Title The effect of MeJA on ethylene biosynthesis and induced disease resistance to Botrytis cinerea in tomato

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#### Abstract

Methyl jasmonate (MeJA), a major derivative of the plant hormone jasmonic acid, plays a critical role in inducing resistance to fungal pathogen. To study the endurance of MeJA-induced resistance and its cause, green mature tomatoes (Solanum esculentum cv. Lichun) were treated with $100 \mu \mathrm{M} \mathrm{MeJA}$ and nordihydroguaiaretic acid (NDGA, LOX inhibitor) at -35 kPa for 0.5 min and incubated at $25 \pm 1^{\circ} \mathrm{C}, 85-90 \%$ RH. Treatment with MeJA reduced disease symptoms in tomato fruit soon after being inoculated with Botrytis cinerea. Lesion size in MeJA-treated fruit was inhibited by $42.5 \%, 27.9 \%$ and $13.9 \%$ respectively $(P<0.05)$ in fruit inoculated 1, 3 and 6 d after treatments. At advanced stages (inoculation carried out 9 and 12 d after treatments), no inhibitory effect of MeJA were found. Ethylene biosynthesis was activated in the response of green mature tomatoes to methyl jasmonate with a rapid (1 d) and enhanced ethylene peak $\left(0.9 \mathrm{ng} \mathrm{kg}^{-1} \mathrm{FW} \mathrm{s}^{-1}\right)$. However the ethylene level was below that of the control from 6 d to 12 d . This rise was closely related with conversion of ACC to ethylene, especially a rise in ACO activity ( 6 h ), which preceded an increase in ACS $(12 \mathrm{~h})$ after MeJA treatment. The development of ethylene biosynthesis was accompanied by a significant increase in LOX activity. Two significant $\mathrm{O}_{2} \cdot{ }^{-}$peaks ( $P<0.05$ ) were detected in MeJA-treated fruit during storage ( $6.18 \mu \mathrm{molg}^{-1} \mathrm{FW} \min ^{-1}$ at 6 h and $5.68 \mu \mathrm{molg}^{-1} \mathrm{FW} \mathrm{min}^{-1}$ at 3 d ). The correlations between LOX, and $\mathrm{O}_{2}{ }^{-}$and ACO activities were $0.75,0.73$ respectively ( $P<0.05$ ). These results indicate that MeJAinduced resistance against $B$. cinerea is durable, MeJA induces LOX and the superoxide radicals formed by LOX may activate ACO and ethylene biosynthesis.


