Biosynthesis and Emission of Terpenoid Volatiles from Arabidopsis Flowers

Feng Chen, a,b,1 Dorothea Tholl, b,1 John C. D’Auria, a Afgan Farooq, b Eran Pichersky, a,2 and Jonathan Gershenzonb

a Department of Molecular, Cellular, and Developmental Biology, University of Michigan, Ann Arbor, Michigan 48109
b Max Planck Institute for Chemical Ecology, Beutenberg Campus, Winzerlaer Strasse 10, D-07745 Jena, Germany

Arabidopsis is believed to be mostly self-pollinated, although several lines of genetic and morphological evidence indicate that insect-mediated outcrossing occurs with at least a low frequency in wild populations. Here, we show that Arabidopsis flowers emit both monoterpenes and sesquiterpenes, potential olfactory cues for pollinating insects. Of the 32 terpene synthase genes in the Arabidopsis genome, 20 were found to be expressed in flowers, 6 of these exclusively or almost exclusively so. Two terpene synthase genes expressed exclusively in the flowers and one terpene synthase gene expressed almost exclusively in the flowers were characterized and found to encode proteins that catalyze the formation of major floral volatiles. A β-glucuronidase fusion construct with a promoter of one of these genes demonstrated that gene expression was restricted to the sepals, stigmas, anther filaments, and receptacles, reaching a peak when the stigma was receptive to cross pollen. The observation that Arabidopsis flowers synthesize and emit volatiles raises intriguing questions about the reproductive behavior of Arabidopsis in the wild and allows detailed investigations of floral volatile biosynthesis and its regulation to be performed with this model plant system.

INTRODUCTION

One of the principal reasons for the success of Arabidopsis as a model system in plant biology is the ability of this species to set copious numbers of seeds under greenhouse and growth-room conditions that arise nearly exclusively from self-pollination (Meyerowitz and Somerville, 1994). This high degree of self-pollination, along with its small flower size (2 to 3 mm long), inconspicuous white petals, and lack of strong scent, suggest that Arabidopsis has not been selected to attract floral visitors for cross-pollination. However, several lines of evidence indicate that insect-mediated cross-pollination may occur in wild populations. First, in the ontogeny of the flower, the stigma becomes receptive and protrudes out of the petals before the stamens mature, providing a brief window for cross-pollination (Jones, 1971). Second, at the base of the stamens, Arabidopsis has floral nectaries that secrete a mixture of sugars that could serve as a reward for floral visitors (Davis et al., 1998). Third, small insects, including hover flies, also have been observed to visit Arabidopsis flowers (Jones, 1971; Snape and Lawrence, 1971). These insects might act as vectors for pollen transfer, although no direct evidence for their role in pollination has been presented. Finally, the frequency of polymorphic loci in Arabidopsis populations is high enough to exclude simple mutation as the source of variation (Loridon et al., 1998). Cross-pollination has been estimated to be as high as 2%, based on segregation for some morphological traits in the progeny of Arabidopsis inbred lines grown outdoors (Snape and Lawrence, 1971), although lower estimates have been obtained from analyses of allozyme loci in wild populations (Abbott and Gomes, 1989).

If Arabidopsis flowers are visited by insect pollinators, this species is likely to possess visual or olfactory cues for pollinator attraction. The emission of volatiles is a major feature of many insect-pollinated flowers (Knudsen et al., 1993; Dudareva and Pichersky, 2000). Mixtures of volatile compounds drawn from several classes of plant metabolites, including terpenes, phenylpropanoids, fatty acid derivatives, and nitrogen- or sulfur-containing compounds (Dudareva and Pichersky, 2000), have been shown to attract a large variety of insects (as well as some mammals) that vector pollen. However, there are no previous reports regarding the detection of volatiles emanating from Arabidopsis flowers either by chemical methods or the human nose.

The detection of floral volatile emission in this species would allow the vast genetic and genomic resources available for this model plant to be applied to studying fundamental questions about the regulation and evolution of volatile

1 These authors contributed equally to this article.
2 To whom correspondence should be addressed. E-mail lex@umich.edu; fax 734-647-0884.
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monoterpenes (C\textsubscript{10}) and geranylgeranyl diphosphate (C\textsubscript{20}) are involved in the production of floral volatiles. Work on the terpene pathway, including dimethylallyl diphosphate (C\textsubscript{10}), farnesyl diphosphate (FPP; C\textsubscript{15}), and geranylgeranyl diphosphate (C\textsubscript{20}), to hemiterpenes (C\textsubscript{5}), monoterpenes (C\textsubscript{10}), sesquiterpenes (C\textsubscript{15}), and diterpenes (C\textsubscript{20}), respectively (Cane, 1999; MacMillan and Beale, 1999; Wise and Croteau, 1999). Nearly all of the products of hemiterpene, monoterpenes, and sesquiterpene synthases are volatile under ambient temperature and atmospheric pressure, and monoterpenes and sesquiterpenes are very typical floral volatiles. However, to date only one characterized terpene synthase, (S)-linalool synthase from Clarkia breweri (Dudareva et al., 1996), has been shown to be involved in the production of floral volatiles. Work on the TPS genes of Arabidopsis also has just begun. Of the 32 predicted functional sequences, only 4 have been characterized. Two monoterpenes synthases have been described, a myrcene/(E)-\beta-cymene synthase (Bohmann et al., 2000) and an (E)-\beta-cymene synthase (Fäldt et al., 2002). Two diterpene synthases also have been reported that are involved in gibberellin formation (Sun and Kamiya, 1994; Yamaguchi et al., 1998).

In this investigation, we searched for volatiles released from flowers and vegetative parts of Arabidopsis using headspace sampling methods. We found a blend of volatile terpenes to be emitted specifically from inflorescences and characterized three TPS genes involved in their biosynthesis.

RESULTS

Arabidopsis Flowers Emit Monoterpenes and Sesquiterpenes

The volatile constituents of Arabidopsis flowers were sampled by passing air over flowering plants and then through a charcoal trap to adsorb organic constituents. A semiopen headspace sampling method was used as well as a sensitive closed-loop stripping method (Donath and Boland, 1995). After desorption of the trap with solvent and analysis by gas chromatography–mass spectrometry (GC–MS), a range of monoterpane and sesquiterpene constituents was detected.

To determine whether these substances were emitted by the flowers, fruits, or vegetative parts, we performed a series of headspace collections in which inflorescences or siliques (fruits) were removed from plants (Figure 1A). By comparing the major volatiles detected, it was determined that the three most abundant monoterpenes (\(\beta\)-myrcene, limonene, and linalool) and most of the major sesquiterpenes, including (\(-\))-(E)-\beta-caryophyllene, (+)-thujopsene, \(\alpha\)-humulene, (E)-\beta-farnesene, (\(+\))-\beta-chammigrene, and (\(-\))-cuparene, were emitted almost exclusively from flowers. Together, monoterpenes and sesquiterpenes represented \(>60\%\) of the total amount of volatiles collected from the flowers (estimated by relative GC–MS peak area; Figures 1B and 1C). The remainder consisted primarily of aliphatic aldehydes or alcohols, including octanal, 2-ethyl-hexanol, nonanal, and decanal. These non-terpenoids also were present in headspace collections from siliques and vegetative parts of flowering plants (data not shown).

Several Arabidopsis TPS Genes Are Expressed Exclusively or Almost Exclusively in the Flowers

To determine which of the predicted functional Arabidopsis TPS genes might be involved in the formation of floral volatiles, the expression of all 32 of these (Aubourg et al., 2002) (Table 1) was determined in various organs of the plant with reverse transcriptase–mediated (RT) PCR. For each gene, specific internal primers designed to span more than one exon were used to discriminate against amplification products from contaminating genomic DNA. No such amplification products were observed, and control PCR experiments omitting reverse transcriptase also resulted in no product. Analysis was performed with RNA isolated from leaves, flowers, siliques, stems, and roots of 6-week-old flowering plants. In these experiments, results for different genes in the same organ are directly comparable, because an identical aliquot of cDNA from the original RT reaction was used in each PCR procedure. To determine whether equal amounts of cDNA were included in the reactions involving different organs, we also performed RT-PCR with primers designed for \(\beta\)-tubulin.

After RT-PCR, amplified fragments from mRNAs of 25 of the 32 TPS genes were obtained from at least one organ (Figure 2). The expression of 11 genes was detected in leaves, 20 in flowers, 11 in siliques, 9 in stems, and 12 in roots. Seven genes (At2g23230, At3g29110, At4g20210, At4g20200, At3g32030, At1g48800, and At1g48820) showed no expression in any of the five organs, whereas eight genes, including the two involved in gibberellin biosynthesis (At1g79460 and At4g02780), were expressed in nearly all of them. Six genes (At3g25810, At4g16730, At1g61680, At5g44630, At3g14490, and At5g23960) showed complete or almost complete flower-specific expression.
Figure 1. Volatile Terpenes Emitted from Arabidopsis (Columbia Ecotype).

(A) Release rates of major terpenes from 6-week-old flowering plants and parts of these plants determined by dynamic headspace sampling. Inflorescences were the source of emission for the monoterpenes and most of the sesquiterpenes. Data represent the mean of four replications ± SE. The emission rate presented is that of a single plant per hour, although collections were made from five plants simultaneously over 8-h periods.

(B) Gas chromatography of monoterpenes collected from 150 cut inflorescences during 12 h of closed-loop stripping (Donath and Boland, 1995). The peak of limonene coeluted with 2-ethyl-hexanol on the (5% phenyl)-polymethylsiloxane column, as shown here, but it was separable from this compound on a polyethylene glycol column.

(C) A later portion of the same chromatogram as in (B), showing the sesquiterpene hydrocarbon region. The inset depicts a portion of the chromatogram of the same sample run on a polyethylene glycol (DB-WAX) column, which shows an improved separation of some components. Compounds marked with dots represent additional sesquiterpene hydrocarbons identified by GC-MS but not yet confirmed by comparison with authentic standards. IS indicates the internal standard, nonyl acetate.

(D) Structures of the compounds identified by numbers in (A), (B), and (C). The number of each compound in (D) corresponds to the numbered columns in (A) and the numbered peaks in (B) and (C). Chirality was determined for all compounds except β-elemene and β-sesquiphellandrene.
Table 1. Primers for RT-PCR Analysis

<table>
<thead>
<tr>
<th>Gene Name</th>
<th>Sequences (5' to 3')</th>
</tr>
</thead>
<tbody>
<tr>
<td>At3g25810</td>
<td>F: TATATTGATGGATCAACAGC&lt;br&gt;R: TTGAAACCAGCATGGAAGAG</td>
</tr>
<tr>
<td>At3g25830</td>
<td>F: ATGATTGATGATGCACTTAAAGAA&lt;br&gt;R: GGAACACCTTAAAGTATTAAAGGT</td>
</tr>
<tr>
<td>At2g24210</td>
<td>F: CTGTTGATGAAAGAGAGAGG&lt;br&gt;R: ATGAAAGATGCAAGGAGAAG</td>
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<tr>
<td>At4g16740</td>
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<td>At1g61680</td>
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<tr>
<td>At2g24210</td>
<td>F: CTGTTGATGAAAGAGAGAGG&lt;br&gt;R: ATGAAAGATGCAAGGAGAAG</td>
</tr>
</tbody>
</table>

F, forward primer; R, reverse primer.

We chose three of these flower-specific genes for further analysis. RNA gel blots (Figure 3) verified that At3g25810 and At5g23960 were expressed exclusively in the flowers. The blots also showed exclusive expression of At1g61680 in the flowers, even though RT-PCR showed that relatively low levels of the transcript of this gene also occurred in the stem, in addition to the much higher levels of transcript in the flowers (Figure 2). This discrepancy likely is attributable to the higher sensitivity of RT-PCR compared with RNA gel blot analysis.

Three Flower-Specific TPS Genes Encode Proteins That Catalyze the Formation of Floral Volatiles

To investigate the catalytic activities of the proteins encoded by At3g25810, At1g61680, and At5g23960, the three flower-specific TPS genes, cDNAs of these obtained by RT-PCR were expressed in Escherichia coli, and the resulting proteins were assayed for terpene synthase activity. Control assays were performed with crude extracts of E. coli (same strain) carrying the same expression vector without any TPS genes inserted.

In plants, the biosynthesis of monoterpenes appears to be localized in plastids, the site of formation of the monoterpeneprecursor GPP, whereas sesquiterpenes are synthesized in the cytosol/endoplasmic reticulum compartment, the site of FPP formation (Lichtenthaler, 1999). Thus, the known monoterpen synthases, all of which are encoded by nuclear genes, possess an N-terminal transit peptide that directs their import into the plastid followed by cleavage of the transit peptide (Turner et al., 1999). Based on sequence analyses, two of the three flower-specific TPS genes, At3g25810 and At1g61680, have transit peptide-like sequences at the N terminus of their proteins (Figure 4) (Aubourg et al., 2002), and they fall into a clade that includes the two previously identified monoterpene synthases of Arabidopsis, At2g24210 and At4g16740 (Bohm et al., 2002; Fäldt et al., 2002). Because it has been shown for other monoterpene synthases that the mature (cleaved) protein often has higher specific activity when expressed for E. coli than the full-length preprotein (Williams et al., 1998), we constructed constructs both with and without the N-terminal sequence.

Both the full-length protein encoded by At3g25810 and its mature version lacking the putative transit peptide catalyzed the formation of eight monoterpenes from GPP, including six monoterpenes, including (E)-α-farnesene, and (E,E)-β-farnesene, (E)-β-farnesene, and trans-α-bergamotene. However, these are not thought to be physiologically relevant products because they are not constituents of the floral volatile blend and the specific activity of the At3g25810 enzyme with GPP was ~40-fold higher than that with FPP (1.7 ± 0.1 pkat/mg protein versus 0.04 ± 0.01 pkat/mg protein).
The complete protein encoded by At1g61680 and its mature version without the putative transit peptide both catalyzed the conversion of GPP to only one product, (+)-3S-linalool (Figure 5B). Incubation of the expressed protein with FPP led to the formation of very low amounts of nerolidol, but this sesquiterpene alcohol is not present in Arabidopsis floral volatiles, and the rate of this reaction was 50-fold lower than the rate with GPP.

The open reading frame of At5g23960 appears to encode a protein with no transit peptide (Figure 3) (Aubourg et al., 2002). When synthesized in E. coli, the At5g23960 protein catalyzed the formation of (−)-E-β-caryophyllene and α-humulene as well as trace amounts of (−)-α-copaene and β-elemene from FPP (Figures 5C and 6). The enzyme did not convert GPP to any monoterpenes.

The Promoter of At3g25810 Is Active in Specific Floral Parts

RT-PCR and RNA gel blot analyses showed flower-exclusive or nearly exclusive expression of At3g25810, At1g61680,
and At5g23960 (Figures 2 and 3) but could not identify the specific part(s) of the flower where gene activity occurs. To investigate the tissue localization of At3g25810 expression, the promoter region (1.4 kb) of At3g25810 was fused to β-glucuronidase (GUS), and the resulting construct was used to transform Arabidopsis of the Columbia ecotype. Flowers and leaves of transgenic plants then were stained for GUS expression. GUS staining was observed only in flowers, confirming the results of the RT-PCR and the RNA gel blot experiments. Among the floral parts, staining was evident in the sepals, stigmas, anther filaments, and receptacles and was visible in mature, but not immature, buds (Figure 7A), in newly opened flowers in which the stigma protrudes above the anthers and petals (Figure 7B), and at a more mature stage of floral development at which the stamens have elongated and the pollen shed (Figures 7C and 7D).

DISCUSSION

Arabidopsis Flowers Release a Mixture of Volatiles Dominated by Terpenes

Arabidopsis inflorescences emit a complex blend of monoterpenes, sesquiterpenes, aliphatic aldehydes, and alcohols. Although the aldehydes and alcohols also are emitted from vegetative parts of mature plants and were detected in volatile collections of younger, rosette-stage plants (Van Poecke et al., 2001), the terpenoids are almost completely specific to the flowers and represent >60% of the total amount of volatiles collected from these organs. The individual terpene constituents are nearly all well known from higher plants. For example, the sesquiterpene hydrocarbon (−)-(E)-β-caryophyllene, the most abundant terpene collected, is very widely distributed in the plant kingdom (Joulain and König, 1998; Cane, 1999) and is a floral volatile of many other plant species (Knudsen et al., 1993). However, the tricyclic α-barbatene, which is common in liverworts, has been reported only rarely from vascular plants (König et al., 1996).

The terpene emission level from Arabidopsis flowers is considerably lower than what is observed from the strongly scented C. breweri, a model wild plant for the study of floral scent. A single mature Arabidopsis plant, which has on average 80 open flowers, emits a total of 23.4 ± 1.4 ng of terpenes per hour. By contrast, a single mature C. breweri plant, which has on average only two flowers in the wild, emits a total of 3.3 to 5.9 µg of volatiles (terpenes, phenylpropanoids, and benzenoids) per hour (calculated from Raguso and Pichersky, 1995), a 140- to 250-fold difference. However, emission from flowers of the closely related species C. concinna is 30- to 60-fold lower than that from C. breweri (Raguso and Pichersky, 1995). Unfortunately, it is difficult to make further interspecific comparisons because absolute floral emission values for wild species generally are not reported. Nevertheless, Arabidopsis flowers clearly emit a terpene-rich blend of volatiles that, although readily detectable by modern gas chromatography, is scentless to the human nose as a result of its high proportion of sesquiterpene hydrocarbons, the low abundance of monoterpenes, and the absence of pungent phenylpropanoids and benzenoids.

The Formation of Arabidopsis Floral Terpenes Is Catalyzed by Terpene Synthases

The volatile terpenes of Arabidopsis inflorescences are biosynthesized by the action of terpene synthases, enzymes that catalyze the conversion of the ubiquitous isoprenoid pathway intermediates, GPP and FPP, to monoterpenes and sesquiterpene products, respectively (Cane, 1999; Wise and Croteau, 1999). Of the 32 intact genes in the Arabidopsis genome that have sequence homology with known TPS genes, we selected three (At3g25810, At1g61680, and At5g23960) for further characterization that were expressed exclusively or almost exclusively in flowers, based on both RT-PCR (Figure 2) and RNA gel blot analysis (Figure 3), and that appeared likely to encode monoterpene and sesquiterpene synthases, based on their sequence similarity to other members of this gene family. When assayed with GPP, At3g25810 catalyzed the formation of multiple monoterpens. Three of these, limonene, β-myrcene, and (E)-β-ocimene, were detected as Arabidopsis floral volatiles in our headspace collections (Figure 1), whereas others, including α-pinene and sabinene, were detected occasionally in low amounts (data not shown). The remainder may be retained in the cell...
and not emitted at detectable levels, or the mixture of products produced by this enzyme in vitro may be different from what occurs in vivo (Jia et al., 1999; Crowell et al., 2002). The ability of a single terpene synthase to form multiple products is a rather common feature of this enzyme family, attributed to the generation of carbocationic intermediates during the reaction that can have more than one metabolic fate (Crock et al., 1997; Cane, 1999; Wise and Croteau, 1999).

Not all terpene synthases make multiple products. At1g61680 catalyzed the formation from GPP of only \((\dfrac{1}{2})\)-\(S\)-linalool, which also was detected in Arabidopsis floral scent (Figure 1). When assayed with FPP, both the At3g25810 and At1g61680 proteins catalyzed the formation of some sesquiterpenes. However, these are likely not to be significant reactions in planta because (1) they occur at significantly lower rates in vitro compared with reactions that form monoterpene from GPP, (2) the sesquiterpene products of such reactions were not found in the blend of floral volatiles, and (3) the proteins themselves may never contact free FPP because they contain transit peptide–like sequences at the N terminus and so are likely targeted to the plastids (Turner et al., 1999), where free FPP is not present (Lichtenthaler, 1999).

The third TPS gene, At5g23960, encodes a protein that catalyzed the formation of five sesquiterpenes. Of these, \((-\dfrac{1}{2})\)E:\(\dfrac{1}{2}\)-\(S\)-linalool, \((-\dfrac{1}{2})\)S-\(S\)-linalool, \((-\dfrac{1}{2})\)E:\(\dfrac{1}{2}\)-\(S\)-linalool, \((-\dfrac{1}{2})\)S-\(S\)-linalool, and \((-\dfrac{1}{2})\)S-\(S\)-linalool, which also was detected in Arabidopsis floral scent (Figure 1). When assayed with FPP, both the At3g25810 and At1g61680 proteins catalyzed the formation of some sesquiterpenes. However, these are likely not to be significant reactions in planta because (1) they occur at significantly lower rates in vitro compared with reactions that form monoterpene from GPP, (2) the sesquiterpene products of such reactions were not found in the blend of floral volatiles, and (3) the proteins themselves may never contact free FPP because they contain transit peptide–like sequences at the N terminus and so are likely targeted to the plastids (Turner et al., 1999), where free FPP is not present (Lichtenthaler, 1999).

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The Members of the Arabidopsis TPS Gene Family Have Different Patterns of Organ Expression

Although many plant terpenes, such as sterols, carotenoids, and gibberellins, have well-known functions in plant growth and development, the roles of the vast majority of this large
class of plant natural products are unknown. The presence
of a large set of terpene synthase genes in the model spe-
cies Arabidopsis affords an excellent opportunity to identify
the functions of plant terpenes. Based on the RT-PCR anal-
ysis presented here (Figure 2), the Arabidopsis TPS
genes
have dramatically different organ expression patterns. Some
are expressed almost exclusively in flowers, roots, or si-
lques, whereas others are expressed to a significant degree
in two or more organs. These results suggest that some TPS
genesis, such as those demonstrated to be involved in floral
volatile production in this study, may produce specialized
products, whereas others may have a more general function.
Further studies to characterize additional Arabidopsis TPS
genesis, manipulate terpene production, and assess pheno-
typic changes should significantly increase our understanding
of the biological raison d’être of plant terpenoid metabolites.

Arabidopsis Floral Volatiles May Function in Pollinator
Attraction and in Other Roles

It is tempting to hypothesize that the biological role of the
terpenes emitted from Arabidopsis inflorescences is to at-
tract insects for cross-pollination. Flowers of many species
emit terpenes and other volatiles for pollinator attraction. Al-
though Arabidopsis readily self-pollinates when grown in-
doors, the presence of floral nectaries (Davis et al., 1998),
the protogynous mode of floral development (early matura-
tion of the stigma while the anthers are still immature)
(Jones, 1971), and several experimental studies (Snape and
Lawrence, 1971; Abbott and Gomes, 1989; Loridon et al.,
1998) all suggest that at least a low degree of cross-pollina-
tion exists in wild populations. Thus, floral volatiles could
provide cues for pollinator attraction. Indeed, anecdotal ob-
servations of various insects visiting Arabidopsis flowers
have been reported (Jones, 1971; Snape and Lawrence,
1971), and we have observed syrphid flies (Diptera: Syr-
phidae) visiting Arabidopsis flowers near Jena, Germany.
However, no information is available regarding whether
these insects facilitate cross-pollination and whether they
can perceive olfactory cues from Arabidopsis. It has been
shown that the antennae of the moth Hyles lineata can de-
tect the monoterpenes linalool and some other C. breweri
floral volatiles at concentrations similar to those produced by
emission from Arabidopsis flowers (Raguso et al., 1996).
Nevertheless, similar experiments need to be conducted

Figure 6. Identification of (E)-β-Caryophyllene as the Major Enzyme Product of At5g23960 Catalysis.

(A) Mass spectrum of (E)-β-caryophyllene produced by the incubation of FPP with a cell-free extract of E. coli BL21 Codon Plus expressing the
At5g23960 protein.

(B) Mass spectrum of authentic (E)-β-caryophyllene standard obtained under the same conditions as in (A).
m/z, mass-to-charge ratio.
with potential insect visitors of Arabidopsis to determine if the compounds emitted from its flowers function as attractants and if the insects so attracted can effect cross-pollination.

The floral volatiles of Arabidopsis also could play a variety of other roles instead of or in addition to pollinator attraction. For example, many terpenes, including β-myrcene, (E)-β-ocimene, linalool, and (E)-β-caryophyllene, react readily with ozone and other reactive oxygen species (Calogirou et al., 1999; Loreto and Velikova, 2001). Thus, Arabidopsis floral volatiles could function to protect the reproductive organs, with their valuable germ line cells, from oxidative damage. A variety of monoterpene and sesquiterpenes also are reported to have antimicrobial activity (Deans and Waterman, 1993). Hence, Arabidopsis floral terpenes also could help defend floral organs from bacterial or fungal infestation. The moist surface of the stigma may be an ideal environment for fungal growth. Finally, it is conceivable that, although Arabidopsis floral terpenes once had a function in the evolutionary past, they no longer do today. Arabidopsis thaliana diverged only recently (~5 million years ago) from a clade made up of A. halleri, A. lyrata, and A. petraea (Koch et al., 2000), a lineage of self-incompatible perennials that presumably relied on insect pollination. Thus, floral terpenes may have been present in a recent ancestor of A. thaliana and simply have not yet been eliminated by genetic drift or natural selection.

Our observations of transgenic plants containing the promoter-GUS fusion construct for the At3g25810 gene also have implications for the function of floral volatiles in Arabidopsis, but these results need to be considered with caution until they are confirmed by additional methods, such as in situ hybridization. GUS activity was observed in four floral

**Figure 7.** Expression Patterns of the At3g25810 Promoter–GUS Fusion Gene during Flower Development.

GUS staining was observed in sepals, stigmas, anther filaments, and receptacles of mature, but not immature, buds (A), newly opened flower (B), and older flower (C). In (D), petals and sepals were removed to show clearly the staining in the stigma of the older flower. The newly opened flower (B) is in the protogynous stage, in which the receptive stigma, protruding above the petals and the immature anthers, is accessible to cross-pollination. The older flower (C) is in the autogamous stage, in which the stamens have elongated to the level of the stigma and dehisced. The proximity of the stamens to the stigma at this stage facilitates self-pollination.
parts: sepals, stigmas, anther filaments, and receptacles (Figure 7). Activity in the stigmas is consistent with the idea that terpene volatiles function to protect the stigma from pathogen attack or oxidative damage, whereas the occurrence of GUS staining in the sepals, filaments, and receptacles suggests a function involving the entire flower, such as pollinator attraction. Interestingly, several genes involved in nonterpenoid floral scent biosynthesis in C. breweri and Antirrhinum majus have been reported to be expressed exclusively in petals, rather than in sepals or other parts of the flower (Wang et al., 1997; Dudareva et al., 1998, 2000). However, the linalool synthase gene of C. breweri also is expressed in the stigma, in addition to showing lower activity in the petals (Dudareva et al., 1996).

The Floral Volatiles of Arabidopsis May Have Implications for the Evolution of This Species and Its Population Genetic Structure

The emission of volatiles from flowers of Arabidopsis may have evolutionary as well as ecological significance. If floral volatiles increase visitation by potential insect pollinators, this will raise the degree of outcrossing and so alter the genetic structure of Arabidopsis populations. Even if the resulting degree of outcrossing is low, as observed by Abbott and Gomes (1989), progeny arising from outcrossing may have greater reproductive fitness because of the genetic polymorphisms they contain (Agren and Schemske, 1993). This may have led to the retention of traits that promote outcrossing even in species that are largely self-fertilizing. Thus, floral volatiles and other traits involved in the attraction of floral visitors may have been critical factors in the evolutionary history of Arabidopsis. From this perspective, the TPS genes involved in the formation of the major floral volatiles may represent valuable markers for the study of the phylogenies of this model species and its nearest relatives.

METHODS

Plant Material

Arabidopsis thaliana of the Columbia ecotype was grown from seeds in a controlled-climate room (22°C, 55% RH, and 100 µmol·m⁻²·s⁻¹ PAR) under long-day conditions (16-h-light/8-h-dark photoperiod) for up to 6 weeks. At this time, plants had developed shoots and flowers (both primary and secondary bolts).

Volatile Analysis

Volatiles from 6-week-old flowering plants were collected in a dynamic headspace sampling system that was installed in a controlled-climate chamber (23°C, 70% RH, and 150 µmol·m⁻²·s⁻¹ PAR). Five plants with intact root balls wrapped in aluminum foil were placed in a 4-L glass chamber that consisted of a flat flange reaction vessel and a flat flange lid with four necks, closed with Teflon stoppers. Charcoal-purified air entered the chamber at a flow rate of 1 L/min from the top through a Teflon hose (whose outlet was positioned in the lower part of the glass vessel). Volatiles were collected for 8 h by pumping air from the chamber through activated charcoal traps (1.5 mg) at a rate of 0.9 L/min. The remaining air was vented through the top of the chamber. For volatile collections from flowers or silicles, these organs were cut from five plants and transferred to a small beaker containing tap water or moistened filter paper.

To collect less abundant volatiles from inflorescences, a high-sensitivity closed-loop stripping method was applied (Donath and Boland, 1995). Ninety or 150 inflorescences were transferred to small glass beakers filled with tap water and were placed in an 1-L bell jar. Under continuous air circulation, emitted volatiles were collected for 12 h on activated charcoal traps that had been fitted into a steel column connected to the circulation pump. Volatiles were eluted with 40 µL of CH₂Cl₂, and 120 ng of nonyl acetate was added as an internal standard.

Samples from volatile collections and from solid-phase microextraction of terpene synthase assays (see below) were analyzed on a Hewlett-Packard 6890 gas chromatograph coupled to a Hewlett-Packard 5973 quadrupole mass selective detector. Separation was performed on (5%-phenyl)-methylpolysiloxane or polyethylene glycol columns (J&W Scientific, Folsom, CA) of 30 m × 0.25 mm i.d. × 0.25 mm thickness. Helium was the carrier gas (flow rate of 2 mL/min), a splitless injection (injection volume of 2 µL) was used, and a temperature gradient of 5°C/min from 40°C (3-min hold) to 240°C was applied. For determination of the enantiomeric composition, a heptakis(2,3-di-O-methyl-6-O-t-butyldimethyl-silyl)-β-cyclodextrin column was used for all of the monoterpenes and the sesquiterpenes (E)-β-caryophyllene, cuparene, and β-bisabolene, whereas a heptakis(2,6-di-O-methyl-3-O-pentyl)-β-cyclodextrin column was used for α-barbatene, α-copaene, thujopsene, and β-chamigrene. The temperature program for the monoterpenes was as follows: 40°C (3-min hold) followed by a ramp of 5°C/min to 125°C. The program for the sesquiterpenes was as follows: 40°C (1-min hold) with a ramp of 30°C/min to 90°C (30-min hold). Mass spectrometry was performed with a transfer line temperature of 230°C, source temperature of 230°C, quadrupole temperature of 150°C, ionization potential of 70 eV, and scan range of 50 to 400 atomic mass units.

The identities of all compounds were determined by comparison of retention times and mass spectra with those of authentic standards and with mass spectra in the National Institute of Standards and Technology and Wiley libraries (Agilent Technologies, Palo Alto, CA). The racemic standards of α-barbatene, (E)-β-caryophyllene, and cuparene were provided by Wilfried A. König (University of Hamburg, Germany). Chirality was determined by direct comparison with standards of both enantiomers for all compounds except β-chamigrene and thujopsene, for which only the (−)-enantiomers were available. For quantification, representative single-ion peaks of each compound were integrated and compared with the equivalent response of the internal standard (single-ion method).

Determination of Gene Expression by Reverse Transcripase–Mediated PCR

Total RNA was isolated with an RNeasy Plant Mini Kit (Qiagen, Valencia, CA), and DNA contamination was removed with DNase (Qiagen)
treatment for 15 min at room temperature. Five micrograms of RNA was reverse transcribed into cDNA in a 33-µL reaction with poly(dT) priming using a First-Strand cDNA Synthesis Kit (Amersham Pharmacia Biotech). The PCR volume was 25 µL containing 100 ng of each primer, 2 mM of each deoxynucleoside triphosphate, 1 µL of cDNA, and 0.75 units of Taq DNA polymerase (Fisher, Pittsburgh, PA). The primers used in the reverse transcriptase–mediated (RT) PCR experiments are shown in Table 1.

A PTC-100 Programmable Thermal Controller (MJ Research, Watertown, MA) was used with an initial denaturation step of 96°C for 1 min, followed by 30 cycles of 94°C for 30 s, 54°C for 30 s, 72°C for 60 s, and a final elongation step of 72°C for 10 min. RT-PCR analysis was performed in duplicate using RNA from two different sets of plants. Results of the two replicates were similar, and only one replicate is depicted in Figure 2. The second RT-PCR was performed using the same reagents as those used for the first PCR except that the RNA was isolated from different plants.

**RNA Gel Blot Analysis**

Total RNA (16 µg per sample) was separated on a 1% agarose gel containing 17% formamide and transferred onto a Hybond-XL nylon membrane (Amersham Pharmacia Biotech). DNA probes were generated by PCR and labeled with 32P-dCTP using a Rediprime II Kit (Amersham Pharmacia Biotech). Standard procedures were followed for prehybridization, hybridization, and washing.

cDNA Cloning by RT-PCR and Protein Expression in *Escherichia coli*

Full-length cDNAs were obtained by RT-PCR as described above. The primers used were 5'-AATGGCTACTCTTTGTATAGGTT-3' (or 5'-ATCCATATGGCGACCTCGGGCAACTCTCAAC-3', containing a Ndel site) and 5'-ATCGGATCTATCTAATGGGATTGGGTC-3' (containing a BamHI site) for At1g61680, 5'-AATGGCCTTTAATAGCTAGCTAC-CAAATAAG-3' (or 5'-AATGATGATGTCATTCAAAGT-3') and 5'-ATTACATTAGAAGTCGGATATG-3' for At5g23960. The second forward primers (listed in parentheses) for At3g25810 and At1g61680 were chosen to eliminate the TG-3 region coding for the transit peptide. The resulting fragments were cloned into the vector pCRT7/CT-TOPO (Invitrogen, Carlsbad, CA), and the truncated cDNA of AtTPS1 was subcloned further into the BamHI and Ndel sites of pET11a (Amersham Pharmacia Biotech). Standard procedures were followed for prehybridization, hybridization, and washing.

**Terpene Synthase Enzyme Assays**

Cells from a 100-mL induced culture were harvested at 4°C, washed with 20 mL of washing buffer (20 mM Tris-HCl, pH 7.0, and 50 mM KCl), and resuspended in 5 mL of extraction buffer (50 mM 3-(N-morpholino)-2-hydroxypropanesulfonic acid, pH 7.0, 10% [v/v] glycerol, 5 mM MgCl2, 5 mM DTT, 5 mM sodium ascorbate, and 0.5 mM phenylmethylsulfonyl fluoride). Cells were disrupted by sonication (2 × 4-min treatment at 50% power with a Bandelin model UW2070 instrument [Berlin, Germany]) and then centrifuged at 20,000 g at 4°C for 30 min. The supernatant was desalted with a Bio-Rad Econo column, and the resulting 4-mL eluate (in assay buffer, containing 10 mM 3-(N-morpholino)-2-hydroxypropanesulfonic acid, pH 7.0, 10% [v/v] glycerol, and 1 mM DTT) was used for the enzyme assay. The volume of each assay was 1 mL, containing 960 µL of enzyme extract, 20 mM MgCl2, 0.2 mM MnCl2, 0.2 mM NaWO4, 0.1 mM NaF, and 60 µM geranyl diphosphate or farnesyl diphosphate (Echelon Research Laboratories, Salt Lake City, UT). The assay was performed in an 8-mL DuPont autosampler vial with a white solid-top polypropylene cap (Alltech, Deerfield, IL), and a solid-phase microextraction PDMS-100 (polydimethylsiloxane) fiber (Supelco, Bellefonte, PA) was inserted into the tube to collect volatiles. The assay was incubated at 30°C for 1 h and then at 42°C for 15 min. After incubation, the solid-phase microextraction fiber was injected into a gas chromatography–mass spectrometry system for analysis. Alternatively, enzyme products were extracted three times with 1 mL of pentane, and the organic extract was concentrated to ~100 µL before gas chromatography–mass spectrometry analysis. Controls included assays with crude extracts of induced *E. coli* (same strain) carrying the expression vector without any insert.

Microscale assays to compare the relative activity of different substrates were performed in a final volume of 500 µL with 450 µL of crude extract and 10 or 30 µM 3-H-geranyl diphosphate (2 or 9 MBq/µmol) or 3-H-farnesyl diphosphate (9 MBq/µmol) Buffer, salt, and incubation conditions were as described above, but 0.4 mM NaWO4 and 0.2 mM NaF were used in assays with At1g61680 enzyme. After a 10-min incubation, the reaction products were extracted two times with 1 mL of hexane. Total radioactivity of the reaction products was determined by scintillation counting. In assays with the At3g25810 enzyme, the hexane extract was treated with silica gel to remove phosphaate products before counting.

**Construction of the At3g25810 Promoter–β-Glucuronidase Reporter Gene Fusion Construct, Arabidopsis Transformation, and histochemical localization of β-Glucuronidase activity**

The At3g25810 promoter region (1.4 kb) was isolated via PCR from Arabidopsis genomic DNA using the primers 5'-CTCCGATTATGG-AATTTAGGGCGATGTCACACC-3' and 5'-GAACTTGGCAATGCATCTAC-3', which contain an EcoRI and an NcoI site, respectively. The resulting PCR product was cloned into the pCRT7/CT-TOPO vector and sequenced. Next, the promoter region of At3g25810 was cut out of the pCRT7/CT-TOPO (Invitrogen, Carlsbad, CA), and the truncated cDNA of AtTPS1 was subcloned further into the BamHI and Ndel sites of pET11a. An *E. coli* BL21 Codon Plus strain, transformed with the appropriate expression construct, was used for protein expression. Induction was performed at 18°C overnight with 1 mM isopropyl-1-thio-β-D-galactopyranoside.

The At3g25810 promoter region was inserted into the binary vector pCAMBIA1303 ([Hajdukiewicz et al., 1994], replacing the 3SS promoter of cauliflower mosaic virus so that the At3g25810 promoter directs the expression of the uidA (β-glucuronidase [GUS] gene). The construct was introduced into *Agrobacterium tumefaciens* strain GV3101, which was used to transform Arabidopsis by floral vacuum infiltration (Bechtold et al., 1993). Transgenic lines transformed with the construct were selected using hygromycin and confirmed by PCR. Enzymatic assays with 5-bromo-4-chloro-3-indolyl-β-D-glucuronide were performed to determine the localization of the enzyme activity of the GUS enzyme (Jefferson et al., 1987). Tissue samples were incubated at 37°C in GUS staining buffer (50 mM sodium phosphate buffer, pH 7.0, 0.1% Triton X-100, and 0.5 mg/mL 5-bromo-4-chloro-3-indolyl-β-D-glucuronide) overnight. After detection of the blue color, chloroform was extracted with 70% ethanol for 24 h. Upon request, all novel materials described in this article will be made available in a timely manner for noncommercial research purposes.
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