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Influence of Pre Harvest and Pre Flower Sprays of Gibberellic acid, Napthalene acetic acid and Ethrel on Flowering Behaviour, Fruit Yield of Phalsa Cultivar Purple Round under Jammu- Sub Tropics

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ABSTRACT

Keywords

Phalsa, Purple Round, Plant growth regulators, Flowering behavior and yield.

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The experiment was conducted at Rainfed Research Sub-Station for Subtropical Fruits (RRSS) Raya, Sher-e-Kashmir University of Agricultural Sciences and Technology of Jammu, during 2015-2016 in Randomized Block Design with three replications and ten treatments; T_1 – 100 ppm GA₃, T_2 - 150 ppm GA₃, T_3 - 200 ppm GA₃, T_4 - 100 ppm NAA, T_5 -150 ppm NAA, T_6 -200 ppm NAA, T_7 - 750 ppm Ethrel, T_8 - 1000 ppm Ethrel, T_9 - 1250 ppm Ethrel and T_{10} - control (water spray). Flowering behaviour like date of flower initiation, date of end of flowering, duration of flowering, average number of flowers per shoot and fruit yield were observed during the investigation. Pre flowering and pre harvest spray of 1250 ppm or 1000 ppm ethrel resulted in early flowering i.e. on 18th March while untreated bushes were last to initiate flowering i.e. on 25th March. Ethrel 1250 ppm resulted into early end date of flowering i.e. on 21st April. GA₃ 150 ppm resulted in maximum average number of flowers and fruit yield in phalsa.

Introduction

Phalsa (*Grewia asiatica* L.) belongs to family Tiliaceae. The phalsa is indigenous to Indian sub-continent South-East and (Chundawat and Singh, 1980). Phalsa is found in wild form all along the foothills of Himalayas. There are 18 genera and 400 species which are mostly distributed in tropical and sub-tropical regions of the world. The genus *Grewia* has 140 species, out of which 40 occur in India. Phalsa is sub-tropical fruit plant, flowering starts from February-March. Phalsa bear flowers on current year growth in the axil of the leaves and the dehiscence of anther takes place before the flowers are completely open (Godara, 1985).

There are three to seven peduncles and each peduncle has three to six flowers of yellow colour. Phalsa is self-compatible, but pollens are not able to reach the stigma to effect the self-pollination due to detraction of stamens away from the stigma causes low fruit set (Randhawa and Dass, 1962). Phalsa fruit ripe in second fortnight of April and continue upto middle of June. Ripe berry contain 50-60 % juice, 10-11 % sugar and 2.0-2.5 % acid, 14.4 % carbohydrates, 1.5 % protein, 0.9 % fat, 129 mg/100g of pulp, 89 mg phosphorous, 3.1 mg iron, 22 mg/100 g of pulp vitamin C and 49 IU vitamin (Aykroyd, 1963). The fruit is non climacteric with extremely short shelf

life. Inconsistent fruit yield, non-uniform ripening are the major bottlenecks in cultivation of phalsa. The use of plant growth regulators has become an important component of agrotechnical procedures for most of the cultivated plants and especially for fruit plants (Monselise, 1979). Plant growth regulators like GA₃, NAA and ethrel has proved effective in increasing the size of fruit, improved fruit yield. The effect of plant growth regulators on growth and fruit yield of phalsa cv. Purple Round is a foregone conclusion, but the beneficial effect of plant growth regulators was yet to be fully explored especially in the Sub-Tropics of Jammu where phalsa bushes were successfully grown.

Materials and Methods

The present investigation entitled "Influence of Pre harvest and Pre flowering Sprays of Gibberellic acid, Napthalene acetic acid and Ethrel on flowering behaviour, fruit yield of phalsa cultivar Purple Round under Jammu-Suib tropics" was carried out at RRSS during the year 2015-16 on thirty years old, healthy phalsa bushes already established in the field. The bushes were pruned in mid of December. Geographically the experimental field is situated at an elevation of 332 m above mean sea level and lies at 32° 39' North latitude and 74° 53' East longitude. The treatments were imposed at two time's first spray of plant growth regulators was done at pre-flowering Stage and second spray was done at preharvesting stage. There were ten treatments; $T_1 - 100 \text{ ppm GA}_3, T_2 - 150 \text{ ppm GA}_3, T_3 - 200$ ppm GA₃,T₄- 100 ppm NAA, T₅-150 ppm NAA, T₆-200 ppm NAA, T₇- 750 ppm Ethrel, T₈- 1000 ppm Ethrel, T₉- 1250 ppm Ethrel T_{10} - control (water spray). observations were recorded for flowering behaviour like date of flower initiation, date of end of flowering, duration of flowering, average number of flowers per shoot and fruit yield were observed during the investigation.

The date of opening of first flower for each treatment was considered as the date of start of flowering. Observations were recorded regularly from tagged branches. End of flowering was recorded when almost all the flowers of bush were opened and there after no more flowering took place. The end of flowering i.e. petal fall was recorded in the field and the date recorded thereof. Duration of flowering was calculated by counting the total number of days from commencement of flowering to end of flowering. Four shoots in four directions (east, west, north and south) of all the treatments were tagged. The number of flowers of each treatment was determined on tagged shoots and expressed as number of flowers per shoot. The fruits harvested from each bush were weighed and considered as total yield and expressed in kg/ bush.

Results and Discussion

Data recorded on flowering behaviour (date of flower initiation, date of end of flowering, duration of flowering, average number of flowers per shoot) and fruit yield of phalsa cv. Purple Round is given in (Table 1). The perusal of data presented in Table 1 revealed that flowering commenced earliest i.e. 18th March in bushes treated with 1000 ppm or 1250 ppm ethrel. The untreated bushes were last to initiate flowering i.e. on 25th March. The earliness in flowering is probably because the ethrel releases ethylene when come in contact with the plant tissue which in turn, triggers the shoots to flower early.

The earlier flowering in ethrel may be due to breaking of dormancy of buds and shoots. The reported results of floral induction as results of ethrel treatments to mango trees agreed with Chacko *et al.*, (1972) and Rath and Das (1979). Similar effect of ethrel on induction of early flowering and initiation of full blooming (opening of flowers) had been reported in Langra mango by (Das *et al.*, 1989), in guava (Brahmachari *et al.*, 1996).

The data regarding effect of plant growth regulators on date of end of flowering is presented in Table 1 indicate that among different treatments, the date of end of flowering was earlier in bushes treated with 1250 ppm ethrel (21st April), while it was prolonged (9th May) in untreated bushes.

Ethylene increases the endogenous level of ethylene production in shoot apices and stimulate the formation of separation layer or abscission zones in flowers by stimulating the activity of hydrolases enzyme. The present observation are in conformity with the findings of (Hussain *et al.*, 2008) in pineapple. The duration of flower in response to different growth regulator treatments is given in Table 1. The minimum duration of flower (35.90 days) was recorded with the application of 1250 ppm and maximum duration of flower (46.00 days) was recorded in control treatment. The present observations are in line with the findings of Ghadage *et al.*, (2016) in cashew.

Table.1 Effect of plant growth regulators on flowering behavior and fruit yield of phalsa cv. Purple Round

Treatments	Date of flower initiation	Date of end of flowering	Durationof flowering (days)	Average number of flowers per shoot	Yield (kg/bush)
T ₁ (100 ppm GA ₃)	20 th March	27 th April	39.00	124.74	2.90
T ₂ (150 ppm GA ₃)	20 th March	26 th April	38.50	128.83	3.00
T ₃ (200 ppm GA ₃)	21 st March	29 th April	40.00	121.25	2.78
T ₄ (100 ppm NAA)	22 nd March	6 th May	45.55	123.12	2.75
T ₅ (150 ppm NAA)	21 st March	2 nd May	43.00	124.66	2.86
T ₆ (200 ppm NAA)	23 rd March	7 th May	45.50	119.10	2.50
T ₇ (750 ppm Ethrel)	18 th March	22 nd April	36.50	112.08	2.18
T ₈ (1000 ppm Ethrel)	18 th March	22 nd April	36.00	110.08	2.02
T ₉ (1250 ppm Ethrel)	18 th March	21 st April	35.90	109.42	1.94
T ₁₀ (Control)	25 th March	9 th May	46.00	106.33	1.88
C.D. (p≤0.05)			0.82	5.17	0.21

It is obvious from the data given in Table 1 that the treatment 150 ppm GA₃ resulted in maximum average number of flowers per shoot (128.83) which was statistically at par with 100 ppm GA₃ (124.74) and 150 ppm NAA (124.66). Whereas, minimum average number of flowers (106.33) were recorded in control treatment. The increase in number of flowers per shoot might be due to the effect of gibberellic accelerating acid in the differentiation of inflorescence and rapid elongation of peduncle, leading to full development of flower buds having functional reproductive parts (Ozgvuen and Kaska, 1990

and Parouissi *et al.*, 2002). The result are in conformity with the findings of (Anwar *et al.*, 1990) in strawberry, (Singh, 2006) in phalsa and (Lal *et al.*, 2013) in mango.

It is clear from the data given in Table 1 that all growth regulator treatments except 1250 ppm and 1000 ppm ethrel effect in increasing the yield of phalsa crop as compared to control. Maximum fruit yield of 3.00 kg/bush was recorded with 150 ppm GA₃ which was statistically at par with 100 ppm GA₃ (2.90 kg/bush) and 150 ppm NAA (2.86 kg/bush). Minimum fruit yield of 1.88 kg/bush was

recorded in control treatment. Among all the treatments GA₃ recorded highest fruit yield, this might be due to increase in fruit set, large number of fruits, low percentage of fruit drop, more retention of fruits, better physiology of developing fruits in turn increased berry size, and berry weight. These results are in line with the findings (Randhawa *et al.*, 1959; Singh *et al.*, 1966; Reddy, 1977; Debnath *et al.*, 2011; Singh *et al.*, 2011 and Singh *et al.*, 2017) in phalsa.

From the above studies it can be concluded that the application of ethrel resulted in earliest flowering, early end date of flowering and GA₃ 150 ppm resulted in maximum average number of flowers and fruit yield in phalsa.

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