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Impact of foliar applied Ethrel, potassic compounds and thiourea on flowering and fruiting attributes of Litchi (*Litchi chinensis* Sonn.)

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Abstract

Plant growth regulators and nutrients play a significant role in altering plant growth subsequently yield and quality. The impact of foliar application of ethrel, potassic compounds [Potassium nitrate (KNO₃), Potassium dihydrogen phosphate (KH2PO4) and Dipotassium hydrogen phosphate (K2HPO4)] and thiourea on flowering and fruiting characteristics of Litchi (Litchi chinensis Sonn.) was explored in an experiment based on a randomized block design with thirteen treatments and three replication at HRC, GBPUA&T Pantnagar (India). The treatments included T₁ [ethrel (400 ppm)], T₂ [KNO₃ (1%)], T₃ [K₂HPO₄ (1%)], T₄ [KH₂PO₄ (1%)], T₅ [thiourea (1%)], T₆ [K₂HPO₄ (2%)], T₇ [KH₂PO₄ (2%)], T₈ $[K_2HPO_4 (1\%) + KNO_3 (1\%)], T_9 [KH_2PO_4 (1\%) + KNO_3 (1\%)], T_{10} [K_2HPO_4 (1\%) + thiourea (0.5\%)],$ T_{11} [KH₂PO₄ (1%) + thiourea (0.5%)], T_{12} [KNO₃ (1%) + thiourea (0.5%)] and T_{13} as control. The foliar spray of these chemicals was done thrice from end of September to November at 20 days interval. Early panicle initiation, advancement in flowering (3 days), maximum number of female (324.63) and total flowers (836.39), highest fruit set per panicle (59.49), minimum fruit drop (75.90%) and highest yield (43. 82 kg/tree) was observed in treatment T₂ [KNO₃ (1%)] whereas, highest fruit weight (20.99g) and aril weight (17.30g) was observed in T₁ [ethrel (400 ppm)] treatment. Maximum flowering intensity and highest pulp recovery was reported in T₉ [KH₂PO₄ (1%) + KNO₃ (1%] and T₄ [KH₂PO₄ (1%)] treatments, respectively.

Keywords: Litchi chinensis, ethrel, potassium, flowering, fruiting, yield

Introduction

Litchi (Litchi chinensis Sonn.) is one of the most important subtropical fruit grown throughout the world for its juicy fleshy aril. It is believed to have originated in Southern China and Northern Vietnam region from where it became widely distributed in the Subtropics (Menzal and Simpson, 1986) [23, 25]. Litchi was introduced to India by the end of the 17th century (Liang, 1981) [20]. India is the second largest litchi producer in the world next to China and has substantially expanded its cultivation in past 50 years. The annual production of litchi in India accounts to 6.86 lakh MT from an area of 0.92 lakh hectare with an average productivity of 7.4 MT/ha (Anonymous, 2018) [3]. Litchi is an excellent source of vitamin C (40 mg/100g) and contains significant amount of protein (0.8-0.9%), fat (0.3%), pectin (0.43%) and mineral content especially calcium, phosphorus and iron (Pandey and Sharma, 1998) [29]. It is among one of the few fruit crops, which is exact to its climatic requirement. Litchi flourishes well in moist subtropical regions with abundant rainfall. Dry winds during summer and frost in winter are detrimental for its growth. Its cultivation is restricted to lower tropics and subtropics where flowering and fruiting is uneven due to lack of prerequisite cool temperature (Hegele et al., 2010) [12]. Irregular flowering in litchi trees during spring season is associated with previous years vegetative flushing in late autumn and early winter which can result in inconsistent yields (Nagao et al., 2000) [27]. Among other problems associated with the production of litchi, the narrow genetic base of litchi results in the availability of litchi for a very short period of time in the market, which is further reduced due to poor shelf life and browning of litchi fruits. The initial fruit set in litchi is very high but a small proportion of fruits finally mature due to

cracking and fruit drop. Chemicals and PGR's play a significant role in altering the plant response from vegetative to reproductive stage in litchi. Foliar application of plant nutrients is helpful in satisfying plant requirements and can be highly effective when nutrient uptake via the root system is limited. Potassium is essential nutrient for numerous physiological and biochemical processes like photosynthesis, phosphorylation, transportation of photo assimilates, enzyme activation, transpiration, stress tolerance (Pettigrew, 2008) [30]. Foliar application of potassium to bearing fruit trees increases fruit size, soluble solids, ascorbic acid, colour, shelf life, and shipping quality (Lester et al. 2007) [19]. Potassic compounds like potassium nitrate (KNO₃) can be used for dormancy breaking in buds (Trewavas, 1983) [40]. Several researchers have confirmed the role of potassic compounds in enhancing flowering in different fruit crops. Valmayor (1962) [42] reported increased internal ethylene concentrations in shoots as well as ethylene forming enzyme (EFE) activity in leaves by KNO₃, which induced flowering. KNO₃ enhances flowering especially in tropical regions where cold temperature for floral induction may not be sufficient (Oosthuyse, 1992). In subtropical regions where winter conditions are usually sufficient for floral induction, flowering enhancement by KNO3 has not been reported. Protacio (2000) [31] hypothesized increase in starch concentration and subsequently floral stimulus in the shoots when the gibberellins levels fall below a threshold level. However, the buds remain quiescent even after sufficient starch accumulation until conditions are favourable for flowering. KNO₃ activates those quiescent buds for floral initiation. Srihari and Rao (1998) [37] reported induced flowering on axillary shoots and increased yield of fruit trees by foliar application of Orthophosphoric acid. Garad et al. (2013) [9] reported increased number of flower percentage, maximum fruits per panicle and maximum cost benefit ratio in mango by spraying KH₂PO₄ (1%) + KNO₃ (1%). Ethylene (H₂C=CH₂) is a simplest olefin that exists in gaseous phase under normal physiological conditions. It is produced from all parts of plants and plays an important role in cellular, developmental and stress related processes in plants. Ethrel or Ethephon (2-chloroethyl phosphonic acid) which is a synthetic plant growth regulator undergoes chemical biodegradation at pH greater than 4.1 in cell cytoplasm to release ethylene (Urwiler and Stutte, 1986; Kasele et al., 1995) [41, 15]. Foliar application of ethrel has been found to increase C/N ratio in shoots and leaves, highest number of flowering panicles and maximum fruits at initial stage and also at final stage in Litchi Cv. Bombai (Mandal et al., 2014) [21]. Thus, well balanced plant nutrition management is essential for maximum yield and fruit quality. Keeping in view the above factors under consideration, the present experiment was conducted to evaluate the effect of different chemicals on flowering and fruiting attributes of litchi cv. Rose Scented.

Material and Methods

The present experiment was conducted at Horticultural Research Centre, Govind Ballabh Pant University of Agriculture and Technology, Pantnagar on 15 years old litchi trees of Cv. Rose Scented planted at 10×10 m planting distance. The centre is located at 29^{0} N latitude, 79.3^{0} E longitudes and 244 m above mean sea level. The region is characterized by humid subtropical climate with dry summers and cold winters. Maximum temperature ranges from 30^{0} C to 43^{0} C in summers and 5^{0} C to 10^{0} C in winters along with

mean rainfall 1400 mm per annum. Monsoon rains occurs from the third week of June to middle of September. Frost can be expected from the last week of December to first week of February. The soil of the experimental site was silty clay loam under the order mollisols. The experiment was laid out in randomized block design with 13 treatments and 3 replications. The treatments T₁ [ethrel (400 ppm)], T₂ [KNO₃ (1%)], T₃ [K₂HPO₄ (1%)], T₄ [KH₂PO₄ (1%)], T₅ [thiourea (1%)], T₆ [K₂HPO₄ (2%)], T₇ [KH₂PO₄ (2%)], T₈ [K₂HPO₄ $(1\%) + KNO_3 (1\%)$], $T_9 [KH_2PO_4 (1\%) + KNO_3 (1\%)]$, T_{10} $[K_2HPO_4 (1\%) + thiourea (0.5\%)], T_{11} [KH_2PO_4 (1\%) +$ thiourea (0.5%)], T_{12} [KNO₃ (1%) + thiourea (0.5%)] were applied. T₁₃ was taken as control and remained untreated during the course of experiment. The foliar spray of chemicals was performed thrice during the last week of September to November at 20 days interval. Date of panicle initiation was recorded by daily field visits during flowering time. Number of male, female and total flowers per panicle was counted in selected panicles of each direction by means of visual counting. Sex ratio was calculated by dividing the number of male flowers with number of female flowers of the same panicle. Flowering intensity was calculated by dividing the number of flowering shoots with total number of shoots in all directions. Flowering advancement was calculated by comparing the date of flowering in different treatments with the date of flowering in control. The time taken to fruit set was determined by counting the number of days between anthesis to the date of fruit set. The fruit set per panicle was calculated by counting the number of fruits in each selected panicle in all the directions by working out the average of data, thus obtained (replication wise). The percentage of fruit drop was calculated on the basis of the total number of fruit dropped from initial fruit set stage to harvesting maturity. Fruit cracking was observed by tagging selected panicles in each direction of the trees. Days taken to fruit maturity were determined by counting the number of days from the day of panicle initiation (bud burst) to the fruit maturity stage. Yield per hectare was recorded by weighing the total harvested fruits in each replication of each treatment by using weighing balance. Average weight of fruits, aril and seed was measured using electronic weighing balance. Pulp recovery per cent was calculated by diving average weight of aril by average weight of fruit. The data was statistically analysed as per the method of Gomez and Gomez (1984) [10]. Data was subjected to ANOVA (Analysis of Variance) using Least Significant Difference at 5% level of significance.

Result and Discussion

Earliest panicle emergence was observed on 16th February in T_1 [ethrel @ 400 ppm] and T_2 [KNO₃ @ 1%] whereas T_8 $[K_2HPO_4 @ 1\% + KNO_3 @ 1\%]$ and T_{13} [control] recorded late panicle emergence on 19th February (Table 1). Ethrel or Ethephon (ethylene releasing substances) are known for leaf expansion (Khan, 2005; Khan et al., 2008) [16, 17] and chlorophyll synthesis at lower concentration (Grewal et al., 1993) [11]. Ethylene releasing compounds have also shown significant effect on leaf photosynthetic rate (Pua and Chi, 1993) [32]. The increased photosynthesis and subsequently high carbohydrate accumulation in shoots might have resulted in early maturation of shoots for inducing early panicle. The results obtained are in close conformity with the findings of Mandal et al. (2014) [21]. Potassium nitrate (KNO₃) has also shown profound effect in stimulating early flowering in trees growing in tropical and subtropical regions (Adam, 1986) [1]. The duration of flowering in all the treatments varied from 17-20 days. The duration was minimum (17 days) in T₂ [KNO3 @ 1%] whereas T₁₃ [control] had maximum flowering duration of 20 days. All the treatments had a significant effect on flowering advancement as compared to control (T₁₃). The maximum advancement of flowering (3 days) was recorded in T2 [KNO3 @ 1%] (Table 1). Similar result of flowering advancement (4-5 days) by application of KNO₃ was reported by Dalal et al. (2005) [4]. The maximum number of female flowers (324.63) and total flowers (836.39) per panicle were observed in T₂ [KNO₃ @ 1%] whereas it was minimum in T₁₃ [control]. Kulkarni (2014) ^[18] reported KNO₃ application increases number of flowers by forcing and sensitizing the flower buds to respond to the floral stimulus which is present in stems. Davenport (2003) [5] observed 100% reproductive shoots and profuse flowering when KNO₃ was applied on older shoots while 2-3 months old shoots resulted in vegetative growth in response to KNO₃. Increase in percent of flowering due to KNO3 might be due to increased concentrations of zeatin or zeatin riboside which are flower induction promoters present in it (Eric Guevara et al., 2012) [6]. The dose and time of application of flower inducing chemicals has a significant effect on the kind of response obtained. In the present study, significant response of increased number of flowers was observed with 1% KNO₃. The results obtained are in support with the findings of Mitra and Sanyal (2001) [26] who reported KNO3 induced flowering in litchi cv. Bombai. Garad et al. (2013) [9] also found increased number of hermaphrodite flowers by application of KNO₃. Significant difference for the sex ratio was observed among all the treatments. The sex ratio (male: female) varied from 1.57 to 1.91 in different treatments. Minimum sex ratio (1.57) was observed in T_2 (KNO₃ @ 1%) whereas T_{13} (control) recorded the maximum sex ratio (1.91).

The flowering intensity was remarkably improved in all the treatments. It was maximum (70.51%) in T₉ [KH₂PO₄ @ 1% + KNO₃ @ 1%] whereas T₁₃ [control] recorded minimum (61.82%) flowering intensity (Table 1). Potassium plays an important role in increasing absorption of Mg, an element that is fundamental in the floral formation and promotes the synthesis of nucleic acids (Feucht, 1982) [8]. Potassium compounds are also known to affect the nitrogen level of the plant and specific products of N metabolism and amino acids plays a direct role in the initiation and differentiation of floral buds in the shoots. Marschner (2002) [22] reported reduced flower bud formation in the shoots in case of a deficiency of P. Increased flowering intensity in the shoots by KH₂PO₄ might be due to the fraction of K in it, that stimulates photosynthesis and transports photo assimilates, which is important for the formation of flowers (Swietlik, 2003) [38] whereas nitrate form (NO3-) in KNO3 increases levels of arginine, compound which can promote flowering (Niu et al. 2007) [28]. Similar results of increased flowering intensity by application of Potassium are reported in other fruits (Agusti, 2003). The results are in close conformity with the findings of Garad et al. (2013) [9] who reported increased flowering intensity by KH₂PO₄ (1%) + KNO₃ (1%). The time taken from flowering to fruit set in all the treatments varied from 16 to 20 days (Table 2). The treatment T₁ [Ethrel @ 400 ppm] took the least time (16 days) to fruit set whereas T₁₃ [control] took maximum time (20 days) for fruit set (Table 2). Similar findings of reduced time period for fruit setting by ethrel were observed by Mandal et al. (2014) [21]. Fruit set per panicle in all the treatments were significantly higher than T_{13} [control].

[KNO₃ @ 1%] followed by T₁ [Ethrel @ 400 ppm]. Similar results regarding increased fruit set per panicle by KNO₃ was reported by Mitra and Sanyal (2001) [26] in litchi and Yeshitela et al. (2004) [44] in mango. The treatment T₂ [KNO₃ @ 1%] had the least fruit drop (75.90%) as compared to other treatments. The results obtained are in support with findings of Sharma et al. (1990) [36] who recorded reduced fruit drop and increased fruit retention by foliar application of KNO3 in mango cv. Langra. Days taken to maturity among all the treatments varied from 51 to 56 days. From fruit set to maturity, growth of litchi fruit occurs in three phase. During first phase, pericarp, seed coat and embryo develop. Second phase is characterized by rapid growth of embryo and beginning of aril development while in last third phase, aril development and maturation takes place (Joubert, 1986; Wang et al., 2017) [14, 43]. Fruits attained the maturity stage when they developed a bright red blush with flattened tubercles. The minimum days for maturity (51 days) were observed in T₁ [ethrel @ 400 ppm] and T₇ (KH₂PO₄ @ 2%) treatments whereas the T₁₃ [control] took maximum days for maturity (56 days). Maximum fruit yield (43.82 kg/tree) was obtained in T₂ [KNO₃ @ 1%] whereas the minimum yield (34.08 kg/tree) was recorded in T₁₃ [control] (Table 2). It is well known fact that yields depends upon reserve food stored by plants as well as many other internal and external factors. Higher fruit set and reduced fruit drop in KNO₃ treated plants subsequently resulted in higher yield. The present results are in accordance with the findings of Sangwan et al. (2008) [34] who reported increased yield by the foliar application of KNO₃. Fruit weight is considered as most important external factor in determining fruit quality as it greatly influences consumers appeal. Foliar application of all the chemicals significantly increased the fruit weight. The maximum fruit weight (20.99 g) was observed in treatment T₁ [ethrel @ 400 ppm] whereas it was minimum (18.46 g) in T₁₃ [control]. The increased fruit weight might be due to more rapid increase in the number of cell during the growth and development of the fruit. The actively proliferating parenchyma cells in the aril are initially elongated and slim, but become irregular in shape during the later phase of isodiametric growth. The involvement of ethylene in the growth of these cells seems probable, since this substance is active in the growth of other fruits. Similar results were observed by Saxena (1994) [35] and Jones et al. (1993) [13] who reported increased fruit weight by foliar application of ethrel. The aril weight in all the treatments was statistically found significant over control. The maximum aril weight (17.30g) was recorded in T_1 [ethrel @ 400 ppm] while minimum aril weight (14.33 g) was observed in T₁₃ [control]. Maximum pulp recovery percent (81.49%) was recorded in T₄ [KH₂PO₄ @ 1%] whereas T₁₃ (control) recorded minimum (77.61%) pulp recovery (Table 2). The K fraction of the KH₂PO₄ is involved in Hill reaction (for generation of NADPH and ATP, together with ionic equilibrium, electron transport, and proton-motive force) and Calvin and Benson cycle (sugar production and transport, partitioning of photoassimilates) of the photosynthesis (Tighe-Neira et al., 2018) [39] whereas phosphate (PO₄) fraction

bounds with the sugars to make different kinds of

carbohydrates and starches, which are then ultimately

transformed into organic compounds of the edible portion,

thus increases the share of edible portion in the fruits (Rice,

The initial fruit set per panicle was maximum (59.49) in T₂

2016) [33].

Table 1: Effect of foliar spray of Ethrel, potassic compounds and Thiourea on flowering attributes of Litchi Cv. Rose scented

Symbols	Treatments	Date of Panicle	Duration of	Advancement in Flowering	Flowers/panicle	Flowers/panicle	Total flowers/	Sex Ratio	Flowering Intensity
Symbols	Treatments	Initiation	-		(Male)	(Female)	panicle	(M/F)	(%)
T_1	Ethrel (400ppm)	16.02.2016	18	2	491.46	307.81	799.27	1.60	65.13
T_2	KNO ₃ (1%)	16.02.2016	17	3	511.76	324.63	836.39	1.57	65.80
T ₃	K ₂ HPO ₄ (1%)	18.02.2016	19	1	414.13	248.48	662.61	1.66	66.84
T_4	KH ₂ PO ₄ (1%)	17.02.2016	19	1	406.34	236.00	642.34	1.74	61.28
T_5	Thiourea (0.5%)	18.02.2016	20	0	396.78	220.09	616.87	1.80	66.26
T_6	K ₂ HPO ₄ (2%)	18.02.2016	19	1	415.96	248.10	664.07	1.67	65.06
T ₇	KH ₂ PO ₄ (2%)	17.02.2016	19	1	368.39	256.16	624.56	1.64	64.21
T_8	$K_2HPO_4(1\%) + KNO_3(1\%)$	19.02.2016	19	1	458.19	244.98	703.18	1.87	65.13
T ₉	$KH_2PO_4(1\%) + KNO_3(1\%)$	18.02.2016	19	1	456.29	246.47	702.76	1.85	70.51
T ₁₀	K ₂ HPO ₄ (1%) + Thiourea (0.5%)	18.02.2016	20	0	430.99	266.06	697.05	1.62	62.21
T ₁₁	KH ₂ PO ₄ (1%) + Thiourea (0.5%)	18.02.2016	18	2	393.21	241.09	634.30	1.64	66.71
T ₁₂	KNO ₃ (1%) + Thiourea (0.5%)	17.02.2016	18	2	462.70	275.75	738.45	1.67	69.10
T ₁₃	Control	19.02.2016	20	0	390.49	203.94	594.43	1.91	61.82
	CD at 5%	-	-	-	NS	36.16	86.88	0.20	3.74
	SEm±	-	-	-	20.01	12.39	29.78	0.06	1.29

Table 2: Effect of foliar spray of Ethrel, potassic compounds and Thiourea on fruiting attributes of Litchi Cv. Rose scented

Symbols	Treatments	Time taken to fruit set (days)	Fruit set per Panicle		Fruit cracking (%)	Days taken to maturity	Yield (Kg/tree)	Fruit weight (g)	Aril weight (g)	Seed weight (g)	Pulp recovery (%)
T_1	Ethrel (400ppm)	16	57.08	81.58	14.6	51	41.29	20.99	17.30	2.00	80.28
T_2	KNO ₃ (1%)	18	59.49	75.90	15.66	53	43.82	20.06	16.16	2.04	80.37
T ₃	K ₂ HPO ₄ (1%)	17	46.07	81.79	14.15	52	40.51	19.61	15.19	2.03	77.95
T ₄	KH ₂ PO ₄ (1%)	19	49.35	82.08	11.46	54	38.16	19.33	15.74	2.16	81.49
T ₅	Thiourea (0.5%)	18	48.76	82.86	12.09	54	35.15	19.57	15.77	2.19	80.58
T_6	K ₂ HPO ₄ (2%)	18	41.79	83.17	15.70	53	40.85	19.08	15.47	2.05	81.05
T 7	KH ₂ PO ₄ (2%)	19	54.90	80.77	16.46	51	37.72	18.60	15.22	2.09	81.08
T ₈	$K_2HPO_4(1\%) + KNO_3(1\%)$	19	52.36	86.48	18.82	54	41.30	20.08	15.70	2.60	78.33
T 9	$KH_2PO_4(1\%) + KNO_3(1\%)$	18	43.99	82.51	19.04	54	41.61	19.66	15.44	2.15	78.50
T_{10}	K_2HPO_4 (1%) + Thiourea (0.5%)	19	55.94	76.27	17.76	55	42.34	18.81	15.20	2.01	80.88
T_{11}	KH_2PO_4 (1%) + Thiourea (0.5%)	18	43.07	91.08	18.86	54	36.53	18.57	14.44	2.24	79.47
T_{12}	KNO ₃ (1%) + Thiourea (0.5%)	17	45.22	77.86	15.82	53	37.56	19.41	15.11	2.30	77.88
T ₁₃	Control	20	39.84	89.24	19.50	56	34.08	18.46	14.33	2.32	77.61
	CD at 5%	-	11.81	7.02	NS	-	5.71	1.36	0.84	NS	2.97
	SEm±	-	4.05	2.40	3.74	-	1.95	0.46	0.28	0.12	1.16

Note: *CD= Critical difference, SEm± = Standard error of mean

Conclusion

The present study indicated that foliar application of chemicals is highly effective for improving flowering and fruiting in case of litchi. The potassic form of chemicals mainly KNO₃ (1%) significantly improved the flowering attributes particularly early and maximum flowering, highest fruit set and subsequently higher yield whereas highest fruit and aril weight was obtained with 400ppm ethrel spray.

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