Effect of 2,4-D pre-treatment on quality during ripening of on-tree longan fruit

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Abstract. The aim of this study was to investigate the effect of 20 mg/L 2,4-D on the quality during ripening of on-tree longan fruits. We investigated external and internal properties of the on-tree longan fruits. The results showed that 2,4-D treatment promotes the growth based on the fruit size and weight. The respiration rate, contents of T SS, total soluble sugar, sucrose, glucose, fructose, and hexose revealed an increasing tendency with advancing the maturity and reached the high status during 110-126DPA. By contrast, the relative electric conductivity and TA content displayed a declining trend during the ripening stage, increase with the senescence. All these pieces of information indicated that 2,4-D treatment could effectively promote the sensory quality of on-tree longan fruit, prolong the harvest time to 118DPA, while CK should harvest before 110DPA.

1 Introduction

Longan (Dimocarpus longan Lour.) is a kind of tropical and subtropical fruit with high business value, it has been cultivated widely in many countries, especially in China, and other Asia-Pacific areas[1-3]. It contains rich nutrients and many other pharmacological materials[4]. Among that sugars play a vital role in fruit quality and are considered as a signal molecule during fruit growth and development which are determined by sugar component and content[5-6]. Sugar of longan fruit mainly consists of sucrose, fructose and glucose with different ratios at different stages[7]. Previous studies have shown that longan fruit sugar receding may occur at unreasonable harvest time, resulting in a change of sugar composition[8], contents of total soluble solids and titratable acidity increase during fruit development, and this is related to the content of sugar[9-11], thus control sugar receding is an important subject to preserve the quality of longan fruit.

Moreover, there are a series of physical and biochemical changes during fruit development which include fruit size and shape and some other alterations inside the longan fruit, for instance, respiration rate, relative electric conductivity and Malondialdehyde (MDA), these factors are also important assessment indicators of fruit quality and shelf-life[8-11]. The diameter in ripening longan fruit is about 15-30 mm, and as a non-climacteric fruit will no longer ripen once harvest, then rapidly deteriorate[12]. Respiration will consume energy if harvest at the non-optimal time, decrease fruit weight, and accelerate senescence[13], meanwhile, during fruit development and ripening stage, cell membrane permeability has changed due to many secondary metabolites produced[10]. Therefore, it is necessary to choose an appropriate harvest time.

It is known that phytohormones are involved throughout the growth and development of plant[16-18]. Some studies have pointed out that auxin plays a significant role in fruit development and ripening[19-21]. 2,4-dichlorophenoxy acid (2,4-D) is a common auxin that could increase fruit size and its effect concentration and date depends on the species[22]. 2, 4-D treatment also could reduce the fruit drop of navel orange[23]. However, it is no more evidence the effect of 2,4-D treatment on the quality of on-tree fruit.

In this work, longan fruit, Dimocarpus longan Lour. cv. Baihuamu was chosen as the material because of its long sugar receding periods to assess the development and quality after 2,4-D treatment before harvesting. It is significant for a better understanding of the effect of 2,4-D, guiding the optimal harvest time.

2 Materials and methods

2.1 Materials and treatments

8-10 years old longan(Dimocarpus longan Lour. cv. Baihuamu) trees with consistent growth and moderate vigor were selected to sprayed 2,4-Dichlorophenoxyacetic acid (2,4-D) before harvesting from an orchard in the Institute of Fruit Tree Research in Guangdong Academy of Agricultural Sciences, Guangzhou, Guangdong Province, China. The spraying time was after full-bloom stage 60 days and 80 days for twice, and the concentration of 2,4-D was 20 mg/kg.

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while the distilled water was control. Three biological replicates were analyzed for each treatment. Fruit samples were taken after 94 days, 102 days (ripening stage), 110 days (full-ripening stage), 118 days, 126 days and 134 days post-anthesis (DPA), more than 50 fruits were harvested from different canopy position. Then transported to the laboratory within 4 h, cut off the stem, fruits with uniformity of maturity, shape and color, and without any diseases and blemishes were selected as materials. Pulps were immediately frozen in liquid nitrogen and stored at -80°C.

2.2 Biological characteristics

Biological characteristics were measured on 30 fruits of each time point. Long diameter, short transverse diameter and vertical diameter were measured by a digital vernier caliper with a precision of 0.01 mm, the analyses were conducted twice. Weight was determined by an electrical scale with a precision of 0.01 g, the analyses were taken 5 times.

2.3 Extraction and assay of titratable acidity (TA), total soluble solids (TSS) and total soluble sugar

Pulp from 20-30 fruits was juiced, then filtered through 4-layer gauze. The filtrate was randomly distributed into 3 groups for analysis of TA, TSS and total soluble sugar. 1 mL of the solution was used to measure TA content by the method of titrating with 0.1 mol/L NaOH. TSS was directly measured using a digital refractometer (PR-32α, ATAGO Co., Ltd., Tokyo, Japan). The content of total soluble sugar was determined according to the method of anthrone sulphuric acid [23].

2.4 Measurement of membrane permeability in pericarp and respiration rate

Membrane permeability was expressed as the relative electric conductivity with % according to the method of Wang et al [25] with small changes. 30 pericarp discs (0.5 mm in diameter) from 10 each fruit, washed twice in deionized water, dried with filter paper then immersed in 25 mL distilled water for 30 min. Initial conductivity was determined with the DDS-307 conductivity meter (Shanghai Scientific Instruments, Shanghai, China). Total conductivity was measured after boiling for 15 min and cooled to 25°C. The relative electric conductivity was expressed as a percentage of the initial value of the total value.

The method of Wang et al [25] was employed to determine the respiration rate with some modifications. 20 fruits were sealed for 2 h and then exhausted air 1 mL, measured by meteorological chromatograph (GC-17A, SHIMADZU, Japan). Respiration rate was expressed as carbon dioxide released per kilogram per hour of fresh weight (mg CO₂·kg⁻¹·h⁻¹). The analyses were carried out for 3 times.

2.5 HPLC analysis of sucrose, glucose and fructose

The assay method of sugar component was reported by Luo et al [8] with a small modification. 1 g pericarp samples were in a microwave oven for 30 s to deactivate the enzymes, and homogenized in a mortar with 2 mL ultrapure water, the homogenate was transferred into a centrifuge tube. The mortar and pestle were rewarshed for 3-4 times, the mixture was diluted to 12 mL. 2 mL aliquot was taken to centrifuge at 13000 rpm for 15 min at ambient temperature. The supernatant was passed through a Sep-Pak C18 Cartridge (Water Corporation, Milford, Massachusetts, USA). Sugar was detected by high-performance liquid chromatography (HPLC) using an Agilent 1200 HPLC system (Agilent Technologies, Waldbronn, Germany) with a G1362A refractive index detector cell maintained at 40°C. A transgenicoc ARB sep Coregel 87C column (CHO-99-5860) With a guard column cartridge (Transgenomic CARB Sep Coregel 87C cartridge) was used. The column was maintained at 80°C with a thermostatted column compartment. The injection volume was 10 μL. Samples were eluted with double-distilled water. The flow rate was 0.6 mL·min⁻¹. Quantification of sugars (sucrose, glucose and fructose) in samples was performed according to external standard solution calibrations (sugar reagents were purchased from Sigma Chemical Co.). The sugar content was calculated according to the peak area of samples and the standard curve of calibrators and expressed as mg·g⁻¹FW.

2.6 Statistical analysis

The variance of data was analyzed Using SPSS software package release 18.0 (SPSS Inc. Chicago, IL, USA). The data were displayed as the means ± standard error.

3 Results and discussion

3.1 Effects of 2, 4-D pre-treatment on external character of on-tree longan fruit

Longan fruit ‘Baihuamu’ development and ripening occurs approximately 110 d after anthesis, the shape of fruits has changed during the time. Some external characters were monitored (Figure 1a,b,c). The transverse diameter (both long diameter and short transverse diameter) showed a tortuous growth trend in all treatments, and the treatment of 2,4-D was higher than CK (Figure 1a,b). It is notable that the transverse diameter value descended at 110 DPA. It may be the errors of sampling. Compared with the transverse diameter, the vertical diameter was increased as a smooth line (Figure 1c), and during the period, a vertical diameter of 2,4-D treatment was higher than CK. This may be because auxin can increase cell size [26]. During the development, fruit weight gradually increased, at the harvest stage, different cultivars weight various from 5 g-20 g [2]. Spraying with 2,4-D promoted the increasing trend, which higher than CK almost 1.261 g and 1.252 g at 102
DPA and 110 DPA, respectively (Figure 1d). These results indicated that 2,4-D could promote longan fruits size and weight. Similar results occurred in the research of auxin on loquat[27].

![Figure 1](https://doi.org/10.1051/e3sconf/202125102031)

### 3.2 Effects of 2, 4-D pre-treatment on content of TA, TSS and total soluble sugar of on-tree longan fruit

The flavor indicators of fruit consist of TSS, total soluble sugar and TA[28]. As Figure 2a shown, the TA contents of both groups displayed a similar pattern: decline firstly and then rising. The 2,4-D treatment reached the lowest at 110 DPA, whereas CK reached the lowest before the ripening stage, and at the later stage of maturity, TA content of 2,4-D treatment was higher than CK. For the TSS contents of CK and 2,4-D treatment, the pattern was opposite to the TA content: increasing firstly then descent(Figure 2b). In comparison to the CK, 2,4-D treatment demonstrated a higher level of TSS content. Figure 2c showed that the total soluble sugar increased with increasing ripening time, after full-ripening time, the total soluble sugar exhibited a downward tendency; While compared with CK, 2,4-D treatment demonstrated a higher level. As figure 2d shown, the tendency of 2,4-D treatment was increasing to 110 DPA and then decreasing; while CK increasing before 102 DPA. Longan is a kind of non-climacteric fruit that will not continue to ripen once harvesting, therefore harvest on time is much important.TSS/TA ratio was applied to determine fruit maturity for most Chinese cultivators[14]. All results indicated that with the development of longan fruit, the flavor and quality gradually became worse, especially after full-ripening stage. 2,4-D treatment could maintain the flavor and prolong the on-tree fruit preservation.

![Figure 2a](https://doi.org/10.1051/e3sconf/202125102031)

![Figure 2b](https://doi.org/10.1051/e3sconf/202125102031)

![Figure 2c](https://doi.org/10.1051/e3sconf/202125102031)

![Figure 2d](https://doi.org/10.1051/e3sconf/202125102031)

**Fig.1.** Effects of 2, 4-D pre-treatment on long diameter(a), short transverse diameter(b), vertical diameter(c) and weight(d) of on-tree longan fruit
3.3 Effects of 2, 4-D pre-treatment on membrane permeability and respiration rate of on-tree longan fruit

Figure 3a illustrated that the relative electric conductivity in CK went up a little at 108 DPA, following by a sharp descent at 110 DPA, then a rapid rise occurred at 118 DPA, descended during the later stage of maturity. However, the relative electric conductivity of 2,4-D treatment showed a rapid descent at 110 DPA, then raised until 126 DPA, finally decreased. The 2,4-D treatment showed a lower level before and after the full-ripening stage. Auxin could promote fruit ripening and softening[29], on the other hand, the fruit reaches maturity and enters senescence caused cell separation[30], all of which leads to higher permeability of 2,4-D treatment at the last maturity stage.

Respiration plays a vital role in the growth and development of plants, and it varies with different stages of fruit development[31]. As shown in Figure 3b, before full-ripening (110 DPA), the fruit respiration rate was raised both in CK and 2,4-D treatment, then gradually decreased. However, the respiration rate of 2,4-D maintained a lower level. As we all know that respiration consumes organic sources resulting in fruit quality worse. Our study showed a lower respiration rate thus retaining the quality of on-tree fruit.

3.4 Effects of 2,4-D pre-treatment on sugar components of on-tree longan fruit

Sugars are kinds of components determining the sensory quality, mainly consist of sucrose, glucose and fructose[32]. The sugar compositions and contents vary with fruit development as well as environmental condition[32,33]. As figure 4a shown, the sucrose content of the CK group increased before 110 DPA then dropped down, in comparison, the sucrose content of 2,4-D treatment ascended before 108 DPA, then decreased after full-ripening. In brief, a higher level of sucrose content revealed in 2,4-D treatment. For the glucose content, fructose content and hexose content, the CK and 2,4-D treatment displayed a similar tendency: decreased at 108 DPA, then ascended until 126 DPA, and finally decline; whereas the contents of 2,4-D treatment peaked at 118 DPA or 126 DPA, then descended, and maintained higher levels after 102 DPA(Figure 4b,c,d). The ratio of sucrose/hexose of 2,4-D treatment decreased during the period of ripening, the ratio of CK increased firstly and then decreased, and remained at a high level all the time(Figure 4e). In a word, before ripening, sucrose, glucose, fructose content of longan fruit increased, afterward decreased. Previous research showed that glucose and fructose contents had a great accumulation coincided with fruit ripening, and dropped in sucrose, the sucrose presented a maximum in the ripening fruit[34]. Sucrose is the major end product in photosynthesis and often hydrolyzed into glucose and fructose[35]. The decrease of sucrose caused the sugar receding in longan fruit which seriously affected the fruit quality[8].
Obviously choosing optimal harvest time is vitally important. In our results that 2,4-D retarded the descent of sucrose, increased the content of glucose and fructose, slowed down the sugar receding rate, maintained the fruit quality.

**Fig.4.** Effects of 2,4-D pre-treatment on sucrose content(a), glucose content(b), fructose content(c), hexose content(d) and ratio of sucrose/hexose(e) of on-tree longan fruit

### 4 Conclusion

In our study, we investigated the properties of 2,4-D treatment during the ripening of on-tree longan fruit. By measuring the appearance trait (size and weight) and some flavor indexes, it has demonstrated that 2,4-D treatment could promote the growth and effectively retain the sensory quality of on-tree longan fruit, prolong the harvest time to 118DPA, while CK should harvest before 110DPA.

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