Optimizing the Storage Temperature and Humidity for Fresh Cranberries: A Reassessment of Chilling Sensitivity

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Abstract. Studies were conducted over three seasons to determine the relationship of temperature and humidity on the storage life of fresh cranberry (Vaccinium macrocarpon Aiton) fruit. Each year, cranberries harvested from four commercial bogs were stored at temperatures ranging from 0 to 10°C in combination with relative humidities (RH) ranging from 75% to 98%. Fruit were stored under these conditions for up to 6 months and were evaluated monthly for marketability, decay, physiological breakdown, weight loss, and firmness immediately after removal and after an additional week at 20°C. The percentage of marketable fruit declined substantially over time in all storage conditions with 41% to 57% becoming unmarketable after 2 months as a result of both decay and physiological breakdown. Relative humidity had a greater effect on fruit storage life than temperature and after 5 months, the amount of marketable fruit stored in high (98%) and medium (88%) RH was 71% and 31% less than that stored in low (75% to 82%) RH. Rates of fresh weight loss increased as RH in storage increased and was 0.41%, 0.81%, and 0.86% per month in fruit stored in high, medium, and low RH, respectively. Fruit firmness was not significantly affected by RH. The effects of storage temperatures ranging from 0 to 7°C on marketable fruit after 2 to 5 months of storage were not significant. Only fruit stored at 10°C consistently had fewer marketable fruit when compared with fruit stored at lower temperatures. Storage temperature had no significant effect on decay incidence. However, physiological breakdown was greatest in fruit stored at 10°C. Rates of fresh weight loss increased with storage temperature, ranging from 0.35% to 1.17% per month for fruit stored at 0 to 10°C, respectively. Contrary to previous reports, no evidence of chilling injury was found in cranberry fruit stored at 0°C. Results suggest that cranberry fruit should be stored at 0 to 7°C and 75% to 82% RH to retain marketable fruit.

There is an increased interest in extending the market life of fresh cranberries as a result of the growing demand for fresh, healthy fruits. To supply this growing demand, fruit quality must be maintained and postharvest losses reduced during extended storage and marketing periods. However, for a variety of known and unknown reasons, successful storage of fresh cranberries has been variable and fruit loss is often excessive.

Unlike most fruit, the optimum conditions for the storage of fresh cranberry fruit are not clearly defined. In various handbooks published in the past 20 years, recommended storage temperatures range from 2 to 7°C (Hardenburg et al., 1986; Kader, 1997; Kasimire and Thompson, 1992; Lidster et al., 1988; Prange, 2004; Spayd et al., 1990). Similarly, the relative humidity (RH) for storage recommended in these handbooks ranges from 80% to 95%. These recommendations are based on a limited number of studies, most being conducted over 60 years ago (Levine et al., 1941; Wright et al., 1937). These variable recommendations reflect the lack of clear differences reported in these studies, which may reflect inherent fruit properties that are influenced by cultural conditions, cultivars, harvest method, postharvest handling, and storage environment.

Postharvest loss of cranberry fruit is primarily the result of physiological breakdown and decay (Forney, 2003). Physiological breakdown is associated with overmature fruit (Doughty et al., 1968), bruising (Patterson et al., 1967), chilling injury (CI) (Hruschka, 1970), freezing (Bristow and Patten, 1995), extended water immersion (Ceposina and Stretch, 1983), and anoxia (Stark et al., 1974) and is characterized by a dull appearance, rubbery texture, and diffusion of red pigment throughout the fruit flesh (Bristow and Patten, 1995). Decay is caused by a complex of fungal organisms, including Allantophomopsis lycopodina (Hohn.) Carris (black rot), Allantophomopsis cytispora (Fr.:Fr.) Petr. (black rot), Strasseria geniculata (Berk.& Br.) Höhnel (black rot), Coleophoma empetri (Rostr.) Petr. (ripe rot), Fusarium putrefaciens Shear. (end rot), Phylllosticta elongata (Shear) Shear (berry speckle), Physalospora vaccinii (Shear) Arx & E. Müller (blotch rot), and Botrytis spp. (yellow rot) (Boone, 1995a, 1995b; Carris, 1995; Caruso, 1995; Oudemans et al., 1998; Pepin and Boone, 1995). Infection of the fruit may occur during bloom or wet harvest in the case of fungi causing black rot. Decay is characterized by external lesions and often only part of the internal flesh is red, whereas the unaffected flesh remains white. Unlike most postharvest decays in other crops, there is little spread of disease from infected to healthy fruit in storage (Oudemans et al., 1998).

Contributing to the uncertainty of optimum storage temperature is the fact that cranberries are reported to be chilling-sensitive and may develop physiological breakdown (CI) when stored at temperatures less than 2.2°C (Levine et al., 1941; Wright et al., 1937). Chilling sensitivity of fruit may be affected by environmental, cultural, or genetic factors, which can alter their response to storage temperature (Patterson and Reid, 1990). In addition, fungal species and races causing fruit decay may vary depending on growing location (Caruso and Ramsdell, 1995). Therefore, decay development may be different in similar storage environments depending on the pathogens present (Bristow and Patten, 1995).

Most berry crops benefit from RH in storage of 95% or greater, which is effective in reducing water loss that would result in shriveling and physiological stress. However, reports indicate that cranberries store better in lower RH with suggested optimum storage RH being 65% to 70% (Stark et al., 1974) or 70% to 75% (Wright et al., 1937). However, in storage handbooks, recommendations are 80% to 90% (Lidster et al., 1988) and 90% to 95% RH (Hardenburg et al., 1986; Kader, 1997; Spayd et al., 1990). These higher recommended RHs are most likely influenced by the positive response of other fruit to high RH.

To clarify storage recommendations, a study was conducted to reassess the effects of temperature and RH on cranberry storage life. The objectives of this study were to determine the relationship among storage temperature, RH, and cranberry fruit storage life and assess the chilling sensitivity of cranberry fruit.

Materials and Methods

Fruit. Storage experiments were conducted over three harvest seasons from 2001 to 2003
to evaluate the effects of temperature and RH on the quality and storage life of fresh cranberries. Each year, mechanically harvested fruit from four commercial bogs were stored for up to 6 months. In 2001, ‘Stevens’ cranberry fruit were harvested from two bogs in Aylesford, Nova Scotia, using a wet-rake harvester and one bog in Aylesford, Nova Scotia, and one bog in Wisconsin using wet-beater harvesters. In 2002, ‘Stevens’ cranberry fruit were harvested from four bogs in Aylesford, Nova Scotia; two bogs were harvested using a wet-rake harvester and two were harvested using a wet-beater harvester from a second grower. In 2003, cranberry fruit were harvested from four bogs in Nova Scotia, representing four commercial growers and three cultivars. Two bogs of ‘Stevens’ were harvested using dry-rake harvesters from commercial bogs in Amherst and Lawrencetown, Nova Scotia, ‘Pilgrim’ fruit were harvested from a commercial bog in Centrela, Nova Scotia, using a wet-rake harvester, and ‘Bergman’ fruit were harvested from a commercial bog in Aylesford, Nova Scotia, using a wet-rake harvester. All fruit were harvested between mid-October and early November. Fruit were transported to the Atlantic Food and Horticulture Research Center in commercial lug bags and held at 5°C and 80% RH for no more than 14 d before being used in storage experiments. Fruit from each bog was randomized and placed into Olympian SP5 5.25-inch square nursery pots (Nursery Supplies, Chambersburg, PA) with a volume of 1.95 L and a depth of 15.3 cm, which is similar to the depth of commercial storage containers. Individual containers held ≈900 g of fruit. Containers were used to hold fruit for storage experiments and served as units for sampling and evaluating fruit.

Storage conditions. For storage experiments, containers of cranberries were held in 60-L plastic tubs. Each tub contained 12 containers of cranberries, six from each of two bogs. Containers of cranberries were randomized and stacked in a four-container × three-level arrangement inside each tub. Containers of cranberries were maintained in the same position throughout the storage experiment. Tubs were sealed with a lid that had a 4-mm diameter hole in the center to provide aeration.

Temperature and humidity. Tubs containing fruit were stored in walk-in controlled-
temperature chambers maintained at 0, 3, 5, 7, or 10°C to study the effects of temperature on fruit storage life. Two chambers for each temperature were used each year with bogs being nested in chambers. Tubs containing fruit from two of the four bogs were stored in one series of chambers at each temperature, whereas tubs containing the fruit from the other two bogs were stored in the second series of chambers.

In the 2001 season, no attempt was made to control RH in the tubs. However, chamber temperatures were monitored hourly and the temperature and RH within tubs was measured after 2, 4, and 6 months. From these observations, it appeared that temperature was maintained within 0.5°C of the target temperature and RH averaged ≈82%. In 2002, “low” or “medium” RH was obtained by placing low RH tubs in the temperature-controlled chambers uncovered as done in 2001, whereas medium RH tubs were sealed in a ventilated plastic garbage bag containing one 4-mm hole near the bottom of the tub. Relative humidity and temperature of the room and each tub were measured immediately before fruit removal using a RH5100 humidity dew point temperature meter (Omega, Stamford, CT) or a 21X Micrologger with a temperature and relative humidity probe (Campbell Scientific, Edmondston, Alberta, Canada). Measurements were obtained by placing the probe ≈30 cm into the tub through a resealable hole in the top of the lid. The temperature in the low RH tubs averaged 0.2 to 0.9°C higher than target storage temperature over the 6 months of storage (Table 1). The RH in the low RH tubs ranged from 77% to 86% and averaged 82%. The temperature in the medium RH tubs averaged 0.2°C higher than temperatures in the low RH tubs. The RH in the medium RH tubs ranged from 81% to 88% and averaged 86%.

In 2003, 1.5 kg of salts was placed in the bottom of each tub to maintain three different RH levels. A 3.5 × 18 × 27-cm plastic tray was used to elevate the containers of berries above the salt. The salts used to obtain an “low,” “medium,” and “high” RH were calcium nitrate (CaNO₃), sodium chloride (NaCl), and potassium nitrate (KNO₃), which have equilibrium RHs of 66%, 76%, and 96%, respectively, across the 0 to 10°C temperature range (O’Brien, 1948). Tubs were placed in the temperature-controlled chambers and sealed in ventilated plastic garbage bags. Temperature and RH of each tub in one of the two chambers at each temperature was monitored using HOBO data loggers (Onset Computer Corp., Bourne, MA) suspended in the center of the tub. Over the 6 months of storage, the temperature in the tubs averaged within 0.2 to 0.8°C of the target storage temperatures (Table 1). The RH in tubs containing the CaNO₃, NaCl, and KNO₃ averaged 75%, 88%, and 98%, respectively, during the storage period.

Fruit quality analysis. At harvest, three containers of fruit from each cranberry bog were evaluated for initial quality. In addition, fruit quality was assessed at monthly intervals for 6 months, at which time one container of cranberries representing each bog and storage treatment was removed from storage. Fruit were evaluated for color, fresh weight loss, firmness, decay, physiological breakdown, and marketable fruit. After fruit showing signs of decay or physiological breakdown were removed from each sample, 340 g of marketable fruit were bagged in perforated polyethylene cranberry bags and held for 1 additional week in air at 20°C. After the week holding period, fruit quality was evaluated as previously described.

Color was assessed as white (less than 50% red) or red (greater than 50% red). Decayed fruit were identified by discoloration and softening. Fruit with physiological breakdown were identified by a dull appearance, rubbery texture, and diffusion of red pigment throughout the flesh. Marketable fruit was calculated as the fruit remaining after all fruit with decay or physiological breakdown were removed from the sample. Fruit identified as white, decay, physiological breakdown, and those considered marketable were all weighed and expressed as a percentage of the total fruit weight.

Firmness was measured using a FirmTech firmness testing instrument (Bio-Works, Stillwater, OK). A subsample of 25 marketable cranberries from each container was placed on its side in the index holes of the FirmTech1 turntable. A 15-mm diameter load plate compressed each fruit at a rate of 10 mm·s⁻¹ to a maximum threshold force of 1.96 N. The average firmness of each cranberry subsample was recorded in N·mm⁻¹ of deformation. In the event that a 25-cranberry subsample was not available for analysis, as a result of low percentages of

Table 1. Recorded temperature (T) and relative humidity (RH) in plastic tubs during storage of cranberry fruit during the 2002 and 2003 storage seasons.¹

<table>
<thead>
<tr>
<th>Season</th>
<th>Humidity</th>
<th>Target temperature</th>
<th>Low T (°C)</th>
<th>Low RH (%)</th>
<th>Medium T (°C)</th>
<th>Medium RH (%)</th>
<th>High T (°C)</th>
<th>High RH (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>2002</td>
<td>Low</td>
<td>0°C</td>
<td>0.5 ± 0.5</td>
<td>86 ± 2</td>
<td>0.7 ± 0.5</td>
<td>88 ± 1</td>
<td>0.0 ± 0.3</td>
<td>72 ± 2</td>
</tr>
<tr>
<td></td>
<td></td>
<td>3°C</td>
<td>3.2 ± 0.3</td>
<td>85 ± 3</td>
<td>3.5 ± 0.3</td>
<td>88 ± 1</td>
<td>2.5 ± 0.2</td>
<td>81 ± 9</td>
</tr>
<tr>
<td></td>
<td></td>
<td>5°C</td>
<td>5.9 ± 0.6</td>
<td>84 ± 4</td>
<td>5.9 ± 0.7</td>
<td>88 ± 1</td>
<td>6.2 ± 0.5</td>
<td>71 ± 3</td>
</tr>
<tr>
<td></td>
<td></td>
<td>7°C</td>
<td>7.5 ± 0.4</td>
<td>80 ± 3</td>
<td>7.8 ± 0.5</td>
<td>86 ± 2</td>
<td>6.8 ± 0.1</td>
<td>76 ± 5</td>
</tr>
<tr>
<td></td>
<td></td>
<td>10°C</td>
<td>10.8 ± 0.3</td>
<td>77 ± 3</td>
<td>11.1 ± 0.5</td>
<td>81 ± 4</td>
<td>9.9 ± 0.3</td>
<td>73 ± 10</td>
</tr>
<tr>
<td>2003</td>
<td>Low</td>
<td>0°C</td>
<td>0.1 ± 0.2</td>
<td>91 ± 1</td>
<td>-0.1 ± 0.3</td>
<td>91 ± 1</td>
<td>-0.4 ± 0.3</td>
<td>93 ± 1</td>
</tr>
<tr>
<td></td>
<td></td>
<td>3°C</td>
<td>2.6 ± 0.2</td>
<td>85 ± 4</td>
<td>2.6 ± 0.2</td>
<td>85 ± 4</td>
<td>2.9 ± 0.0</td>
<td>97 ± 2</td>
</tr>
<tr>
<td></td>
<td></td>
<td>5°C</td>
<td>5.5 ± 0.2</td>
<td>90 ± 2</td>
<td>5.5 ± 0.2</td>
<td>90 ± 2</td>
<td>5.7 ± 0.4</td>
<td>99 ± 2</td>
</tr>
<tr>
<td></td>
<td></td>
<td>7°C</td>
<td>6.5 ± 0.1</td>
<td>84 ± 6</td>
<td>6.5 ± 0.1</td>
<td>84 ± 6</td>
<td>6.7 ± 0.1</td>
<td>91 ± 6</td>
</tr>
<tr>
<td></td>
<td></td>
<td>10°C</td>
<td>10.0 ± 0.2</td>
<td>88 ± 4</td>
<td>10.0 ± 0.2</td>
<td>88 ± 4</td>
<td>9.7 ± 0.3</td>
<td>98 ± 2</td>
</tr>
</tbody>
</table>

¹In the 2002 season, the “low” and “medium” RH were maintained by sealing the “medium” RH tubs in a ventilated plastic bag. In the 2003 season, the “low,” “medium,” and “high” RHs were maintained by placing CaNO₃, NaCl, and KNO₃ salts, respectively, in the bottom of tubs and sealing the tubs in ventilated plastic bags.

Mean ± SD; n = 20 for 2002, n = 138 for 2003.
marketable fruit, an average was obtained from the available fruit.

Statistical analysis. Data from all 3 years were combined for analysis. The RH treatments were grouped as low, medium, and high. The low RH group was comprised of all 2001 data, the 2002 low RH treatment, and the 2003 CaNO₃ treatment; the medium RH group was comprised of the 2002 medium RH treatment and the 2003 NaCl treatment; and the high RH group was comprised of the 2003 KNO₃ treatment. To account for the unbalanced design, data were analyzed using the Restricted Maximum Likelihood (REML) procedure of GenStat (Payne, 2003) to determine fixed effects among temperature (0, 3, 5, 7, 10 °C), RH (low, medium, high), storage time (1, 2, 3, 4, 5, 6 months), and their interactions. The REML procedure tested variation in the effects of temperature and RH among years and cultivars and within years to account for unbalanced treatments and treatment combinations. Year was assigned as a random factor in the model, which removed variation differences resulting from year. The model used the common cultivar (‘Stevens’) among years to adjust year-to-year variation and variation resulting from the additional cultivars in year 3. For each temperature, berries from four cranberry bogs were randomly assigned to two chambers. Within a chamber, the humidity treatments were randomized to tubs and removals were randomized across levels within a tub. Data expressed as a percentage were transformed using an angular transformation to normalize the distribution. Pairwise t-probabilities among the levels of the main effect and among treatments within the table of interactions were calculated to determine mean separation. Evaluations made immediately after storage and after the 7-d period at 20 °C were analyzed separately.

Results

Initial quality. Initial fruit quality is described in Table 2. On average over the three seasons, 92% of the fruit was judged to be marketable before being placed into storage. Of the unmarketable fruit, ≈2.7% showed signs of decay and 4% had physiological breakdown. The amount of decay and physiological breakdown varied by bog, which is reflected in the large standard deviations associated with these values. After 7 d at 20 °C, 79% of the fruit remained marketable. During these 7 d, ≈7% of the fruit decayed and 14% developed physiological breakdown. In addition, most of the white fruit turned red. Fruit firmness declined by 4% and fresh weight by 1.4%.

Storage humidity. Relative humidity in storage affected the quantity of marketable cranberry fruit to a greater extent than temperature did with the greatest losses of marketable fruit occurring in the high RH (Table 3; Fig. 1A). This effect of RH on marketable fruit increased during storage. After 1 month in storage, marketable fruit stored in the three levels of RH were not significantly different. However, after 2 months, the high RH stored fruit had 37% fewer marketable fruit than the low RH stored fruit. This difference increased further during storage and after 5 months, the medium and high RH storages produced 31% and 71% fewer marketable fruit than the low RH storage, respectively.

The loss of marketable fruit continued at a higher rate in the high RH stored fruit after fruit were removed from storage and marketable fruit held an additional 7 d at 20 °C (Table 3; Fig. 1B). After 1 month plus 7 d, high RH stored fruit had 13% less marketable fruit than those stored in low RH, whereas after 3 months plus 7 d, the medium and high RH stored fruit had 10% and 41% fewer marketable fruit, respectively, than the low RH stored fruit.

The different levels of RH affected both decay and physiological breakdown (Table 3; Fig. 1C, E). After 1 month of storage, decay was greatest in fruit stored in high RH at 14% compared with 5% and 9% decay in fruit from the medium and low RH storage, respectively. However, as the percentage of decayed fruit increased during the first 4 months of storage, these differences were no longer significant. Physiological breakdown was not significantly different after 1 month of storage from the three storage RHs. However, after 2 months, differences between high and low or medium RH stored fruit were observed, which increased through 5 months of storage. After 5 months, physiological breakdown comprised 52% of the fruit stored in high RH compared with 36% and 31% for the medium and low RH stored fruit, respectively.

Decay and physiological breakdown of marketable fruit held 7 d at 20 °C after storage was also affected by the storage RH (Table 3; Fig. 1D, F). Decay was greatest in fruit previously stored in the high RH for the first 4 months, reaching 34% after 3 months plus 7 d. At this time, decay of fruit from the medium RH reached 16%, whereas that from the low RH was 9%. Rates of physiological breakdown in fruit that had been stored in the low RH remained fairly constant ranging from 10% to 13%. However, physiological breakdown of fruit stored in high RH was elevated and ranged from 16% to 22% after 2 to 6 months.

Fresh weight loss was reduced by elevated RH (Table 3; Fig. 1G). Rates of weight loss, determined by linear regression, for fruit stored in the low, medium, and high RH were

### Table 2. Average quality of cranberry fruit immediately after harvest (initial) and after 7 d at 20 °C.

<table>
<thead>
<tr>
<th>Fruit quality</th>
<th>Initial</th>
<th>Mean</th>
<th>sd</th>
<th>Plus 7 d at 20 °C</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Marketable fruit (%)</strong></td>
<td>92.1</td>
<td>3.9</td>
<td>79.0</td>
<td>10.1</td>
</tr>
<tr>
<td><strong>Decay (%)</strong></td>
<td>2.7</td>
<td>4.0</td>
<td>7.4</td>
<td>7.9</td>
</tr>
<tr>
<td><strong>Physiological breakdown (%)</strong></td>
<td>4.4</td>
<td>3.9</td>
<td>13.7</td>
<td>6.5</td>
</tr>
<tr>
<td><strong>Fruit less than 50% white (%)</strong></td>
<td>1.6</td>
<td>1.8</td>
<td>0.1</td>
<td>0.4</td>
</tr>
<tr>
<td><strong>Firmness (N-mm⁻¹)</strong></td>
<td>6.6</td>
<td>0.7</td>
<td>6.3</td>
<td>0.7</td>
</tr>
<tr>
<td><strong>Weight loss (%)</strong></td>
<td>—</td>
<td>—</td>
<td>1.4</td>
<td>0.5</td>
</tr>
</tbody>
</table>

*Values are the average of fruit from 12 commercial bogs (four bogs in each of three seasons).
When marketable fruit were bagged and held an additional 7 d at 20 °C, fruit previously stored at higher temperatures maintained more marketable fruit than those previously stored at 0 or 3 °C (Table 3; Fig. 2B). The bagged fruit previously stored at 7 or 10 °C for 1 month had 75% marketable fruit compared with 66% from the 0 °C storage. This difference was accentuated after 6 months of storage, where marketable fruit was maintained at greater than 70% in fruit previously stored at 7 or 10 °C compared with only 47% when stored at 0 °C. Maintenance of fruit quality after storage at 7 or 10 °C, and to a lesser extent 5 °C, was maintained during the additional 7-d holding period throughout the 6-month experiment at levels similar to those measured for the initial levels. However, marketability of fruit that had been stored at either 0 or 3 °C declined with increasing storage duration.

Loss of marketable fruit was caused by increases in decay and physiological breakdown (Fig. 2C, E). When fruit were evaluated immediately after storage, storage temperature did not significantly affect decay (Table 3). Decay increased from 9% after 1 month to 46% after 6 months of storage. Physiological breakdown was affected by storage temperature and was least at 0 or 3 °C and greatest at 10 °C. After 1 month of storage, 14% and 25% of fruit developed physiological breakdown when stored at 0 and 10 °C, respectively. These differences were maintained through 5 months of storage at which time 33% and 49% developed physiological breakdown. However, after 6 months, these differences were lost as a result of increases in decay.

Both decay and physiological breakdown were affected by prior storage temperature when marketable fruit were bagged after storage (Table 3). Decay was greater in fruit previously stored at the low temperatures of 0 and 3 °C and increased during the first 4 months of storage (Fig. 2D). Similarly, physiological breakdown was also greater in 0 and 3 °C stored fruit (Fig. 2F). However, quantities of fruit developing physiological breakdown remained fairly constant during the first 4 months of storage, ranging from 8% to 18%, which were similar to the physiological breakdown observed with the initial levels. After 5 months of storage, physiological breakdown of fruit stored at 0 °C increased to ~28%. This increase coincided with a decrease in decay resulting in a similar drop in marketable fruit.

Fresh weight loss increased as storage duration and temperature increased (Table 3; Fig. 2G). Rates of fresh weight loss, determined by linear regression, were 0.35%, 0.40%, 0.59%, 1.13%, and 1.17% per month for fruit stored at 0, 3, 5, 7, and 10 °C, respectively. After storage, marketable fruit lost 1.5% to 2.5% fresh weight during the additional week at 20 °C and this rate of loss remained constant regardless of the prior storage temperature or duration (Fig. 2H).

Fruit firmness followed a similar trend decreasing with storage duration; however,
the effect of temperature, although significant, was more variable (Table 3; Fig. 2). Fruit stored at temperatures from 3 to 10 °C lost firmness at rates averaging 0.31 to 0.35 N-mm⁻¹ per month. Fruit stored at 0 °C lost firmness at a similar rate for the first 4 months, but then softened at a higher rate for the remaining 2 months. Fruit firmness continued to decrease during the 7 d at 20 °C and averaged 0.39 to 0.78 N-mm⁻¹ less than fruit measured immediately after storage (Fig. 2). Fruit previously stored at higher temperatures tended to lose more firmness.

Discussion

The storage life of cranberry fruit, as assessed by marketability, was affected more by storage RH than by temperature. The greater loss of cranberry fruit stored at high RH was also reported in other studies. Wright et al. (1937) found that cranberries better maintained quality when stored in 70% to 75% RH compared with 90% to 95% RH. In extended storage of cranberries under a nitrogen atmosphere, Stark et al. (1974) found that fruit stored in 65% to 70% RH had less decay than fruit stored in 90% to 95% RH. Likewise, decay and physiological breakdown were higher in fruits stored in polyethylene bags or liners that maintained high RH than in fruit stored in boxes, well-ventilated bags, or unlined cartons (Anderson et al., 1963; Hruschka, 1970). In a series of experiments that looked at the storage of fresh cranberries in bulk storage containers, Norton (1982) concluded that storage rot was minimized when adequate ventilation was maintained through the containers and RH was held ≈70%. He further suggested that storage rot was minimized when fruit weight loss was maintained at 1% every 12 d.

Reducing RH in the storage tubs increased fresh weight loss of the fruit reflecting the differences in RH. Weight loss provides a better integrated measure of RH and vapor pressure deficit (VPD) than direct measure of these parameters as a result of variation of RH and temperature and error associated with their measurement (van den Berg and Lentz, 1978). However, the rate of weight loss in the low RH treatment was approximately three times less (0.86% versus 2.5% every month) than the optimum rate suggested by Norton (1982). Although the RH in the low RH treatments was similar to that reported by Norton, the difference in air movement around the fruit may account for the large difference in weight loss. Interestingly, the higher rates of weight loss seen at low RH did not result in softer fruit. This contrasts with other fruits such as bell peppers that have a higher incidence of soft fruit when stored in low RH (Polderdijk et al., 1993). Cranberry fruit softened more rapidly as storage temperature increased suggesting that firmness loss may be the result of metabolically controlled changes in cell wall structure or other processes rather than turgor pressure loss. Turgor pressure loss, which is associated with dehydration and texture loss, may have been minimized by osmotic adjustment within the cells (Pomper and Breen, 1997) or by modification of cellular structure (Levitt, 1986).

The driving force of water loss in stored fruit is the VPD between the storage atmosphere and the fruit. The VPD is a function of both the temperature and the RH of the storage environment. Assuming the RH of the interior of the fruit is near saturation, the VPD can be calculated based on the temperature and RH of the storage environment and the fruit temperature. When the VPD, calculated from the average temperature and RH in each of the storage treatments, was plotted against the average marketable fruit, a positive correlation was found with VPD accounting for approximately half of the variation in marketable fruit (Fig. 3). Both fruit decay and physiological breakdown were greater at low VPD, but this relationship was strongest for decay.

These effects of humidity on cranberry storage life are contrary to the response of most fresh fruits and vegetables. Generally, high RH minimizes weight loss and water stress on fresh produce, which often delays senescence and decay (Hardenburg et al., 1986; Paul, 1999; van den Berg and Lentz, 1978). However, in cranberries, high RH resulted in increased rates of both decay and physiological breakdown. Water stress induced by low RH may impart resistance to other stresses by affecting growth regulator levels (Grierson and Wardowski, 1978). Low RH may also alter gas diffusion into and out of the fruit, which may affect storage life (Lidster, 1990). The optimum RH favoring pathogen development may vary among pathogens (Grierson and Wardowski, 1978),
and fruit water potential may influence the development of decay (Cook and Papendick, 1978). The unique complex of fungal pathogens that is responsible for cranberry fruit decay may respond differently to storage RH than the more common pathogens found in other fruits and vegetables. Understanding the physiological response of cranberry fruit to VPD and water loss may lead to new strategies to reduce postharvest deterioration of these fruit.

Cranberry fruit were not found to be chilling-sensitive in the current study. As storage temperatures increased, both decay and physiological breakdown remained unchanged or increased and there was no increase in physiological breakdown associated with chilling temperatures contrary to previous reports (Anderson et al., 1963; Hruschka, 1970; Levine et al., 1941; Wright et al., 1937). When fruit were held for an additional week at 20 °C, greater amounts of decay and physiological breakdown were observed in fruit previously held at lower temperatures, i.e., 0 and 3 °C. It has been reported that CI may not be expressed until after fruit is warmed (Morris, 1982). However, when the cumulative effects of both cold storage and the additional 7 d at 20 °C were plotted for total marketable fruit (Fig. 4A), total decay (Fig. 4B), and total physiological breakdown (Fig. 4C), very little effect of temperature was observed (Fig. 4). It appears that cooler storage temperatures prolonged the storage life of fruit that are more susceptible to decay or physiological breakdown, but when these fruit were warmed to 20 °C, they rapidly deteriorated. The fact that the total quantity of fruit developing physiological breakdown or decay was not enhanced by prior storage at 0 °C indicates that no CI was induced.

Unlike many fruits and vegetables, cranberry fruit do not respond strongly to changes in storage temperature between 0 and 10 °C. In this study, there was little significant difference among marketable fruit stored at temperatures ranging from 0 to 7 °C over 6 months. Similarly, Levine et al. (1941) found rates of spoilage differed by only 0.5% to 4% when ‘Early Black’ and ‘Howes’ fruit were stored at temperatures ranging from 1.7 to 7.2 °C for 1.5 to 4 months. Most other fruits and vegetables respond more dramatically to storage temperature (Paull, 1999). For example, blueberries stored for 4 weeks at 4.4 °C had 67% more unmarketable fruit than those stored at 0 °C (Hruschka and Kushman, 1963).

The classification of cranberries as chilling-sensitive originates back to several studies, the first being that of Wright et al. (1937) in which large amounts of “sterile breakdown” (physiological breakdown) occurred when ‘Early Black’ and ‘Howes’ fruit were held at –1.1 °C and to a lesser extent 0 °C. Because the freezing point of cranberries ranges from –1.4 to –0.9 °C (Whiteman, 1957), the physiological breakdown that occurred at –1.1 °C was most likely freezing and not CI. The cause of physiological breakdown that occurred at 0 °C is less clear. These fruit may also have experienced freezing if temperature fluctuations in the refrigeration system occurred. Wright et al. (1937) describe the breakdown that occurred at –1.1 °C and 0 °C to have a “taste and appearance not unlike those of frozen berries.” The next coldest temperature tested was 2.2 °C at which the fewest unmarketable cranberries were found. Levine et al. (1941) also reported low temperature breakdown when ‘Early Black’ and ‘Howes’ fruit were stored at –1.1 °C and concluded that the best storage temperature was 1.7 °C, although rates of spoilage did not differ greatly when fruit
were stored at temperatures ranging from 1.7 to 7.2 °C after 7 to 18 weeks of storage. Temperatures between –1.1 and 1.7 °C were not tested. Anderson et al. (1963) reported little difference in the spoilage of ‘Howes’ cranberries stored at 0 or 3.3 °C every 2 or 4 weeks for 1 d reduced both decay and physiological breakdown. These reductions were most apparent after 4 or more months of storage after the additional 1 week at 20 °C (unpublished data).

The apparent chilling sensitivity of cranberries in these past studies could be a response to RH rather than temperature. With the exception of the study by Wright et al. (1937), RH was not controlled in these studies. Under most storage conditions where RH is not actively controlled, the VPD would most likely be lowest at the lowest storage temperature and increase as storage temperature increased. As a result, greater rates of physiological breakdown or decay that were attributed to CI could have been a response to high RH (low VPD).

Cultivar differences also could be responsible for the differences in apparent CI. Wang and Stretch (2001) compared the effects of storage temperatures ranging from 0 to 20 °C on the antioxidant capacity of 10 cranberry cultivars. Although decay and physiological breakdown of the fruit were not reported, some of the cultivars exhibited symptoms of CI after 3 months of storage at 0 °C (Wang, pers. comm.).

The wide variation in recommended storage temperatures of fresh cranberries appears to be the result of the lack of response of fresh cranberries to storage temperatures ranging from 0 to 7 °C and the unexpected effects of RH on decay and physiological breakdown. An increased understanding of the effects of humidity, VPD, and weight loss on fruit physiology and fruit–pathogen interactions would lead to improved methods for the storage of fresh cranberries. In addition, the chilling response of cranberry fruit of various cultivars should be reassessed under controlled humidity conditions.

### Literature Cited


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