IMPACT OF POSTHARVEST HANDLING PROCEDURES ON SOFT ROT DECAY OF BELL PEPPERS¹

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Abstract. Bacterial soft rot or stem-end decay caused by Erwinia carotovora (L R. Jones) Holland can be a devastating postharvest problem for bell peppers (Capsicum annuum L). Studies were conducted at commercial packinghouses to de termine the impact of several postharvest handling proced ures on the severity of soft rot decay after a simulated transit and market period. In all cases inoculation with Erwinia greatly increased decay. An overnight delay before packing inoculated fruits resulted in a 23% decrease in marketable peppers after the simulated transit period, compared to only a 4% decrease for uninoculated fruits. In one test, chlorina tion of packinghouse spray-wash water resulted in a 12% in crease in marketable peppers. In another study comparing peppers handled dry to others spray-washed with water or water plus chlorine, washing (water or water plus chlorine) of inoculated peppers resulted in a 17% increase in market able fruits. Vacuum cooling peppers immediately after pack ing reduced decay by about 10% compared to uncooled fruits.

Florida is the leading state in the production of fresh market bell peppers. During the 1981-82 season, Florida growers produced 7.9 million bushels of peppers with a total shipping point value of \$55.6 million (3). Severe postharvest losses may be incurred from bacterial soft rot or stem-end decay (1, 2, 7). Florida shippers have not adopted the hot water treatment recommended for commercial control of bacterial soft rot in peppers (6). The objective of this study was to examine the impact of several postharvest handling practices on soft rot decay of bell peppers.

Materials and Methods

Commercially grown and harvested 'Early Calwonder' peppers were used in 2 experiments during May and June 1982. Both experiments were randomized complete blocks with 4 replications. Freshly harvested peppers were sorted to eliminate damaged fruits. Twenty peppers of medium size $(\approx 130g)$ were selected, jumble-packed in waxed corrugated cartons, and randomly assigned to a treatment.

In the first experiment treatments were a complete fac torial arrangement of 2 inoculations [stem and calyx dipped in sterile buffered saline solution (control) or stem and calyx dipped in a buffered saline solution containing 1.0×10^6 cells ml⁻¹ E. carotovora]; 2 chlorination levels (15-sec spraywash with tap water or tap water plus 150 ppm free chlorine from NaHOCl); and 2 cooling methods (none or vacuum cooled). Vacuum cooling consisted of a 20 min cooling cycle with a minimum absolute pressure of 5.8 mm Hg.

Treatments in the second experiment were a complete factorial arrangement of 2 inoculations (as described above); 2 packing times (packed immediately or packed after an overnight delay at 23° C); and 3 washing methods (none, a

30 sec spray-wash with well water, or a 30 sec spray-wash with well water plus 150 ppm free chlorine from NaHOCl). All peppers in this experiment received a 20 min vacuum cooling following treatment.

Peppers from both experiments were transported to Gainesville and placed in storage at 13°C for 4 days with an additional 2 days at 20°C to simulate transit and market conditions. Following storage at each temperature, peppers were rated for decay with the following scale: $l = none$; $2 =$ slight, not objectionable; $3 =$ moderate, objectionable; $4 =$ severe, decay extended into walls of fruit; 5=extreme, completely decayed. The number of fruits receiving a decay rat ing of 1 was used to calculate the $%$ decay-free. The number of fruits receiving a decay rating of 1 or 2 was used to cal culate the % marketable. An analysis of variance was per formed for each factor using the Statistical Analysis System (4) and means for main effects and interactions were sep arated with orthogonal comparisons where appropriate (10) .

Results

Inoculation with E. carotovora significantly increased decay and reduced the marketable peppers after 6 days of simulated transit and market conditions (Table 1). An overnight delay between Erwinia inoculation and further handling and packing significantly increased decay and re duced the marketable peppers (Table 2). The effect of the handling delay was evident on all peppers after the 2 addi tional days storage at 20°C when the mean decay ratings were 1.8 and 2.0 for immediate and delayed handling, re spectively. This resulted in 9% less marketable peppers for fruits receiving the delay before handling $(73\%$ and 64% marketable for immediate and delayed handling, respectively).

Table 1. Effect of inoculation with Erwinia carotovora on pepper de cay rating, decay-free peppers, and marketable peppers after 6 days of simulated transit and market conditions.

 z Decay rating: $l =$ none; $2 =$ slight, not objectionable; $3 =$ moderate, objectionable; $\frac{3}{4}$ = severe, decay extended into wall of fruit; 5 = extreme, completely decayed.

 y Significantly different from the inoculated treatment at 0.1% level by F test.

The addition of 150 ppm free chlorine to the spraywash water increased the decay-free and marketable peppers in the first experiment (Table 3). The effect of chlorination on decay was most pronounced for peppers inoculated with E. carotovora (decay ratings of 1.3 and 1.2 for control peppers compared to ratings of 2.9 and 2.2 for inoculated peppers spray-washed in water or water plus chlorine, re spectively). There were no significant differences between spray-washing peppers in water or water plus chlorine in the second experiment. However, spray-washing peppers inoculated with $E.$ carotovora in water or water plus chlorine significantly reduced decay and increased the marketable peppers when compared to peppers not washed (Table 4).

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Table 2. Effect of immediate and delayed handling and inoculation with Erwinia carotovora on the decay rating and marketable peppers after 4 days storage at 12.5 °C.

²Decay rating: l = none; 2 = slight, not objectionable; 3 = moderate, objectionable; $\tilde{4}$ =severe, decay extended into wall of fruit; 5 =extreme, completely decayed.

ySignificantly different from the delayed and inoculated treatment at 5% level by \dot{F} test.

^Significantly different from the delayed and inoculated treatment at 1% level by F test.

Table 3. Effect of chlorinating packinghouse spray-wash water on the decay-free and marketable peppers after 6 days of simulated transit and market conditions.

²Significantly different from the chlorinated water treatment at 1% level by F test.

Table 4. Effect of washing treatment and inoculation with Erwinia carotovora on the decay rating and marketable peppers after 6 days of simulated transit and market conditions.

^zDecay rating: $l = none$; $2 = slight$, not objectionable; $3 = moderate$, objectionable; $\check{4}$ = severe, decay extended into wall of fruit; 5 = extreme, completely decayed.

yOrthogonal comparison for the interaction of inoculation x washing (dry vs. [water + water plus chlorine]) was significant at 5% level.

Vacuum cooling reduced decay and increased the decayfree peppers (Table 5). Wetting the peppers prior to vacuum cooling enhanced the removal of heat during the cooling cycle. Pepper temperatures averaged 24.5°C prior to cooling. For dry fruits pepper wall temperatures averaged 23°C after vacuum cooling; calyx and placenta temperatures were about 3°C lower. For wet fruits pepper wall temperatures averaged 20.5°C; calyx and placenta temperatures were about 3°C lower.

Table 5. Effect of vacuum cooling on the decay rating and decay-free peppers after 6 days of simulated transit and market conditions.

zDecay rating: $l = none$; $2 = slight$, not objectionable; $3 = moderate$, objectionable; $\bar{4}$ =severe, decay extended into wall of fruit; 5 =extreme, completely decayed.

ySignificantly different from the vacuum cooled treatment at 1% level by F test.

 x Significantly different from the vacuum cooled treatment at 5% level by F test.

Discussion

The devastating effect of inoculation with Erwinia carotovora (Table 1) underscores the need for strict sanitation measures during pepper harvesting and handling pro cedures. Fundamentally these measures should begin with the exclusion of decayed fruit during harvest and clean picking and transport containers. Fruits should be handled carefully to avoid mechanical injuries which are easily inoculated. Harvest crews should be instructed not to over fill the transport containers.

The fact that harvest needs to be coordinated with packinghouse operations is evident from the detrimental effect that an overnight delay before subsequent handling had on marketable fruits (Table 2). Harvest operations should be curtailed before the packing capacity is exceeded.

Chlorination of vegetable packinghouse water has been recommended for many years (5). However, data on the effectiveness of chlorine for decay reduction are conflicting (8). Our own results are also conflicting. In the first experi ment there was clearly a benefit from adding chlorine to the spray-wash water (Table 3). In the second experiment wash ing with water alone was as effective in reducing decay as washing with chlorinated water (Table 4). Both washing treatments were far better than not washing inoculated peppers. Chlorination of the wash water is relatively in expensive and serves to prevent the buildup of inoculum on the rollers, brushes, and conveyors that the peppers contact further along the packing line. Therefore, it would seem most practical to include chlorination of spray-wash water as a routine sanitation practice for bell peppers.

Vacuum cooling was clearly beneficial when compared to not cooling peppers (Table 5). This result was consistent with an earlier study (9) where thoroughly cooling peppers to 10°C soon after harvest effectively delayed soft rot decay. Forced-air cooling is another effective cooling method for peppers (9), but the portability of vacuum cooling equip ment may make it more widely available to Florida pepper shippers.

None of the handling procedures studied in this paper provided absolute control of postharvest decay caused by Erwinia carotovora. However, packers and shippers can re duce their losses by careful attention to detail during han dling. Modern pepper harvesting and packing should in clude strict sanitation procedures to prevent fruit inocula tion, prompt careful handling to prevent mechanical dam age, and thorough cooling soon after harvest.

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EFFECT OF SCRUBBING ETHYLENE DURING STORAGE OF TOMATOES¹

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Abstract. In 3 storage tests, Florida mature-green tomatoes were stored at 55°F for 1, 2 and 3 wk in sealed chambers with and without the circulating air being filtered through potassium permanganate/aluminum oxide to remove ethylene from the circulating air. After 55°F storage, the toma toes were evaluated each day for color and held at 70°F until each tomato reached full red color, at which time firm ness and decay were also recorded. Generally, the removal of ethylene from the storage atmosphere only increased the number of days for the tomatoes to reach full red color after 1 wk of storage and they were firmer than tomatoes stored without ethylene removal after all storage periods. Removal of ethylene during storage did not affect the amount of de cay.

Florida is the largest producer of fresh market tomatoes and for many years shippers and receivers have been inter ested in exporting Florida tomatoes to Europe. Florida tomatoes generally do not meet commercial trade standards on arrival in Europe. The most successful shipments have been of vine-ripened fruit or large-sized mature-green toma toes that have been treated with ethylene. Many European receivers have indicated that they would import larger quantities if tomatoes arrived consistently in good condition and of uniform color and size.

Recommendations for the transit and storage of ethylenetreated (ET), mature-green tomatoes generally specify that temperatures be maintained at 55° to 70°F, and for ripe tomatoes at 45° to 50° F (1, 3, 6, 8, 9). There are many studies which describe the climacteric (respiration) cycle of tomatoes during the ripening process (7), and studies de scribing the roles of ethylene $(4, 5)$ and $CO₂$ (2) activity during ripening. Generally, Florida producers harvest their tomatoes during the mature-green stage and then treat them with ethylene for the initiation of the ripening process. However, there is very little known about the effects of scrubbing (removal) endogenously produced ethylene from the storage area of ethylene-treated mature-green toma toes during the ripening process. The objectives of this study were to determine the effects of scrubbing ethylene from the storage area on tomato ripening and quality.

Materials and Methods

An initial experiment with 3 replications was conducted to determine if Ethysorb® (potassium permanganate/alu minum oxide) absorbs ethylene during storage, air samples were obtained every 24 hr from each of 2 chambers and analyzed for presence of ethylene. Each chamber contained 60 gasses ET mature-green tomatoes (cultivar FTE-12), 20 each of 3 ripening stages (turners, pinks and light reds). In both chambers, the air was circulated, in one chamber through 0.53 oz of Ethysorb® and in the other chamber without Ethysorb® The gas samples were analyzed on a gas chromatograph.

Mature-green tomatoes (cultivar FTE-12) were obtained from a Dade County packinghouse. Three tests were con ducted at ca. 5-wk intervals starting in January 1983. The mature-green tomatoes were commercially packed but not gassed (ET to initiate ripening) at the packinghouse. To matoes (size 6 x 6) were brought to the Orlando USDA Laboratory immediately after packing. Immediately on ar rival in Orlando, the tomatoes were gassed for ca. 72 hr with 50 ppm ethylene at 70°F (21°C). After gassing, fruit were color sorted into lots of breakers and pinks in accordance with United States Standards for Grades of Fresh Tomatoes (10).

For each treatment, 20 (6.5 lb.) tomatoes were placed on polystyrene foam trays. Each tray was placed in a sealed chamber (28 x 28 x 14 inches) where the contained air was continuously circulated by means of an electric pump and tubing. The temperature of all chambers was maintained at 55°F and 80-85% relative humidity. One tray of each of the following treatments was stored for 1, 2 or 3 wk. The follow ing is a breakdown of the treatments:

- Treatment 1 $(B+)$ —1 tray of breaker tomatoes in a sealed chamber and circulated air.
- Treatment 2 $(B-)$ –1 tray of breaker tomatoes in a sealed chamber and circulated air filtered through 0.53 oz of Ethysorb®.
- Treatment 3 $(P+)$ —1 tray of pink tomatoes in a sealed chamber and circulated air.
- Treatment 4 $(P-)$ -1 tray of pink tomatoes in a sealed chamber and circulated air filtered through 0.53 oz of Ethysorb®.

Tomatoes were removed from the test chambers after each specific storage period of 1, 2, or 3 wk, and were then stored at 70°F (21°C) and 88-92% relative humidity. Each fruit was inspected daily and removed on reaching the full red color stage. All color measurements were made with a Hunter® colorimeter (Model D25-9) signal processor with a Model D25-L optical sensor. The tomatoes were called red when the external 'a' value of readings on the colorim eter averaged 32.0 which was judged to be fully red and matched the USDA standards. External color readings were

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