Relationship between a Reduced Aroma Production and Lipid Metabolism of Apples after Long-term Controlled-atmosphere Storage

A. Brackmann¹, J. Streif, and F. Bangerth
Institut für Obstbau, Universität Hohenheim, 7000 Stuttgart-70, Germany

Abstract. ‘Golden Delicious’ apples (Malus domestica Borkh.) harvested at the preclimacteric and climacteric stages of ripening were stored for up to 8 months at 1C in air and under various controlled atmosphere(s) (CA), including ultralow oxygen (ULO) storage conditions. Aroma volatiles were measured at 2-month intervals in fruit ripened for 10 days at 20C. Fruits harvested at the climacteric stage produced more volatiles during all storage conditions than preclimacteric fruit. All CA storage treatments suppressed aroma production compared to cold storage. The greatest reduction was found under ULO (1% O₂ and high CO₂,3%) conditions. A partial recovery of aroma production was observed when CA fruits were subsequently stored for 14 days under cold storage conditions. Suppression of aroma production under ULO conditions seems to be related to low fatty acid synthesis and/or degradation, and is restricted to volatiles having a straight C chain. Production of branched C-chain aroma compounds was suppressed by high CO₂ concentrations. The reduced capacity of aroma production during shelf life after ULO storage is confined to apple cultivars producing mainly ester compounds with a straight C-chain, e.g., ‘Golden Delicious’.

CA storage, in particular ULO, diminishes the loss of firmness, acidity, chlorophyll, and sugar of the fruit and reduces the incidence of some physiological disorders in comparison with cold storage of apples (Bohling and Hansen, 1985; Schouten, 1988; Streif, 1985). However, CA conditions suppress flavor development in apples (Shatat et al., 1978; Streif and Bangerth, 1988). The suppression depends on both the composition of the storage atmosphere and the length of storage. In general, suppression of volatile production is particularly marked under conditions that delay ripening (Hatfield and Patterson, 1975). The phenomenon of decreasing aroma production has been observed in ‘Cox Orange’ (Knee and Hatfield, 1981), ‘Golden Delicious’ (Streif and Bangerth, 1988), and ‘McIntosh’ (Lidster et al., 1981).

Reasons for the suppression of aroma production after CA storage are not fully understood. Patterson et al. (1974) attribute the decline in aroma volatiles of long-term-stored apples to a loss of substrates or enzymes essential for the formation of esters. Hatfield and Patterson (1975) suggest a connection between inhibition of respiration after CA storage and suppression of aroma production. According to Knee and Hatfield (1981), the production of esters by ‘Cox Orange’ in low O₂ atmosphere is limited by the availability of alcohols from which the aroma esters are derived. Bangerth and Streif (1987) suggest that changes in the sensitivity of fruits to C₂H₄ might be responsible for the decline in aroma production by ‘Golden Delicious’ because inhibition of CH₃biosynthesis and/or action under cold storage conditions gave similar effects on volatile production (Bangerth and Streif, 1987; Halder-Doll and Bangerth, 1987). High CO₂ concentrations in combination with ULO conditions further suppress aroma production in shelf life. DePooter et al. (1987) indicated that the antagonistic effect of high CO₂ concentrations on aroma production might not only interfere with alcohol dehydrogenase (EC 1.1.1.1), which catalyses the synthesis of alcohols from aldehydes, but also with carboxylic acid metabolism.

The pathway of ester synthesis in apples is not completely understood. Studies of lipid metabolism in apples indicated an enhanced turnover of lipids during ripening (Bartley, 1985), and the long chain liberated fatty acids are thought to be precursors for some of the esters (Bartley, 1986). Also, amino acids serve as precursors for the synthesis of aroma volatiles in apples (Tressler et al., 1970).

The purpose of this investigation was to determine the effect of various storage conditions, especially ULO, on the production of aromatic volatiles and fatty acids during ripening. Further, this investigation evaluated the effect of several poststorage treatments on aroma production after long-term ULO storage.

Materials and Methods

Apples from ‘Golden Delicious’ trees grown on M 9 rootstocks were used for the experiments. The fruits were harvested either at the preclimacteric or climacteric ripening stages, as determined by internal ethylene concentrations (<0.05 µl·liter⁻¹ and >0.6 µl·liter⁻¹, respectively) and CO₂-production. Immediately after picking, fruits were selected for uniformity and stored at 1C. For CA storage, 40 kg of apples were placed in 240-liter containers. The concentrations of CO₂ and O₂ were continuously monitored by gas analysers connected to a process computer. To achieve CO₂ absorption in the low CO₂ atmospheres (<1%CO₂) it was necessary to place Ca(OH)₂ in the container. Excess CO₂ in the other CA containers was removed by N₂ flushing. Storage conditions were: air storage: 3% CO₂+3% O₂;3% CO₂+1%O₂;5%CO₂+1%O₂. During 8 months of storage, 1-kg samples of fruit in each of two replicates were removed from the storage containers at 2-month intervals and placed in separate gas-tight glass jars at 20C for 10 days. These were continuously ventilated with air at a rate of 11.3 liter·h⁻¹. After 3, 6, and 9 days of ripening, 2 liters of air from the jars was passed through Pasteur pipettes in which activated charcoal adsorbed the

Received for publication 16 Sept. 1991. Accepted for publication 3 Sept. 1992. This research was supported in part by a grant from the German Academic Exchange Service (Deutscher Akademischer Austauschdienst). The cost of publishing this paper was defrayed in part by the payment of page charges. Under postal regulations, this paper therefore must be hereby marked advertisement solely to indicate this fact.

¹Versuchsstation für Obstbau, 7980 Ravensburg-Bavendorf, Germany.

Abbreviations: CA, controlled atmosphere; ULO, ultralow oxygen.
volatiles. The pipettes were directly inserted into the injection port of a gas chromatograph and the desorbed volatiles analysed on a packed Carbowax 20M column (Streif, 1981). Retention indices of authentic standards (Roth, FRG) were used to identify the 15 most important volatiles. Restriction to these 15 compounds was possible because earlier methodological experiments had shown that most of the volatile substances produced in smaller amounts were similar in production trends as the major compounds.

The wax and fatty acid composition of the fruit peel was determined after 7 months of storage in one of the two above replicates. For these determinations, 200 disks 1.2 cm in diameter from five fruits were taken for each storage treatment. Wax and fatty acids were extracted with petroleum ether in a Soxhlet apparatus for 3 h. Gas chromatography of the lipids and fatty acids was done according to Neubeller (1971).

In an additional poststorage treatment, six samples (seven fruits each) from the 3 % CO₂ + 1 % O₂ atmosphere were removed after 8 months of storage and stored in air at 1°C for 2 weeks. The fruit was then ripened as stated above.

To evaluate the effect of exogenous aroma precursors, fruit (three randomized samples of five apples each) from the 3% CO₂, + 1% O₂, atmosphere were taken after 7 months of storage, enclosed in 5-liter jars, and held at 20°C. The various alcohols, aldehydes, organic acids, and esters (Table 1) were directly injected onto a filter paper in the jars, the total final concentration in the jar being 160 µl·liter⁻¹. After 24 h, the samples of apples were transferred to ventilated jars for 10 days of ripening and then volatiles analysed as described by Streif (1981). Production was determined after 1, 5, and 9 days of ripening.

Results

The main constituents in the aroma volatiles of ‘Golden Delicious’ were esters of which butyl and hexyl acetate represented = 60% of the total volatile production of ripe fruits (Fig. 1). Fruits harvested at the climacteric stage produced more volatiles than preclimacteric fruits under all storage conditions and removal dates (Fig. 2). CA storage conditions suppressed the aroma production in comparison to air storage. In general, the production of volatiles was negatively affected by low O₂ (1%) and high CO₂ (3%) concentrations. The greatest suppression of volatiles was caused by ULO (3% CO₂ + 1% O₂), and suppression was accentuated with increasing storage time. ULO especially suppressed compounds with a straight C-chain, such as butyl acetate (Fig. 3).

Production of compounds with a branched C-chain, such as 2-methyl butyl acetate and 2-methyl butanol, was not suppressed by low O₂ but by high CO₂ after long-term storage (Figs. 4 and 5).

The amount of fatty acids in the peel increased with storage time, e.g., in air-stored fruits, from 18 to 45 µg·cm⁻² over 7 months. Production, however, was inhibited by low O₂ and high CO₂, a possible reason for the lowest concentration being found in ULO (Fig. 6). Synthesis of linoleic acid was suppressed the most. In contrast, the wax content of the skin was not influenced by storage conditions. After 8 months in store, the wax content of all treatments was ≈ 400 µg·cm⁻².

Air storage at 1°C for 2 weeks immediately after 8 months of ULO storage greatly improved production of volatiles during ripening. This treatment, however, failed to allow a complete recovery of the potential for aroma production in comparison with air storage (Fig. 7).

In the experiment in which aroma precursors were added to long-term ULO-stored fruits, the apples showed an intensive turnover of these substances as indicated by changes in the concentration of compounds produced from them (Table 1). One mechanism by which the precursors could be transformed is β-oxidation forming compounds having two fewer C atoms. Exogenous butyrate, e.g., was reduced to butanol, but mainly degraded by β-oxidation to ethanol, which was then partly esterified to ethyl

![Fig. 1](image1.png) Relative amount of the 15 most important aroma volatiles of ‘Golden Delicious’ apples (see Material and Methods) at harvest and ripened for 9 days at 20°C. Peak identity was done by comparison of retention times with reference standards.

![Fig. 2](image2.png) Total volatile production of (A) preclimacteric and (B) climacteric ‘Golden Delicious’ apples during 8 months of storage at 1°C: air storage; CA 3% CO₂ + 3% O₂; CA 3% CO₂ + 1% O₂ (ULO); CA <1% CO₂ + 1% O₂.

![Fig. 3](image3.png)
butyrate. The aldehyde butanal was oxidized or partly reduced to its corresponding acid and alcohol, which were subsequently combined in ester compounds such as ethyl isobutyrate, butyl- or pentyl butyrate. The supplied ester, methyl acetate, was hydrolized, and the formed acid was reduced to alcohol and then esterified with another acid preferentially forming ethyl isobutyrate. The three most prominent volatile compounds, viz. butyl acetate, 2-methyl butyl acetate, and hexyl acetate, were surprisingly little or not at all affected by the precursors applied. Since unlabeled precursor compounds were used for these experiments, our conclusions have to be viewed as tentative.

**Discussion**

It is generally accepted that apple volatiles are synthesized enzymatically by coupling the respective acid and alcohol moieties (Salunkhe and Do, 1976). The straight-chain fatty acids can be formed by P-oxidation of long chain fatty acids (Bartley, 1986) and the branched-chain acid moieties from amino acids (Tressl et al., 1970).

Butyl and hexyl acetate, the most abundant volatiles produced by ‘Golden Delicious’ (Fig. 1), are formed from butanol and hexanol combining with acetate derived from the metabolism of fatty acids (Bartley, 1986). Among the 15 aroma volatile compounds evaluated in this study it was evident that production of esters and alcohols with straight chains was strongly suppressed by long storage times under ULO. However, esters with branched-chains were not suppressed under these conditions. Only under high CO₂ (3%) concentrations did the production of these substances decline.

The suppression of volatiles with straight C-chains under ULO could be related to the influence of low O₂ concentrations on lipid metabolism and/or synthesis. Measurements of the fatty acid content and composition in the fruit peel, which probably reflect fatty acid composition in the cortex, confirms that low O₂, and high CO₂ concentrations reduce the levels of all fatty acids, especially linoleic acid. This finding agrees with that of Schmitz (1968), who
observed inhibition of unsaturated fatty acid synthesis during CA storage of ‘Golden Delicious’ apples. The unsaturated fatty acids, such as linoleic acid, need O₂ for their synthesis (Stumpf, 1980). This fact could partly explain the inhibition of unsaturated fatty acid synthesis under low O₂ concentration. However, low O₂ and high CO₂ concentrations could inhibit respiration and general metabolism, thus reducing the supply of energy equivalents, such as NADPH, that are needed for the synthesis of fatty acids during the ripening of the fruit (Drawert, 1975). This explanation is relevant because inhibition of ethylene biosynthesis and/or action also inhibits respiration and the production of unsaturated fatty acids and aroma volatiles, irrespective of O₂ concentration in the storage atmosphere (Halder-Doll and Bangerth, 1987).

The experiment with aroma precursors shows that fruits are capable of synthesizing alcohols and esters from exogenously supplied precursors after 8 months of ULO storage. This means that enzymes in the last steps of β-oxidation, viz. alcohol dehydrogenase and esterase, are active. This leads to the conclusion that low O₂ limits the availability of alcohols from which esters are derived (Knee and Hatfield, 1981). Inhibition of fatty acid biosynthesis and/or degradation could be another reason for the lack of alcohol precursors, thus low volatile production after ULO storage. If impairment of fatty acid degradation is the reason, then the limiting step would be expected to be early in the catabolism of these compounds, since precursors of the size of octanol and smaller are easily metabolized by ULO fruits (see Table 1).

Oxidation of, e.g., linoleic acid by lipoxigenase (EC 1.13.11.12), which requires O₂ and yields hexanal besides three larger molecules (aldehydes, acids) (Drawert, 1975), is a more likely alternative for these first steps than β-oxidation.

Since high CO₂ concentrations suppress the production of aroma compounds with branched and unbranched C-chains, then high CO₂ probably affects the metabolism of amino acids and fatty acids. Rolle (1968) concluded that apple cultivars synthesizing comparatively large amounts of “branched volatiles” produce the alcohol precursors for these substances mainly from amino acids. The tricarboxylic acid cycle, from which most amino acid precursors are derived, is inhibited by elevated CO₂ (Frenkel and Patterson, 1973). Thus, the influence of CO₂ on the production of volatiles is one of suppressed amino acid metabolism.

Drawert et al. (1972) classified apple cultivars by their volatile composition into two types: the “ester-type” and the “alcohol-type.” Whereas the alcohol-type cultivars produce most of their volatile precursors from amino acids, the precursors of the ester-type cultivars originate mainly from fatty acids. Taking this information into account, we hypothesize that the suppression of aroma volatiles during shelf life, after ULO storage, is important only for the ester-type apple cultivars, such as ‘Cox Orange’, ‘Golden Delicious’, ‘Jonathan’, etc. Therefore, attempts to improve the volatile aroma production of “ester-type” apple cultivars under ULO conditions would be important.

Literature Cited


Bartley, I.M. 1986. Changes in sterol and phospholipid composition of...


