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Original article

Tree age affects postharvest attributes and mineral content in Amrapali mango (Mangifera indica) fruits

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ABSTRACT

The study was carried out to investigate the effect of tree age on postharvest attributes and mineral content of Amrapali mango fruits. Effect of 3 different tree ages (6, 18 and 30 years) on functional components, including the antioxidant activity (AOX), total phenols, total carotenoids, ascorbic acid and minerals like Ca, K, Mg, Fe, Zn, Cu, Mn and B along with total sugars, total soluble solids (TSS) and titratable acidity (TA), respiration rate, polygalacturonase (PG) and pectin methylesterase (PME) activities in Amrapali cultivar were studied. With tree ageing total phenols, ascorbic acid and antioxidant activity decreased whereas total carotenoids increased. Ca diminished and K elevated with the tree age progression while, B, Fe, Cu, Zn, and Mn showed an indefinite pattern. Total soluble solids and total sugars were recorded higher in 18 year old tree fruits. Fruit respiration rate, polygalacturonase and pectin methylesterase activities showed an upward trend with tree ageing. The study

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tree fruits. Fruit respiration rate, polygalacturonase and pectin methylesterase activities showed an upward trend with tree ageing. The sturindicates that fruit produced from middle age group mango orchard (18 year old) suits to the requirement of consumers as well as industry.

Keywords: Mango (Mangifera indica); Tree age; Functional parameter; Mineral

1 1. Introduction

Mango is the most important tropical fruit crop belonging to
the botanical family Anacardiacae. Due to its delicate taste, pleasant aroma and high nutritional value it is considered as King
of fruits. India is the largest mango producing country in the
world. Mango contributes around 20.7% of total fruit production
in the country with annual production of 18.43 million tonnes
(Anonymous, 2015).

9 There are many pre-harvest factors which affect produc-10 tion, quality and storage life of mango fruits. These are culti-11 vars (Seymour et al., 1990), orchard soil management, irrigation water (Duran-Zuazo et al., 2004), rootstock (Dayal et al., 2016), 12 foliar application of nutrients (Sarker and Rahim, 2013; Kare-13 mera and Habimana, 2014; Taha et al., 2014), canopy manage-14 ment (Lal and Mishra, 2007; Asrey et al., 2013), micro leaf area 15 near fruits, bagging (Wu et al., 2013; Haldankar et al., 2015), veg-16 etative vigor of tree, position of fruit on tree (Kawphaitoon et al., 17 2016), harvesting stage (Baloch and Bibi, 2012), use of growth reg-18 ulators (Tandel and Patel, 2011), and insect pests of mango fruit 19

(Whitney et al., 1990; Ketsa et al., 1992; Whiley and Schaffer, 1994;20Maqbool and Mazhar, 2007). The different influences of these pre-
harvest factors have been investigated for enhancing mango pro-
duction by researchers world over.21

Above reviews reflect that in the past majority of researchers 24 were focused on enhancing production and productivity of 25 mango crop. In the era of consumer awareness toward healthy 26 foods, the quantity aspect has taken backseat and quality food 27 production with enhanced functionality is now remaining a key 28 challenge for researchers. 29

The sound postharvest management is one of the weak clinch 30 in Indian mango supply chain and above all quality aspects have 31 virtually remained neglected. There are few reports that tree age 32 also affects the nutritive properties and storage physiology of 33 harvested fruits. Old trees of guava showed decline in mineral 34 absorption and accumulation which significantly affected the 35 physicochemical properties and mineral content of guava fruits 36 (Asrey et al., 2007). Khalid et al. (2012) reported that Kinnow fruits 37 harvested from young (3 year old) tree were rich in high rag mass, 38 rind mass, ascorbic acid, non-reducing sugar, rind manganese 39

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Nirmal Kumar Meena, Ram Asrey

40 and iron content as compared to 6, 18 and 35 year old trees. Tree age factors influenced the pomological parameters of olive fruits 41 and physiochemical characteristics of virgin olive oil (Bouchaala 42 et al., 2014). Young trees of orange produced poor quality fruits 43 than the older tree (Hearn, 1993). Fruit tree age also affected 44 45 fruit yield and physical characteristics of pummelo (Nakorn and 46 Chalumpak, 2016) and apple fruits (Arshad et al., 2014). Ozeker (2000) reported that fruits of 20 year old "Marsh Seedless" grape-47 fruit produce heavier fruit with thin rind of superior quality 48 (juice content, total soluble solids and acidity) as compared to 49 fruits of 34 year old trees. Tree age affected acid content of juice 50 and total soluble solids of Satsuma mandarin (Matsumato et al., 51 1972). 52

53 In Asian nations; particularly in South Asian countries, around 30% mango orchards are too old and in senile condition 54 55 which ultimately affect productivity and quality (Baba et al., 2011). Amrapali is the first commercialized mango hybrid having 56 distinctly dwarf stature, regular and prolific bearer, precocious 57 in nature which is highly suitable for high-density planting 58 59 (Majumdar et al., 1982; Singh, 1996). The pulp of Amrapali is deep 60 orange-red color which may be used for preparing mango nectar and juice and has about 2.5-3.0 times higher carotene content 61 than its parents (Singh et al., 2001, 2012). This study attempts to 62 find the effect of tree age on the postharvest quality attributes 63 and mineral content of Amrapali mango fruits. 64

65 2. Materials and methods

66 2.1. Experimental site

The study was conducted during July-August (peak harvesting time of Amrapali mango) at the experimental farm of Indian Agricultural Research Institute, New Delhi, India. The experimental farm is situated at 28°8′ N and 77°12′ E at an elevation of 229 m above mean sea level.

72 2.2. Selection of tree age group

The mango trees were selected from three different age 73 groups (6, 18 and 30 years) grafted on local seedling rootstock 74 75 (sourced from a single mother plant) and planted at the recommended spacing of 2.5 m \times 2.5 m in the experimental orchard. 76 77 These three tree age groups are important for mango crop as trees start to bear fruits at the age of 5-6 years and their productivity 78 reaches at peak between 15-18 years afterwards it enters into de-79 clining phase after 25-30 years. 80

81 2.3. Fruit sample

82 Fruits were randomly harvested from ten selected trees at commercial maturity stage having total soluble solids (TSS) 83 84 \sim 10%. From each age group uniform size (average weight 230 g per fruit) and healthy fruits free from diseases and insect infes-85 tation were selected; thoroughly de-sapped and surface cleaned 86 with tissue paper. Fruits from each age group were divided into 87 three separate lots, each having 160 fruits for analysis of various 88 functional and nutritional parameters. 89

90 2.4. Study parameters

91 2.4.1. Physical parameters

The specific gravity of mango fruits/stone was calculated by dividing the weight of the fruit/stone by the volume of the fruits/stone as recorded by water displacement method. Mango94stones were obtained by removing peel and pulp of the fruits fol-95lowed by rinsing in potable water and wiping with tissue paper96in order to remove free moisture.97

Fruit firmness was determined by using a texture analyzer 98 (Model: TA+Di, Stable microsystems, UK) under compression test 99 adopted by Jha et al. (2010). Each fruit was compressed using a 100 cylindrical probe (2 mm diameter) having a programmed setting 101 for speed as pre-test, test and post-test speed as 5, 2 and 10 mm/s 102 respectively with probe distance of 10 mm. It was measured at 103 three places (top, mid and bottom) of the individual fruit and ex-104 pressed as mean of three values. First peak force Newton (N) in 105 the force-deformation curve was taken as firmness of the sample. 106

For computing stone/pulp ratio, the pulp was separated from 107 both peel and stone with the help of knife and peeler. The mango 108 pulp and stone were individually weighed. The ratio was deter-109 mined by dividing the weight of stone by weight of pulp. Peel 110 thickness of fresh fruits was measured by digital vernier caliper 111 (Precision 150 Digital Caliper, India). For this, the pulp was care-112 fully removed from the peel with minimum damage or scrap-113 ping to the peel. Fruit peel thickness was measured at three 114 places (top, mid and bottom) and mean value was expressed in 115 millimeter. 116

2.4.2. Biochemical parameters

Total soluble solids (TSS) content was determined by using 118 hand refractometer (Model: PAL-3, ATAGO, Japan) as suggested 119 by Ranganna (1999). Hand refractometer prism was carefully 120 washed with double distilled water and wiped with tissue paper. 121 Juice of mango was extracted by straining of pulp with the muslin 122 cloth. Then two drops of juice were placed on the prism and cor-123 responding refraction index was read and expressed as %. Percent 124 titratable acidity was determined by taking 10 g mango pulp. Ten 125 milliliter of filtered fruit juice was titrated with standard sodium 126 hydroxide (0.1 mol L⁻¹) using phenolphthalein as an indicator for 127 each sample (AOAC, 2006). 128

Similarly, total sugars (%) were determined by taking 50 mL 129 aliquot filtered mango juice titrated with boiling Fehling's solution using methylene blue indicator till brick red color appeared 131 (AOAC, 2006). 132

2.4.3. Physiological parameters

 $\begin{array}{ll} \mbox{The fruit respiration rate was estimated by adopting the static 134 headspace technique using gas analyzer (Model: Checkmate 9900 135 O_2/CO_2, PBI Dansensor, Denmark) followed by Barman and Asrey 136 (2014), and results were expressed as mL CO_2 kg^{-1} h^{-1}. 137 \\ \end{array}$

 $\begin{array}{ll} \mbox{Pectin methylesterase (PME) activity was measured by follow-138 ing the method of Hagerman and Austin (1986), and expressed 139 as μmol min^{-1} g^{-1}$ FW. Polygalacturonase (PG) activity was determined by following the method of Lazan et al. (1995) and expressed as \$\mu\$g galacturonic acid g^{-1} h^{-1} FW. 142

2.4.4. Functional parameters

The analysis of total phenols was carried out by Folin-144 Ciocalteu spectrophotometric method suggested by Singleton 145 and Rossi (1965). Total carotenoids were estimated by the col-146 orimetric method on the spectrophotometer (Model: Jasco V-670 147 UV-VIS-NIR spectrophotometer, Japan) as suggested by Roy 148 (1973). Total phenols and total carotenoids were calculated and 149 expressed in μ g gallic acid g⁻¹ and mg kg⁻¹ pulp respectively. 150 The antioxidant activity was determined by cupric reducing 151 antioxidant capacity method followed by Apak et al. (2004) and 152

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