Methyl Jasmonate Promotes Apple Fruit Degreening Independently of Ethylene Action

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Abstract. Climacteric ‘Fuji’ apples (Malus ×domestica Borkh.) were treated with water, 0.45 mmol·m⁻³ 1-methylcyclopropene (MCP), 2 mmol·L⁻¹ methyl jasmonate (MJ), or both MCP and MJ. Fruit were kept at 20 °C for 17 days after treatment. Ethylene production, respiration, and color change were all inhibited following MCP treatment. Ethylene production following MJ treatment fluctuated below and above that of controls, but was representative of postclimacteric apples at all times. Rates of respiration and color change were enhanced by MJ, even when fruit were previously treated with MCP. The results indicate that MJ can enhance rate of color change and respiration in apple fruit independently of ethylene action.

Jasmonates [jasmonic acid and its methyl ester, methyl jasmonate (MJ)] are cyclopentanone compounds that modulate a wide range of plant responses (Creehan and Mullet 1997; Sembdner and Parthier, 1993), including promotion of chlorophyll degradation (Abeles et al., 1989; Hung and Kao, 1996; Ueda and Kato, 1980). Exogenous MJ promotes chlorophyll degradation and β-carotene accumulation in tomato (Saniewski and Czapski, 1983) and chlorophyll degradation in apple fruit (Perez et al., 1993). Jasmonates also stimulate ethylene biosynthesis, and the endogenous jasmonate concentration rises coincident with the onset of ripening in apple and tomato fruit (Fan et al., 1998b). Exogenous jasmonates accelerate degreening and stimulate ethylene production in apple fruit in a concentration- and developmental stage-dependent manner (Fan et al., 1997, 1998c). Methyl jasmonate is more effective in stimulating degreening than is ethephon (an ethylene releasing agent) on a molar basis. However, it is unclear whether the effect of MJ on degreening is related to its effect on ethylene action. The ethylene action inhibitor MCP (Sisler and Serek, 1997) delays senescence of cut flowers and potted plants at very low concentrations (Serek et al., 1994; Sisler et al., 1996). In this study, MCP was used to investigate whether enhanced degreening of apple fruit by MJ requires ethylene action.

Materials and Methods

Climacteric ‘Fuji’ apples with a mean internal ethylene concentration of 32.5 µL·L⁻¹ (range 10.1–67.9) were harvested from a commercial orchard near Orondo, Wash. Fruit were treated with water (control), 2.0 mmol·L⁻¹ MJ (Bedoukian Research, Danbury, Conn.), 0.45 mmol·m⁻³ MCP (BioTechnology for Horticulture Inc., Burr Ridge, Ill.), or a combination of the two chemicals. There were four replicates of five apples each per treatment. Methyl jasmonate was applied in water as described by Fan et al. (1998c). For MCP treatment, each replicate group of apples was placed in a 20-L glass jar and MCP was injected into the jar through a rubber septum in the lid. After 12 h, fruit were removed from the jar and dipped in water for 2 min. For the combination MCP + MJ treatment, fruit were dipped in 2 mmol·L⁻¹ MJ for 2 min after the MCP treatment. All fruit were enclosed in 20-L jars for 12 h and dipped in 0.02% Tween 20 (v/v) and 1% (v/v) ethanol whether or not MJ or MCP was present. After treatment, fruit were kept at 20 °C, and ethylene production, respiration, and color were measured 0, 1, 2, 3, 5, 7, 9, 11, 13, 15, and 17 d after treatment. Measurement of fruit color, internal ethylene concentration, ethylene production, and respiration rates were as described previously (Fan et al., 1998c). The nonblushed area of each individual fruit was marked, and color on the same spot was recorded as CIEL*a*, b* with a chromometer (model CR-200; Minolta, Japan) using CIE illuminant C and an 8-mm measuring aperture. The chromometer was calibrated with a standard white plate (CR-A43) and d/0 was the illuminant/viewing geometry. Hue and chroma values were calculated from a* and b* (McGuire, 1992).

Results and Discussion

Ethylene production was reduced by MJ 2, 3, and 5 d after treatment (Fig. 1A) but remained at a rate indicative of postclimacteric apple fruit (Burg and Burg, 1962). Ethylene production recovered 7 d after treatment, and was higher than the controls thereafter. There was no detectable ethylene production by MCP-treated fruit throughout the 17-d period, suggesting that continued ethylene biosynthesis by apple fruit requires continuous ethylene action. Another ethylene action inhibitor, diazocyclopentadiene (DACP), also inhibits ethylene production in postclimacteric apple fruit, but the inhibition is only ≈50% and lasts <13 d at 20 °C, even at a concentration of 83 mmol·m⁻³ (Fan et al., 1998a). Blankenship and Sisler (1993) reported that DACP at the same concentration inhibited ethylene production for at least 20 d at 21 °C.

Treatment with MJ overcame some of the inhibition of ethylene production by MCP, but not until 9 d after treatment. Responses elicited by exogenous MJ are dependent on the fruit’s stage of development (Fan et al., 1998b, 1998c; Saniewski et al., 1987). Exogenous jasmonate stimulates ethylene production in preclimacteric fruit but inhibits it in postclimacteric fruit. Ethylene production was not detectable following MCP treatment of climacteric fruit in this study. Renewed ethylene production 9 d after MJ treatment of MCP-treated fruit was similar to that following MJ treatment of preclimacteric fruit (Fan et al., 1997). 1-Methylcyclopropene presumably binds to ethylene receptor(s) and blocks ethylene action (Sisler and Serek, 1997), but the mechanism of action of MJ is not known.

Treatment with MJ stimulated respiration (Fig. 1B), while treatment with MCP inhibited it. The respiration rate of MCP-treated fruit was ≈50% that of controls, suggesting that climacteric respiration requires continuous ethylene action. Ethylene stimulates respiration in almost all fruit tested (Abeles et al., 1992). Although MJ promoted respiration, it inhibited ethylene production immediately after treatment. Furthermore, MJ promoted respiration even when ethylene action was blocked by MCP. This effect does not appear to involve ethylene action, and MJ and MCP appear to have opposing effects on apple fruit respiration.

Treatment with MJ hastened the decrease in hue and the increase in chroma relative to controls (Fig. 2). By definition, chroma indicates the degree of departure from gray toward a pure chromatic color, and hue is the visually perceived color (McGuire, 1992). In this study, the decline in hue angle indicated the change from green to yellow, and the increase in chroma reflected increasing intensity of yellow color. Both chroma and hue values increased following MJ treatment, a change indicative of the development of the deep yellow ground color that develops as ‘Fuji’ apples ripen. In contrast, MCP treatment inhibited color change, suggesting that the change that occurs during normal apple fruit ripening requires ethylene action. Exogenous MJ pro-
Promoted color change even when ethylene production was inhibited by MCP, and the magnitude of the color change was not affected by MCP (Fig. 2). This suggests that MJ promoted color change independently of ethylene action. Emery and Reid (1996) reported that MJ promoted chlorophyll degradation in the absence of ethylene action. Abeles et al. (1989) observed that MJ was as effective as ethylene in inducing chlorophyll degradation in excised cucumber (Cucumis sativus L.) cotyledons. Hung and Kao (1996), however, observed that silver thiosulfate, an inhibitor of ethylene action (Veen, 1985), inhibited MJ-promoted chlorophyll degradation in detached maize (Zea mays L.) leaves, and concluded that MJ-promoted chlorophyll degradation is mediated through an increase in ethylene sensitivity. Considering the evidence for ethylene promotion of chlorophyll degradation (Abeles et al., 1992), jasmonates may modulate color changes in two ways: 1) by promoting ethylene biosynthesis, ethylene in turn promoting color change and 2) independently of ethylene action as evidenced by color changes after treatment with MCP followed by MJ. The color change (yellowing) during apple fruit ripening is associated with chlorophyll degradation, carotenoid synthesis, and metabolism of other pigments. Both ethylene and jasmonate can promote chlorophyll degradation and β-carotene synthesis (Perez et al., 1993; Saniewski and Czapski, 1983), but whether MJ and ethylene mediate pigment metabolism in similar ways is not known.

In summary, MCP blocked ethylene production, and inhibited respiration and color changes in apple fruit, whereas MJ promoted color change and enhanced respiration even in fruit that produced no detectable ethylene. Our results indicate that MJ can act independently of ethylene action on color change and respiration of apple fruit, and may promote color change in two ways, one ethylene-dependent and the other ethylene-independent.

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