to 90%) favors dormancy breaking and early bud outgrowth and temperature above 35°C can immediately break dormancy inducing a condition called psychopathy or heat sprouting [11].

Considering the storability of potato the term dormancy is significantly important. The tuber dormancy duration gives information on how long the potato can be stored. It helps in selecting varieties for short to long term storage. Regarding process and fresh market potatoes, detrimental quality concerns raised if sprouting begins which causes changes in carbohydrate status, weight loss, increase in respiration rate and storage management issues such as impeded airflow. Therefore controlling the length of the dormancy period is of considerable economic importance especially in case of seed potato. New strategies are needed because current cold storage techniques are often problematic. Seed growers may need to accelerate or retard sprout development depending upon the time of year and intended seed market. When working with seed production programs it is essential to standardized different chemicals and their concentrations needed for dormancy breaking for routine applications because different types of genetic materials (varieties) react differently to various chemicals that promote sprouting. Usually late maturing clones have a long dormancy that is difficult to break than that of early maturing clones [13].

Early potato planting requires dormancy breaking. Various methods for chemical regulation of dormancy period has been developed which includes tuber treatment with Rindite [14], bromoethane [15], carbon disulfide [16], sugar [17], thiourea [18], GA₃ [19], ethanol [20], or bromoethane favors dormancy breaking [15]. Moreover, cytokinins and GA₃ (gibberellins) reactivate meristematic and sprouting activity shortly before dormancy termination [21,22]. The response of potato genotypes to different chemical treatments and storage methods may also be differed due to genetic factors, type of chemicals and storage conditions [23].

In seed production programs it is indispensable to standardize different protocols and chemicals used for dormancy breaking. The common way of producing pre-basic seed of potato is the production of mini-tubers in the greenhouse from invitro plantlets produced through tissue culture. Generally for certified seed production cycle the minitubers harvested during the month of May have to be planted usually in the first week of June in Hilly areas suitable for seed production. So there is not enough time between harvest and planting to break dormancy naturally. Therefore, it is anticipated to develop an effective technique easy to apply in order to induce seed-tuber dormancy breakage in different potato varieties. The aim of this study is to develop an effective protocol in order to analyze the performance of various chemical treatments in interaction to certain environmental factor (light;dark) and plant genotypes to find a potential relation between treatments applied in the storage phase and to record the dormancy breakage duration of each variety and to evaluate potato crop performance after plantation.

2. MATERIALS AND METHODS

This study was conducted in the greenhouse at Hazara Agriculture Research Station, Abbottabad. The research experiment was carried out in two phases: In the first phase (at storage level minitubers were treated with different chemicals while in the second phase the same treated minitubers were planted in the green house and evaluated for plant growth and yield performance.

This research is based on 3 factorial randomized complete design. Experimental factors were chemical treatments as factor one with five levels (control, 50ppm gibberellic acid (GA₃), 1% thiourea, thiourea +GA₃ (1% thiourea and 50ppm

GA₃) and Sugar treatment (100g sugar in 200ml water), potato genotypes as the second factor with two levels (SM Kaghan and Roko) and storage condition as the third factor with two levels (light, dark conditions). Freshly harvested 100 minitubers of uniform shape and size of each variety was selected. The weight of each selected tuber was between 30-40 gram.

The tubers were soaked in the 1% Thiourea solution for 1 hour and air-dried. For GA₃ treatment tubers were soaked in 50ppm solution for 30 minutes and air dried. For sugar treatment 1:2 ratio sugar solution (dissolving 100g of table sugar in 200ml of water) was prepared according to the procedure described by Zaghum et al., 2021 and tubers were sprayed thoroughly with this solution and air-dried. For thiourea+GA₃ treatment first tubers were soaked in thiourea (1%) solution for 1 hour then transferred to 50ppm GA₃ solution and soaked for 30 minutes, removed and air dried while for control the tuber were treated with distilled water. After each treatment application the treated minitubers were placed in plastic trays and kept in the dark and light conditions at 25±2°C and 80±5% RH. Data were recorded on sprouting (%), number of sprouts per tuber and any tuber rotting (%).

In the second phase of the experiment the same treated tubers that showed sprouting were sown in earthen pots (18x20cm) filled with 4 kg of sterilized peat moss soil and data on plant growth in terms of plant height, number of branches, average tuber number and tuber yield per plant were recorded. The recorded data were analyzed by Analysis of Variance (ANOVA) for the F-test and Least significance difference test (LSD) at 95% level of significance was performed to assess significance of results. All the statistical analyses were done using computer software Statistix 8 Version 8.1 [24].

3. RESULTS

Potato minitubers of both varieties i.e SM Kaghan and Roko for each respective treatment were observed regularly to ensure the emergence of any sprout. Only the sprout that reached a length of about 2mm was counted as the first sprout from that particular tuber. Data pertaining to sprouting percentage of each respective treatment interaction showed that after 22 days of treatments application only 10% sprouting was first observed in potato variety Roko minitubers placed in Dark and treated with Thiourea + GA₃ (Fig. 1). Any other treatment

interaction has not shown sprouting in both varieties.

After 25 days figure 2 showed 10% sprouting in Roko variety treated with thiourea under dark conditions followed by 20% sprouting in GA₃ treatment . 30% sprouting was recorded in Thiourea + GA₃ treatment in dark environment while only 10% tubers showed sprouting in the same treatment but under light conditions in Roko minitubers. Whereas variety SM Kaghan showed 10% sprouting in GA₃ treatment under light conditions (Fig. 2). In the sugar treatment and control no sprouting was observed.

The analyzed data for 28 days after treatments application showed a significant interaction effect of experimental factors on sprouting percentage and considerable increase in sprouting percentage was observed in both varieties for Thiourea + GA₃ treatment under dark conditions in which 70% sprouting occurred in Roko followed by 50% in SM Kaghan (Fig. 3). In GA₃ treatment under light conditions both varieties showed sprouting with 30% and 10% in SM Kaghan and Roko, respectively. However, no sprouting was recorded in sugar treatment and in control.

It was observed that Thiourea along with GA₃ significantly affected the tubers dormancy as compared to other treatment combinations after 31 days and 100 % tubers were sprouted in Roko and 80% in SM Kaghan under dark conditions (Fig. 4). Second highest sprouting percentage (80%) was recorded in GA₃

treatment in dark for potato variety Roko followed by 60% sprouting in thiourea+GA₃ treatment under light conditions (Fig. 4). Moreover, breaking of dormancy in SM Kaghan variety was also observed in sugar treatment and 10% tubers showed sprouting under dark conditions in this interaction. No sprouting was observed in both varieties for control (Fig. 4).

Analyzed data presented in Figure 5 revealed a significant increase in sprouting (100%) in GA₃ treatment alone and in thiourea+GA₃ treatment in dark environment in Roko variety after 34 days. However SM kaghan minitubers showed 90% sprouting in thiourea+GA₃. The lowest sprouting i.e 20% was recorded in sugar treatment under dark conditions and in thiourea treatment in light conditions in SM kaghan (Fig. 5). It was observed that tubers treated with sugar treatment did not showed good response in both varieties under dark and light conditions even after 34 days of treatment application. No sprouting was observed in control for both varieties during this period (Fig. 5).

By comparing the varietal response to the chemical treatments applied it was found that minitubers of potato variety Roko showed comparatively better sprouting % whereas the dormancy breakage and the respective sprouting percentage in potato variety SM Kaghan was comparatively low in all the interactions except in sugar treatment in which only SM Kaghan showed 20% tuber sprouting only under dark conditions (Fig. 5).

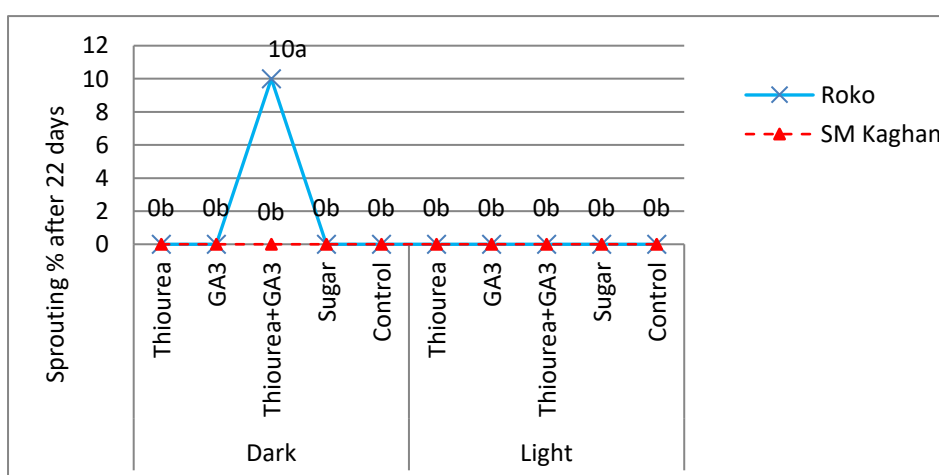


Fig. 1. Mean comparison of sprouting % in potato minitubers under interactive effect of genotype (variety) x chemical treatments x storage condition after 22 days of treatment application

Mean values accompanied by different letters are significantly different at $P \leq 0.05$

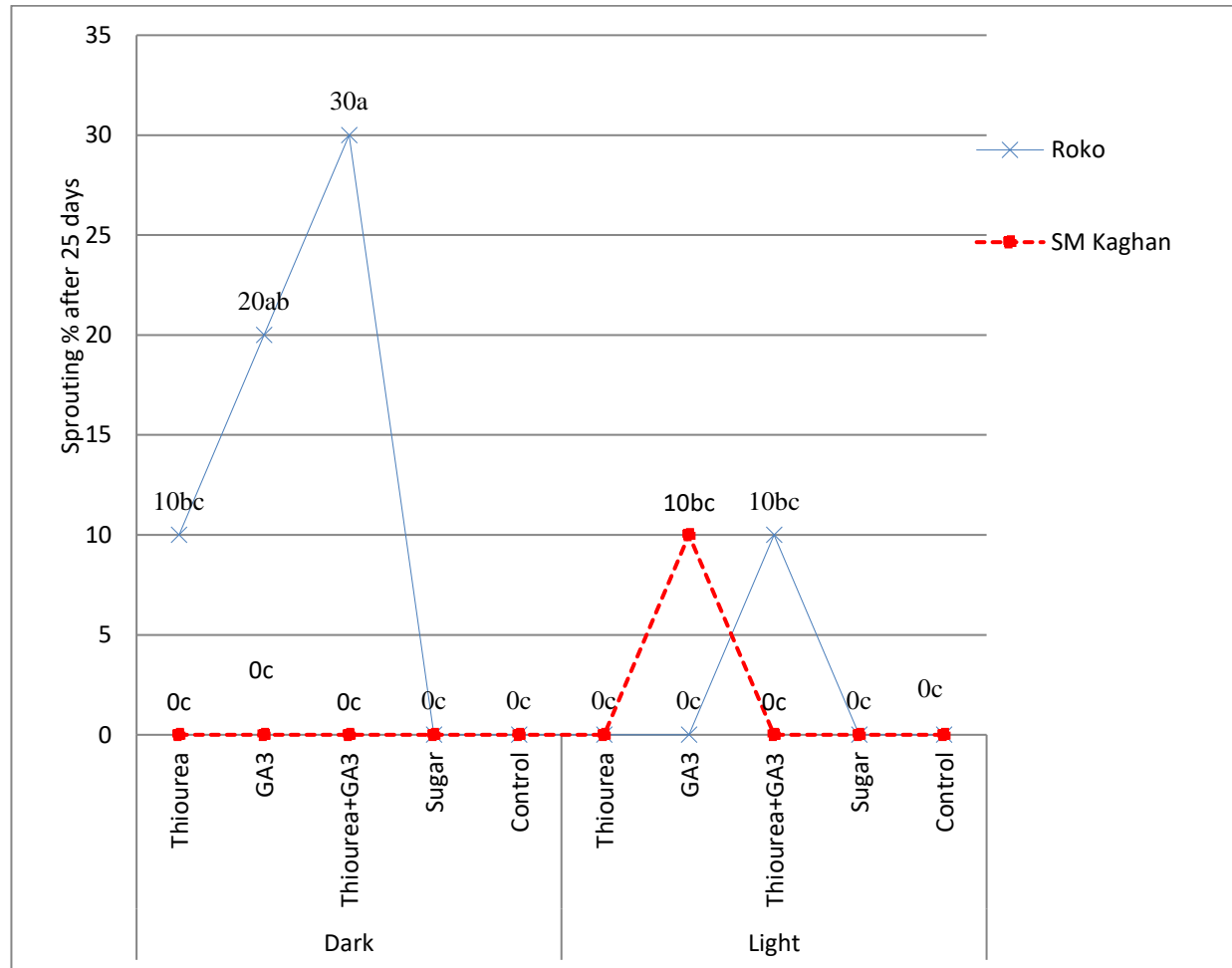


Fig. 2. Mean sprouting % in potato minitubers under interactive effect of genotype (variety) x chemical treatments x storage condition after 25 days of treatment application

Mean values followed by different letters are significantly different at $P \leq 0.05$

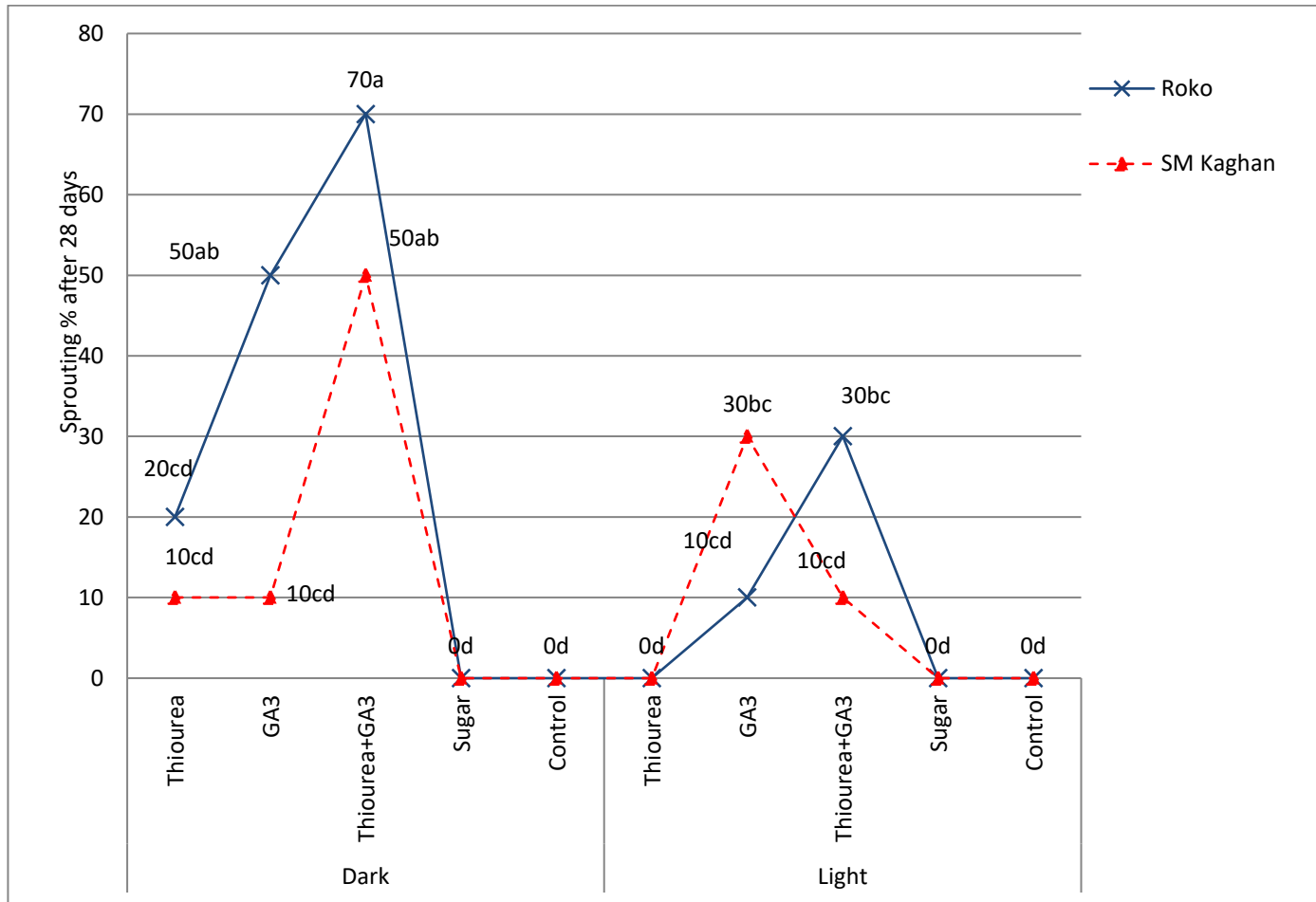


Fig. 3. Mean sprouting % in potato minitubers under interactive effect of genotype (variety) x chemical treatments x storage condition after 28 days of treatment application

Means followed by different letters are significantly different at $P \leq 0.05$

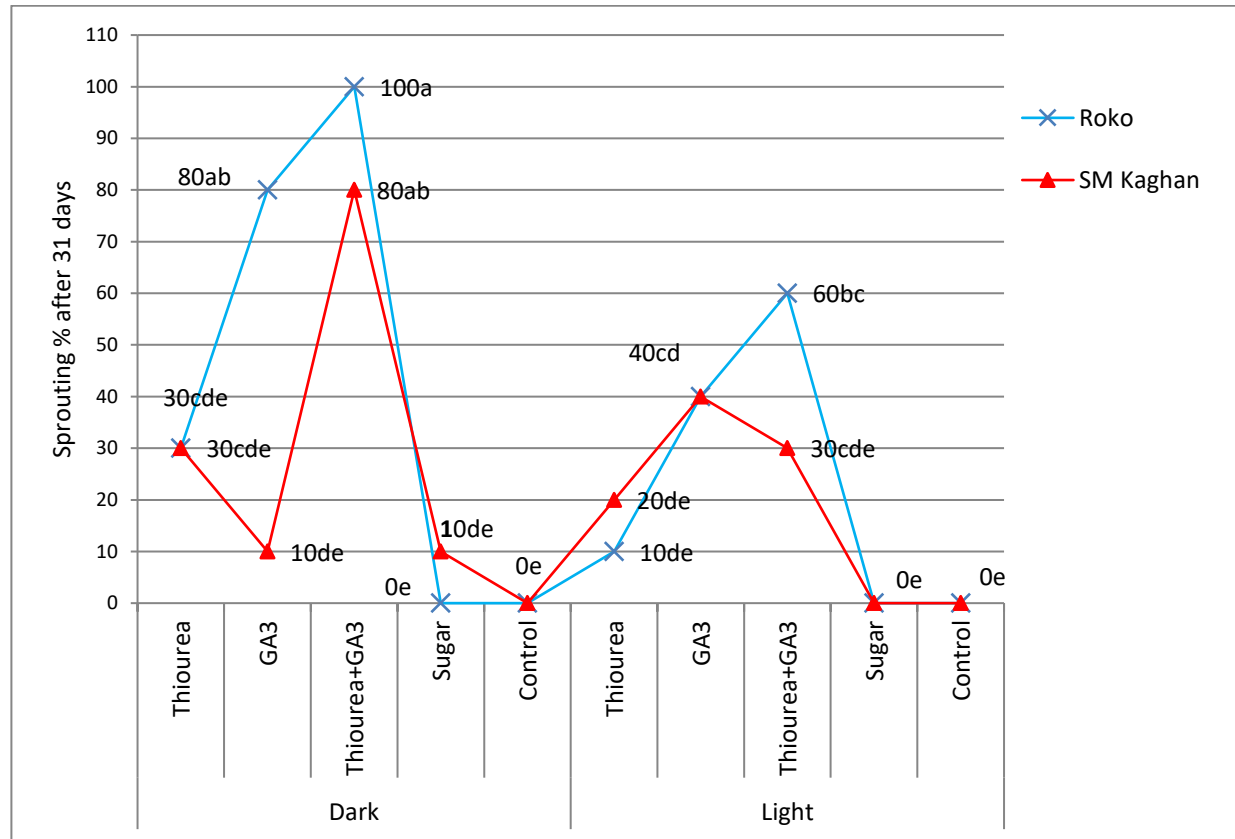


Fig. 4. Mean sprouting % in potato minitubers under interactive effect of genotype (variety) x chemical treatments x storage condition after 31 days of treatment application

Means followed by different letters are significantly different at $P \leq 0.05$

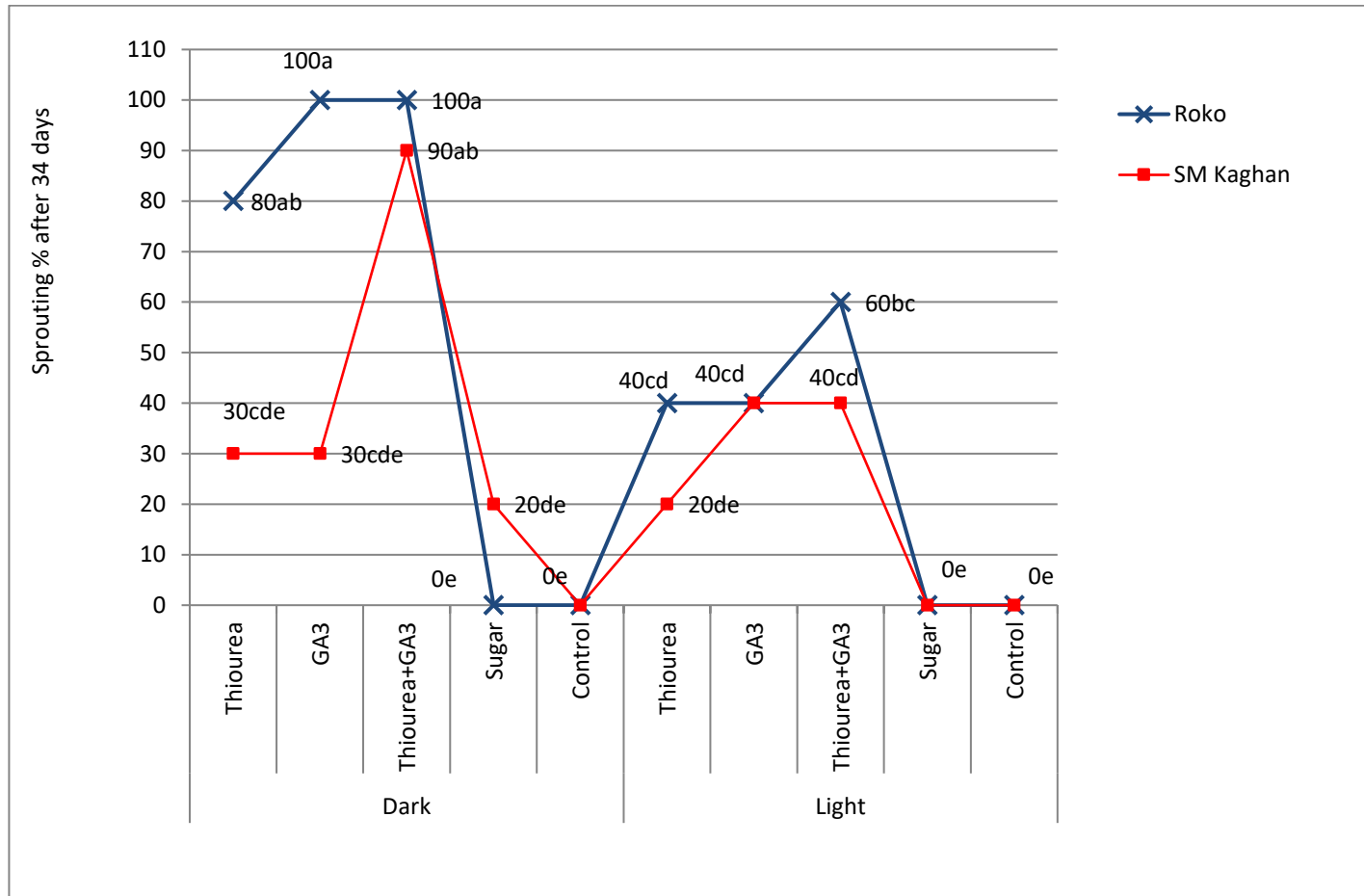


Fig. 5. Mean sprouting % in potato minitubers under interactive effect of genotype (variety) x chemical treatments x storage condition after 34 days of treatment application
Means followed by different letters are significantly different at $P \leq 0.05$

Table 1. Mean number of sprouts per minituber under interactive effect of genotype (variety) x chemical treatments x storage condition after 34 days of treatment application

Variety	Treatments									
	Thiourea		GA ₃		Thiourea + GA ₃		Sugar		Control	
	Dark	Light	Dark	Light	Dark	Light	Dark	Light	Dark	Light
Roko	1.83abcd	1.17def	2.30ab	1.17def	2.50a	2.00abc	0g	0g	0g	0g
SM Kaghan	1.00ef	1.50cde	1.50cde	1.67bcde	1.50cde	1.00ef	1.00ef	0g	0g	0g

Means followed by different letters are significantly different at P≤0.05

Table 2. Mean emergence % in potato minitubers under interactive effect of genotype (variety) x chemical treatments x storage condition after 10 and 15 days of plantation

No. of days	Variety	Treatments									
		Thiourea		GA ₃		Thiourea + GA ₃		Sugar		Control	
		Dark	Light	Dark	Light	Dark	Light	Dark	Light	Dark	Light
10 days	Roko	40b	10cd	30bc	10cd	30bc	10cd	0d	0d	50ab	25bcd
	SM Kaghan	0d	0d	0d	0d	10cd	0d	0d	0d	50ab	75a
15 days	Roko	100a	100a	100a	100a	100a	100a	0c	0c	100a	100a
	SM Kaghan	100a	100a	100a	100a	100a	100a	100a	0c	75b	100a

Means followed by different letters are significantly different at P≤0.05

The results in Table 1 showed that the application of growth regulators on minitubers of different varieties have a significant effect on the total number of sprouts. The highest mean number of sprouts i.e 2.50 were found in roko minitubers in thiourea+GA₃ treatment followed by 2.30 mean sprouts number in GA₃ treatment alone under dark conditions (Table 1). The least mean number of sprouts (1.00) was recorded in SM kaghan variety minitubers in thiourea and sugar treatment under dark and in thiourea+GA₃ treatment under light conditions (Table 1).

In the current research project after dormancy breaking of minitubers of both potato varieties, the sprouted tubers were planted in pots in the green house to study the interactive effect of various experimental treatments (factors) including an untreated sprouted control on the plant growth and yield. The tubers of Roko and SM kaghan in sugar treatment under light conditions were not sprouted (Fig. 5) and were not planted however, the SM Kaghan tubers of sugar treatment under dark conditions showed 20% sprouting (Fig. 5) and were also planted. It was observed that after 10 days of plantation in both varieties the control showed higher emergence percentage (Table 2). In potato variety roko 40% emergence occurred in Thiourea treatment under dark conditions whereas 10% emergence in SM kaghan minitubers occurred in thiourea+GA₃ treatment under dark conditions (Table 2).

Data recorded for percent emergence after 15 days of plantation showed a considerable increase in emergence % i.e 100% in both varieties including control (Table 2) except 75% emergence observed in minitubers of SM kaghan control under dark conditions. The SM Kaghan tubers in sugar treatment under dark conditions also showed 100% emergence after 15 days of plantation (Table 2).

Statistical analysis of the data after 30 days of emergence showed no significant difference among different treatment interactions including control in potato variety Roko. However, Thiourea+GA₃ dark conditions treatment showed highest mean shoot length (19.67 cm). The lowest mean value (15.13 cm) was observed in Thiourea treatment with light conditions (Table 3). In SM Kaghan variety the highest plant height (20.47cm) was recorded in Thiourea+GA₃ treatment interacted with dark conditions followed by 19.90cm in the same treatment but under light conditions (Table 3). In the sugar

treatment significantly lowest plant height (8.40cm) was observed in SM kaghan variety plants developed from minitubers placed in dark conditions whereas no data was recorded in the sugar treatment of roko variety as no tubers were found sprouted and were not sown (Fig. 5; Table 2). In the control no significant difference was found in plant height between the two varieties however plants developed from tubers of light condition have comparatively higher shoot length after 30 days (Table 3).

The interaction effect among varieties x chemical treatment x storage conditions on plant height after 60 days of emergence showed that the potato variety Roko showed highest plant height in thiourea+GA₃ followed by GA₃ treatment (46.91cm) and (43.42cm), respectively under dark storage conditions (Table 3), whereas in SM kaghan the thiourea+GA₃ treatment of light conditions has highest shoot length i.e 38.58 cm followed by 36.65 cm in GA₃ with dark conditions. The lowest mean plant height (29.65cm) was recorded in sugar treatment dark conditions of SM kaghan variety (Table 3).

The data collected on mean number of branches per plant after 30 days (Table 3) showed that no significant difference found in mean number of branches of variety Roko in all the interactions including control, however the thiourea+GA₃ treatment in both light and dark conditions showed highest mean number of branches i.e 8.80 and 8.60 followed by 7.60 and 7.30 in GA₃ treatment under dark and light conditions respectively (Table 3). In the potato variety SM kaghan highest mean branch number (8.75; 8.42) was recorded in GA₃ and thiourea+GA₃ treatments with dark conditions, respectively. The least number of branches (4.00) were recorded in sugar treatment (Table 3). An increase in mean number of branches of both varieties was observed after 60 days of emergence in all the treatment interactions in both potato varieties (Table 3).

Data presented in Table 4 showed that highest mean tuber number i.e 5.16 was recorded from thiourea+GA₃ treatment of SM kaghan variety under light conditions followed by 5.00 and 4.83 mean tuber number in control and in GA₃ interacted with light conditions, respectively of the same variety SM Kaghan. Whereas, in variety Roko highest mean tuber number (4.25) was obtained in control under light conditions followed by GA₃ under dark condition. The lower mean tuber number i.e 3.00 was recorded in

Table 3. Mean plant growth parameters under interactive effect of genotype (variety) x chemical treatments x storage condition after 30 and 60 days of emergence

No. of days	Varieties	Treatments										
		Mean Growth parameter	Thiourea		GA ₃		Thiourea + GA ₃		Sugar		Control	
			Dark	Light	Dark	Light	Dark	Light	Dark	Light	Dark	Light
30 days	Roko	Shoot length (cm)	15.51 def	15.13 ef	18.00 abcde	19.16 abcd	19.67 abc	18.03 abcde	0h	0h	17.22 abcde	17.48 abcde
		No. of branches	5.80abcd	6.65abcd	7.60abc	7.30abcd	8.60a	8.80a	0e	0e	5.60abcd	4.75cd
	SM Kaghan	Shoot length (cm)	12.57 f	18.00 abcde	19.40 abc	17.13 abcde	20.47a	19.90 ab	8.40g	0h	16.05 cdef	16.16 bcdef
		No. of branches	5.00cd	6.50abcd	8.75a	5.15bcd	8.42ab	7.30abcd	4.00d	0e	5.50abcd	5.00cd
60 days	Roko	Shoot length (cm)	42.23ab	39.25 abcde	43.42ab	39.70abcde	46.91a	41.26abcd	0g	0g	41.77abc	39.56abcde
		No. of branches	8.95bcde	8.65bcde	10.50abc	10.50abc	11.50ab	12.50a	0f	0f	11.37ab	9.95abcd
	SM Kaghan	Shoot length (cm)	33.97 cdef	35.30 bcdef	36.65 bcdef	32.06 ef	33.38 def	38.58 bcde	29.65f	0g	36.56 bcdef	33.85 cdef
		No. of branches	7.00e	8.00cde	10.20abcd	9.00bcde	10.65abc	9.35 bcde	7.60de	0f	9.81abcde	8.40cde

Means followed by different letters are significantly different at $P \leq 0.05$

individual Thiourea and GA₃ treatments under light condition (Table 4). Regarding tuber size the data in Table 4 showed that the number of small size tubers is greater in both varieties than medium and large size tubers with highest mean small tubers (4.00) produced in thiourea treatment under light conditions of potato variety SM kaghan.

Statistically no significant difference was found among various experimental factors regarding mean tuber yield per plant in both potato varieties (Table 5). However, among the variety, treatment and light/dark interaction, the maximum average tuber weight (42.37g) was recorded in potato variety SM kaghan for thiourea+GA₃ treatment under light conditions.

4. DISCUSSION

Major problem in potato seed production cycle is the in time availability of sprouted seed. Seed dormancy lead to poor sprouting which results in delayed planting and poor emergence. In the present study a significant percentage of minitubers of potato varieties i.e Roko and SM kaghan showed sprouting earlier when treated with GA₃ either individually or in combination with thiourea when placed in dark. By comparing dormancy period of tubers treated with Thiourea and GA₃ individually it was noticed that GA₃ was found superior and just after 28 days of treatments application higher percentage of minitubers started sprouting (Fig. 3). This may be due to the fact that the exogenous application of GA₃ enhances the endogenous level of gibberellins which affects the synthesis of amylase enzyme thereby enhances starch break down causing quick sprout growth as demonstrated by [25,26]. Carrera et al., [27] studied the effect of gibberellins on potato tuber dormancy and reported that internal growth regulators such as gibberellic acid, abscissic acid and cytokinins are responsible for dormancy termination and commencement of sprouting under invitro conditions and Demo [28] found that the maintenance of seed quality in terms of seed health and vigor are related to gibberellins application. The previous scientific research showed that tuber treatment with GA₃ caused breakdown of starch and accumulation of renewable sugars in potato tubers that can stimulate germination and consequently dormancy breaking [29].

Our results are consistent with the findings of Rehman et al., [19] who showed that number of

days required for 50% sprouting of potato minitubers can be reduced by the application of GA₃ under laboratory conditions. It is clear from the data presented in Fig. 5 that Roko variety showed greater sprouting % than SM kaghan variety in treatments that include GA₃ either alone or in combination with thiourea. These results are in line with the findings of Shibairo et al., [30] who reported differences among genotype response to GA₃ and found that potato seed tubers of genotype Kenya Sifa showed delayed in dormancy breaking and had least sprouting %. Carrera et al., [27] elaborated the reasons for such differences as that ectopic expression of gene coding for GA biosynthetic enzyme GA sub 20-oxidase resulted in high GA content of potato tuber and premature sprouting. Genotypic differences could therefore be attributed to inherent factors such as amounts of growth regulators present in each genotype. It is possible that the quantities of GA₃ present in some genotypes may be too low and therefore external supplementation is needed to prop up successive sprout growth.

The tubers of both varieties expressed very poor response to sugar treatment under both dark and light conditions and only 20% tubers showed sprouting in SM Kaghan variety under dark conditions. These findings are in contradiction to the results reported by Zaghum et al., [17] who demonstrated that treating potato tubers with 300grams sugar in 600ml water solution breaks the dormancy of potato tubers within 12-16 days at open light conditions. Contrary to this it was noticed in the present study that the tubers treated with sugar solution caused tuber decay mostly by fungal deterioration of tuber eyes.

The variations observed in dormancy period of different genotypes i.e Roko and SM kaghan in each treatment under the effect of dark and light conditions showed that after 31 days dormancy was broken in maximum tubers under dark. Demo et al., [31] demonstrated that seed storage under diffused light conditions can delay dormancy breaking by delaying tuber ageing process and reduce apical dominance. It was evident from the results of our study that tuber storage under dark and light conditions in interaction with the use of chemical treatments affected the dormancy period of potato varieties Roko and SM Kaghan which normally take 80-90 and 100-120 days for complete tuber dormancy elimination respectively. Greater number of minitubers sprouted in Roko than SM Kaghan cultivar in just 28 days which could be stated due

Table 4. Mean number of tubers per plant at the time of harvesting under interactive effect of genotype (variety) x chemical treatments x storage condition

Variety	*Tuber category	Treatments									
		Thiourea		GA ₃		Thiourea + GA ₃		Sugar		Control	
		Dark	Light	Dark	Light	Dark	Light	Dark	Light	Dark	Light
Roko	Large	0.60ab	0.17b	0.20b	0.17b	0.40ab	0b	0b	0b	1.00a	0.25ab
	Medium	0.87abc	1.50ab	1.10ab	1.83a	1.10ab	1.33ab	0c	0c	0.75bc	1.25ab
	Small	1.87abcd	1.33bcd	2.90abc	1.00cd	2.60abc	2.67abc	0d	0d	1.75abcd	2.75abc
	Mean Total	3.33ab	3.00b	4.20ab	3.00b	4.10ab	4.00ab	0c	0c	3.50ab	4.25ab
SM	Large	0.50ab	0.50ab	0.50ab	0.17b	0.32ab	0b	0b	0b	0b	0b
Kaghan	Medium	1.25ab	0c	1.50ab	0.83abc	1.37ab	1.33ab	0.50bc	0c	1.00abc	1.50ab
	Small	1.25bcd	4.00a	1.00cd	3.83a	1.93abcd	3.83a	3.00abc	0d	2.50abc	3.50ab
	Mean Total	3.00b	4.50ab	3.00b	4.83ab	3.62ab	5.16a	3.50ab	0c	4.75ab	5.00a

Means followed by different letters are significantly different at $P \leq 0.05$. *Tuber Category on the basis of weight (Large >20grams; Medium 10-20grams; Small < 10grams)

Table 5. Average tuber yield per plant (grams) under interactive effect of genotype (variety) x chemical treatments x storage condition

Variety	Treatments									
	Thiourea		GA ₃		Thiourea + GA ₃		Sugar		Control	
	Dark	Light	Dark	Light	Dark	Light	Dark	Light	Dark	Light
Roko	34.60a	38.10a	39.98a	34.88a	41.29a	39.35a	0b	0b	40.73a	38.75a
SM Kaghan	37.91a	30.10a	41.62a	42.35a	40.84a	42.37a	24.85a	0b	37.95a	36.52a

Means followed by different letters are significantly different at $P \leq 0.05$

to different physiological characteristics of different varieties; may be one is long duration (120 days crop) (SM Kaghan) whereas Roko has normal 90 days crop maturity. Differences in this characteristic have been observed in different varieties in the other experiments in literature. Salimi et al., [16] observed that regarding the genotypes, potato cultivars Burren and Agria achieved longest and shortest dormancy period after various chemical treatments.

Throughout the experiment, we can observe the significant effect of Thiourea +GA₃ treatment in Dark on dormancy breaking of tubers. The dormancy period was shorter in the tubers of both varieties treated with Thiourea + GA₃ and placed in Dark conditions as compared to other treatment interactions particularly tubers placed under light, while in control no sprouts were emerged in all treatment combinations of the two varieties. During the experiment it was also observed that sprouts developed from the tubers of the two varieties treated with gibberellic acid and placed in dark conditions were found slender and elongated as compared to untreated control. This might be due to the fact that breaking dormancy naturally requires greater physiological age, which affects membrane integrity [32] and metabolic activity of cell division and elongation [33]. Alexopoulos et al., [34] reported that potato tubers treated with gibberellic acid produced seedlings that were thinner and longer than those that sprouted naturally.

It was also observed in this study that in different experimental treatment interactions the sprouted tubers after sowing showed 100% germination, good growth and production performance. Barrani et al. [35] showed that when seed potatoes were treated either dormant or sprouted with various concentrations of gibberellic acid, emergence of plants from treated seed was more rapid than from untreated seed tubers.

Statistically plant height was non-significantly ($p \geq 0.05$) affected by cultivar, GA₃ and thiourea and storage conditions i.e light and dark (Table 3). However, data recorded on growth parameters of potato minitubers revealed that plants developed from tubers treated with GA₃ individually and along with thiourea showed increase in plant height of the two cultivars i.e. Roko, and SM kaghan and the response did not vary significantly between light and dark storage conditions. The process of plant growth occurred as cell division and cell elongation. The

comparatively increase in plant height might be due to GA₃ effect on the elongation of internodes and shoot growth. Reza et al. [36] demonstrated that the GA₃ causes increase in cell number and cell size resulting in a significant effect on growth. Chindi and Tsegaw [37] reported that GA₃ applied on the foliage increased plant growth (8%) as compared to the control, and it may be related to the effect of internodes and shoot growth. But in this study no significant difference was found between treatments \times Cultivar \times light/dark storage interaction which can be due to the reason that no foliar application was done in this experiment. Plant height in sugar treatment was found lower than control in SM Kaghan variety, the possible reason for this might be that the sugar is not a hormone or growth promoting agent and therefore showed no effect.

Non-significant differences observed regarding mean number of branches of both varieties in all the interactions including control, however effect of GA₃ was strongly enhanced by thiourea, since the most efficient plant growth and higher number of branches were recorded in thiourea+GA₃ \times dark/light conditions in both varieties (Table 3).

Non-significant differences observed in the yield parameters between varieties, chemical treatments and storage environment. GA₃, thiourea and sugar treatment didn't cause any significant change in number of tubers and total yield in two potato cultivars (Tables 4 & 5). However, the utilization of thiourea and GA₃ individually and with other hormones in solving dormancy problem has not been optimized for all potato genotypes and there are still many constraints on the information of potential yields obtained after breaking dormancy.

5. CONCLUSION AND RECOMMENDATION

On the basis of results it was concluded that Thiourea along with GA₃ shortens the dormancy period of tubers as compared to other treatment combinations. GA₃ can break the tuber dormancy to about 31 days if stored in dark environment. Tubers placed in light conditions (diffused light) can take much longer time to sprout even treated with GA₃ or Thiourea or both. It was also concluded that the dormancy breaking treatments did not affect the plant growth and yield in comparison with control. Visual observation concluded that storage in light

conditions produced shoots with better quality than ones in dark conditions.

It is recommended that the different concentrations of GA3 can be tested alone and with different concentrations of thiourea in combination with different environmental conditions temperature, humidity and ventilation of storage place for different potato varieties to find a comprehensive protocol to shorten the dormancy duration in certified seed potato production cycle for those areas where there is only one potato crop per year.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

- Ekin Z. Some analytical quality characteristics for evaluating the utilization and consumption of potato (*Solanum tuberosum* L.) tubers. *Afr J Biotechnol.* 2011; 10: 6001–6010.
- King JC, Slavin JL. White potatoes, human health, and dietary guidance. *Adv Nutr.* 2013; 4:393S–401S. DOI:<https://doi.org/10.3945/an.112.003525>
- Hawkes J. The potato evolution, biodiversity and genetic resources. 1990; pp.viii + 259 pp. ref.7 pp. Re-trived from <https://www.cabdirect.org/cabdirect/abstract/19901615687>
- FAOSTAT. 2008. Food and Agriculture Organization of the United Nations Statistical Database. Retrived from <http://faostat.fao.org>
- Tiwari JK, Buckseth T, Zinta R, Bhatia N, Dalamu D, Naik S, Poonia AK, Kardile HB, Challam C, Singh RK, Luthra SK, Kumar V and Kumar M. Germplasm, Breeding, and Genomics in Potato Improvement of Biotic and Abiotic Stresses Tolerance. *Front. Plant Sci.* 2022; 13: 805671. doi: <https://doi.org/10.3389/fpls.2022.805671>
- Gul Z, Qureshi F, Jamal Z. Growth and minituber yield response of potato plantlets in micropropagation to different plant spacing under greenhouse conditions. *Int J Agric Environ Food Sci.* 2020; 4 (3): 271-277. DOI: <https://doi.org/10.31015/jaefs.2020.3.5>
- Bashir KM, Ali A, Farrukh MU, Alam M.. Estimation of economic and production efficiency of potato production in Central Punjab. Pakistan. *Custos e @gronegocio on line* 2021; 17: 2–23.
- Khan IA. PARS-70: an interspecific potato hybrid suitable for long storage and autumn-to-autumn seed multiplication. *Potato Res.* 2004; 47: 187–193. doi: <https://doi.org/10.1007/BF02735984>.
- Haider MW, Ayyub CM, Malik AU, Ahmad R. Plant growth regulators and electric current break tuber dormancy by modulating antioxidant activities of potato. *Pakistan J. Agri. Sci.* 2019; 56: 867–877. doi: <https://doi.org/10.21162/PAKJAS/19.7428>
- Suttle JC. Dormancy and sprouting, In: D. Vreugdenhil, J. Bradshaw, C. Gebhardt, F. Govers, M. Taylor, D. MacKerron, and H. Ross (eds.). *Potato physiology and biotechnology. Advances and perspectives.* Elsevier, New York, NY. 2011; p. 288–308.
- Suttle JC. Dormancy and sprouting. In *Potato Biology and Biotechnology. Advances and Perspectives*, 1st ed.; Vreugdenhil, D., Bradshaw, J., Gebhardt, C., Govers, F., MacKerron, D.K.L., Taylor, M.A., Eds.; Elsevier: Amsterdam, The Netherlands. 2007; pp. 287–309.
- TeperBamnolker P, Buskila Y, Lopesco Y, Ben Dor S, Saad I, Holdengreber V, Belausov E, Zem ach H, Ori N, Lers A, Eshel D. Release of Apical Dominance in Potato Tuber Is Accompanied by Programmed Cell Death in the Apical Bud Meristem. *Plant Physiol.* 2012; 158: 2053–2067.
- Bryan JE. Breaking dormancy of potato tubers. *CIP Research Guide 16.* International Potato Center, Lima, Peru. 1989; 12 p.
- Kim HS, Joen JH, Choi KH, Joung YH, Joung H. Effect of rindite on breaking dormancy of potato microtubers. *Am. J. Potato Res.* 1999; 76: 5-8.
- Alexopoulos AA, Aivalakis G, Akoumianakis KA, Passam HC. Bromoethane Induces Dormancy Breakage and Metabolic Changes in Tubers Derived from True Potato Seed, *Postharv. Biol. Technol.* 2009; 54: 165–171. doi: <https://doi.org/10.1016/j.postharvbio.2009.07.004>
- Salimi K, Afshari RT, Hosseini MB, Struik PC. Effects of gibberellic acid and carbon disulphide on sprouting of potato

- minitubers. *Sci. Hortic.* 2010; 124: 14–18. doi: <https://doi.org/10.1016/j.scienta.2009.12.026>.
17. Zaghum MJ, Zohaib A, Khalid MN, Muhy-Ud-Deen G, Ahmad A, Zaidi SSH. Effect of Different Chemicals on Eliminating the Dormancy Period of Freshly Harvested Seed Potatoes. *Acta Scientific Agriculture.* 2021; 5(3): 2-7.
 18. Mani F, Bettaieb T, Doudech N, Hannachi C. Effect of hydrogen peroxide and thiourea on dormancy breaking of microtubers and field-grown tubers of potato. *Afr. Crop Sci. J.* 2013; 21: 221–234.
 19. Rehman F, Lee SK, Kim HS, Jeon JH, Park J, Joung H. Dormancy breaking and effects on tuber yield of potato subjected to various chemicals and growth regulators under greenhouse conditions. *J. Biol. Sci.* 2001;1: 818–820.
 20. Claassens MMJ, Verhees J, van der Plas LH, van der Krol AR, Vreugdenhil D. Ethanol Breaks Dormancy of the Potato Tuber Apical Bud. *J. Exp. Bot.* 2005, 56: 2515–2525.
 21. Hartmann A, Senning M, Hedden P, Sonnewald U, Sonnewald S. Reactivation of meristem activity and sprout growth in potato tubers require both cytokinin and gibberellin. *Plant Physiol.* 2011; 155: 776–796.
 22. Suttle JC. Physiological regulation of potato tuber dormancy. *Am. J. Potato Res.* 2004; 81: 253.
 23. Mustefa G, Mohammed W, Dechassa N, Gelmesa D. Effects of different dormancy-breaking and storage methods on tuber tuber sprouting and subsequent yield of two potato (*Solanum tuberosum* L.) varieties. *Open Agriculture.*, 2017; 2: 220–229.
 24. Analytical software. Statistix 8.1. User's Manual. Analytical Software, Tallahassee, Florida; 2005.
 25. Krochko JE, Abrams GD, Loewen MK, Abrams SR, Cutler A J. (+)-Abscisic acid 8'-hydroxylase is a cytochrome P450 monooxygenase. *Plant Physiol.* 1998; 118: 849–860. doi: <https://doi.org/10.1104/pp.118.3.849>.
 26. Zhang H, Hou J, Liu J, Xie C, Song B. Amylase analysis in potato starch degradation during cold storage and sprouting. *Potato Res.* 2014;57:47–58. doi: <https://doi.org/10.1007/s11540-014-9252-6>
 27. Carrera E, Bou J, García-Martínez J L, Prat S. Changes in GA 20-oxidase gene expression strongly affect stem length, tuber induction and tuber yield of potato plants. *Plant J.* 2000; 22: 247–256. doi: <https://doi.org/10.1046/j.1365-313x.2000.00736.x>
 28. Demo P. Strategies for seed potato (*Solanum tuberosum* L.) production using rooted apical stem cuttings and tubers in Cameroon. 2002; Ph.D Thesis, Department of Agronomy, University of Ibadan, Nigeria.
 29. Alexopoulos AA, Akoumianakis KA, Vemmos SN, Passam HC. The effect of postharvest application of gibberellic acid and benzyl adenine on the duration of dormancy of potatoes produced by plants grown from TPS. *Postharvest Bio. Tech.* 2007; 46: 54-62.
 30. Shibairo, SI, Demo P, Kabira JN, Gildemacher P, Gachango E, Menza M, Nyankanga R O, Chemining'wa GN, Narla RD. Effects of gibberellic acid (GA3) on sprouting and quality of potato seed tubers in diffused light and pit storage conditions. *Journal of biological sciences.* 2006; 6(4): 723-733.
 31. Demo P, Akoroda MO, El-Bedewy R, Asiedu R. Monitoring storage loses of seed potato (*Solanum tuberosum* L.) tubers of different sizes under diffused light conditions. Proceedings, 6th triennial congress of the African Potato Association (APA), 5-10 April, 2004, Agadir Morocco. 2004; 363-370.
 32. Knowles LO, Knowles NR. Changes in fatty acid composition of phospholipids from different ages of potato seed-tubers during sprouting. *Ann. Bot.* 1990; 65: 217-223.
 33. Taiz L, Zeiger E. *Plant Physiology.* 3rd ed. Sinauer Associates, Sunderland, MA. 2002; 690pp.
 34. Alexopoulos AA, Aivalakis G, Akoumianakis KA, Passam HC. Effect of gibberellic acid on the duration of dormancy of potato tubers produced by plants derived from true potato seed. *Posth. Biol. & Tec.* 2008; 49: 424-430.
 35. Barani M, Akbari N, Ahmadi H. The effect of gibberellic acid (GA3) on seed size and sprouting of potato tubers (*Solanum tuberosum* L.) . *African journal of Agricultural Research.* 2013; 8(29): pp. 3898-3903, 1 August, 2013. DOI: <https://doi.org/10.5897/AJAR09.419>.

36. Reza M, Islam M, Hoque A, Sikder RK, Mehraj H, Jamal Uddin AFM. Influence of Different GA Concentrations on Growth and Yield of Broccoli. American-Eurasian Journal of Scientific Research. 2015; 10 (5): 332-335.
37. Chindi A, Tsegaw T. Effect of gibberellic acid on growth, yield and quality of potato (*Solanum tuberosum* L.) in central highlands of Ethiopia. Journal Horticulture Sci. 2019;1(2):1-10.

© 2023 Gul and Iqbal; This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Peer-review history:
The peer review history for this paper can be accessed here:
<https://www.sdiarticle5.com/review-history/109232>