

Optimum O₂ or CO₂ Atmosphere for Storing Broccoli Florets at Various Temperatures

Hidemi Izumi¹, Alley E. Watada², and Willard Douglas

Belts vine Agricultural Research Center, Agricultural Research Service, U.S. Department of Agriculture, Beltsville, MD 20705

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Abstract. 'Marathon' broccoli (*Brassica oleracea* L. var. *italica*) florets were stored in air, low O₂ (0.25%, 0.5%, and 1%) or high CO₂ (3%, 6%, and 10%) at 0, 5, and 10C. Oxygen consumption and CO₂ production were reduced under low O₂ or high CO₂ atmosphere, the reduction being greater at lower O₂ and higher CO₂ levels. No differences were found in ethylene production among the different atmospheres. Low O₂ and high CO₂ retained color of broccoli florets to about the same extent at 10C but had no effect at 0 and 5C. Development of soft rot and browning was suppressed by low O₂ or high CO₂, but offensive off-odor occurred in 0.25% O₂ at all temperatures and 0.5% O₂ at 10C. These results indicate that the best O₂ and CO₂ levels seem to be 0.5% O₂ and 10% CO₂ at 0 and 5C, and 1% O₂ and 10% CO₂ at 10C.

A controlled atmosphere with reduced O₂ and/or elevated CO₂ reduces respiration (Kasmire et al., 1974; Lebermann et al., 1968), ethylene production (Wang, 1979), weight loss (Anelli et al., 1984; Makhlof et al., 1989a), and decay (Lipton and Harris, 1974; Makhlof et al., 1989a) and retards yellowing (Kasmire et al., 1974; Lebermann et al., 1968; Lipton and Harris, 1974; Wang, 1979; Makhlof et al., 1989a) of broccoli heads. Based on these results, lowering O₂ to 1% to 2% and/or increasing CO₂ to 5% to 10% is recommended for storing broccoli heads to maintain quality at 0 to 5C (Saltveit, 1993). With the increasing demand for lightly processed vegetables, controlled or modified atmosphere for broccoli florets also has been researched over the past few years (Ballantine et al., 1988; Barth et al., 1993; Bastrash et al., 1993; Berrang et al., 1990). However, these studies were conducted only at a single temperature, which differed among the studies.

Since the optimum storage atmosphere can differ with storage organ-commodity and storage temperature, our objective was to determine the optimum treatment atmosphere for storage of broccoli florets at 0, 5, and 10C. The 10C was included because the temperature of a few commercial holding rooms approaches 10C. Optimum atmosphere was based on physiological activity and visual quality of the florets. Oxygen levels below 1% were examined in this study because the internal O₂ content in broccoli heads probably drops substantially below 1% when using 1% to 2% O₂ atmosphere. In broccoli heads, compared to the florets, the gas diffusion is more limited, which causes a larger gradient in gas concentration from the external to internal tissue (Brecht, 1980).

Materials and Methods

Heads of 'Marathon' broccoli were obtained from the Wholesale distribution center in Jessup, Md., and separated with a knife into florets of about 15 g each. A 150-g sample was placed on an elevated screen in a 2-liter glass jar with 100 ml of distilled water in the bottom to maintain high relative humidity. Three replicated samples were stored at 0, 5, and 10C under a continuous stream of

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¹Present address: Kinki Univ., Naga, Wakayama, 649-64 Japan.

²Research leader, Horticultural Crops Quality Laboratory. To whom reprint requests should be addressed.

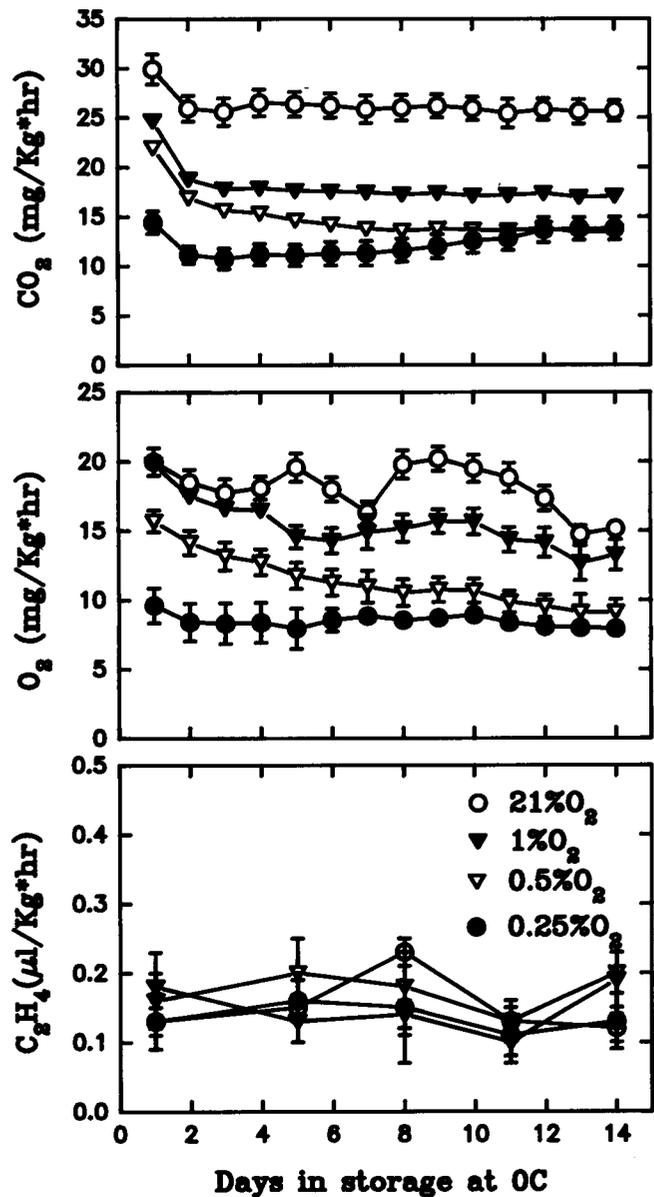


Fig. 1. Rates of CO₂ production, O₂ consumption, and ethylene production of broccoli florets during storage at 0C under air or low O₂ atmospheres. Vertical lines represent SE. SE bars not shown when within symbols.

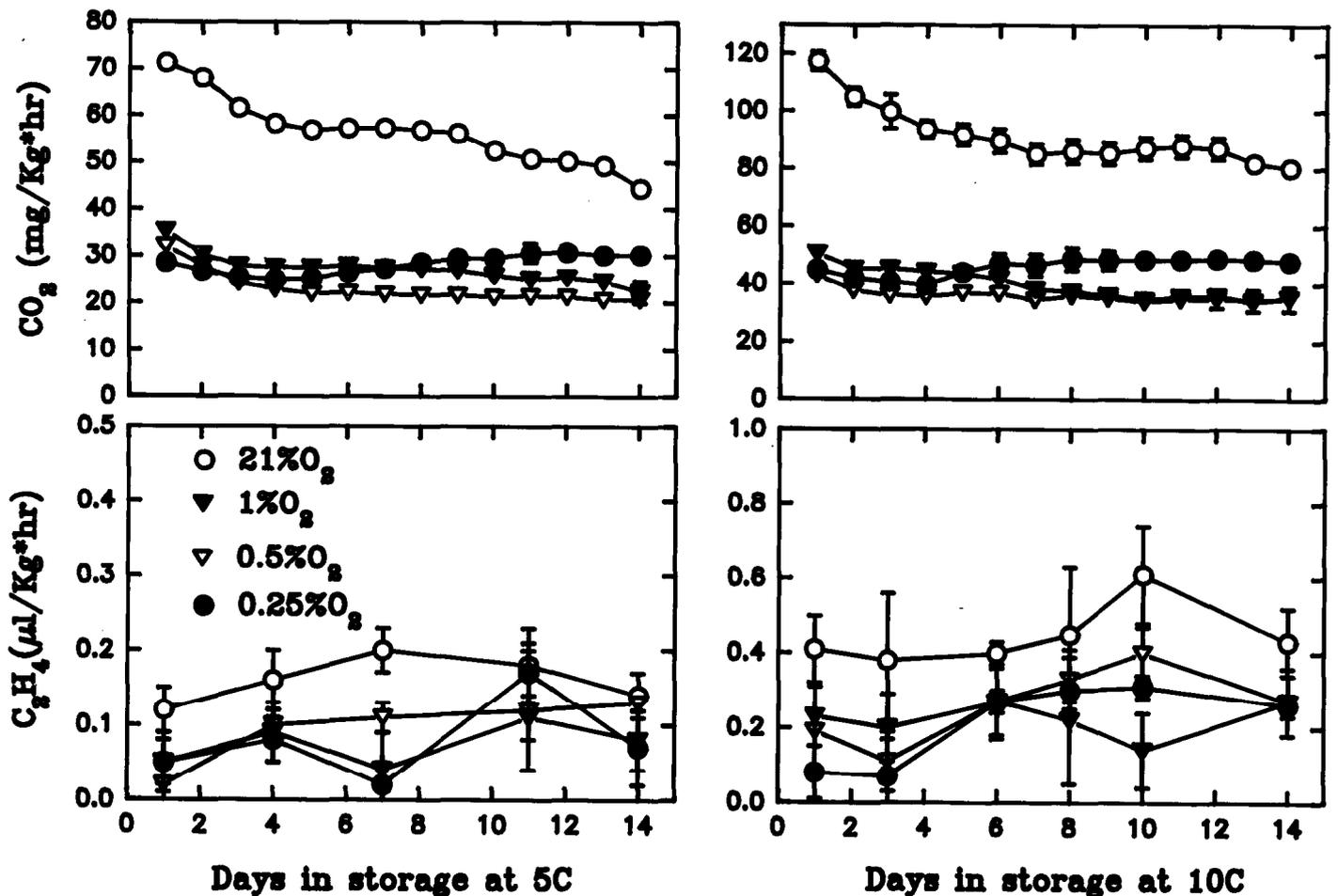


Fig. 2. Rates of CO_2 production and ethylene production of broccoli florets during storage at 5 and 10C under air or low O_2 atmospheres. Vertical lines represent SE. SE bars not shown when within symbols

air, low O_2 (0.25%, 0.5%, and 1%) or high CO_2 (3%, 6%, and 10%) atmosphere at 10, 20, and 40 $\text{ml}\cdot\text{min}^{-1}$, respectively. The balance of the mixture was N_2 for the low O_2 atmosphere, and N_2 and air to give a minimum of 19% O_2 for the high CO_2 atmosphere. Different lots of broccoli were used for the low O_2 and the high CO_2 studies.

Oxygen and CO_2 contents of inlet and outlet streams of each jar were monitored with an oxygen analyzer and carbon dioxide analyzer (model S-3A/I and model CD-3A; Ametek, Pittsburgh). Ethylene levels in 5-ml gas samples taken from each jar were measured with an analytical gas chromatography (Model AGC-211; Carle, Tulsa, Okla.), equipped with a flame ionization detector. Color, decay, and odor were evaluated after storage. The color [hue angle, $\tan^{-1}(b^*/a^*)$] of the central part of the inflorescence was determined using a chroma meter (model CR-300; Minolta, Japan), with a 1-cm aperture. Incidence of soft rot and browning were expressed as a percentage of the total number of florets inspected. Raw and cooked odor of florets that had been placed in a microwave oven for 2 min were rated on a scale of 0 to 4, with 0 = normal and 4 = severely objectionable.

Results

Carbon dioxide production and O_2 consumption of florets held at 0C were lower with decreasing O_2 levels (Fig. 1). However, CO_2 production of florets at 0.25% O_2 decreased on day 2, and then increased gradually thereafter and approached the rate of those held at 0.5% O_2 by day 10, with samples held at 0C. In comparison, the CO_2 production rates of samples in 0.25% and 1% O_2 merged

on day 7 at 5C and day 5 at 10C (Fig. 2). Carbon dioxide production of samples held in 0.25% O_2 was higher than those held in 0.5% and 1% O_2 after day 11 at 5C, and day 8 at 10C (Fig. 2). No major differences were found in ethylene production between samples in air and low O_2 atmospheres at all temperatures (Figs. 1 and 2).

Green color, expressed as hue angle, was retained in florets stored in all atmospheres at 0 and 5C (127.1 ± 1.1), whereas it decreased by 80% from the initial value for florets in air at 10C (104.7 ± 1.6). Low O_2 levels suppressed the development of soft rot on flower buds and browning of cut surfaces at 0C (Table 1). Browning was only suppressed at 5 and 10C. Soft rot incidence was not affected at 5C, but increased when florets were held in 0.25% and 0.5% O_2 atmospheres at 10C. An offensive off-odor was produced by raw florets held in low O_2 at all temperatures except 1% O_2 at 0C. After cooking, however, the odor was not detected with samples held at 0 and 5C in 0.5% and 1% O_2 and was minimal with samples at 10C in 1% O_2 .

High CO_2 atmosphere inhibited CO_2 production of florets held at 0C, with the inhibition being greater in 10% or 6% CO_2 (Fig. 3). This pattern of differences among treatments was also observed at 5 and 10C (Fig. 4). Oxygen consumption tended to be lower in high CO_2 atmosphere than in air at 0C, although the difference was not as defined as that in CO_2 production (Fig. 3). There were no significant differences in ethylene production between air and high CO_2 atmospheres at all temperatures, except for the initial day at 0C (Figs. 3 and 4).

High CO_2 had no effect on retention of green color at 0 and 5C, but did retard its loss at 10C, as noted with low O_2 atmosphere (data

Table 1. Incidence of soft rot and browning and development of objectionable odor during storage of broccoli florets at 0, 5, and 10C under air or low O₂ atmospheres.

Storage (°C)	Treatment	Odor ^y			
		Soft rot ^t	Browning ^z	Raw	Cooked
0	21% O ₂	20 a ^x	15a	0c	0b
	1% O ₂	4b	0b	0c	0b
	0.5% O ₂	0b	0b	1.5b	0b
	0.25% O ₂	0b	0b	2.0a	3.0a
5	21% O ₂	17a	44a	1.0d	0b
	1% O ₂	21a	15b	1.7c	0.7b
	0.5% O ₂	21a	10b	2.0b	0.7b
	0.25% O ₂	11a	0b	3.0a	3.0a
10	21% O ₂	37b	33a	1.0c	0d
	1% O ₂	45ab	19b	1.0c	1.0c
	0.5% O ₂	59a	13b	2.3b	2.0b
	0.25% O ₂	61a	0c	3.0a	3.0a

^t(Number of injured florets/ 100 number of observed florets) × 100.

^yRated on a scale of 0 to 4, with 0 = normal and 4 = severely objectionable.

^zMean separation within each temperature by Duncan's multiple range test, *P* = 0.05.

not shown). Soft rot, browning, and off-odor were minimal in high CO₂ atmosphere at all temperatures, while the incidence of soft rot in air at 10C was about five times greater than that in high CO₂ at 10C (Table 2). No off-odor occurred after cooking samples held in high CO₂ atmospheres at all temperatures.

Discussion

Respiration rate of broccoli florets was reduced by both low O₂ and elevated CO₂. The effect was greater at 5 and 10C when florets were held in O₂ levels of 1% or less compared to CO₂ levels of up to 10%. Lebermann (1968) reported that respiration of broccoli heads was reduced when held in 2% O₂ or 20% CO₂ at 7C, but the reduction was similar in both atmospheres. With florets in our study, the greater effect of O₂ relative to CO₂ may have been due to a change in the mechanism by which low O₂ reduced the respiration rate, because respiratory quotient (RQ) was higher in florets held in 0.25% O₂ than in air, but lower in those held in high CO₂ atmospheres (data not shown). As the O₂ level was reduced to 0.25%, respiration may have changed from the aerobic to anaerobic pathway, and this shift would occur more readily at 5 and 10C than at 0C. High CO₂ has been shown to reduce respiration rate, the mechanism of action is unknown, but may be associated with changes in the Krebs cycle (Kader, 1986).

Ethylene production was minimal and green color was retained with florets held in low O₂ at 10C. Others have reported that the rate of ethylene production is correlated positively with the rate of chlorophyll degradation (Aharoni et al., 1985; Makhoul et al., 1989b; Wang, 1979). However, in our study with the high CO₂ atmosphere at 10C, ethylene production had increased but the green color was retained. High CO₂ probably caused a stress condition which induced ethylene production but did not have a deteriorative effect on color (Aharoni et al., 1985). However, with extended storage, ethylene can cause degradation of chlorophyll (Makhoul et al., 1989b). We found that high CO₂ and low O₂ retained the green color of florets to about the same degree at 10C. However, others have indicated that elevated CO₂ was more effective than reduced O₂ in retarding the yellowing of broccoli (Kasmire et al., 1974; Lebermann et al., 1968; Makhoul et al., 1989a). The differences in the results may be due to the differences

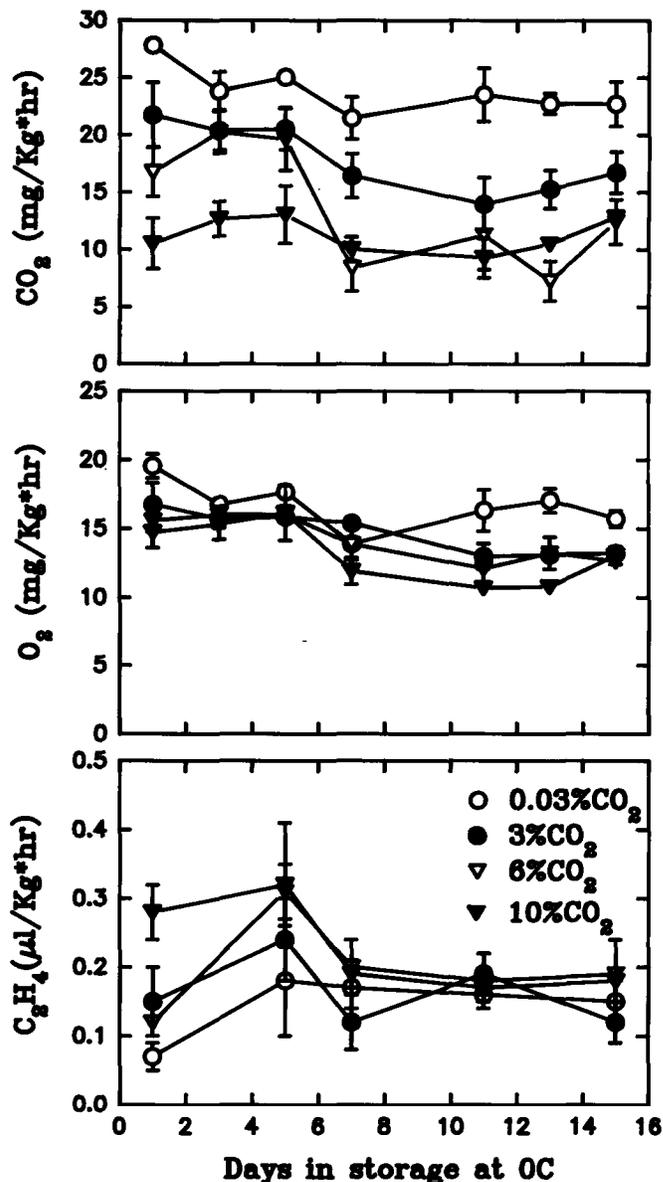


Fig. 3. Rates of CO₂ production, O₂ consumption, and ethylene production of broccoli florets during storage at 0C under air or high CO₂ atmospheres. Vertical lines represent SE. SE bars not shown when within symbols.

in the temperature and length of storage periods.

High CO₂ was more effective than low O₂ in reducing the incidence of soft rot at 10C, and of undesirable odor at all temperatures. However, this comparison may not be valid because the incidence of soft rot of samples held in air (control) was lower in the CO₂ experiment (Table 2) than in the O₂ experiment (Table 1), possibly due to the differences in maturity of broccoli and the degree of field and packing house contamination.

A 0.5% O₂ level at 0 and 5C and 1% O₂ level at 10C is recommended for broccoli florets to prevent the development of off-odor induced by low O₂. Others have reported that off-odor occurred with raw broccoli heads held in O₂ below 1% at 2.5C (Kasmire et al., 1974) and by cooked broccoli heads that were held in O₂ below 0.25% at 5 and 7.5C (Lipton and Hams, 1974).

Our results indicate that 0.5% O₂ and 10% CO₂ at 0 and 5C, and 1% O₂ and 10% CO₂ at 10C seemed to be the best atmosphere for preservation of broccoli florets without risking the development of undesirable odors and disorders. These O₂ levels were lower than

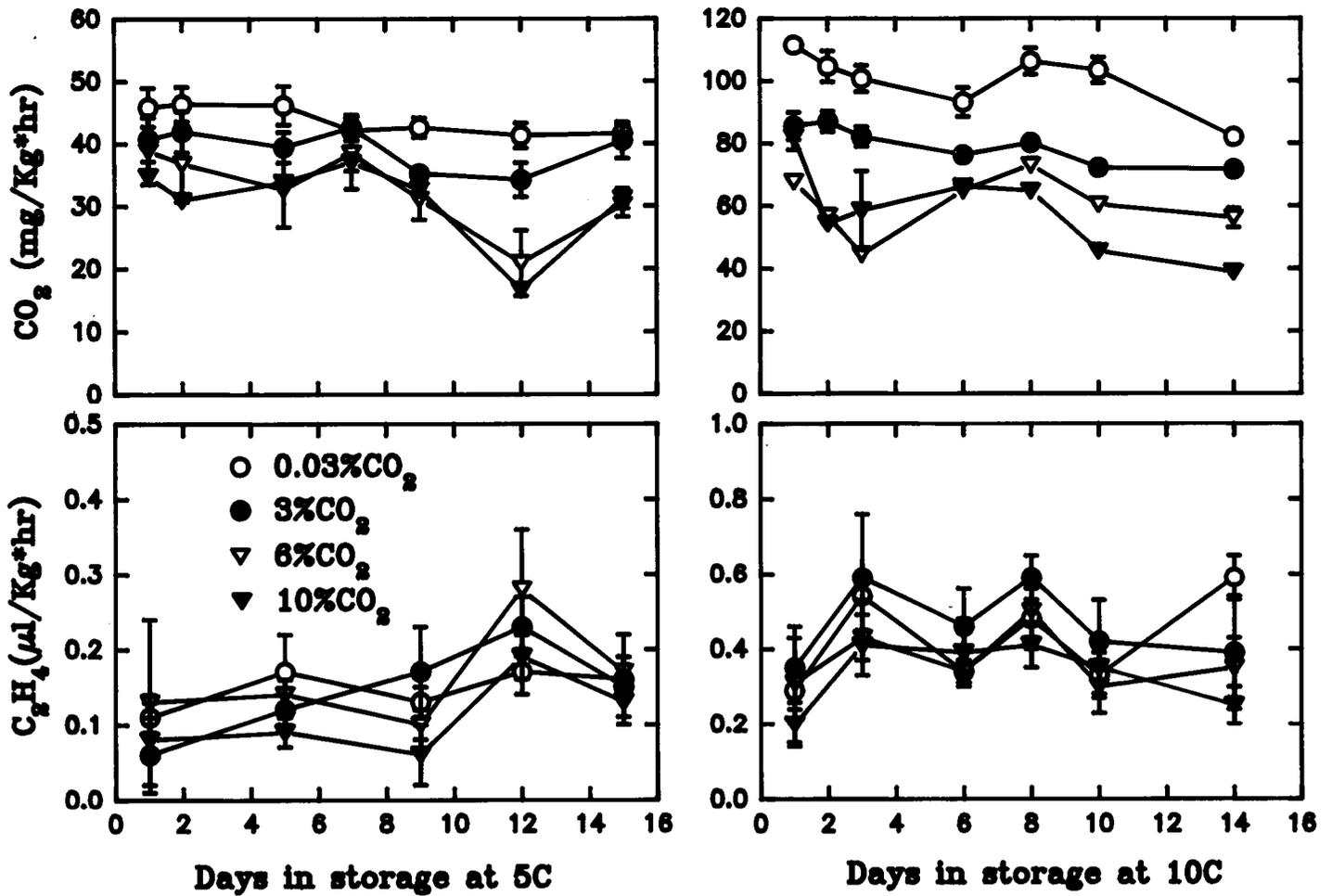


Fig. 4. Rates of CO₂ production and ethylene production of broccoli florets during storage at 5 and 10C under air or high CO₂ atmospheres. Vertical lines represent SE. SE bars not shown when within symbols.

Table 2. Incidence of soft rot and browning and development of objectionable odor during storage of broccoli florets at 0, 5, and 10C under air or high CO₂ atmospheres

Storage (°C)	Treatment	Soft rot ^a	Browning	Odor ^b	
				Raw	Cooked
0	0.03% CO ₂	0 a	0 a	2.0 a	0.7 a
	3% CO ₂	0 a	0 a	1.0b	0a
	6% CO ₂	0 a	0 a	1.0b	0a
	10% CO ₂	0 a	0 a	1.0b	0.7 a
5	0.03% CO ₂	0 a	0 a	1.7a	0.3 a
	3% CO ₂	0 a	0 a	1.0a	0.7 a
	6% CO ₂	0 a	0 a	1.0a	0.7 a
	10% CO ₂	2 a	2 a	1.0a	0.3 a
10	0.03% CO ₂	17 a	6 a	1.0a	0a
	3% CO ₂	4 b	0 a	1.3a	0a
	6% CO ₂	2 b	0 a	1.3a	0a
	10% CO ₂	2 b	0 a	1.7a	0a

^aNumber of injured florets × 100 number of observed florets.
^bRated on a scale of 0 to 4, with 0 = normal and 4 = severely objectionable.
^cMean separation within each temperature by Duncan's multiple range test, *P* = 0.05.

those recommended for broccoli heads at 0 to 5C (Saltveit, 1993), suggesting that a smaller gradient of O₂ concentration from the external to internal tissue had occurred in florets than in heads due to the less distance of gas diffusion in florets. When florets were held at 10C, it is important to keep O₂ levels above 1%. The results

could be beneficial for storage of packaged broccoli florets, because ambient O₂ of packaged florets can become very low due to the high respiration rate of florets as compared with heads (Bastrash et al., 1993; Rushing, 1990).

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