To study the effect of different growth regulators at various concentration on fruit drop control in mango cv. Lagra

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DOI: https://doi.org/10.22271/chemi.2020.v8.i2v.8962

Abstract
Mango (Mangifera indica L.) is the most important among the tropical fruits of India. It is the premier and choice fruit of India and undoubtedly is one of the best fruits of the world. Mango occupies a pre-eminent place amongst the fruit crops grown in India and is acknowledged as the king of the fruit of the country. The Langra, also known as Banarasi Langra, is a mango cultivar primarily grown in Banaras, Northern India, Bangladesh and Pakistan. This cultivar retains a greenish tinge while ripening. It is normally harvested during the last half of July. Around 2006, it was known to be gaining popularity on the international market. It is considered suitable for slicing and canning. Fruit drop is the serious problem of central and north India. There is a lot of problematic factor for fruit drop like low stigmatic receptivity, defective perfect flowers (having defective embryo sac development at anthesis), poor pollen transference, occurrence and extent of self-incompatibility, competition between developing fruitlets and drought or lack of irrigation and incidence of insect pest and diseases. NAA is a plant growth regulator. The effect of growth regulator i.e. naphthalene acetic acid (NAA) on fruit drop, yield and quality in mango, cultivar ‘Langra’ was studied. Full grown mango plants were sprayed with 10, 20, 30 and 40 ppm NAA at different stages of fruit development. It shows the positive response to all yield attribute. 2,4-D is a also plant growth regulator and also applied in 5, 10, 15 and 20 ppm. The effect of growth regulator i.e. naphthalene acetic acid (NAA) on fruit drop control in mango cv. Lagra was studied. Full grown mango plants were sprayed with 10, 20, 30 and 40 ppm NAA at various concentration on fruit drop control in mango cv. Lagra.

Keywords: Mango cultivar langra, NAA, 2,4-D, GA3

Introduction
Mango (Mangifera indica L.) is the most important among the tropical fruits of India. It is the premier and choice fruit of India and undoubtedly is one of the best fruits of the world. Mango occupies a pre-eminent place amongst the fruit crops grown in India and is acknowledged as the king of the fruit of the country. Mangoes have been cultivated for an estimated 4000 years in India and active selection of superior cultivars has taken place (Mukherjee, 1972; Singh, 1978). It belongs to family Anacardiaceae and is the native of Indo-Burma region (De Candole, 1904). Besides its delicious taste, excellent flavor and attractive fragrance, it is an excellent source of vitamin A (6375-20750 μg /100g ß-carotene) and Vitamin C (6.8-38.8 mg /100g) and mineral contents (Chandra and Chandra, 1997) [2].

Mango is very well adapted to tropical and sub-tropical climate and it thrives well in almost all regions of India from sea level to an altitude of 1500 m. Mango can be grown successfully at a temperature between 24ºC to 27ºC. The tree is hardy in nature and it can endure even the temperature as high as 48º C. It can be grown successfully in low rainfall area of as low as 25 cm and as high as 375 cm per annum. It is gaining more popularity as one of the important dry land horticultural crops. Hence, there is a lot of scope for large scale planting of mango in the vast low rainfall areas of the country provided the temperature and soil are ideal for its growth. Mango is being grown in more than 87 countries of the world but India ranks first in the world with respect to 1.60 million hectares area and 10.78 million tones production (Gandhi, 2012) [3]. India contributes to more than 70 per cent of the total world’s production and this offers bright prospects for further boosting exports. Madhya Pradesh is one of the important mango-growing states of India and the crop occupies an area of 25764 hectare with production of 113801.8 metric tonnes (Anonymous, 2012-13) [1]. Most of the area of Madhya Pradesh is rainfed and vast acreage has an immense potential to improve mango production.
Mango can be used in all the stages of immature, mature and unripe fruits, because of its excellent flavor, attractive fragrance, and beautiful shade of colour, delicious taste and healthful value. Immature and green fruits are used for preparing curries, pickles and chutneys and ripe fruits are utilized in preparing squash, nectar, jam, cereal flakes, custard powder, baby food, mango leather, toffee and several other products.

Mango takes about three months time for fruit set to maturity and ripening. Some varieties may take four months for the same. In the northern region mango is harvested during May to July, while it is November to January in southern region. In South India, the fruits come to harvest earlier by two months as compared to north. Due to multiplicity of cultivars, the harvesting season is spread for over three months. In Madhya Pradesh, ripe mango fruit of North Indian cultivars are available 2-3 weeks earlier, whereas South Indian cultivars are available 2-3 weeks later in the season. This variation has great scope of utilization of mango fruit for export (state and abroad), fresh consumption and processing purpose. Therefore, knowledge of developmental stages of mango fruits is necessary to meet out the demand of standard quality fruits both for processing industry and market supply of the region.

Fruit drop of mango is a serious problem causing heavy losses to the orchardists. Sometimes, 99 per cent crop is lost due to drop of hermaphrodite flowers and immature fruits (Mukherjee, 1949) [1]. The intensity of drop is maximum within 15 days after pollination/anthesis, in which about 60-70 per cent hermaphrodite flowers and immature fruit, drop. The second drop (about 30%) occurs between 28-35 days, when the fruits are at the marble stage. The third drop, which occurs irregularly from 40 days to maturity of fruits is quite low (3-5%), but it is important because immature fruits at the advance stage of growth drop during this period.

Several attempts have been made in the past to control fruit drop in mango cultivars by various workers using auxins (NAA, planofix, 2, 4-D, 2, 4, 5-T, and MENA), gibberellins (GA3), cytokinin, growth retardants (alar, cycocel) and other chemicals (urea, sevin, silver nitrate). However, increase in fruit retention with these chemicals varied between 5.75 and 50.00 per cent (Shukla, 2011) [8]. Major cause attribute to this phenomenon are lack of pollination, low stigmatic receptivity, defective perfect flowers (having defective embryo sac development at anthesis), poor pollen transference, occurrence and extent of self-incompatibility, competition between developing fruitlets and drought or lack of irrigation and incidence of insect pest and diseases (Singh, 1960). Number of research works have been noticed the performance of different chemicals in mango fruit retention and some of the best chemicals and their concentration are 10 ppm 2, 4-D (Singh et al., 1910), 10-20 ppm NAA (Singh et al., 1994) [10], 40 ppm NAA (Gupta and Brahmachari, 2004), 200 ppm GA3 (Rani and Brahmachari, 2013) [7] and 3 per cent urea (Tripathi, 2012), 200 ppm Sevin (Ram and Sirohi, 1909).

### Method and materials

This paper deals with a concise description of the materials used and the techniques adopted during the course of the investigation. The present investigation entitled “Effect of Plant Growth Regulators on Fruit Drop and Physico-Chemical Composition of Mango (Mangifera indica L.) cv. Langra” was conducted at the Department of Crop Sciences, MGCGV, Chitrakoot Satna (M.P.)

### Experimental site

Ten years old mango trees, planted at 10 x 10 m spacing were used for experiment, which was carried out at the research field of Horticulture, Tulsi Krush Vigyan Kendra Ganeewa Rajapur, Dist- Chitrakoot (U.P.) during the year 2001

### Choice of variety

Madhya Pradesh and Uttar Pradesh are reputed for producing early maturing and best quality of Langra mango. Therefore, it was selected for the present investigation.

### Selection of trees

Thirty-nine healthy, vigorous and uniform disease free, bearing trees of about Ten years of age were selected for the experiment. Selected trees were kept under uniform cultural practices, i.e. irrigation, weeding and hoeing etc.

### Preparation of solution

Required quantity of 2, 4-D and GA3 were dissolved in a small quantity of ethyl alcohol but NAA was dissolved in a few drops of NH4OH to avoid precipitation. Then the final volume was made upto one litre with distilled water. The stock solution of the chemicals was diluted with water for preparing the required strength of the foliar spray. Spraying was done with fixed nozzle having mist droplets with sprayer.

### Time of spray

The spraying was done on 20th February and 5th March 2014 from 8.00 a.m. to 5.00 p.m. on a sunny day. Plant growth regulators in different concentrations were sprayed at pin head stage and marble stage. In all the treatments, solutions were sprayed on panicle/fruit and foliage of the tree.

### 3.9 Irrigation

Experimental trees were irrigated whenever necessary.

### 3.10 Observations recorded

To find out the relative performance of the individual treatment the following characters were recorded:

<table>
<thead>
<tr>
<th>Treatments</th>
<th>15 DAS</th>
<th>30 DAS</th>
<th>45 DAS</th>
<th>60 DAS</th>
<th>75 DAS</th>
</tr>
</thead>
<tbody>
<tr>
<td>T0: Without application of plant growth regulators (control)</td>
<td>*35.63a (33.94)</td>
<td>22.35a (14.46)</td>
<td>16.71a (8.27)</td>
<td>13.30a (5.29)</td>
<td>12.56a (4.73)</td>
</tr>
<tr>
<td>T1: 10 ppm foliar application of NAA</td>
<td>42.65cdef (45.90)</td>
<td>25.70abc (18.81)</td>
<td>20.31b (12.05)</td>
<td>17.02bc (8.57)</td>
<td>15.42bc (7.08)</td>
</tr>
<tr>
<td>T2: 20 ppm foliar application of NAA</td>
<td>42.63cdef (45.87)</td>
<td>27.71bc (21.62)</td>
<td>21.67b (13.64)</td>
<td>16.91bc (8.46)</td>
<td>17.02bcde (8.57)</td>
</tr>
<tr>
<td>T3: 30 ppm foliar application of NAA</td>
<td>44.01def (48.27)</td>
<td>28.79c (23.19)</td>
<td>23.53cd (15.94)</td>
<td>19.78ef (11.45)</td>
<td>19.41f (11.04)</td>
</tr>
<tr>
<td>T4: 40 ppm foliar application of NAA</td>
<td>45.51f (50.89)</td>
<td>34.86d (32.67)</td>
<td>30.34e (25.51)</td>
<td>25.38e (18.37)</td>
<td>24.86g (17.67)</td>
</tr>
<tr>
<td>T5: 5 ppm foliar application of 2, 4-D</td>
<td>43.57def (47.51)</td>
<td>26.23abc (19.53)</td>
<td>21.02bc (12.87)</td>
<td>17.67cde (9.21)</td>
<td>17.19e (8.73)</td>
</tr>
<tr>
<td>T6: 10 ppm foliar application of 2, 4-D</td>
<td>(47.51)</td>
<td>(46.27f)</td>
<td>(19.53)</td>
<td>(29.51)</td>
<td>(12.87)</td>
</tr>
<tr>
<td>T7: 15 ppm foliar application of 2, 4-D</td>
<td>41.68bcde (44.22)</td>
<td>27.63bc (21.51)</td>
<td>23.67cd (16.12)</td>
<td>17.85cde (9.40)</td>
<td>17.11cde (8.65)</td>
</tr>
</tbody>
</table>

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increase the fruit development. It can also be concluded that the foliar application of 40 ppm NAA increased fruit retention at 30, 45, 60 and 75 days after spray. The yield and quality of mango cv. Langra and Dashehari. M.Sc. (Ag.) thesis Indira Gandhi Agricultural University, Raipur (2011); 32(2):7-9.

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References