

Regulation of Monoterpene Accumulation in Leaves of Peppermint¹

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Plants synthesize numerous classes of natural products that accumulate during development and are thought to function as constitutive defenses against herbivores and pathogens. However, little information is available about how the levels of such defenses are regulated. We measured the accumulation of monoterpenes, a model group of constitutive defenses, in peppermint (*Mentha × piperita* L.) leaves and investigated several physiological processes that could regulate their accumulation: the rate of biosynthesis, the rate of metabolic loss, and the rate of volatilization. Monoterpene accumulation was found to be restricted to leaves of 12 to 20 d of age, the period of maximal leaf expansion. The rate of monoterpene biosynthesis determined by ¹⁴CO₂ incorporation was closely correlated with monoterpene accumulation, as determined by gas chromatographic analysis, and appeared to be the principal factor controlling the monoterpene level of peppermint leaves. No significant catabolic losses of monoterpenes were detected throughout leaf development, and monoterpene volatilization was found to occur at a very low rate, which, on a monthly basis, represented less than 1% of the total pool of stored monoterpenes. The composition of volatilized monoterpenes differed significantly from that of the total plant monoterpene pool, suggesting that these volatilized products may arise from a separate secretory system. With the demonstration that the rate of biosynthesis is the chief process that determines monoterpene accumulation in peppermint, efforts to improve production in this species can now focus on the genes, enzymes, and cell differentiation processes that regulate monoterpene biosynthesis.

Plants produce an enormous variety of natural products that are thought to play a critical role in defense against herbivores and pathogens (Wink, 1999). These metabolites may be synthesized constitutively in specific organs or at specific stages of development, or their production may be induced by herbivore or pathogen attack. Considerable information is available about the mechanisms by which plant damage induces the synthesis of defensive metabolites (Karban and Baldwin, 1997). By comparison, much less is known about what controls the formation of constitutive defenses. Compounds such as monoterpenes (Gambliel and Cates, 1995), naphthoquinones (Brigham et al.,

1999), pyrrolizidine alkaloids (Hartmann and Dierich, 1998), and glucosinolates (Blake-Kalff et al., 1998) accumulate during normal root or shoot development in thousands of plant taxa, but the physiological and molecular mechanisms that regulate the production of these natural products have seldom been examined.

One of the best-studied examples of constitutive plant defenses are the monoterpenes, the C₁₀ members of the terpenoid (isoprenoid) family of natural products. Monoterpenes are colorless, lipophilic, volatile substances that have been implicated as defenses against a variety of herbivores and pathogens (Langenheim, 1994). Known from species of Pinaceae, Lamiaceae, Rutaceae, Myrtaceae, Asteraceae, and many other plant families (Charlwood and Charlwood, 1991), they are responsible for many of the characteristic odors of plants (Hay and Waterman, 1993). Monoterpenes are also frequent constituents of oils and resins, and their accumulation is often associated with complex secretory structures such as glandular trichomes, secretory cavities, or resin ducts (Fahn, 1979).

From an economic standpoint, the most important monoterpene-producing species is peppermint (*Mentha × piperita* L.), a perennial herb of the Lamiaceae that produces high levels of *p*-menthane monoterpenes (Fig. 1) in glandular trichomes found on the surfaces of leaves, young stems, and parts of the inflorescence (Amelunxen, 1965). During leaf development, the total content of monoterpenes increases with age (Burbott and Loomis, 1969; Croteau and Martinkus, 1979), and the composition of monoterpenes is significantly altered. Limonene and menthone are the major monoterpenes present in the youngest leaves. The proportion of limonene declines rapidly with development, while menthone increases in prominence and declines only at later stages as menthol becomes the dominant monoterpene constituent (Burbott and Loomis, 1969; Croteau and Martinkus, 1979; Brun and Voirin, 1991).

The pathway of monoterpene biosynthesis in peppermint has been well established by *in vivo* and cell-free studies (Kjonaas and Croteau, 1983; Kjonaas et al., 1985; Croteau and Venkatachalam, 1986), and all of the enzymes involved have been described (Kjonaas et al., 1982, 1985; Croteau and Venkatachalam, 1986; Karp et al., 1990; Croteau et al., 1991; Alonso et al., 1992; Rajaonarivony et al., 1992; Colby et al., 1993). In addition, the site of monoterpene biosynthesis has been specifically localized to the secretory cells of the glandular trichomes (Gershenzon et

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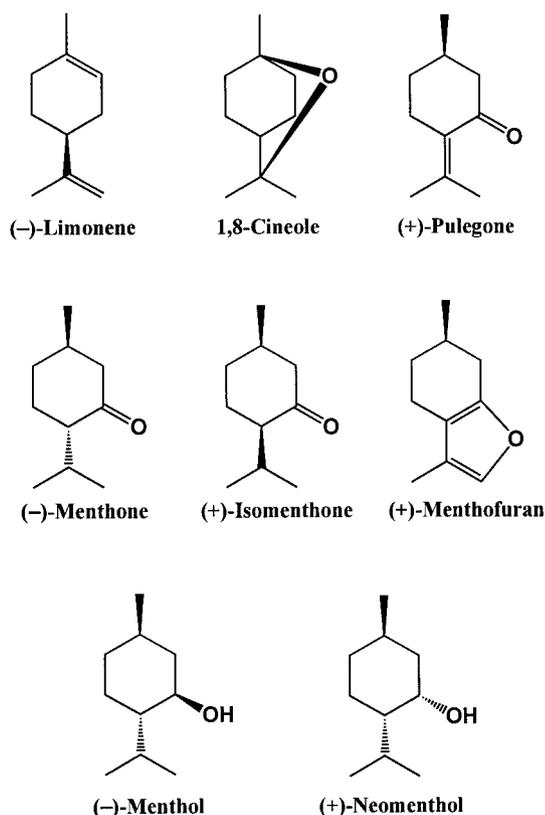


Figure 1. Major monoterpene constituents of peppermint leaves.

al., 1989; McCaskill et al., 1992). However, the physiological factors regulating monoterpene accumulation are poorly known. The accumulation of any metabolite is controlled by the balance between the rate of formation and the rate of loss, a consequence of direct release into the environment and/or catabolism. For volatile compounds such as monoterpenes, emission into the atmosphere may be a major route of loss from plants (Lerdau et al., 1997). However, catabolism must also be considered, because, while short-term monoterpene turnover in peppermint has been ruled out (Mihaliak et al., 1991), a pathway for the long-term degradation of monoterpenes in mature leaves of this species has been previously described (Croteau, 1988).

In this study, the pattern of monoterpene accumulation in the developing leaves of peppermint was measured, and the rates of monoterpene synthesis, loss, and volatile emission were determined at various stages during development to evaluate the influence of these processes on monoterpene yield. The results define the physiological factors that control the accumulation of a major class of defensive compounds in plants, and provide the necessary foundation for further regulatory studies at the biochemical and molecular levels. Given the economic importance of monoterpenes for the fragrance, flavor, and pharmaceutical industries, knowledge of the processes that control monoterpene accumulation in plants can be of value in increasing the yields of these commercially valuable natural products.

MATERIALS AND METHODS

Plant Material

Peppermint (*Mentha × piperita* L.) was propagated from rhizomes and raised in a plant growth chamber equipped with a mixture of fluorescent and incandescent lights (16-h photoperiod, $350 \mu\text{mol m}^{-2} \text{s}^{-1}$ of photosynthetically active radiation [PAR] at plant height) and a temperature cycle of $22^\circ\text{C}/10^\circ\text{C}$ (day/night). Plants were grown in peat moss:pumice:sand (55:35:10, v/v), watered daily, and fertilized on alternate days with a complete fertilizer (N:P:K, 20:20:20, v/v) plus iron chelate and micronutrients. To study developmental changes in monoterpene accumulation and the rate of monoterpene biosynthesis, a cohort of leaves was utilized that was initiated on 3-week-old stems. Samples of this cohort were removed at ages ranging from 5 to 55 d, at which time the majority of leaves of this group had senesced or abscised.

Monoterpene Extraction and Analysis

Fresh leaves of different ages were soaked in 5 mL of diethyl ether for 1 h, and then again in a second portion of diethyl ether for 1 h at room temperature. A mixture of $1.25 \mu\text{mol}$ of isobutyl benzene and $1.25 \mu\text{mol}$ of camphor was added to the combined extract for each age group as an internal standard for the quantification of monoterpene olefins and oxygenated monoterpenes, respectively, followed by concentration under nitrogen to approximately 3 mL and treatment with 25 mg of activated charcoal. After filtration and washing with 1 mL of water, the organic extract was passed through a short column of anhydrous sodium sulfate and silica gel in a Pasteur pipette and concentrated further to 0.5 mL.

Gas chromatography was performed on a model HP5890 gas chromatograph (Hewlett-Packard, Palo Alto, CA) with an AT-1000 column (polyethylene glycol ester, 30-m \times 0.25-mm i.d., 0.2- μm film thickness, Alltech, Deerfield, IL) operated with hydrogen (1.5 mL min^{-1}) as a carrier, a split injector (injector temperature 220°C , injection volume $2 \mu\text{L}$, split ratio 75:1) flame ionization detector (300°C), and a temperature program from 45°C (5-min hold) to 150°C at $10^\circ\text{C min}^{-1}$, and to 220°C at $50^\circ\text{C min}^{-1}$ (with a 10-min hold). Components were identified by comparison of retention times and mass spectra with authentic standards from our own collection (Kjonaas et al., 1985), and were quantified by comparison of detector response with that of the appropriate internal standard. Gas chromatography-mass spectrometry analysis was performed on a Hewlett-Packard 5840A-5985B system at 70 eV, with the same column and separation conditions described above.

Rate of Monoterpene Synthesis

Pulse-labeling experiments were conducted using $^{14}\text{CO}_2$ with rooted plants in a 40-L plexiglass chamber. A pulse of 37 MBq of $^{14}\text{CO}_2$ was administered by acidification (with 1 mL of perchloric acid) of $\text{Na}_2^{14}\text{CO}_3$ (20 GBq mmol^{-1} ; DuPont/NEN, Wilmington, DE) dissolved in 0.5 mL of water. After the plants were placed in the chamber and the door

was sealed, the acid was added to a beaker containing the $\text{Na}_2^{14}\text{CO}_3$ solution by injection through a septum inlet in the chamber wall. Plants were exposed to $^{14}\text{CO}_2$ for 5 min under incandescent lights providing $250 \mu\text{mol m}^{-2} \text{s}^{-1}$ PAR. Temperature was maintained at 22°C by the use of water-filled trays placed on top of the chamber through which the light was filtered. $^{14}\text{CO}_2$ concentration in the chamber was measured by sampling air through the septum with a gastight syringe. Air samples were transferred to septum-capped, glass scintillation vials containing 0.2 mL of 1 N KOH and allowed to stand for 30 min to permit the trapping of $^{14}\text{CO}_2$ as carbonate. After the addition of scintillation cocktail (10 mL of 0.4% [w/v] DuPont/NEN Omnifluor in toluene/ethanol, 7:3, v/v), samples were analyzed in a liquid scintillation counter (Tricarb 460 CD, Packard Instruments, Meriden, CT) with a ^{14}C counting efficiency of 91%. Plants absorbed 20% to 25% of the administered $^{14}\text{CO}_2$ pulse. After the pulse, unincorporated $^{14}\text{CO}_2$ was exhausted from the chamber into a 10 N KOH trap, and the plants were left under the incandescent lights in a fume cabinet for 6 h. Replicate samples of leaves of the various ages were then harvested for analysis. Each sample, consisting of four to 20 leaves depending on leaf size, was immediately weighed and frozen at -20°C . At least three samples were analyzed for each age group.

Radiolabeled monoterpenes were extracted from peppermint leaves by simultaneous steam distillation-pentane extraction using a Likens-Nickerson apparatus (J&W Scientific, Folsom, CA) equipped with a standard condenser that was cooled with ice water. The leaves were heated to reflux in a flask with 30 mL of distilled water and $3 \mu\text{mol}$ of camphor as an internal standard. The organic phase consisted of 10 mL of pentane. Both solvents were heated for 30 min after refluxing had begun, and the pentane layer was then collected, dried over anhydrous sodium sulfate, and concentrated to 4 mL under a stream of nitrogen. A portion was removed for liquid scintillation counting, and the remainder analyzed by gas chromatography as described above to evaluate losses during extraction and concentration.

Rate of Monoterpene Loss

A 5-min pulse of $^{14}\text{CO}_2$ similar to that described above was administered to 5-week-old peppermint stems. Plants were then kept in a fume cabinet under lights for 3 d and then transferred to a confined greenhouse, where they were allowed to grow for an additional 6 weeks. The greenhouse had supplemental light (16-h photoperiod, $350\text{--}550 \mu\text{mol m}^{-2} \text{s}^{-1}$ of PAR at plant height) and a $30^\circ\text{C}/15^\circ\text{C}$ (day/night) temperature cycle. Samples were taken periodically from a group of leaves that were approximately 2 weeks old at the time of pulsing, and thus (according to measurements of the rate of monoterpene biosynthesis; see Fig. 2) at the stage of maximal monoterpene production. This was the same cohort of leaves used for measurements of monoterpene accumulation and the rate of biosynthesis. At least three samples were harvested at each time point, with each sample consisting of a pair of leaves from a single stem. To determine whether there were significant differences between time points, statistical

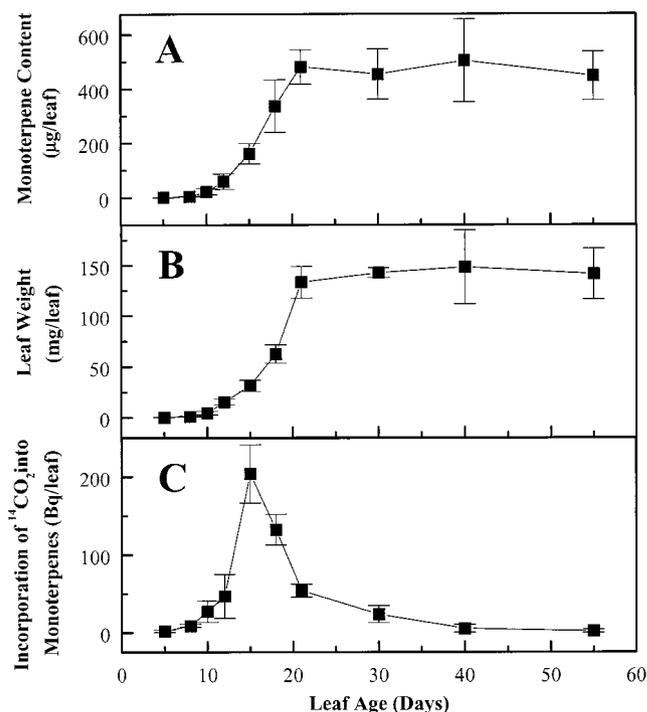


Figure 2. Changes in monoterpene content (A), leaf weight (B), and rate of monoterpene biosynthesis (C) during peppermint leaf development. Monoterpenes were extracted with diethyl ether and analyzed by gas chromatography. To determine the rate of monoterpene biosynthesis, leaves of various ages were exposed to a 5-min pulse of $^{14}\text{CO}_2$, and the incorporation of ^{14}C into monoterpenes was measured after a 6-h chase period. Each data point is the mean of three to six independent measurements. Bars indicate sd.

analyses were performed using SAS software (SAS Institute, Cary, NC).

Rate of Monoterpene Volatilization

The monoterpenes volatilized from peppermint were collected by headspace sorption from intact plants enclosed in a 40-L plexiglass chamber lined with a 0.005-cm layer of transparent polyvinylfluoride film (Tedlar, DuPont/NEN). A constant stream of air flowing at 1.1 L min^{-1} was drawn through the chamber with two small diaphragm vacuum pumps (no. 8803, Welch Vacuum Technology, Skokie, IL) connected in parallel. Flow was carefully regulated by controlling the pump speed and adjusting a valve on the inlet side of the chamber. The air exiting the chamber was passed through an adsorbent trap consisting of a $200 \times 7\text{-mm}$ glass tube containing 150 mg of Tenax GC (a polymer of 2,6-diphenyl-*p*-phenylene oxide, 60/80 mesh, Alltech) and 150 mg of Super Q (a polymer of divinylbenzene, 80/100 mesh, Alltech) held in place with plugs of silanized glass wool. Preliminary trials showed that the combination of these two adsorbents trapped the full spectrum of peppermint monoterpenes with higher efficiency than either one alone.

An additional adsorbent trap of identical construction was placed at the inlet of the chamber to purify incoming air. Trials with two such traps connected in series at the

chamber exit indicated that there was no detectable "break-through" (i.e. no loss of volatile monoterpenes from the first trap due to overloading, even when collections were carried out for periods of up to 8 h). Prior to initial use, the adsorbents were extracted exhaustively with diethyl ether and pentane, and before every subsequent use they were washed free of residual material with 50 mL of diethyl ether and dried with a stream of compressed air. In addition to the adsorbent traps, all fittings and connecting tubes were of glass. The entire volatile collection apparatus was contained in a controlled environment room that was adjusted such that light (16-h photoperiod) inside the plant chamber was maintained at an intensity of $350 \mu\text{mol m}^{-2} \text{s}^{-1}$ PAR with a constant temperature of 24°C .

Plants were placed in the chamber prior to volatile collection and left undisturbed with the chamber cover open for at least 2 h to ensure that any volatiles released by handling had dissipated. After the chamber was sealed and the airflow initiated, actual collection was not begun for an additional 2 h to ensure that an equilibrium concentration of volatiles had been reached in the chamber. Collection was then initiated for 3 to 6 h to accumulate sufficient material for accurate analysis. Trapped volatiles were desorbed from the trap with 25 mL of diethyl ether, and the sample was concentrated to 1 mL under a stream of nitrogen, and analyzed by gas chromatography as described above. Internal standards (75 μg each of isobutyl benzene and fenchone) were added prior to desorption to adjust for losses during sample processing. Tared vials of 2-carene and camphor (monoterpenes not found in peppermint) were also placed directly in the plant chamber to assess recovery of standard monoterpenes during the collection process. Collections made without plants (with pots and soil, with empty pots, or with an empty chamber) established that the background of monoterpenes present was negligible, and that the "carryover" from one collection to the next was insignificant.

RESULTS AND DISCUSSION

Monoterpenes in Peppermint Leaves Accumulate with Leaf Development

The pattern of monoterpene accumulation in peppermint leaves was measured by following a single cohort of leaves from initiation to senescence. Leaves were harvested at nine different stages during development, and the monoterpenes were extracted and analyzed by gas chromatography. The cohort of leaves chosen was initiated on 3-week-old (10- to 15-cm) stems and reached full expansion 21 d later when stems were 20 to 25 cm in length. Flower buds first appeared at the stem apex when leaves were 30 to 35 d old (stems 30–35 cm tall), and flowering commenced at approximately 45 to 50 d. By the time leaves were 55 d old, they had begun to senesce and abscise.

The monoterpene content of young leaves increased rapidly for the first 21 d of leaf development, then leveled off and was stable for the remainder of leaf life (Fig. 2A). Leaf weight showed a nearly identical trend (Fig. 2B). Similar profiles of monoterpene accumulation have been described

for leaves and fruits of other species, including dill (*Anethum graveolens*) (Porter et al., 1983), garden sage (*Salvia officinalis*) (Croteau et al., 1981), lemongrass (*Cymbopogon flexuosus*) (Singh et al., 1989), and caraway (*Carum carvi*) (Bouwmeester et al., 1998). In all of these taxa, the monoterpene content increases during the early stages of organ development and then remains relatively constant over the rest of organ life. In contrast, several studies of peppermint and other Lamiaceae have reported that monoterpene content declines as leaves age. These results are probably attributable to unusual growth conditions, such as extensive overhead irrigation (Croteau, 1977a), or to sampling schemes in which leaves of different ages were all harvested from the same stem at the same time (Srivastava et al., 1990; Srivastava and Luthra, 1991). The latter sampling method does not represent a true developmental gradient, since it is known that peppermint leaves initiated at early growth stages never attain monoterpene levels as high as leaves initiated at later stages (Burbott and Loomis, 1969).

The developmental changes in monoterpene accumulation in peppermint were accompanied by alterations in the monoterpene composition. The proportions of limonene, menthofuran, and pulegone declined as leaves aged, while those of 1,8-cineole, menthol, and neomenthol increased substantially (Table I). The major constituent, menthone, which was present in 5-d-old leaves at 36% of total monoterpenes, increased to approximately 75% at 15 d and then declined to 10% by the end of the study. Among the minor constituents, β -pinene, myrcene, and linalool showed increased percentages during development. Some of these compositional shifts have been documented in previous studies (Grahle and Holtzel, 1963; Brun and Voirin, 1991; Court et al., 1993; Voirin and Bayet, 1996; Rohloff, 1999).

Monoterpene Biosynthesis Is Restricted to a Brief Period Early in Leaf Development

The ontogenetic profile of monoterpene accumulation in peppermint may be influenced by both monoterpene synthesis and loss. To examine the rate of monoterpene biosynthesis, rooted plants were exposed to a 5-min pulse of $^{14}\text{CO}_2$. Leaves of different ages from the same cohort as that used to study changes in monoterpene content were harvested 6 h after $^{14}\text{CO}_2$ exposure, and the monoterpenes were isolated by simultaneous steam distillation-pentane extraction. The ^{14}C content of each extract was determined by liquid scintillation counting, and radio-gas chromatography was used to measure the percentage of radioactivity attributable to monoterpenes. There was a sharp peak of biosynthetic activity centered at 15 d, when leaves were still expanding, but only very low rates of biosynthesis were observed in leaves younger than 12 or older than 20 d (Fig. 2C). The rapid decline in biosynthetic rate between 15 and 20 d coincides with the leveling off of leaf monoterpene content (Fig. 2A) and the cessation of leaf expansion (Fig. 2B).

These results are consistent with those of several previous investigations on plant terpene formation. The biosynthesis of monoterpenes in *S. officinalis* leaves (Croteau et al., 1981), *Majorana hortensis* leaves (Croteau, 1977b), *C. carvi*

Table 1. Changes in monoterpene composition during leaf development in peppermint

Diethyl ether extracts prepared from leaves of different ages were analyzed by gas chromatography. Each value is the mean of at least three separate analyses. "tr" indicates that <0.1% was detected.

Compound	Leaf Age (d)								
	5	8	12	15	18	21	30	40	55
	% of total monoterpenes								
α -Pinene	1.5	1.2	1.1	0.8	1.5	1.4	1.8	1.6	2.0
β -Pinene	1.1	1.3	1.1	1.0	1.7	1.9	2.4	2.3	2.9
Sabinene	1.0	0.3	0.6	0.5	0.8	1.0	1.3	1.3	1.6
Myrcene	tr ^a	tr	0.4	0.3	0.4	0.5	0.6	0.6	0.7
Limonene	26.8	19.9	13.0	5.0	2.4	1.4	2.4	1.3	1.8
1,8-Cineole	tr	0.5	0.8	2.4	6.0	8.5	12.0	11.7	15.6
Menthone	36.4	53.6	66.0	75.5	58.5	63.5	40.2	33.3	10.0
Menthofuran	3.9	1.9	2.8	1.4	4.6	1.5	tr	tr	tr
Isomenthone	3.5	3.5	4.2	4.5	3.7	4.2	3.4	3.7	3.6
Linalool	0.4	0.2	0.2	0.2	0.2	0.3	0.8	0.6	1.0
Neomenthol	tr	tr	tr	tr	tr	tr	0.8	2.6	4.6
Menthol	tr	tr	tr	tr	1.6	2.6	12.0	35.2	54.0
Pulegone	24.9	16.9	8.5	7.1	17.5	11.8	21.1	4.6	0.8
α -Terpineol	tr	tr	0.1	0.1	0.2	0.3	0.4	0.3	0.2
Piperitone	0.4	0.8	1.2	1.1	0.8	1.0	0.7	0.9	1.2

^a tr, Trace (<0.1% detected).

fruits (Bouwmeester et al., 1998), *C. flexuosus* blades (Singh et al., 1989), and maritime pine (*Pinus pinaster*) foliage (Bernard-Dagan et al., 1982) is also restricted to a short interval during organ ontogeny. Among other terpenes, the biosynthesis of sesquiterpenes in *Heterothecca subaxillaris* (Mihaliak and Lincoln, 1989), diterpenes in *Newcastelia viscida* (Dell and McComb, 1978), and triterpenes in *Euphorbia lathyris* (Koops and Groeneveld, 1991) is also confined to early development. A ¹⁴CO₂ pulse-labeling experiment previously conducted with peppermint showed similar trends (Srivastava and Luthra, 1991). However, these latter results were deemed unreliable because cuttings rather than rooted plants were used (peppermint cuttings pulsed with ¹⁴CO₂ exhibit an artifactual turnover of monoterpenes [Mihaliak et al., 1991]), and because the monoterpene extracts were not examined for radiochemical purity.

The peak period of monoterpene biosynthesis in peppermint coincides with the time when the secretory cells of the glandular trichomes are metabolically active. Monoterpene synthesis in this species is localized to the secretory cells of glandular trichomes (Gershenzon et al., 1989; McCaskill et al., 1992), and the monoterpenes are discharged into a surmounting subcuticular storage compartment formed by expansion of the cuticle (Amelunxen, 1965). Anatomical studies indicated that the formation of Lamiaceae glands and filling of the subcuticular space occur only in actively growing, protodermal regions of the leaf surface (Werker et al., 1993); tracking the distribution of various gland developmental stages during peppermint leaf development has shown that 2-week-old leaves have especially high proportions of filling glands (G. Turner, J. Gershenzon, and R. Croteau, unpublished data). The occurrence of monoterpene biosynthesis in other species is also associated with the metabolic activity of glandular trichomes or other specialized secretory structures, such as secretory cavities and resin ducts, in which monoterpenes are synthesized and

sequestered. Such structures commonly differentiate in young, expanding tissue (Werker and Fahn, 1981; Charon et al., 1986; Russin et al., 1988), which may account for the fact that monoterpene biosynthesis is often highest in immature tissue.

Rate of Monoterpene Loss Is Negligible throughout Leaf Development

To determine the rate of monoterpene loss, similar pulse-labeling experiments were employed, except that plants were allowed to grow for 6 more weeks following the initial 5-min pulse of ¹⁴CO₂. Samples of a single, marked group of leaves (2 weeks old at the time of pulsing) were harvested periodically for monoterpene extraction and determination of ¹⁴C content. There was a rapid incorporation of ¹⁴C into monoterpenes during the first 2 d following the pulse, but no further significant changes in the radioactivity of extracted monoterpenes were observed over the remainder of the time course (Tukey's studentized range test, *P* > 0.05) (Fig. 3A). Thus, the metabolic pools of monoterpenes in peppermint leaves appear to be stable, and do not exhibit any detectable turnover. In contrast, the weight and total monoterpene content of these leaves increased steadily over the period of measurement (Fig. 3, B and C).

These results extend those of an earlier study in which peppermint plants exhibited no significant losses of radio-labeled monoterpenes over 29 h following a 5-min pulse of ¹⁴CO₂ (Mihaliak et al., 1991). They are also in accord with recent experiments conducted to examine terpenoid turnover conducted with lodgepole pine (*Pinus contorta*), Australian tea tree (*Melaleuca alternifolia*), garden sage (*S. officinalis*), and common tansy (*Tanacetum vulgare*) (Gershenzon et al., 1993). None of these species, which span a range of taxonomically distant plant families, contain different types of secretory structures, and include plants that con-

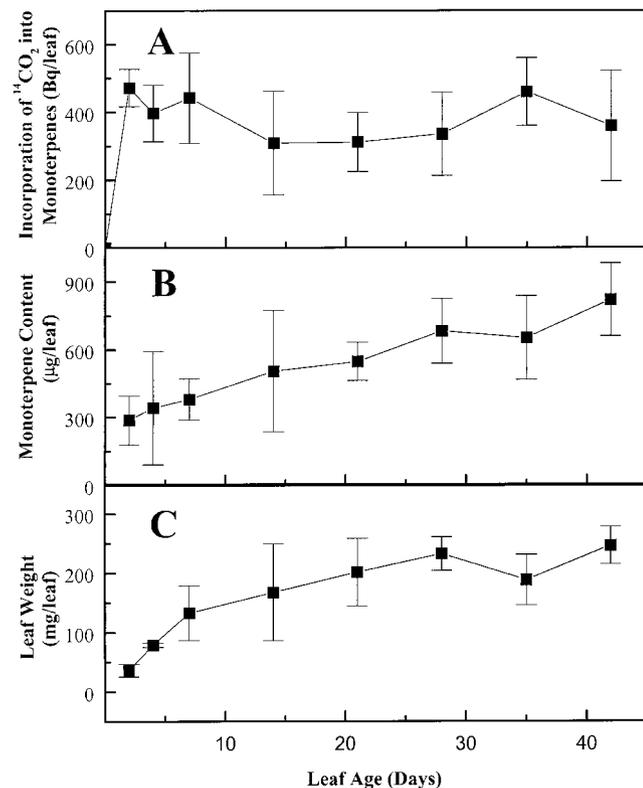


Figure 3. Lack of monoterpene turnover in peppermint leaves. Plants were exposed to a 5-min pulse of $^{14}\text{CO}_2$, and samples were harvested over the next 6 weeks for determination of monoterpene content. Incorporation of ^{14}C into monoterpenes (A) did not change significantly (Tukey's studentized range test, $P > 0.05$) over the time course of the experiment, indicating the lack of detectable monoterpene loss. In contrast, total monoterpene content (B) and leaf weight (C) increased steadily over the time course of the experiment. Each data point represents the mean of at least three independently measured samples, each consisting of a pair of leaves from a single stem. Bars indicate SD.

tain sesquiterpenes and diterpenes as well as monoterpenes, exhibited significant losses of terpenoids over a period of 10 to 14 d following an initial $^{14}\text{CO}_2$ pulse. In contrast, detached stems of peppermint studied under similar conditions displayed pronounced monoterpene turnover, an artifact also observed in numerous other investigations with detached tissues of other monoterpene-containing species (Francis and O'Connell, 1969; Croteau et al., 1972; Njar et al., 1989).

In the current study, peppermint leaves showed no significant loss of monoterpenes during an interval of 6 weeks after pulse labeling. At the end of this period, the group of marked leaves under study was 8 weeks old, and the experiment was terminated because most of the remaining marked leaves had begun to senesce. Previous investigations have indicated that peppermint monoterpenes can be metabolically degraded at later stages of leaf development (Croteau, 1988). A catabolic pathway was described involving the sequential reduction and glucosylation of menthene to neomenthol glucoside (Croteau and Martinkus, 1979), which is transported to the rhizome and degraded

(Croteau et al., 1984; Croteau and Sood, 1985). Although the present study provided no evidence for monoterpene catabolism, the large variances in monoterpene incorporation after pulse labeling (Fig. 3C) may have prevented its detection. Alternatively, the monoterpene degradation enzymes previously described may function not to degrade stored monoterpenes, but to detoxify monoterpenes that have come into contact with living cells after damage to the secretory structures. Monoterpenes such as limonene, linalool, and isomenthone have been demonstrated to be toxic to plant tissues (Brown et al., 1987), and disruption of monoterpene-containing secretory structures has been reported to cause damage to surrounding cells (Shomer and Erner, 1989; Loveys et al., 1992).

Rate of Monoterpene Volatilization Is Low and Does Not Significantly Influence Accumulation

Many species of plants release volatile monoterpenes into the atmosphere (Lerdau et al., 1997). Therefore, any consideration of the factors that regulate monoterpene accumulation in peppermint would be incomplete without an assessment of the rate of volatilization. The volatilization of peppermint monoterpenes was quantified by dynamic headspace sampling (Dobson, 1991) under a controlled-environment regime. Potted plants were placed in a chamber through which a stream of filtered air was passed. The air exiting the chamber passed through a cartridge packed with adsorbents to trap volatile organic compounds. Monoterpenes and other adsorbed substances were desorbed from the trap with organic solvent and analyzed by gas chromatography. Intact plants rather than cut leaves or stems were used, since detached tissues may have altered volatilization rates (Mookherjee et al., 1989; Nielsen et al., 1995).

Preliminary observations indicated that the rate of monoterpene volatilization from peppermint plants varied with light, temperature, time of day, and stage of development. For the purposes of this study, nonflowering, 6-week-old plants were measured under light and temperature conditions that were virtually identical to those used in the other experiments described here. The volatilization rate during the light period was $1.22 \mu\text{g h}^{-1} \text{ plant}^{-1}$, which can be expressed as $4.95 \mu\text{g h}^{-1} \text{ m}^{-2}$ leaf area or $0.254 \mu\text{g h}^{-1} \text{ per g}^{-1}$ dry weight. During the dark period, the volatilization rate was slightly higher at $1.73 \mu\text{g h}^{-1} \text{ plant}^{-1}$, or $7.02 \mu\text{g h}^{-1} \text{ m}^{-2}$ leaf area and $0.36 \mu\text{g h}^{-1} \text{ per g}^{-1}$ dry weight. According to the calculations in Table II, the monoterpenes emitted as volatiles represent only a small fraction of the total pool of monoterpenes present in the plant. Extrapolation over a typical 6-month growing period leads to the conclusion that $<5\%$ of total monoterpenes would be released to the atmosphere. This proportion could be somewhat higher under conditions of elevated temperature (Dement et al., 1975; Loreto et al., 1996) or higher humidity (Croteau, 1977a).

The low rate of monoterpene volatilization measured is consistent with the results of the $^{14}\text{CO}_2$ pulse experiments (Fig. 3), which showed no significant loss of labeled monoterpenes over 6 weeks of leaf development. Peppermint monoterpenes are stored in glandular trichomes within a

Table II. Comparison of monoterpene content and monoterpene volatilization rate of peppermint shoots

Measurements were performed on a set of six plants that were 6 weeks old, 25 cm tall, and had not yet begun to flower. Stored monoterpenes were extracted by soaking in diethyl ether and analyzed by gas chromatography. Volatiles were collected by headspace sorption from intact plants at 24°C (see "Materials and Methods" for details). Each plant was measured three times during the light period and once during the dark period for a span of 3 to 6 h at a time. All values are given as mean \pm SD.

Parameter	Measurement
Shoot monoterpene content	
Monoterpene content of average leaf ($n = 50$)	658 \pm 115 μg
Average number of leaves per plant ($n = 6$)	185 \pm 66.9 leaves
Monoterpene content of average stem ($n = 50$)	2.78 \pm 0.47 mg m^{-1}
Average length of total stems per plant ($n = 6$)	3.59 \pm 0.77 m
Total shoot monoterpene content per plant = (Ia \times Ib) + (Ic \times Id)	132 μg
Rate of monoterpene loss	
Average daytime volatilization rate per plant ($n = 18$)	1.22 \pm 0.64 $\mu\text{g h}^{-1}$
Average nighttime volatilization rate per plant ($n = 6$)	1.73 \pm 0.60 $\mu\text{g h}^{-1}$
Average monoterpene loss over 24-h period per plant = (IIa \times 16) + (IIb \times 8)	33.4 μg
Percentage	
Total shoot monoterpenes volatilized per day = (IIc)/(Ie)	0.025%
Total shoot monoterpenes volatilized per month = (IIc \times 30)/(Ie)	0.759%

subcuticular compartment that remains intact unless the leaf is damaged (Amelunxen, 1965; G. Turner, J. Gershenson, and R. Croteau, unpublished results), and therefore their persistence is not surprising. The rate of monoterpene volatilization from peppermint foliage is also lower than rates reported for most other monoterpene-emitting species, including *Salvia mellifera* (Dement et al., 1975), *Citrus* sp. (Winer et al., 1992), *Eucalyptus globulus* (Evans et al., 1982), *Quercus ilex* (Street et al., 1997), and various conifers (Janson, 1993; Staudt et al., 1997).

The composition of the monoterpenes emitted from peppermint is quite different from the composition of the total monoterpenes present in the plant (Fig. 4). Compared with the total pool, the mix of emitted monoterpenes contains higher proportions of menthofuran (12.4% versus 3.9%) and three olefins, β -pinene (10.1% versus 1.3%), sabinene (13.5% versus 0.7%), and limonene (16.4% versus 1.5%). In contrast, the two major monoterpenes found in mature leaves, menthone and menthol, are found in much lower proportions in the emitted mix than in the total pool (menthone at 10.1% versus 43.5%, and menthol at 19.5% versus 29.0%). If peppermint monoterpenes are ranked in order of volatility based on vapor pressure at 25°C (a ranking roughly approximating elution order on gas chromatography as indicated on the "compound" axis of Fig. 4), it can be seen that the more volatile substances are better represented in the mixture of emitted monoterpenes than in the mixture of total monoterpenes. Therefore, emission could occur directly from the total stored pool with greater representation of those compounds with greater volatility. However, two compounds of relatively low volatility, menthofuran and the α,β -unsaturated ketone, pulegone, are actually more abundant in the emitted mixture than in the total pool, suggesting that the membranes of the storage compartment might be selectively more permeable to some monoterpenes or that the emitted substances may be associated with an entirely different secretory compartment than the stored monoterpenes.

The existence of a separate compartment for the synthesis of emitted (as distinct from stored) monoterpenes has

been inferred from previous studies with other plant species. For example, herbivore damage to cotton (*Gossypium hirsutum*) foliage results in the immediate release of α -pinene, β -pinene, myrcene, limonene, and some sesquiterpenes from stored pools located in subepidermal glands (Loughrin et al., 1994). Herbivory to cotton also induces the release of a second set of monoterpenes and sesquiterpenes, including β -ocimene and linalool, that are largely absent from the stored pools and whose emission is not observed until 2 to 4 d after initial damage (Loughrin et al., 1994; Röse et al., 1996). An elegant series of [^{13}C]CO $_2$ tracer

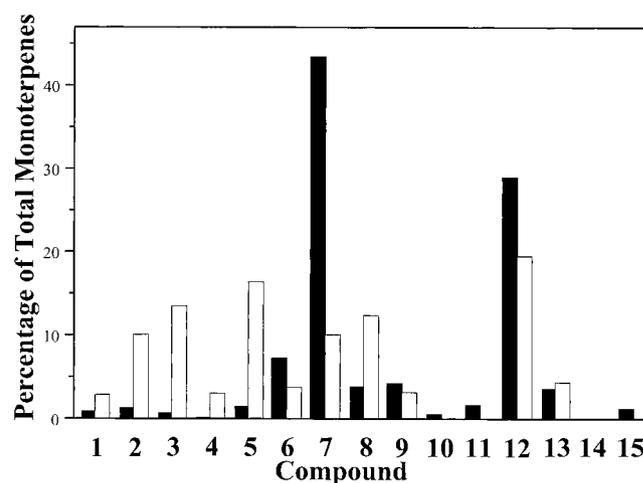


Figure 4. Comparison of the composition of monoterpenes stored (black bars) and emitted (white bars) by peppermint shoots. Aerial parts of 6-week-old plants were examined. Stored monoterpenes were extracted with diethyl ether and analyzed by gas chromatography. Emitted monoterpenes were collected by headspace sorption from intact plants (see "Materials and Methods" for details). Each value is the mean of five determinations. Key to compounds (in order of elution on gas chromatography, which approximates the range from most to least volatile): 1, α -Pinene; 2, β -pinene; 3, sabinene; 4, myrcene; 5, limonene; 6, 1,8-cineole; 7, menthone; 8, menthofuran; 9, isomenthone; 10, linalool; 11, neomenthol; 12, menthol; 13, pulegone; 14, α -terpineol; and 15, piperitone.

studies conducted following herbivore damage to cotton (Pare and Tumlinson, 1997) revealed that the immediately emitted terpenes arise from stored pools, whereas most of the later-emitted compounds are synthesized *de novo* just prior to their release. These late-emitted terpenes, like the monoterpenes released from peppermint, may be part of a secretory system that is distinct from that producing the stored terpenes, and under the control of different physiological factors. Thus, their emission would have no effect on the monoterpenes of stored pools. In the case of peppermint, it is possible that the emitted monoterpenes derive from the smaller, sparsely distributed capitate glands, whereas the bulk of the monoterpene pool is produced and permanently stored in the peltate glandular trichomes (Fahn, 1979).

CONCLUSIONS

The accumulation of monoterpenes in developing peppermint leaves could, in theory, be influenced by both the rate of monoterpene synthesis and the rate of monoterpene loss. However, no evidence for significant loss of monoterpenes during leaf development as a result of volatilization, metabolic degradation, or other routes was observed. The pattern of monoterpene accumulation can be explained solely by changes in the rate of monoterpene biosynthesis. The large increase in monoterpene content of 12- to 20-d-old leaves coincides with the peak period of monoterpene biosynthesis. Prior to this stage, monoterpene biosynthesis and accumulation are negligible, while after this stage, the rate of synthesis declines precipitously and monoterpene accumulation ceases. The preeminent role of biosynthesis in controlling monoterpene accumulation in peppermint has stimulated further interest in glandular trichome development (G. Turner, J. Gershenzon, and R. Croteau, in preparation) and in those factors that regulate the rate of monoterpene biosynthesis, including the activities of individual pathway enzymes and the expression of the corresponding structural genes, as described in the following paper (McConkey et al., 2000).

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