

REVIEW

View Article Online
View Journal | View Issue



The volatile chemistry of orchid pollination

Cite this: *Nat. Prod. Rep.*, 2023, 40, 819

James Perkins,^a Tobias Hayashi,^a Rod Peakall,^{ab} Gavin R. Flematti^b and Björn Bohman^c  ^{*abc}

Covering: up to September 2022

Orchids are renowned not only for their diversity of floral forms, but also for their many and often highly specialised pollination strategies. Volatile semiochemicals play a crucial role in the attraction of a wide variety of insect pollinators of orchids. The compounds produced by orchid flowers are as diverse as the pollinators they attract, and here we summarise some of the chemical diversity found across orchid taxa and pollination strategies. We focus on compounds that have been experimentally demonstrated to underpin pollinator attraction. We also highlight the structural elucidation and synthesis of a select subset of important orchid pollinator attractants, and discuss the ecological significance of the discoveries, the gaps in our current knowledge of orchid pollination chemistry, and some opportunities for future research in this field.

Received 1st September 2022

DOI: 10.1039/d2np00060a

rsc.li/npr

1	Background
2	Fatty acid derivatives
2.1	Alkanes and alkenes
2.2	Alcohols
2.3	Aldehydes
2.4	Carboxylic acids
2.5	Esters
2.6	Lactones
2.7	Tetrahydrofuran derivatives
2.8	Chiloglottones
3	Isoprenoids
3.1	Monoterpenes
3.1.1	Acyclic monoterpenes
3.1.2	Cyclic monoterpenes
3.2	Sesquiterpenes
3.2.1	Acyclic sesquiterpenes
3.2.2	Cyclic sesquiterpenes
3.3	Other terpenoids
4	Benzenoids
4.1	C ₆ -C ₀
4.2	C ₆ -C ₁
4.3	C ₆ -C ₂
4.4	C ₆ -C ₃
4.5	C ₆ -C ₄
5	Other compounds
5.1	Nitrogenous compounds

5.2	Sulfurous compounds
6	Methods
6.1	Isolation
6.2	Identification and data analysis
6.3	Synthesis
7	Final remarks
7.1	The importance of behavioural bioassays
7.2	Considering absolute configuration
7.3	Lessons from sexual deception
8	Conclusion
9	Conflicts of interest
10	Acknowledgements
11	References

1 Background

The orchid family (Orchidaceae) is one of the most species-rich and diverse of all plant families, with over 29 000 species.¹ Orchids are also well known for the extraordinary diversity of their interactions with animal pollinators, making use of unusual and often highly specialized pollination strategies that go well beyond the typical exchange of nectar or pollen as food to pollinators for the transfer of gametes. Some orchids provide unusual rewards, such as perfumes to male euglossine bee pollinators, which collect them to attract mates.² Many other orchids employ deceptive pollination strategies. For example, many 'food deceptive' orchids mimic the visual and olfactory signals of rewarding flowers, but do not provide a tangible reward. Others have evolved even more complex means to attract pollinators, such as mimicking sites for insects to lay

^aResearch School of Biology, The Australian National University, Australia

^bSchool of Molecular Sciences, The University of Western Australia, Australia

^cDepartment of Plant Protection Biology, Swedish University of Agricultural Sciences, Sweden. E-mail: bjorn.bohman@slu.se



eggs,³ or even the mimicry of insects themselves. Some insect-mimicking species emit alarm pheromones of aphids⁴ or honey bees⁵ to attract predators of those species as pollinators, and others emit aggregation pheromones to attract large swarms of potential pollinators.⁶ By far the most common insect-mimicry pollination strategy amongst orchids is 'sexual deception', whereby orchids mimic the sex pheromones and appearance of specific female insects to sexually lure conspecific males as pollinators. In these cases, pollination often occurs during attempted copulation with the flower, highlighting the extreme effectiveness of this bizarre trickery.^{7,8}

Whether it is by honest signalling of reward or by deception, orchid pollinator attraction is often dependent on volatile or semi-volatile compounds. Indeed, orchids are the most well-represented family in studies of plant volatiles, and are the source of many unusual natural products.⁹ Given the great diversity of orchids and their numerous chemically mediated pollinator interactions, orchids likely offer a treasure trove of natural products remaining to be discovered.

While much progress has been made towards cataloguing the diversity of known floral volatiles within orchids,^{9–12} relatively few studies have confirmed the ecological activity of the compounds, and fewer still have elucidated the chemical structures of potentially new-to-science compounds. When done systematically, such studies provide valuable insights into the diversity of compounds used by both plants and insects for communication, and aid our understanding of the evolution of plant-animal interactions.

In this review, we highlight the diversity of volatile and semi-volatile natural products from orchid flowers that may be relevant to pollination. We broadly follow the groupings established in prior reviews of plant volatiles,^{9,12,13} which class compounds based on their likely biosynthetic origin into three main groups – fatty acid derivatives, isoprenoids, and benzenoids. We prioritise compounds for which credible evidence for biological activity has been established. We regard experimental confirmation of pollinator attraction in field bioassays with reference compounds as the most compelling evidence for the biological activity of a compound, while we recognise that experiments with wind tunnels and/or olfactometers in the laboratory may also provide such evidence. Many studies report findings on the electrophysiological activity of the compounds extracted from orchids to one or more species of pollinator, based on coupled GC-Electroantennography Detection (GC-EAD), or less commonly *via* electroantennography (EAG) or GC-Single Sensillum Recordings (GC-SSR).¹⁴ While electrophysiological activity of a compound does not always translate to attractiveness in field experiments (see final remarks, 7.1), antennal electrophysiology is often critical for narrowing down the number of compounds of interest for further experiments in the field. Therefore, we highlight electrophysiological results where possible. Finally, we briefly explore the methods used to elucidate and synthetically prepare some orchid pollinator attractants, before discussing what we see as some exciting opportunities in this expansive field in the coming years.

2 Fatty acid derivatives

Fatty acid derivatives are often shorter in chain length than their precursor and may or may not retain a carbonyl moiety. They are formed by oxidative cleavage and/or decarboxylation of the fatty acid precursor.¹⁵ The biosynthesis of fatty acids from condensation of acetic acid (C2) units is well known and is catalysed by the enzyme fatty acid synthase. In animals and fungi, this is a single large multifunctional protein, however in plants and bacteria the individual steps are catalysed by an assembly of discrete enzymes, allowing for much more diversity in structure.¹⁶ These differences in biosynthesis make the plant and bacterial enzymes attractive targets for herbicides and antibacterials, respectively.^{17,18}

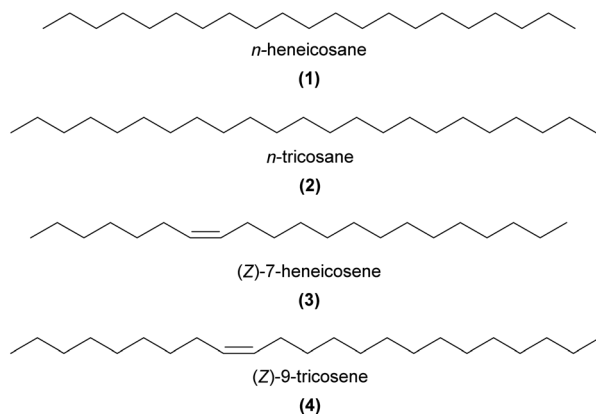
2.1 Alkanes and alkenes

Alkanes and alkenes are biosynthesised from fatty acid intermediates through reduction to fatty aldehydes, followed by decarbonylation *via* aldehyde decarbonylase enzymes that have been found in both plants and insects.^{19,20} This pathway results in the production of mostly odd-numbered hydrocarbons from even-numbered fatty acid precursors. Unsaturation to form alkenes is achieved earlier in the biosynthetic sequence by desaturase enzymes acting on fatty acid derivatives, yielding predominantly (*Z*)-configured compounds in both plants and insects.²¹

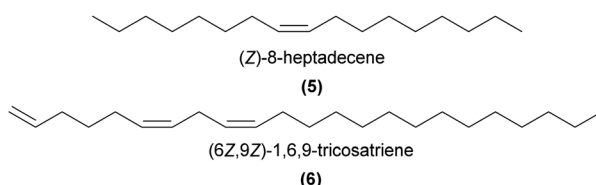
Unbranched saturated hydrocarbons (*n*-alkanes), and unsaturated hydrocarbons (alkenes), are important constituents of the cuticular waxy layer of plants, which has a critical primary function of protection against water loss, UV light, pathogens, and pests.^{22,23} Cuticular alkanes and alkenes are also widely found in insects, where they function as protection against desiccation and also as signalling molecules.^{24,25} Specific alkanes and alkenes or combinations are widely known as sex pheromones and mate recognition cues across a range of insect groups,²⁶ including the Hymenopteran pollinators of some orchids. It is perhaps unsurprising, then, that alkanes and alkenes have been demonstrated to play key roles in the pollination of sexually deceptive orchids.

Using GC-EAD in combination with field bioassays, Schiestl *et al.*^{27,28} were the first to convincingly demonstrate female insect sex pheromone mimicry in sexually deceptive *Ophrys* orchids. The initial breakthrough was made in *O. sphegodes*, pollinated by sexually attracted male *Andrena nigroaenea* bees. A blend of 14 hydrocarbons (C₁₉–C₂₉, *e.g.* 1 and 2) that had been found to be both EAD-active to the male bee pollinator and shared between the orchid labellum and female bees, were also sexually attractive in field bioassays.²⁸ Later, in *O. exaltata*, a specific blend of EAD-active alkenes and *n*-alkanes was found to be attractive to male *Colletes cunicularius* bee pollinators, but alkenes were also attractive alone (*e.g.* 3 and 4).²⁹ In both orchid systems, alkene double bond configuration is pivotal for the specific bee pollinator attraction ((*Z*)-9, 11, and 12 in *O. sphegodes*, and (*Z*)-7 and 9 in *O. exaltata* but with the (*Z*)-7-alkenes being the most attractive).^{27–29}





Alkanes and alkenes are now widely implicated as key components of the sex pheromone mimicry across bee-pollinated European *Ophrys* (see detailed reviews in ref. 7 and 30–32). In particular, C_{21} – C_{33} *n*-alkanes and specific alkenes, always in (Z) configuration, have been demonstrated to be EAD-active to the bee pollinators of multiple *Ophrys* species.^{27–29,33–37} Shorter chain hydrocarbons, (Z)-8-heptadecene (5) and *n*-pentadecane have been shown to be EAD-active in the wasp-pollinated *O. insectifera*, although their role in pollinator attraction remains unknown.³⁸ Alkadienes, typically C_{25} – C_{33} and all (Z), have also been detected in some *Ophrys* spp.,^{31,39,40} including some that are EAD-active,²⁹ although their role in pollinator attraction is also yet to be demonstrated.



Perhaps one of the most unusual examples of hydrocarbon sex pheromone mimicry is found in the Australian orchid *Pterostylis orbiculata*, which attracts its male *Mycomya* sp. fungus gnat pollinator with a tri-unsaturated hydrocarbon.⁴¹ Laboratory bioassays with captive male gnats revealed that males responded to a GC-derived fraction containing two unsaturated hydrocarbons subsequently identified as (6Z,9Z)-1,6,9-tricosatriene (6) and (6Z,9Z)-6,9-tricosadiene. Field bioassays confirmed that male gnats were sexually attracted to a synthetic blend of five compounds in common between orchid labella and female gnats, comprising the two unsaturated hydrocarbons and three C_{21} – C_{25} *n*-alkanes. The triene alone was attractive, although gnats displayed weaker sexual behaviour than to the blend, and the blend without the triene was unattractive. In related species, (3Z,6Z,9Z)-3,6,9-tricosatriene (*P. concava*) and an unidentified tricosatriene and tricosatetraene (*P. vittata*) were detected, suggesting polyenes may be key to pollinator attraction in other *Pterostylis* species.⁴¹ Polyenes are not as common in nature as monoenes or dienes, but are well known as type II moth sex pheromones⁴² and have been occasionally reported in other insects⁴³ and plants.⁴²

Alkenes are also components of the pollinator-attracting blend in the South American orchid *Telipogon peruvianus*, pollinated by male *Eudejeania* tachinid flies. Only pre-

copulatory behaviour is observed at the orchid flower in this system,⁴⁴ and a more complex mechanism than sex pheromone mimicry alone may be involved.⁴⁵ Nonetheless, the EAD-active blend is dominated by (Z)-9-tricosene (4), along with other C_{20} – C_{25} saturated and monounsaturated hydrocarbons and the aldehyde tetradecanal (14) in similar ratios to the female fly.⁴⁵ Field bioassays with synthetic blends in both flower and female fly ratios elicited pre-copulatory behaviour from male flies, but attempted copulation was not observed.⁴⁵

Alkenes and *n*-alkanes (C_{23} – C_{31} , alkenes in (Z)-9 configuration) also dominate the solvent extracts of another sexually deceptive South American orchid *Maxillaria lineolata* (previously *Mormolyca ringens*).⁴⁶ Compared to fresh flowers, lower amounts of C_{25} and C_{27} (Z)-9-alkenes and (9Z)-9,17-octadecadienal were present in pollinated flowers that were unattractive to male (drone) bee pollinators. Solvent extracts elicited electrophysiological activity from antennae of male bees, although it is unknown which specific compounds are EAD-active.⁴⁶

Alkenes have also been detected, but not yet shown to be biologically active, in a range of other non-sexually deceptive orchids (e.g. *Maxillaria*, *Anacamptis*, *Gymnadenia*, *Serapias*, *Dactylorhiza*, *Himantoglossum*, *Platanthera*, *Neotinea*^{47–49}) suggesting alkene production in orchids is not unusual.

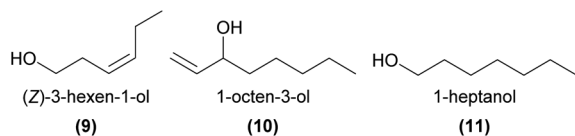
2.2 Alcohols

Several unbranched primary and secondary alcohols, and short chain unsaturated alcohols have been found across a variety of orchids, with evidence that some are involved in pollination. In the compelling case of *Dendrobium sinense*, which is pollinated by the predatory hornet *Vespa bicolor*, a synthetic mixture of EAD-active 1-octadecanol (7), 1-eicosanol, and (Z)-11-eicosen-1-ol (8), as well as 8 on its own, were attractive to the hornets in a flight cage and in Y-tube choice tests.⁵ This study was the first to report 8 as a plant volatile, and further noted that this molecule is a major component of the Asian and European honey bee alarm pheromones. These findings, coupled with the unusual observations of hornets 'pouncing' on the flowers and thus picking up pollinia, led the authors to conclude that the rewardless *D. sinense* mimics the honeybee alarm pheromone to deceptively attract its hornet pollinator.



In another case of orchid pollination by predatory wasps, a synthetic mixture of EAD-active (Z)-3-hexen-1-ol (9) with two other green leaf volatiles, the esters hexyl acetate and (Z)-3-hexenyl acetate, was strongly attractive in Y-tube choice tests to the *Vespa* pollinators of the nectar rewarding *Epipactis helleborine*.⁵⁰ It is thought that the orchid emits these green leaf volatiles, usually associated with wounding by herbivorous insects, as a deceptive signal to mimic the presence of herbivorous prey species and attract its predatory wasp pollinator.





In the mushroom-scented *Dracula lafleurii*, the mushroom alcohol, 1-octen-3-ol (10), is emitted by the attractive labellum. This compound is predicted to be a key attractant of the small fly pollinators – although single compound bioassays have yet to be conducted.⁵¹

In some food rewarding and food deceptive systems, the short chain 1-heptanol (11), 1-octanol, and 1-decanol were EAD-active,^{52,53} while the longer chain 1-hexadecanol, 1-octadecanol (7), and their unsaturated analogs were EAD-active components of attractive orchid extracts in two sexually deceptive *Ophrys* species.³⁴ However, the precise roles of these primary alcohols in pollination remain to be more fully tested.

2.3 Aldehydes

Across the plant kingdom, aliphatic aldehydes are ubiquitous floral and vegetative volatiles. It is not surprising that these compounds have been reported across a wide range of orchid species (for example ref. 54–59), yet a decisive role for aldehydes in orchid pollination has rarely been confirmed. Lahondère *et al.*⁵³ showed that the short chain aldehydes heptanal (12), octanal, and nonanal (13) were emitted by *Platanthera obtusata* and that 13 was critical to the attraction of its mosquito pollinators.⁵³ Mosquito pollination is unusual in this orchid genus, with other species predominantly pollinated by moths, where terpenes often play a key role in pollinator attraction (see Section 3 for further details).



Aldehydes (C_7 – C_{26}) have been found to be present in many species of sexually deceptive *Ophrys* species, and EAD-active in several.^{28,34–37,60–62} Cuervo *et al.*⁶³ found 20 EAD-active compounds in common between orchids and associated female *Eucera kullenbergi* bees, including seven aldehydes (C_7 – C_{16}), as well as alcohols, fatty acids, and hydrocarbons.⁶³ Synthetic blends of all EAD-active compounds in females and orchids, as well as polar compounds only, aldehydes only, and alkanes only, elicited more sexual attraction than the control in field bioassays, but were not as attractive as female bee and orchid extracts. The presence of aldehydes, in both females and orchids, their EAD-activity, and some sexual attraction indicate a possible role as sex pheromone components, although their importance in the overall blend remains unclear.

Sexually deceptive *Ophrys sphegodes* exhibit within-inflorescence variation in the relative amount of aldehydes and esters produced, with the uppermost flowers producing less

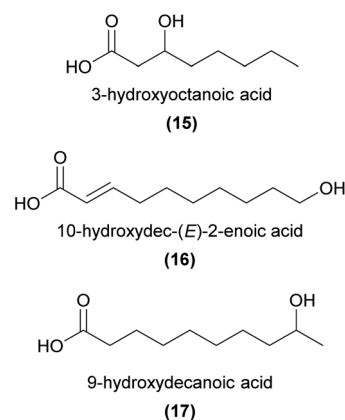
of these compounds than the second uppermost flower and consequently attract fewer *Andrena nigroaenea* pollinators.³⁹ Ayasse *et al.*,³⁹ however, showed that this reduction in visitation could be avoided by applying aldehydes and esters to the uppermost flowers, suggesting that bees may learn the scent of individual flowers in part due to aldehydes and esters.

In *Telipogon peruvianus*, tetradecanal (14) is present in both orchids and female *Eudejeania* sp. aff. *browni* tachinid flies, and is the only non-hydrocarbon EAD-active to males.⁴⁵ The blend of hydrocarbons and tetradecanal is sexually attractive in field bioassays, although the role of tetradecanal alone has not been reported.

2.4 Carboxylic acids

Carboxylic acids, with chain lengths between C_6 to C_{18} , are relatively common floral volatiles,⁹ and have frequently been found in orchids.^{57,64–66} Despite their prevalence, carboxylic acids have rarely been shown to function in orchid pollinator attraction, with two notable exceptions.

In a fascinating pollination system, *Cymbidium floribundum* attracts swarms of the Japanese honeybee, *Apis cerana japonica*, by emitting bee aggregation pheromones.⁶ Sugahara *et al.*⁶ identified 3-hydroxyoctanoic acid (15) and 10-hydroxydec-(*E*)-2-enoic acid (16) as attractive components from the flowers. They also noted that 15 and similar unsaturated acids are known mandibular gland components of *A. cerana*,⁶⁷ and that 16 is a major component of royal jelly in *A. mellifera*.⁶⁸ Interestingly, neither compound was attractive alone, but instead a specific blend of both compounds was required to attract the bees.



Ophrys speculum is a sexually deceptive, scoliid wasp-pollinated species in an otherwise mostly bee-pollinated genus. Eight EAD-active compounds are shared between the flowers and associated female wasps.^{31,60} These comprised saturated (ω -1)-hydroxy acids (e.g. 17) and (ω -1)-oxo acids, aldehydes and ethyl esters. In combination, at blend ratios comparable to those found in the orchid and wasp, these compounds elicited high rates of attempted copulation in field bioassays.⁶⁰ Notably, the enantiomeric composition of 7-hydroxyoctanoic acid and 9-hydroxydecanoic acid (17) was similar in orchids and wasps ($R:S = 6:4$), and this enantiomeric ratio was essential for strong sexual attraction in the field bioassays.



2.5 Esters

Esters are reported as floral volatiles across a range of orchids with different pollination strategies, although few studies have conclusively linked these molecules to orchid pollinator attraction. One important exception is the non-photosynthetic *Gastrodia similis*, which attracts its specific drosophilid fly pollinator, *Scaptodrosophila bangi*, by emitting ethyl acetate (**18**), ethyl isobutyrate (**19**), and methyl isobutyrate (**20**).⁶⁹ The flies were strongly attracted to a synthetic mix of the three esters, and each compound on its own was also partially attractive.



Another interesting case of specific chemical attraction of pollinators by an ester is found in the Japanese orchid *Luisia teres*, which sexually attracts male *Protaetia pryri pryri* beetles.^{70,71} Wakamura *et al.*⁷¹ showed that male beetles were exclusively attracted to (*R*)-2,3-dihydroxypropyl isovalerate (**21**), but not to the (*S*)-enantiomer or the racemate. The (*R*)-enantiomer was also found in virgin female beetles but did not occur in males, strongly suggesting that this compound is related to sex signalling in this species.

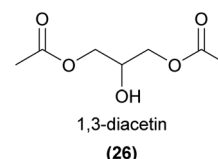


Several esters with relatively long parent alcohol and acid chains are EAD-active in sexually deceptive *Ophrys*. Examples include 2-nonyl dodecanoate (**22**) and dodecyl tetradecanoate in *O. sphegodes*,^{28,39} octyl and nonyl pentadecanoate (**23**) in *O. aymoninii*,³³ and 2-nonyl hexadecanoate reported in *O. iricolor*,⁶² *O. lupercalis*,³⁶ and *O. fabrella*.³⁷



Shorter ethyl- and methyl esters and acetates are reported in several food rewarding and food deceptive species, but their role in pollination also remains unresolved. Examples include C₁₀, C₁₂, and C₁₄ EAD-active alkyl and alkenyl acetates in the food rewarding *Gymnadenia conopsea*,⁷² a series of C₆ to C₁₆ alkyl acetates (*e.g.* **24**) in *G. conopsea*⁷³ and the food deceptive *Cypripedium calceolus*,⁵² methyl octanoate (**25**) and methyl decanoate in the food deceptive *Orchis mascula*,⁷⁴ a series of C₁₀ to C₁₆ methyl and ethyl esters in the food rewarding *Cremastra appendiculata*,⁷⁵ and methyl hexadecanoate in the rewarding *Platanthera bifolia*.^{76,77}

Oil rewarding plants partake in highly specialized interactions with 'oil bees', by secreting oils comprised of various acylglycerols and free fatty acids as a reward for pollination. The bees collect and use these oils for larval provisioning, brood cell water-proofing, or food for adults.⁷⁸ Due to the low volatility of floral oils, we do not focus on these here, however a related volatile acetylated glycerol has been identified as a common compound associated with oil rewarding plants that may function as a signal of oil reward. 1,3-Diacetin (**26**) and both enantiomers of 1,2-diacetin have been detected from many oil rewarding plants, including several orchid species.^{79,80} Interestingly, these compounds were shown to be electrophysiologically and behaviourally active in oil bees, but not detectable to co-occurring non-oil bee species, suggesting that these compounds might represent a 'private communication channel' between oil flowers and oil bees.⁷⁹



2.6 Lactones

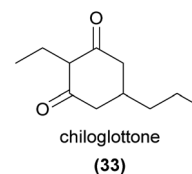
In a remarkable demonstration of the potential of chemical ecology, Cohen *et al.*⁸¹ were able to successfully identify the sexual attractant from a single flower of the exceedingly rare South African orchid *Disa forficaria*. Using GC-EAD and HRMS, the authors identified a single active compound, (16*S*,9*Z*)-16-ethyl hexadec-9-enolide (termed disalactone, **27**), that was highly attractive and elicited prolonged copulatory behaviour from male *Chorothyse hessei* longhorn beetle pollinators in field bioassays. Further bioassays with individual isomers showed that the (16*R*,9*Z*)-stereoisomer was only weakly attractive, and (*rac*,9*E*)-16-ethyl hexadec-9-enolide and (*rac*,8*Z*)-16-ethyl hexadec-8-enolide were also weakly attractive.⁸¹ While this is the only macrolide presently known to play a role in pollination, several macrolides are known insect pheromones,⁸² suggesting similar compounds may be more broadly exploited. We further note that (ω -1)-hydroxy acids (*e.g.* **17**) such as those identified by Ayasse *et al.*⁶⁰ may readily cyclise to form macrolides.





In the rare Australian orchid *Drakaea micrantha*, a blend of three compounds, likely from two separate biosynthetic pathways, elicit strong sexual behaviour from male *Zebleboria* sp. thynnine wasp pollinators. Similar to other *Drakaea* spp., two hydroxymethylpyrazines are part of the blend, however the pyrazines are barely attractive on their own and a β -hydroxylactone (4-hydroxy-3-methyl-6S-(pentan-2S-yl)-5,6-dihydro-2H-pyran-2-one, termed drakolide (28)), is required to elicit strong sexual behaviour.⁸³ Further investigations showed that both the naturally occurring stereoisomer of 28 and a blend of four stereoisomers prepared from racemic reagents elicited strong sexual behaviour from male wasps in field bioassays when presented in combination with the two hydroxymethylpyrazines.⁸⁴ Field bioassays with six structural analogues of drakolide featuring substituents at positions 3 and 6 revealed reduced sexual behaviour and varying levels of attractiveness.⁸⁴

Several other lactones, such as the 'Aerangis lactone' ((5S,6S)-6-pentyl-5-methyltetrahydropyran-2-one, 29)^{10,85} and the jasmine lactone ((Z)-6-(pent-2-en-1-yl)tetrahydro-2H-pyran-2-one, 30),⁸⁶ have been identified as orchid volatiles. Noted for their pleasant scents to humans, these may be important for pollination, but their biological activities have not yet been reported.



2.7 Tetrahydrofuran derivatives

Using bioassay-guided fractionation in combination with field bioassays, Bohman *et al.*⁸⁷ demonstrated that (S)-2-(tetrahydrofuran-2-yl)acetic acid (31) produced by the sexually deceptive Australian orchid *Cryptostylis ovata*, is attractive to male *Lissopimpla excelsa* wasp pollinators.⁸⁷ The ester derivatives methyl (S)-2-(tetrahydrofuran-2-yl)acetate (32) and ethyl (S)-2-(tetrahydrofuran-2-yl)acetate, present in small amounts in the orchid, were similarly attractive to male wasps. However, few landings and no attempted copulations were observed in response to either orchid solvent extracts, 31, the ester derivatives, or a combination of all three, indicating that additional compounds or other non-chemical cues are required to induce the strong copulatory behaviour observed at the flower.

2.8 Chiloglottones

In a first step towards understanding the chemical basis of sexual pheromone mimicry in the many sexually deceptive orchids of Australia, Schiestl *et al.*⁸⁸ demonstrated that the sole EAD-active compound in the male thynnine wasp pollinated *Chiloglottis trapeziformis* orchid was 2-ethyl-5-propylcyclohexan-1,3-dione, termed chiloglottone (33),⁸⁸ which was also confirmed to be the female sex pheromone of the pollinator species. In field bioassays, synthetic 33 was equally as attractive as orchid extracts, whole flowers, and female wasps. Subsequent investigations revealed the presence of five additional related compounds in other species of *Chiloglottis*, with compounds differing in the alkyl or alkenyl chains at positions 2 and 5.^{89,90} Each of eleven *Chiloglottis* species investigated was found to produce one or two chiloglottones, with some sharing of chiloglottones among allopatric taxa and occasional sharing of chiloglottones among sympatric taxa.^{90,91} Chiloglottones have also been detected in the allied *Arthrochilus* and *Paracaleana*, and are EAD-active in the latter genus, although their role in pollination is yet to be confirmed.⁹⁰

3 Isoprenoids

Isoprenoids are a common group of hydrocarbon compounds based on condensations of five-carbon (isoprene) building blocks derived from dimethylallyl diphosphate (DMAPP) and isopentenyl diphosphate (IPP).⁹² Isoprenoids represent the largest class of natural products⁹³ and can be classified by the number of isoprene units as monoterpenes (C₁₀), sesquiterpenes (C₁₅), diterpenes (C₂₀), triterpenes (C₃₀), *etc.* Terpenoids are terpenes that have been modified in some way, usually through oxidation. They constitute a wide and structurally diverse group of specialised metabolites produced by plants and have many roles in defence and pollinator attraction.¹⁵ Below we include terpenoids within the respective group of terpenes.

3.1 Monoterpenes

Monoterpenes are among the most common of all floral volatiles, being found across the plant kingdom and commonly



reported in orchid floral volatile profiles. Here we focus on compounds shown *via* bioassays to be directly involved in orchid pollination, but we note that given their prevalence, monoterpenes likely play roles in pollination of many other orchid species.

Of particular note, while limonene (34), (*E*)-ocimene (35), and myrcene (36) are the three most commonly occurring floral volatiles across studied plant families⁹ and are also common in orchid flowers, they have rarely been confirmed to be involved in pollination. One exception is in the perfume rewarding pollination systems, where monoterpenes frequently appear in detailed lists of compounds that are attractive to perfume-collecting male euglossine bee pollinators.^{94,95} We discuss some of these compounds here, but refer interested readers to these reviews and references therein for a comprehensive account of the compounds potentially⁹⁶ or known to be^{94,95} involved.



3.1.1 Acyclic monoterpenes. Linalool (37), geraniol (38) and nerol (39) are common orchid floral volatiles for which EAD activity has been found in multiple cases.^{52,53,73–75,97} Linalool (37) has also been confirmed to be active in field bioassays to the euglossine bee pollinators of several perfume rewarding orchids.^{94,98–100} Terpenoids 38 and 39 have also been confirmed as behaviourally active to the male *Eulama* euglossine bee pollinator of the orchid *Sarcoglottis acaulis* in field studies. Interestingly, in this case the typical perfume collecting behaviour was not observed at either the orchid flower or the chemical baits, and instead pollinators appeared to forage for nectar, despite the specific attraction of males.⁹⁷



Linalool (37) is also one of very few monoterpenoids to be implicated in pollination by sexual deception. Borg-Karlson *et al.*¹⁰¹ first showed that the (*S*)-enantiomer was attractive to male *Colletes cunicularius* bees, the pollinator of *Ophrys exaltata*, whereas the racemate was less attractive. Mant *et al.*²⁹ also found racemic linalool rarely induced attempted copulation. However, in blends with 12 EAD-active hydrocarbons (odd chain lengths of C₂₁–C₃₁ alkanes, alkenes, and one alkadiene), contacts increased,

indicating linalool acts as a long-range attractant, while the hydrocarbons are essential to induce sexual behaviour.

To our knowledge, the only other monoterpene shown to play a key role in sexual deception is (*S*)- β -citronellol (40).¹⁰² In *Caladenia plicata*, this compound in an optimal 1 : 4 blend with 2-hydroxy-6-methylacetophenone, secures pollination by male *Zebleboria* sp. thynnine wasps. Field bioassays revealed neither compound was active on its own,¹⁰² while further field tests confirmed replacements with (*R*)- β -citronellol and alternative regioisomers of the acetophenone derivative were barely attractive.¹⁰³



Ipsdienol (41) has also been found from several perfume rewarding orchids, and is known to play a key role in specific attraction of perfume-collecting bees.^{99,104} Schorkopf *et al.*¹⁰⁵ further showed that the (*R*)-enantiomer was attractive, but the (*S*)-enantiomer was not.

3.1.2 Cyclic monoterpenes. Some early comprehensive work on the chemical basis of pollinator attraction was conducted on the food rewarding *Platanthera bifolia*, which is pollinated by the silver Y moth, *Autographa gamma*.^{76,77} Several benzenoids and terpenoid lilac aldehydes (42) were confirmed to be EAD-active. However, despite the dominance of benzenoid compounds in the headspace, a mixture of lilac aldehyde isomers was equally attractive to the moths as the full blend of EAD-active compounds, indicating one or multiple of these isomers are the key attractive compounds. Indeed, while several other benzenoids were partially attractive, the next most attractive compound, methyl benzoate, elicited only half as many contacts with the source of volatiles in behavioural bioassays as did the mixture of lilac aldehyde isomers.

Nearly two decades later, another detailed and innovative study focused on the pollination chemistry of *Platanthera* species. Lahondère *et al.*⁵³ showed that in *P. obtusata*, the unusual attraction of mosquitoes was achieved primarily *via* the enhanced emission of nonanal (13) relative to lilac aldehydes (42), and that related species not pollinated by mosquitoes had much lower levels of 13 and much greater amounts of 42 and linalool (37), as well as several other EAD-active monoterpenes. The same study confirmed through behavioural and physiological experiments that increasing the amount of 42 (a mixture of three isomers) in a synthetic blend approximating the orchid scents reduced attractiveness to mosquitoes.

In tandem, these studies of *Platanthera* species highlight the complexity of plant–pollinator interactions mediated by chemistry, wherein one compound can be attractive to some pollinator species while deterring others. It remains unclear which of the eight possible lilac aldehyde stereoisomers were important to attraction or deterrence in each study, which may add

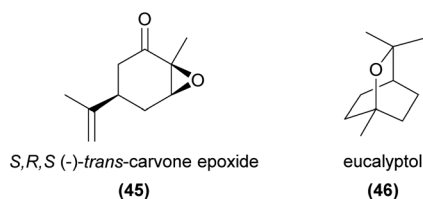


further complexity given that pollinator antennae may respond differently to each isomer.¹⁰⁶



Both α - and β -pinene (**43**, **44**) are key components of the oviposition site mimicry employed by *Epipactis veratrifolia*, which mimics the volatile compounds of aphids to attract several species of hoverflies as pollinators.⁴ These hoverflies seek to lay eggs near aphid colonies, as their larvae feed on aphids. Stökl *et al.*⁴ found that the floral volatiles of *E. veratrifolia* were very similar to those emitted as alarm pheromones from the aphid *Megoura viciae*, comprising primarily **43**, **44**, myrcene (**36**), and β -phellandrene, inducing the pollinator to lay eggs near the orchid. EAD experiments showed that each of these compounds were physiologically active in *Episyrphus balteatus* antennae, and further showed that when these compounds were added to bean plants, *E. balteatus* females laid significantly more eggs on the dosed plants compared to controls. A later study confirmed that a synthetic mixture of **43** and **44** alone was sufficient to attract hoverflies in field bioassays.¹⁰⁷ The monoterpene **44** also attracted the hoverfly pollinators of *Cypripedium subtropicum* in field trapping experiments.¹⁰⁸

Carvone epoxide (**45**) stands out as one of the more unusual monoterpenoids to be experimentally confirmed as a key to the attraction of the perfume-collecting euglossine bee pollinators of various orchids.^{109,110} Recently, Brandt *et al.*¹¹¹ investigated the relative attractiveness and prevalence of all four isomers of carvone epoxide, across five *Catasetum* species pollinated specifically by *Eulama* bee species. Only the (*S,R,S*)-(-)-*trans*-isomer was produced in these species and this isomer elicited both the strongest bee antennal response and the strongest attraction in field bioassays when compared to the other isomers.



Eucalyptol (1,8-cineole, **46**) has been known to be attractive to euglossine bees for over 50 years.² It has also been confirmed as an EAD-active component of attractive synthetic blends in several food rewarding and food deceptive orchids,^{53,74,112} and thus may warrant further attention in field bioassays.

3.2 Sesquiterpenes

Sesquiterpenes are composed of three isoprene units, and can be further modified to produce a great diversity of floral volatile

compounds. However, fewer sesquiterpenes have been shown to be involved in orchid pollination than monoterpenes, and their roles are often less well defined.

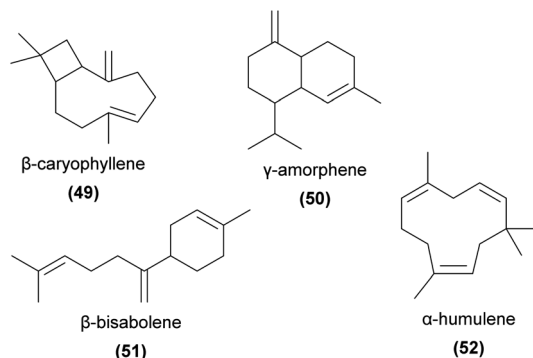
3.2.1 Acyclic sesquiterpenes. (*E*)- β -Farnesene (**47**) is a key attractant of the hoverfly pollinators of the intriguing aphid mimicking *Cypripedium subtropicum*.¹⁰⁸ This orchid produces white, hair-like tufts that visually mimic an aphid colony and also act as a nutrient reward. Hoverfly attraction to this visual mimic is facilitated by the emission of several terpenoids, including geranyl acetone (**53**), **47**, β -citronellol (**40**), β -caryophyllene (**49**), α -humulene (**52**), and β -pinene (**44**). All sampled aphid species co-occurring with *C. subtropicum* produced (*E*)- β -farnesene (**47**) as the dominant compound. Subsequent trapping experiments, using different compounds detected in the flower headspace as baits, showed that a synthetic mixture of all detected compounds was nearly as attractive as the flower itself, and that **47** alone was only slightly less attractive than the full mix of compounds. β -Pinene (**44**) also attracted significantly more hoverflies than the pentane control. Various isomers of farnesene have been reported in other orchid species,^{52,74,113} as has the related (*E,E*)-farnesol.^{28,39,66,86,114}



Farnesyl hexanoate (**48**) is one of few terpenoids known to play a role in sexual deception, where it acts as a repellent rather than an attractant. Schiestl and Ayasse¹¹⁵ showed that *Ophrys sphegodes* flowers exhibited a large increase in **48** after pollination. Furthermore, addition of **48** to unpollinated flowers substantially reduced their attractiveness to the male *Andrena nigroaenea* pollinator,¹¹⁵ in line with previous observations that production of this compound by mated female bees lowers their attractiveness to males.¹¹⁶

3.2.2 Cyclic sesquiterpenes. The common cyclic sesquiterpene β -caryophyllene (**49**) has been found to be EAD-active and/or a component of an attractive blend in several rewarding orchid species, including *Satyrion microrrhynchum*,¹¹⁷ *Chamorchis alpina*,¹¹² and *Cypripedium subtropicum*.¹⁰⁸ Outside the orchids, β -caryophyllene is more commonly associated with herbivore and pathogen resistance than pollination,^{118,119} which may also be the case for orchids as there are no reports of a definitive attractive role of β -caryophyllene. Several other cyclic sesquiterpenes have also been identified as orchid volatiles. Some have been shown to be EAD-active to pollinators, such as γ -amorphene (**50**),¹¹⁷ β -bisabolene (**51**),¹¹³ and α -humulene (**52**).¹⁰⁸ For each of these compounds, bioassays will be required to determine whether they play any role in pollinator interactions.



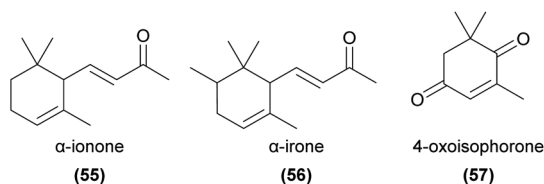


3.3 Other terpenoids

Geranylacetone (53) has been found to be a major floral volatile component of the aphid mimicking *Cypripedium subtropicum* and is weakly attractive to its hoverfly pollinators in field trapping experiments.¹⁰⁸ This compound has also been identified from several other orchids.^{80,120} 6-Methyl-5-hepten-2-one (sulcatone, 54) has also been found in many orchid species.^{57,59,66,120–125} It is EAD-active to the pollinators of *Gymnadenia conopsea* and *Orchis mascula*,^{73,74} but appears not to have been tested in field bioassays.



Apocarotenoids are degradation products of C₄₀-carotenoids, usually formed through oxidative cleavage of pigment compounds.¹²⁶ The products often play important signalling roles (for example, the phytohormones abscisic acid and strigolactones), and several are known as floral volatiles. Relevant examples of volatile apocarotenoids are ionones (e.g. 55) and their derivatives, which are known to be responsible for the smell of violets¹²⁷ and are common in some orchid species.¹⁰ α-Ionone, β-ionone and α-irone (56) are known to be attractive to perfume-collecting bees,^{94,128} and β-ionone elicits strong antennal responses from some perfume-collecting bees.⁹⁸ 4-Oxoisophorone (57) has been found to be EAD-active in pollinators of *Cypripedium calceolus*⁵² and has been detected in several other orchid species,^{122,129–132} as have several ionone isomers,^{59,64,129,133–140} though their functions in pollination are generally unknown.



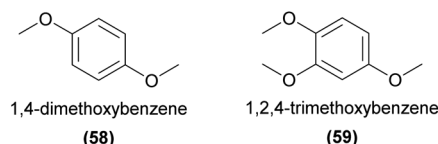
4 Benzenoids

Benzenoids are widespread plant volatile compounds and are usually biosynthesized from the aromatic amino acid phenylalanine, which itself is derived from the shikimate pathway.¹⁶ They can be classified based on the length of the main side chain, (C₆–C_{0–4}, for side chains comprising 0–4 carbons) with modifications such as methylation, hydroxylation, and acetylation leading to considerable structural diversity.¹³

Several C₆–C₁ benzenoids, such as benzyl alcohol (60), benzaldehyde (62), and benzyl benzoate (63) are amongst the most common of all floral volatiles, being found in more than half of the angiosperm families studied.⁹ Unsurprisingly, these compounds and many other benzenoids are well-represented across the orchids, and some have been shown to be relevant to pollination.

4.1 C₆–C₀

Some relatively common C₆–C₀ compounds found in orchid flowers are the dimethoxybenzenes (e.g. 58). While each of the isomers are known as orchid floral volatiles, to our knowledge only the *ortho* and *para* compounds have been confirmed as pollinator attractants in field bioassays. For example, 1,4-dimethoxybenzene (58) has been confirmed to be attractive in field bioassays to various perfume-collecting bee species and is reported to be an abundant floral volatile in members of the orchid genera *Embreea*, *Gongora*, and *Mormodes*.⁹⁴ More recently, this isomer has also been shown to elicit strong antennal responses from many other species of perfume-collecting bees across several genera.⁹⁸ The related 1,2,4-trimethoxybenzene (59) has also been shown to elicit antennal responses in several perfume-collecting bee species.^{98,113}



Outside of perfume rewarding systems, Salzmann *et al.*¹⁴¹ found that the volatile profile of the bumblebee-pollinated *Anacamptis coriophora* was dominated by benzenoids, most notably 58 and 4-methoxybenzaldehyde. These two compounds were the only compounds emitted by the orchid found to elicit a response from *Bombus terrestris* queen antennae.

4.2 C₆–C₁

Benzyl alcohol (60) is emitted from many orchid species, and has been shown to be attractive to various perfume-collecting bee species.^{94,100} It has also been shown to be EAD-active to pollinators of perfume rewarding,⁹⁸ food rewarding,^{73,77,142} food deceptive,⁵² and sexually deceptive⁶³ orchids, as well as to



pollinators of the honeybee alarm pheromone mimicking *Dendrobium sinense*.⁵



Similarly, the common benzenoids benzaldehyde (61),^{50,53,73,77,142,143} benzyl acetate,^{5,52,73,76,98,99,142,143} benzyl benzoate (62),^{77,99,143} and methyl salicylate (63)^{77,98,99,117} have been found to be EAD-active in several orchid pollinators.

4.3 C₆-C₂

In an interesting study, Nunes *et al.*¹⁴⁴ found that *Dichaea pendula* almost exclusively emitted 2-methoxy-4-vinylphenol (64). They further showed that this compound was not only attractive to several perfume-collecting bee pollinator species, but that it was even more strongly attractive to florivorous weevils, potentially representing a major evolutionary trade-off between beneficial and detrimental ecological interactions.

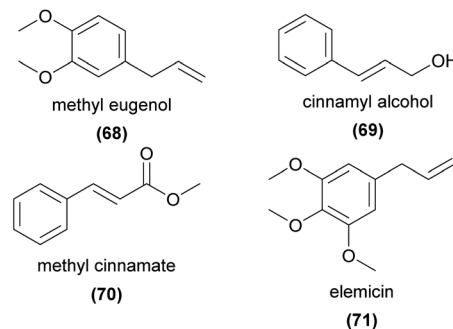
Phenylacetaldehyde (65) has been demonstrated to be attractive to moth pollinators of *Gymnadenia odoratissima* as a trap bait in field tests¹⁴³ and has been shown to be EAD-active in several other lepidopteran orchid pollinators.^{73,142}



Other common C₆-C₂ compounds have also been shown to be attractive to perfume-collecting bees in the field,⁹⁴ while 2-phenylethanol (66)^{52,73,142} and 2-phenylethyl acetate (67)^{52,73,142} are EAD-active in moth and bee pollinators. More recently, Chapurlat *et al.*⁷³ also demonstrated that phenylacetaldehyde (65), 66, 60, 67, and the phenylpropanoids methylisoeugenol and eugenol, are EAD-active components of the floral scent of *Gymnadenia conopsea* to two moth pollinators, *Deilephila porcellus* and *Aglaia urticae*.

4.4 C₆-C₃

Methyl eugenol (68) is strongly attractive to the fruit fly pollinators of several *Bulbophyllum* orchid species.¹⁴⁵ Isomers of 68 and eugenol have also been shown to be EAD-active in moth pollinators of various food rewarding orchids,^{72,73,143} beetle pollinators of *Satyrium microrrhynchum*,¹¹⁷ and bee pollinators of various perfume rewarding orchid species.^{94,98,99,128}



Cinnamyl alcohol (69) and its methyl ester, methyl cinnamate (70), are relatively common orchid volatiles. Both compounds are known as attractants of perfume-collecting bees and found in perfume rewarding flowers.^{94,128} The alcohol 69 is weakly attractive to the moth pollinators of *Platanthera bifolia*,⁷⁷ and is EAD-active in the moth pollinators of *Gymnadenia conopsea*⁷² where emission rates are correlated with pollinator activity.¹⁴⁶ The ester 70 is also EAD-active in the bumblebee pollinators of *Orchis mascula*.⁷⁴ Elemicin (71) is known to attract the fruit fly pollinators of *Bulbophyllum* orchids¹⁴⁷ as well as perfume-collecting bees,⁹⁴ and is EAD-active in moth^{72,73} and beetle¹¹⁷ pollinators.

4.5 C₆-C₄

Several phenylbutanoid compounds have been shown to be key attractants in the well-studied 'fruit fly orchids'. In a compelling early study, Nishida *et al.*¹⁴⁸ showed that male *Zeugodacus cucurbitae* flies were attracted to and fed from a single spot on a developed TLC plate separating the crude extract of *Dendrobium superbum* petals. They further showed that the attractive compound, 'raspberry ketone' (4-(4-hydroxyphenyl)-2-butanone, 72), was sequestered in the rectal glands of the male flies, and was only present in rectal gland extracts of males that fed on the flower. Raspberry ketone (72) was later found in other *Bulbophyllum* species,¹⁴⁹⁻¹⁵² and a methoxylated derivative, 'zingeron' (4-(4-hydroxy-3-methoxyphenyl)-2-butanone, 73), has been identified as a fruit fly attractant from several *Bulbophyllum* species as well.¹⁵¹⁻¹⁵⁴ Other studies have found these and related phenylbutanoids such as anisyl acetone (74), zingerol, rhododendrol, and 4-(4-methoxyphenyl)-2-butanol as significant components of the floral scent of *Bulbophyllum* species,^{149,152} all of which were also attractive to some extent to male *Zeugodacus cucurbitae* flies in laboratory bioassays.¹⁴⁹





5 Other compounds

5.1 Nitrogenous compounds

The nitrogen atom in organic compounds usually originates from an amino acid.¹⁶ In many alkaloids, building blocks from the acetate, shikimate, and terpene pathways are often also incorporated leading to a diverse group of natural products.

Indole (75) is a particularly common nitrogenous plant volatile, being derived from the same pathway as the amino acid tryptophan.¹⁵ While known as an attractant and modifier of perfume rewarding orchid fragrances,^{94,100} 75 has not been directly shown to be responsible for pollinator attraction in any other orchid species. However, it has been found to be EAD-active in several orchid pollinators,^{52,73,99} and is present widely in orchids with a variety of pollination strategies.^{54,55,131,135,155,156} It has been implicated in oviposition site mimicry³ and has been detected in a few orchids with this strategy.^{56,157} However, a direct role in orchid pollinator attraction has not yet been established.



Other nitrogenous compounds commonly found in orchids include benzyl nitrile (76),^{55,66,131,136} 2-aminobenzaldehyde (77),⁷³ and several aldoximes,^{55,132,136,155,158,159} but to our knowledge their roles in orchid pollination have not been established. However, these and many other nitrogenous compounds are known to be EAD- and behaviourally active in various other, non-orchid pollination systems (*e.g.* ref. 160–162), which indicates likely roles in orchid pollination as well.

Pyrazines are volatile heterocyclic nitrogenous compounds predicted to be derived from oxidative dimerisation of α -aminocarbonyls derived from amino acids. They are widely distributed in plants, insects, fungi and bacteria, but to date, their biosynthesis has only been established in bacteria.¹⁶³

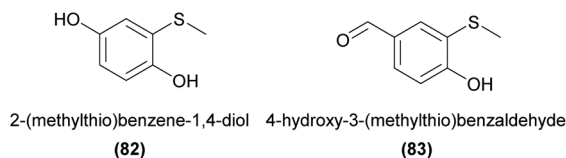
The sexually deceptive orchid genus *Drakaea* attracts male thynnine wasp pollinators primarily using various tetrasubstituted pyrazines. In *Drakaea glyptodon*, three alkyl pyrazines and one hydroxymethylpyrazine are EAD-active to its male *Zaspilothynnus trilobatus* thynnine wasp pollinator. Only two of these compounds, 2-butyl-3,5-dimethylpyrazine (78) and 2-hydroxymethyl-3,6-diethyl-5-methylpyrazine (79), in a 3 : 1 ratio, are required to elicit rates of sexual behaviour in field bioassays comparable to the flower.¹⁶⁴ The related species *D. micrantha* elicits sexual behaviour from its male *Zeleeboria* sp. thynnine wasp pollinator with a combination of drakolide (28) and two hydroxymethylpyrazines.⁸³

In a *D. livida* ecotype pollinated by male *Zaspilothynnus nigripes* thynnine wasps, 2-hydroxymethyl-3-(3-methylbutyl)-5-methylpyrazine (80) is EAD-active and present in both orchids and female wasps.¹⁶⁵ In another *D. livida* ecotype pollinated by male *Catocheilus* sp. thynnine wasps, four tetrasubstituted pyrazines and (2,5-dimethylpyrazin-3-yl)methyl 3-methylbutanoate (81) have been shown to be EAD-active.¹⁶⁶

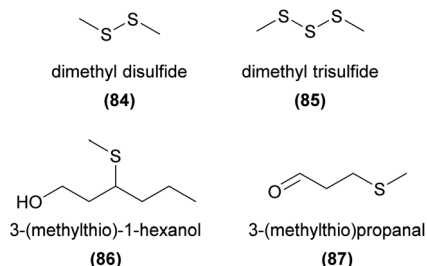
5.2 Sulfurous compounds

Natural compounds containing sulfur are generally derived from cysteine or methionine amino acid building blocks with cysteine being the major sulfur donor for thiosulfates and glucosinolates. A large portion of volatile sulfurous compounds are the result of enzymatic decomposition of these latter biosynthetic products.¹⁶⁷ (Methylthio)phenols have been identified from *Caladenia* and *Drakaea* orchids and found to act as pollinator attractants in two sexually deceptive *Caladenia* spider orchids. A set of four (methylthio)phenols were found to be used by *C. crebra* to sexually deceive its thynnine wasp pollinator *Campylothynnus flavopictus*. Three of the four semiochemicals had not previously been found as natural products.¹⁶⁸ Two of these compounds (82 and 83) have also been confirmed to attract another *Campylothynnus* sp. pollinator to *Caladenia attingens*.¹⁶⁹ More recently, all three previously unknown (methylthio)phenols have also been identified in one of the ecotypes of *Drakaea livida*, although no function has yet been reported.¹⁷⁰





Dimethyl disulfide (84) and dimethyl trisulfide (85) are strongly correlated with oviposition site mimicry, and are frequently emitted from flowers utilizing this pollination strategy.³ Surprisingly, to our knowledge these compounds have only twice been reported in orchids, without any confirmation of function.^{10,157} Several other sulfurous compounds have also been identified from orchids, again without any confirmed function, such as 3-(methylthio)-1-hexanol (86)¹³⁴ and 3-(methylthio)propanal (87).¹³⁹



6 Methods

Deciphering the chemical basis of pollinator attraction is no easy task. The floral volatile signals involved are often a complex blend with many of the components stored and emitted in sub-microgram amounts against a background of tens to hundreds of other compounds. For headspace sampling, detection may be hampered by slow release rates, while solvent extractions may fail to find compounds that are emitted as soon as they are produced. Even in highly specialised pollination systems, where just one or a few active compounds are involved in pollinator attraction, pinpointing candidate compounds can be challenging. Furthermore, in such cases the compounds involved are often unusual, making them challenging to identify without detailed structural elucidation.

Here, we highlight studies that illustrate some of the key methods that have facilitated new chemical discoveries and broadened the field of pollination chemistry. We commence with an overview of methods for isolation, before covering identification and data analysis. We conclude this section by covering some of the synthetic methods that have made it possible to confirm the identity and activity of unusual specific compounds in selected studies.

6.1 Isolation

To date, most studies of wild orchids have not attempted isolation of pure compounds directly, due to limited availability of flowers, and the generally low abundance of semiochemicals. Much more commonly, gas chromatography with mass spectrometry (GC-MS) is used to separate and analyse the compounds, with gas chromatography coupled with

electroantennographic detection (GC-EAD) frequently used in parallel to narrow down the candidate compounds.

Sampling of volatiles has generally been done with solvent extraction or headspace collection methods. Solvent extractions involve direct immersion of floral tissue in solvents of various polarities, and are particularly useful for analysing less volatile compounds, stored compounds, and to quickly answer questions about tissue specificity of volatiles. Headspace sampling captures volatile compounds from the air surrounding flowers by adsorption or absorption onto one of a variety of stationary phases, and is particularly useful for assessing the composition of volatiles emitted by flowers and that might realistically be encountered by pollinators in the environment. Headspace methods utilise either static sampling (e.g. Solid Phase Micro-extraction, SPME), or dynamic sampling, where volatiles are actively pumped onto adsorbents, and subsequently eluted with solvents or desorbed in a heated GC inlet.

The effectiveness of these techniques depends on the volatility, solubility, and polarity of each compound, which is often unknown at the outset. There are often large qualitative and quantitative differences between headspace analysis and solvent extraction. Generally, headspace analysis is most effective when relatively large amounts of volatiles are released from the plants, while solvent extraction is superior for plants that have large storage of semiochemicals and low release rates. For example, Gervasi *et al.*¹⁷¹ isolated a range of electrophysiologically active long-chain hydrocarbons by dichloromethane extraction of *Ophrys insectifera*, while Bohman *et al.*³⁸ employed SPME to extract shorter C₁₅- and C₁₇-hydrocarbons from the same species that were also found to be EAD-active to the pollinator *Argogorytes mystaceus*. Similarly, studies utilising both headspace and solvent extractions often find limited overlap between the results of each method.^{47,172} On the other hand, SPME has been unsuccessful for isolation of bioactive compounds from Australian *Drakaea* flowers despite multiple attempts (B. Bohman, pers. comm.), possibly due to low release rates of semiochemicals, while solvent extraction with dichloromethane has yielded the compounds of interest for several species.^{83,164,166} Another option to sample less volatile floral compounds with a non-destructive method is to sample floral tissue surfaces by rubbing with SPME fibres.^{40,173}

Adsorption of pheromone components from glass surfaces followed by desorption by a solvent such as diethyl ether is another effective way of isolating semiochemicals emitted at low rates and quantities. Wakamura *et al.*⁷¹ used this method to isolate sufficient amounts of 2,3-dihydroxypropyl isovalerate (21) to obtain 2D-NMR data using a cryoprobe following column chromatography, and identified this ester as the sex pheromone of the sexually deceived scarab beetle *Protaetia pryeri pryeri*.⁷¹ In short, the beetles were placed in a stainless steel net cage inside a closed glass beaker not allowing the beetles to touch the glass. Later, the cage was removed and the beaker rinsed with diethyl ether. Some studies of orchids pollinated by oil collecting bees have also used traditional natural product isolation procedures with column chromatography and NMR spectroscopy to identify less volatile compounds such as triterpenes and glycosides, although whether these compounds are involved in pollination remains to be confirmed.¹⁷⁴



For sexually deceptive orchids in particular, there are excellent examples of detailed chemical studies involving preparative chromatography methods.¹⁷⁵ One example is the study by Cuervo *et al.*⁶³ that fractionated extracts of the European *Ophrys leochroma*, pollinated by *Eucera* bees. These authors found that polar fractions eluted from a silica column with chloroform, dichloromethane, and diethyl ether, were more attractive to the bee pollinators than non-polar fractions of hexane elutions. This finding, together with the work of Ayasse *et al.*⁶⁰ on wasp-pollinated *Ophrys*, show that both these orchids lure their sexually deceived pollinators with polar compounds. In contrast, the vast majority of studies of *Ophrys* have only used non-polar solvents such as hexanes, with hydrocarbons predominantly identified as pollinator attractants.⁷

Three recent Australian studies report using semi-preparative gas chromatography to isolate bioactive floral compounds for NMR analysis (using a microprobe) and bioassays. These studies involve three diverse pollinator taxa; fungus gnats,⁴¹ ichneumonid wasps,⁸⁷ and thynnine wasps.⁸³ Hayashi *et al.*⁴¹ collected fractions from *Pterostylis orbiculata* for laboratory choice-bioassays with *Mycomya* fungus gnats, in order to successfully isolate the unsaturated hydrocarbons (6Z,9Z)-6,9-tricosadiene and (6Z,9Z)-1,6,9-tricosatriene (**6**) present in the flowers that elicited attraction. Bohman *et al.*⁸⁷ used SPME and semi-preparative gas chromatography concurrently to determine the set of long-range tetrahydrofuran acid derivatives **31** and **32** as attractants for the ichneumonid wasp *Lissopimpla excelsa* that pollinates *Cryptostylis ovata*. Thynnine wasp attractants have also been isolated with semi-preparative GC methods. Here, the hydroxymethylpyrazine **79** was initially identified as a candidate from *Drakaea micrantha* to attract its *Zebleboria* sp. pollinator, while the second component of the blend was unknown. By testing a combination of **79** and different GC-fractions of the orchid extract until a single fraction containing one main compound was confirmed as attractive to the thynnine pollinator, the first drakolide (**28**) was isolated.⁸³

6.2 Identification and data analysis

In most cases, especially for less specific pollination systems, candidate compounds are tentatively identified by GC-MS data comparisons with increasingly comprehensive databases. Commercially available reference libraries now provide EI-MS and retention index data for several hundred thousand compounds, and synthetic standards can often be readily obtained for confirmation. Nonetheless, it can still be difficult to distinguish between compounds with very similar properties based only on mass spectrometric fragmentation and retention time data. Furthermore, recently discovered compounds, including known pollinator attractants, will not be present in the commercial databases. Thus, their identification is often only possible by careful interpretation of GC-MS data after detection with electroantennography or isolation with preparative chromatography. For example, Bohman *et al.*^{164–166} tentatively identified novel hydroxymethylpyrazines from *Drakaea livida* and *D. glyptodon*, attracting thynnine wasp pollinators, by extrapolating mass fragmentations from other known

pyrazines, in combination with GC retention data. These identifications were then confirmed by comparison with synthesised compounds. Similarly, Cohen *et al.*⁸¹ identified a novel macro-lide, **27**, that attracts the cerambycid beetle pollinator of the exceptionally rare South African orchid *Disa forficaria* with GC-MS alone, and confirmed this identification by synthesis.

Additional techniques such as GC-FTIR^{89,176} are useful to differentiate between isomers while purification and NMR^{6,41,147} are helpful in elucidating new natural products when a sufficient amount of floral material is available – although this can be particularly challenging when the orchid species is rare. For example, the amount of sample required for a ¹H NMR is at least a thousand-fold higher than for a routine GC-MS analysis, and more comprehensive 2D-experiments require even higher amounts of isolated compound.

Differentiating between many possible structural isomers of certain compounds often requires additional methods to unambiguously assign a structure. Derivatisation with dimethyl disulphide (DMDS) is frequently used to determine alkene double bond positions, and while the studies that developed these techniques for volatile and semi-volatile compounds (ref. 177 and 178) are regularly cited, often very little experimental detail about the analysis is described. It should be noted that in the original papers, each candidate compound was synthesised and derivatised to enable direct comparison with derivatised biological extracts. The adduct formation upon treatment of alkenes is far from straightforward, particularly for complex extracts and for compounds containing more than one double bond¹⁷⁹ or other functional groups, such as alcohols or carbonyls.¹⁸⁰ Consequently, it would be advisable to provide more detailed experimental data when these experiments are carried out, such as clearly stating whether synthetic compounds have been derivatised and co-injected,¹⁸¹ and whether other isomers were ruled out. Including mass spectra and retention indices of adducts on which the identification is based would also provide the reader greater confidence in the identifications, as is common practice for identification of natural products by GC-MS. If semi-preparative GC is available, the analysis can be simplified by fractionating before treating each fraction with DMDS.³⁸

More generally, it is also important to be aware that obtaining synthetic standards and confirming retention times and mass spectra may not be sufficient for unambiguous identification if other structurally similar compounds cannot be ruled out. For example, in the identification of drakolide (**28**),⁸³ several structural analogues were synthesised and exhibited virtually identical GC-MS retention and mass spectral data. Furthermore, two of these isomers (differing in branching within a sidechain) required comparison across four GC-columns before they could be separated and the natural product confirmed. (Ref. 83 and B. Bohman pers. comm). Unfortunately, not even biological activity is always proof of a correctly identified natural product, as structural analogues may also show very similar activity.^{84,182} Overall, multiple GC-columns of differing stationary phase and carefully optimised methods should be employed in order to adequately compare retention data and mass spectra between natural products and synthetic standards.



Chiral semiochemicals also require additional identification efforts. Wakamura *et al.*⁷¹ used HPLC with a chiral-phase column to separate the two enantiomers of 2,3-dihydroxypropyl isovalerate, while chiral monoterpenes, pyrazines and drakolides have been easily separated by chiral-phase GC.^{4,83,165} In order to determine the absolute configuration, chromatographic separation of stereoisomers (usually using a chiral stationary phase) in combination with methods such as X-ray crystallography or measurement of the optical rotation or

electronic circular dichroism (ECD) of synthetic standards can be applied. Then, the peak of the natural product can be aligned chromatographically with the corresponding peak of the fully characterised synthetic standard. This methodology was used in the confirmation of the absolute configuration of the sexually attractive drakolide from *Drakaea micrantha*.⁸³

Chemical analysis may also involve comparative analysis of the recorded data. (Semi)automated analysis of GC-MS data with peak-deconvolution and alignment software, statistical



Scheme 1 Examples of synthetic routes to orchid pollinator attractants: (A) (8Z,20Z)-nonacosadiene from *Ophrys exalata*, (B) chiloglottone from *Chiloglottis* spp., (C) hydroxymethylpyrazines from *Drakaea livida*, (D) drakolide (**28**) from *Drakaea micrantha*, (E) (methylthio)phenols from *Caladenia crebra*, (F) disalactone (**27**) from *Disa forficaria*.

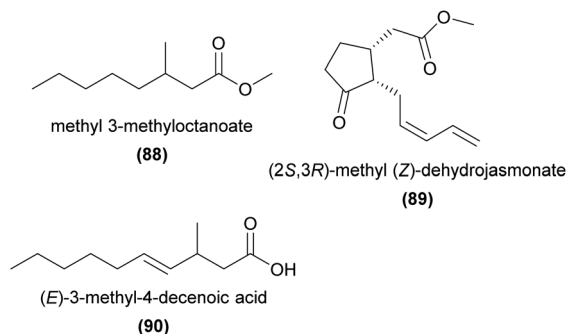


treatment, and graphical visualisation of data have undergone rapid development over the last decade, largely due to the expansion of the field of metabolomics.^{183,184} These developments have facilitated comparative studies of comprehensive floral volatiles, which complement structural elucidation of new specific semiochemicals to enable new insights into the roles of chemical communication. Such “chemical phenotyping” studies have, for example, revealed relationships between chemical composition and the phylogeny in taxa of perfume rewarding orchids,^{59,125,135} disentangled visual and olfactory signals in mushroom-mimicking *Dracula* orchids,⁵¹ showed correlations between floral cuticular hydrocarbon compositions and fly pollinators in *Neotinea*,⁴⁷ and distinguished chemotypes of sexually deceptive *Drakaea* orchids.¹⁷⁰ Here, it is important to note that to identify any specific compounds revealed by such analyses as potentially biologically relevant, comparison of retention data and mass spectral fragmentations with those of fully characterised reference compounds is critical.

6.3 Synthesis

Many of the unusual compounds found in orchids, particularly from the highly specialised sexually deceptive orchid taxa, have required the development of new routes of synthesis.

Excellent examples of earlier work in preparing orchid natural products by enantioselective methods include the suite of papers by Kitahara and coworkers,^{185–188} who synthesised candidate stereoisomers of a range of natural products from orchids, and confirmed the absolute configuration with chiral-phase GC. Both enantiomers of ethyl 3-methyloctanoate (**88**) were prepared in nine steps from methyl 3-hydroxy-2-methylpropanoate. Comparing both enantiomers by chiral-phase GC revealed that the natural product in the African *Aerangis* spp. orchid studied had *S*-configuration.¹⁸⁶ Methyl *cis*-(*Z*)-dihydrojasmonate (**89**) was determined to have (2*S*,3*R*)-configuration in the Asian *Cymbidium goeringii* after synthesis of all four stereoisomers from an in-house available intermediate in eight steps.¹⁸⁷ Similarly, (*E*)-3-methyl-4-decenoic acid (**90**) and derivatives were prepared to determine the configuration of a range of European and Indo-Australian orchids¹⁸⁵ and both enantiomers of *cis*-3-methyl-4-decanolide were synthesised in eight steps from (1*S*,5*R*)-2-oxabicyclo[3.3.0]oct-6-en-3-one (**29**).¹⁸⁸



In the case of the many European *Ophrys* studies, few have reported the detailed synthesis of alkenes. The more unusual (8*Z*,20*Z*)-nonacosadiene, found in *O. exaltata*, was however

prepared in five steps from the commercially available 10-bromo-1-decanol, *via* Lindlar hydrogenation of the corresponding alkyne (Scheme 1A). Chromatography on silica provided the (*Z*)-alkene in 94% purity.²⁹ Another unusual unsaturated hydrocarbon, (6*Z*,9*Z*)-1,6,9-tricosatriene (**6**), found in Australian *Pterostylis orbiculata*, was identified, synthesised in five steps from propargyl alcohol, and confirmed to be a sexual attractant to the male fungus gnat pollinator.⁴¹

A nice example where orchid chemistry has motivated the modification of existing methods is provided by Delle-Vedove *et al.*,¹⁵⁸ who developed a new stereoselective method for synthesis of (3*E*)-4,8-dimethyl-1,3,7-nonatriene (DMNT). An aqueous Wittig–Horner–Emmons reaction in combination with a Cu-catalysed decarboxylation to complement the classic Wittig-approach yielded a 38/62 (*Z/E*) ratio of the product (as determined by NMR) allowing the determination of the configuration of the natural product.¹⁵⁸

Chiloglottones, which were the first orchid pollinator attractants to be identified in Australia,⁸⁸ have been synthesised by multiple methods. Initially, a condensation approach between ethyl 3-oxohexanoate and ethyl (2*E*)-hex-2-enoate was taken to prepare chiloglottone (**33**).⁸⁹ This protocol was later replaced by a more general route from 1,3-dimethoxybenzoic acid, *via* resorcinol derivatives, allowing a greater flexibility of substituents (Scheme 1B).¹⁸⁹

Several semiochemicals from *Drakaea* orchids have also been synthesised *via* alternative routes, with the new refinements improving flexibility. For hydroxymethylpyrazines, two synthetic methods have been reported. In the earlier work, Kumada–Corriu cross-couplings were combined with Boekelheide hydroxylations to prepare for example 2-hydroxymethyl-3-(3-methylbutyl)-5-methylpyrazine (**80**) from 2,5-dimethylpyrazine in six steps.¹⁶⁵ Later, a more convenient four-step method was used based on Minisci-type chemistry (Scheme 1C).^{164,190} Drakolide (**28**), was prepared from (*S*)-2-methylpentanal and ethyl-2-methyl-3-oxobutanoate (Scheme 1D), with the resulting diastereoisomers separated with HPLC and enantiomeric purity enhanced with chiral-phase HPLC. The absolute configuration of **28** was determined with X-ray crystallography of the synthetic stereoisomer and confirmed to match the natural product by chiral-phase HPLC and GC.⁸³ Another recent example from Australia is the synthesis of three (methylthio)phenols, used as wasp pollinator attractants by spider orchids such as *Caladenia crebra*. 4-Hydroxy-3-(methylthio)benzaldehyde (**83**) was prepared from 2-methoxythiophenol through an *S*-methylation, Vilsmeier–Haack formylation, and *O*-demethylation. The corresponding alcohol (**82**) was formed through a sodium borohydride reduction. 2-(Methylthio)benzene-1,4-diol was prepared by oxidising the commercially available 2-methylthiophenol using Fremy's salt (Scheme 1E).¹⁶⁸

Most recently, Katte *et al.*¹⁴⁹ have provided good examples of orchid semiochemical synthesis by preparing ‘syringerone’ (4-(4-hydroxy-3,5-dimethoxyphenyl)-2-butanone), ‘raspberry ketone’ (**72**) and its acetate from *Bulbophyllum* fruit fly orchids of Asia. In South Africa, Cohen *et al.*⁸¹ synthesised the macrolide



27 found in *Disa forficaria* in five steps from (9Z)-16-hydroxy-octadec-9-enoic acid (Scheme 1F).

7 Final remarks

7.1 The importance of behavioural bioassays

There are no shortcuts in establishing the biological role of particular compounds in specific orchid–pollinator interactions. Instead, behavioural bioassays are required to determine which compounds are relevant, and to what extent.

Often there are tens to hundreds of compounds present in floral extracts, and so for simplicity it may be tempting to draw conclusions about the function of chemical compounds in one system based on their function in another. Specifically, when compounds are known to be key to pollinator attraction in one orchid, finding them in another might suggest a similar role. In some cases this may indeed be a valid inference, but there are also many examples in which attractive compounds in one system are seemingly unimportant or play contrasting roles in another. As a case in point, two studies on *Platanthera* species show that shared compounds do not always have the same function for pollinator attraction even in related orchids. Plepys *et al.*⁷⁷ showed that lilac aldehydes (**42**) were key floral attractants to the hawkmoth pollinators of *P. bifolia*, while Lahondère *et al.*⁵³ showed that lilac aldehydes (**42**) reduced attraction of the mosquito pollinators of *P. obtusata*. These studies highlight the need for behavioural investigations of pollination chemistry to determine function, and both provide good examples of behavioural and physiological investigations to understand the chemical basis of pollinator attraction.

Measuring electroantennographic physiological activity (EAG or GC-EAD) has proven a powerful tool for pinpointing only the compounds perceptible to pollinators, but does not in itself provide information about the function of those compounds. For example, in the hornet pollinators of *Dendrobium sinense*, benzyl acetate and benzyl alcohol (**60**) are the most strongly EAD-active compounds from the orchid volatile profile, but are not required to attract pollinators.⁵ Similarly, Huber *et al.*¹⁴³ found that benzaldehyde (**61**), phenylacetaldehyde (**65**), 2-phenylethyl acetate (**67**), benzyl acetate, eugenol, and 1-phenyl-2,3-butanedione were emitted from *Gymnadenia odoratissima* and were EAD-active to their pyralid moth pollinators, but only phenylacetaldehyde (**65**) was attractive in the field.

It is worth noting that EAD-active compounds that do not appear to be involved in pollinator attraction may still warrant closer research attention, since they may play roles beyond pollinator attraction. For example, the alkanes and alkenes may be particularly interesting in this context. Given their prevalence in orchid extracts and their very wide use as insect signalling compounds, it is plausible that some chemical information could be encoded by these compounds. For example, honey bees and bumblebees are known to be able to detect hydrocarbon ‘footprints’ from conspecifics present on flowers and avoid visiting these flowers, thereby saving time that might be wasted visiting a flower that had recently had nectar or pollen resources collected.^{191,192} While there is

currently no evidence that hydrocarbons play this role in orchids, it serves as a reminder that there may be many subtle cues perceptible to pollinators that are as yet unknown.

7.2 Considering absolute configuration

The absolute configuration of semiochemicals, including orchid pollinator attractants, is often fundamental for biological activity. As a result, it is often important to analyse samples enantioselectively and to test stereoisomers separately in bioassays in order to accurately determine any biological activity. For example, Schorkopf *et al.*¹⁰⁵ assessed pollinator discrimination between enantiomers of ipsdienol, a monoterpene alcohol known from several perfume rewarding species.^{104,113} Male *Euglossa cyanura* bees were attracted to the (*R*)-enantiomer (**41**) and the racemate, but showed virtually no attraction to the (*S*)-enantiomer. They further showed that male *E. cyanura* antennae were consistently more responsive to the (*R*)-enantiomer, potentially contributing to the difference in behavioural response.

Similar observations, but with the (*S*)-enantiomer and racemate rather than the (*R*)-enantiomer and racemate being active have been reported in an analysis of the field activity of β -citronellol (**40**) to the thynnine wasp pollinator of *Caladenia plicata*.¹⁰³ In other cases, while one enantiomer is attractive, the presence of the other reduces or completely prevents attraction to the racemate. For example, Williams and Whitten⁹⁴ described baiting with both enantiomers of α -pinene, and found that while the (–)-enantiomer (1*S*,5*S*) was attractive to *Eulama nigrita*, the (+)-enantiomer (1*R*,5*R*) was not. Furthermore, the racemate was also unattractive, suggesting that the (+)-enantiomer was either actively repellent or masked the effect of the (–)-enantiomer to this species. More recently, Wakamura *et al.*⁷¹ showed that male *Protaetia* beetles were exclusively attracted to the (*R*)-enantiomer of 2,3-dihydroxypropyl isovalerate (**21**), but were not attracted to the (*S*)-enantiomer or to the racemate, again suggesting that the presence of the (*S*)-enantiomer prevents attraction to the (*R*)-enantiomer. These findings highlight the critical importance of not ruling out the attractiveness of compounds based on bioassays using mixtures of stereoisomers. Instead, pure stereoisomers are required for unambiguous evaluation of biological activity.

7.3 Lessons from sexual deception

The chemistry of pollination by sexual deception has been frequently highlighted throughout this review. Indeed, there has been remarkable progress on this topic in the 20+ years since Schiestl *et al.*²⁷ first confirmed the chemical sexual mimicry of female *Andrena* bees by *Ophrys* orchids, and this research continues to grow. Since 2016, the chemistry of sexual deception has been confirmed in diverse Australian genera,^{41,102,168} and for the first time in Africa,⁸¹ Asia,⁷¹ and South America.⁴⁵ These recent chemical discoveries include the first examples from sexually deceptive systems pollinated by Diptera^{41,45} and Coleoptera.^{71,81} Notably these ‘solved’ cases represent a small fraction of the hundreds of known cases of sexual deception, meaning that there are many more discoveries to be made.



The ongoing progress in this area is likely because sexually deceptive systems are particularly amenable to investigations of pollination chemistry. They often have somewhat simpler floral volatile profiles than more generalist systems, and in some studies the dominant volatiles are the active ones (e.g. ref. 41 and 102). The strong sexual responses of males to sex pheromones also make them particularly amenable to experimental bioassays in the laboratory and field, as there are typically many rapid responses to the 'correct' compound or mixture of compounds. Finally, investigations of sexually deceptive orchids often integrate multiple lines of evidence and methods to establish active chemical compounds. For example, the ability to reduce the number of candidate compounds by comparing chemical extracts of active and non-active floral tissues with extracts of the female of the pollinator being mimicked has proven powerful. The addition of GC-EAD and/or semi preparative techniques, particularly when combined with iterative field and/or lab bioassays further helps to pinpoint the compounds involved.¹⁷⁵

While sexual deception is undoubtedly among the most highly specialised of all plant pollination strategies, the dependence of an orchid species on just a few pollinator species is common.¹⁹³ It follows that the strategies used to disentangle the chemistry of sexual deception can be applied to many hundreds or even thousands of other orchid species. Other types of mimicry-based pollination strategies are particularly ripe for investigation. Indeed, this review has highlighted several cases where the chemistry of unusual plant–pollinator interactions has been solved by integrating information from the orchid, the pollinator, and the putative model (e.g. mimicry of honeybee alarm pheromones,⁵ mimicry of green leaf volatiles⁵⁰ etc.).

In less specific pollination systems, where a wider variety of species are attracted to general scents indicative of a food reward, the particular compounds responsible for attraction can be more challenging to reliably elucidate than in more specialised cases. Here, attraction is often due to a large 'bouquet' of compounds, each contributing partially to pollinator attraction. For this reason, studies attempting to disentangle the effects of individual compounds in food rewarding and food deceptive systems are comparatively rare (but see ref. 53, 77 and 143). We predict that a chemical phenotyping approach, where taxonomic units are compared, and diagnostic compounds are subsequently identified, could be the first steps towards a more comprehensive understanding of the role of chemistry in diverse pollination systems. Examples of this approach have already been successfully applied in perfume rewarding and sexually deceptive pollination systems.^{47,59,135,170} Chemical phenotyping may also provide important clues about the evolution of pollination strategy shifts and the evolution of novel semiochemicals. Thus, we recommend this approach as a potentially fruitful area for further research.

8 Conclusion

This review has demonstrated that orchids are a rich source of interesting and unusual natural products. Yet, despite many

decades of interdisciplinary studies, we have most likely only identified a very small fraction of the compounds pivotal for pollination of orchids. Given the huge number and great diversity of orchid species, the opportunities for new discoveries in this space are near endless.

9 Conflicts of interest

There are no conflicts to declare.

10 Acknowledgements

This work was supported by the Australian Research Council DP210100471 (R. P. and G. R. F.) and an Australian Government Research Training Program Scholarship to J. P.

11 References

- 1 R. Govaerts, P. Bernet, K. Kratochvil, G. Gerlach, G. Carr, P. Alrich, A. M. Pridgeon, J. Pfahl, M. A. Campacci, D. Holland Baptista, H. Tigges, J. Shaw, P. Cribb, A. George, K. Kreuz and J. Wood, *World Checklist of Orchidaceae. Facilitated by the Royal Botanic Gardens, Kew. Published on the Internet*, (accessed August 28, 2022).
- 2 C. H. Dodson, R. L. Dressler, H. G. Hills, R. M. Adams and N. H. Williams, *Science*, 1969, **164**, 1243–1249.
- 3 A. Jürgens, S.-L. Wee, A. Shuttlesworth and S. D. Johnson, *Ecol. Lett.*, 2013, **16**, 1157–1167.
- 4 J. Stökl, J. Brodmann, A. Dafni, M. Ayasse and B. S. Hansson, *Proc. R. Soc. B*, 2011, **278**, 1216–1222.
- 5 J. Brodmann, R. Twele, W. Francke, L. Yi-bo, S. Xi-qiang and M. Ayasse, *Curr. Biol.*, 2009, **19**, 1368–1372.
- 6 M. Sugahara, K. Izutsu, Y. Nishimura and F. Sakamoto, *Zool. Sci.*, 2013, **30**, 99–104.
- 7 B. Bohman, G. R. Flematti, R. A. Barrow, E. Pichersky and R. Peakall, *Curr. Opin. Plant Biol.*, 2016, **32**, 37–46.
- 8 F. P. Schiestl, *Naturwissenschaften*, 2005, **92**, 255–264.
- 9 J. T. Knudsen, R. Eriksson, J. Gershenzon and B. Ståhl, *Bot. Rev.*, 2006, **72**, 1–120.
- 10 R. Kaiser, *The Scent of Orchids. Olfactory and chemical investigations*, Elsevier, Amsterdam, 1993.
- 11 R. Kaiser, *Meaningful Scents Around the World: Olfactory, Chemical, Biological, and Cultural Considerations*, Wiley, 2006.
- 12 J. T. Knudsen and J. Gershenzon, in *Biology of Plant Volatiles*, ed. E. Pichersky and N. Dudareva, CRC Press, 2020, pp. 57–79.
- 13 J. K. Muhlemann, A. Klempien and N. Dudareva, *Plant, Cell Environ.*, 2014, **37**, 1936–1949.
- 14 F. P. Schiestl and F. Marion-Poll, in *Analysis of Taste and Aroma*, ed. J. F. Jackson and H. F. Linskens, Springer, Berlin, Heidelberg, 2002, pp. 173–198.
- 15 E. Pichersky, J. P. Noel and N. Dudareva, *Science*, 2006, **311**, 808–811.
- 16 P. M. Dewick, *Medicinal Natural Products: A Biosynthetic Approach*, John Wiley & Sons, 2009.



- 17 X. Yang, I. A. Guschina, S. Hurst, S. Wood, M. Langford, T. Hawkes and J. L. Harwood, *Pest Manage. Sci.*, 2010, **66**, 794–800.
- 18 J. W. Campbell and J. E. Cronan, *Annu. Rev. Microbiol.*, 2001, **55**, 305–332.
- 19 A. Bernard, F. Domergue, S. Pascal, R. Jetter, C. Renne, J.-D. Faure, R. P. Haslam, J. A. Napier, R. Lessire and J. Joubès, *Plant Cell*, 2012, **24**, 3106–3118.
- 20 Y. Qiu, C. Tittiger, C. Wicker-Thomas, G. Le Goff, S. Young, E. Wajnberg, T. Fricaux, N. Taquet, G. J. Blomquist and R. Feyereisen, *Proc. Natl. Acad. Sci. U. S. A.*, 2012, **109**, 14858–14863.
- 21 P. M. Schlüter, S. Xu, V. Gagliardini, E. Whittle, J. Shanklin, U. Grossniklaus and F. P. Schiestl, *Proc. Natl. Acad. Sci. U. S. A.*, 2011, **108**, 5696–5701.
- 22 S. B. Lee and M. C. Suh, *Plant Cell Rep.*, 2015, **34**, 557–572.
- 23 L. Samuels, L. Kunst and R. Jetter, *Annu. Rev. Plant Biol.*, 2008, **59**, 683–707.
- 24 G. J. Blomquist, C. Tittiger and R. Jurenka, in *Hydrocarbons, Oils and Lipids: Diversity, Origin, Chemistry and Fate*, ed. H. Wilkes, Springer International Publishing, Cham, 2018, pp. 1–32.
- 25 H. Holze, L. Schrader and J. Buellesbach, *Heredity*, 2021, **126**, 219–234.
- 26 H. Chung and S. B. Carroll, *BioEssays*, 2015, **37**, 822–830.
- 27 F. P. Schiestl, M. Ayasse, H. F. Paulus, C. Löfstedt, B. S. Hansson, F. Ibarra and W. Francke, *Nature*, 1999, **399**, 421.
- 28 F. P. Schiestl, M. Ayasse, H. F. Paulus, C. Löfstedt, B. S. Hansson, F. Ibarra and W. Francke, *J. Comp. Physiol. A*, 2000, **186**, 567–574.
- 29 J. Mant, C. Brändli, N. J. Vereecken, C. M. Schulz, W. Francke and F. P. Schiestl, *J. Chem. Ecol.*, 2005, **31**, 1765–1787.
- 30 M. Ayasse, in *Biology of Floral Scent*, ed. N. Dudareva and E. Pichersky, CRC Press, 2006, pp. 219–241.
- 31 M. Ayasse, J. Stöckl and W. Francke, *Phytochemistry*, 2011, **72**, 1667–1677.
- 32 R. Peakall, D. C. Wong, B. Bohman, G. R. Flematti and E. Pichersky, in *Biology of Plant Volatiles*, ed. E. Pichersky and N. Dudareva, CRC Press, 2020, pp. 271–295.
- 33 D. D. L. Gervasi, M.-A. Selosse, M. Sauve, W. Francke, N. J. Vereecken, S. Cozzolino and F. P. Schiestl, *Ecol. Evol.*, 2017, **7**, 6023–6034.
- 34 J. Göglér, J. Stöckl, A. Sramkova, R. Twele, W. Francke, S. Cozzolino, P. Cortis, A. Scrugli and M. Ayasse, *Evolution*, 2009, **63**, 2222–2234.
- 35 J. Stöckl, H. Paulus, A. Dafni, C. Schulz, W. Francke and M. Ayasse, *Plant Syst. Evol.*, 2005, **254**, 105–120.
- 36 J. Stöckl, P. M. Schlüter, T. F. Stuessy, H. F. Paulus, G. Assum and M. Ayasse, *Am. J. Bot.*, 2008, **95**, 472–481.
- 37 J. Stöckl, P. M. Schlüter, T. F. Stuessy, H. F. Paulus, R. Fraberger, D. Erdmann, C. Schulz, W. Francke, G. Assum and M. Ayasse, *Biol. J. Linn. Soc.*, 2009, **98**, 439–451.
- 38 B. Bohman, A. M. Weinstein, R. Mozuraitis, G. R. Flematti and A.-K. Borg-Karlson, *Int. J. Mol. Sci.*, 2020, **21**, 620.
- 39 M. Ayasse, F. P. Schiestl, H. F. Paulus, C. Löfstedt, B. Hansson, F. Ibarra and W. Francke, *Evolution*, 2000, **54**, 1995–2006.
- 40 N. Joffard, V. Arnal, B. Buatois, B. Schatz and C. Montgelard, *Plant Biol.*, 2020, **22**, 881–889.
- 41 T. Hayashi, B. Bohman, A. Scaffidi, R. Peakall and G. R. Flematti, *Curr. Biol.*, 2021, **31**, 1954–1961.
- 42 C. Löfstedt, N. Wahlberg and J. G. Millar, in *Pheromone Communication in Moths*, ed. J. Allison and R. T. Cardé, University of California Press, 2016, pp. 43–78.
- 43 A. El-Sayed, *The pherobase: database of insect pheromones and semiochemicals*, <https://www.pherobase.com>, (accessed August 29, 2022).
- 44 C. Martel, L. Cairampoma, F. W. Stauffer and M. Ayasse, *PLoS One*, 2016, **11**, e0165896.
- 45 C. Martel, W. Francke and M. Ayasse, *New Phytol.*, 2019, **223**, 1989–2001.
- 46 A. Flach, A. J. Marsaioli, R. B. Singer, M. D. C. E. Amaral, C. Menezes, W. E. Kerr, L. G. Batista-Pereira and A. G. Corrêa, *J. Chem. Ecol.*, 2006, **32**, 59–70.
- 47 C. Martel, D. Rakosy, P. E. Romero, J. Jersáková and M. Ayasse, *J. Systemat. Evol.*, 2021, DOI: [10.1111/jse.12812](https://doi.org/10.1111/jse.12812).
- 48 F. P. Schiestl and S. Cozzolino, *BMC Evol. Biol.*, 2008, **8**, 27.
- 49 N. J. Vereecken, C. A. Wilson, S. Hötling, S. Schulz, S. A. Banketov and P. Mardulyn, *Proc. R. Soc. B*, 2012, **279**, 4786–4794.
- 50 J. Brodmann, R. Twele, W. Francke, G. Hölzler, Q.-H. Zhang and M. Ayasse, *Curr. Biol.*, 2008, **18**, 740–744.
- 51 T. Policha, A. Davis, M. Barnadas, B. T. M. Dentinger, R. A. Raguso and B. A. Roy, *New Phytol.*, 2016, **210**, 1058–1071.
- 52 H. Braunschmid, B. Mükisch, T. Rupp, I. Schäffler, P. Zito, D. Birtele and S. Dötterl, *Arthropod-Plant Interact.*, 2017, **11**, 363–379.
- 53 C. Lahondère, C. Vinauger, R. P. Okubo, G. H. Wolff, J. K. Chan, O. S. Akbari and J. A. Riffell, *Proc. Natl. Acad. Sci. U. S. A.*, 2020, **117**, 708–716.
- 54 J. M. R. B. V. Aguiar, G. D. S. Ferreira, P. A. Sanches, J. M. S. Bento and M. Sazima, *Phytochemistry*, 2021, **182**, 112591.
- 55 M. G. Balducci, T. Van Der Niet and S. D. Johnson, *Ann. Bot.*, 2020, **126**, 1155–1164.
- 56 L. Humeau, C. Micheneau, H. Jacquemyn, A. Gauvin-Bialecki, J. Fournel and T. Pailler, *J. Trop. Ecol.*, 2011, **27**, 591–599.
- 57 N. Joffard, I. Le Roncé, A. Langlois, J. Renoult, B. Buatois, L. Dormont and B. Schatz, *J. Evol. Biol.*, 2020, **33**, 1028–1038.
- 58 F. S. Robustelli della Cuna, J. Calevo, M. Bazzicalupo, C. Sottani, E. Grignani and S. Preda, *Plants*, 2021, **10**, 1718.
- 59 D. W. Roubik and J. T. Knudsen, *Flora*, 2017, **232**, 117–127.
- 60 M. Ayasse, F. P. Schiestl, H. F. Paulus, F. Ibarra and W. Francke, *Proc. R. Soc. B*, 2003, **270**, 517–522.
- 61 J. Mant, R. Peakall and F. P. Schiestl, *Evolution*, 2005, **59**, 1449–1463.
- 62 J. Stöckl, R. Twele, D. H. Erdmann, W. Francke and M. Ayasse, *Chemoecology*, 2007, **17**, 231–233.



- 63 M. Cuervo, D. Rakosy, C. Martel, S. Schulz and M. Ayasse, *J. Chem. Ecol.*, 2017, **43**, 469–479.
- 64 J. Calevo, M. Bazzicalupo, M. Adamo, F. S. Robustelli della Cuna, S. Voyron, M. Girlanda, K. J. Duffy, A. Giovannini and L. Cornara, *Diversity*, 2021, **13**, 550.
- 65 A. Manzo, S. Panzeri, I. Vagge and A. Giorgi, *Molecules*, 2014, **19**, 7913–7936.
- 66 C. I. Peter and S. D. Johnson, *Ann. Bot.*, 2014, **113**, 277–288.
- 67 C. I. Keeling, G. W. Otis, S. Hadisoelilo and K. N. Slessor, *Apidologie*, 2001, **32**, 243–252.
- 68 G. Lercker, P. Capella, L. S. Conte, F. Ruini and G. Giordani, *Lipids*, 1981, **16**, 912–919.
- 69 F. Martos, M.-L. Cariou, T. Pailler, J. Fournel, B. Bytebier and S. D. Johnson, *New Phytol.*, 2015, **207**, 225–234.
- 70 N. Arakaki, K. Yasuda, S. Kanayama, S. Jitsuno, M. Oike and S. Wakamura, *Appl. Entomol. Zool.*, 2016, **51**, 241–246.
- 71 S. Wakamura, N. Arakaki, D. Moriyama, S. Kanayama, M. Oike, A. Kimura, S. Wajima, H. Ono and H. Yasui, *Chemoecology*, 2020, **30**, 49–57.
- 72 J. Jersáková, S. Castro, N. Sonk, K. Milchreit, I. Schödelbauerová, T. Tolasch and S. Dötterl, *Evol. Ecol.*, 2010, **24**, 1199–1218.
- 73 E. Chapurlat, J. Ågren, J. Anderson, M. Friberg and N. Sletvold, *New Phytol.*, 2019, **222**, 2009–2022.
- 74 L. Dormont, T. Fort, J.-M. Bessière, M. Proffit, E. Garcia Hidalgo, B. Buatois and B. Schatz, *Acta Oecol.*, 2020, **107**, 103600.
- 75 R. Kubo and M. Ono, *Can. Entomol.*, 2017, **149**, 372–376.
- 76 D. Plepys, F. Ibarra, W. Francke and C. Löfstedt, *Oikos*, 2002, **99**, 75–82.
- 77 D. Plepys, F. Ibarra and C. Löfstedt, *Oikos*, 2002, **99**, 69–74.
- 78 S. L. Buchmann, *Annu. Rev. Ecol. Syst.*, 1987, **18**, 343–369.
- 79 I. Schäffler, K. E. Steiner, M. Haid, S. S. van Berkel, G. Gerlach, S. D. Johnson, L. Wessjohann and S. Dötterl, *Sci. Rep.*, 2015, **5**, 12779.
- 80 M. Castañeda-Zárate, S. D. Johnson and T. van der Niet, *Curr. Biol.*, 2021, **31**, 238–246.
- 81 C. Cohen, W. R. Liltved, J. F. Colville, A. Shuttleworth, J. Weissflog, A. Svatoš, B. Bytebier and S. D. Johnson, *Curr. Biol.*, 2021, **31**, 1962–1969.
- 82 S. Schulz and S. Hötling, *Nat. Prod. Rep.*, 2015, **32**, 1042–1066.
- 83 B. Bohman, M. M. Y. Tan, R. D. Phillips, A. Scaffidi, A. N. Sobolev, S. A. Moggach, G. R. Flematti and R. Peakall, *Angew. Chem., Int. Ed.*, 2020, **59**, 1124–1128.
- 84 B. Bohman, M. M. Y. Tan, G. R. Flematti and R. Peakall, *J. Chem. Ecol.*, 2022, **48**, 323–336.
- 85 D. Bartschat, D. Lehmann, A. Dietrich, A. Mosandl and R. Kaiser, *Phytochem. Anal.*, 1995, **6**, 130–134.
- 86 R. Luyt and S. D. Johnson, *Plant Syst. Evol.*, 2001, **228**, 49–62.
- 87 B. Bohman, A. M. Weinstein, R. D. Phillips, R. Peakall and G. R. Flematti, *J. Nat. Prod.*, 2019, **82**, 1107–1113.
- 88 F. P. Schiestl, R. Peakall, J. G. Mant, F. Ibarra, C. Schulz, S. Franke and W. Francke, *Science*, 2003, **302**, 437–438.
- 89 S. Franke, F. Ibarra, C. M. Schulz, R. Twele, J. Poldy, R. A. Barrow, R. Peakall, F. P. Schiestl and W. Francke, *Proc. Natl. Acad. Sci. U. S. A.*, 2009, **106**, 8877–8882.
- 90 R. Peakall, D. Ebert, J. Poldy, R. A. Barrow, W. Francke, C. C. Bower and F. P. Schiestl, *New Phytol.*, 2010, **188**, 437–450.
- 91 F. P. Schiestl and R. Peakall, *Funct. Ecol.*, 2005, **19**, 674–680.
- 92 Z. Yin and J. S. Dickschat, *Nat. Prod. Rep.*, 2022, DOI: [10.1039/D2NP00020B](https://doi.org/10.1039/D2NP00020B).
- 93 J. Gershenzon and N. Dudareva, *Nat. Chem. Biol.*, 2007, **3**, 408–414.
- 94 N. H. Williams and W. M. Whitten, *Biol. Bull.*, 1983, **164**, 355–395.
- 95 S. Ramírez, R. L. Dressle and M. Ospina, *Biota Colomb.*, 2002, **3**, 7–118.
- 96 G. Gerlach and R. Schill, *Bot. Acta*, 1991, **104**, 385–391.
- 97 N. S. L. Albuquerque, P. Milet-Pinheiro, D. D. Cruz, D. M. A. F. Navarro and I. C. Machado, *Plant Biol.*, 2021, **23**, 719–727.
- 98 K. Brandt, S. Dötterl, S. R. Ramírez, F. Etl, I. C. Machado, D. M. do A. F. Navarro, D. Dobler, O. Reiser, M. Ayasse and P. Milet-Pinheiro, *Front. Ecol. Evol.*, 2021, **9**, 727471.
- 99 P. Milet-Pinheiro, D. M. Do Amaral Ferraz Navarro, S. Dötterl, A. T. Carvalho, C. E. Pinto, M. Ayasse and C. Schlindwein, *Phytochemistry*, 2015, **116**, 149–161.
- 100 E. R. Pansarin and M. C. E. Amaral, *Flora*, 2009, **204**, 238–249.
- 101 A.-K. Borg-Karlson, J. Tengö, I. Valterová, C. R. Unelius, T. Taghizadeh, T. Tolasch and W. Francke, *J. Chem. Ecol.*, 2003, **29**, 1–14.
- 102 H. Xu, B. Bohman, D. C. J. Wong, C. Rodriguez-Delgado, A. Scaffidi, G. R. Flematti, R. D. Phillips, E. Pichersky and R. Peakall, *Curr. Biol.*, 2017, **27**, 1867–1877.
- 103 B. Bohman, A. Karton, G. R. Flematti, A. Scaffidi and R. Peakall, *J. Chem. Ecol.*, 2018, **44**, 436–443.
- 104 W. M. Whitten, H. G. Hills and N. H. Williams, *Phytochemistry*, 1988, **27**, 2759–2760.
- 105 D. L. P. Schorkopf, L. Mitko and T. Eltz, *J. Chem. Ecol.*, 2011, **37**, 953.
- 106 S. Dötterl, D. Burkhardt, B. Weißbecker, A. Jürgens, S. Schütz and A. Mosandl, *J. Chromatogr. A*, 2006, **1113**, 231–238.
- 107 X.-H. Jin, Z.-X. Ren, S.-Z. Xu, H. Wang, D.-Z. Li and Z.-Y. Li, *BMC Plant Biol.*, 2014, **14**, 63.
- 108 H. Jiang, J.-J. Kong, H.-C. Chen, Z.-Y. Xiang, W.-P. Zhang, Z.-D. Han, P.-C. Liao and Y.-I. Lee, *New Phytol.*, 2020, **227**, 1213–1221.
- 109 N. Lindquist, M. A. Battiste, W. Mark Whitten, N. H. Williams and L. Strekowski, *Phytochemistry*, 1985, **24**, 863–865.
- 110 P. Milet-Pinheiro and G. Gerlach, *Perspect. Plant Ecol. Evol. Systemat.*, 2017, **27**, 23–34.
- 111 K. Brandt, S. Dötterl, R. Fuchs, D. M. A. F. Navarro, I. C. S. Machado, D. Dobler, O. Reiser, M. Ayasse and P. Milet-Pinheiro, *J. Chem. Ecol.*, 2019, **45**, 464–473.
- 112 F. P. Schiestl and F. Glaser, *Alpine Bot.*, 2012, **122**, 1–9.



- 113 P. Milet-Pinheiro, D. M. do A. F. Navarro, S. Dötterl, A. T. Carvalho, C. E. Pinto, M. Ayasse and C. Schindwein, *Phytochemistry*, 2015, **116**, 149–161.
- 114 A.-K. Borg-Karlson, *Phytochemistry*, 1990, **29**, 1359–1387.
- 115 F. P. Schiestl and M. Ayasse, *Oecologia*, 2001, **126**, 531–534.
- 116 F. P. Schiestl and M. Ayasse, *Behav. Ecol. Sociobiol.*, 2000, **48**, 303–307.
- 117 S. D. Johnson, A. Ellis and S. Dötterl, *Am. J. Bot.*, 2007, **94**, 47–55.
- 118 H. Bouwmeester, R. C. Schuurink, P. M. Bleeker and F. Schiestl, *Plant J.*, 2019, **100**, 892–907.
- 119 M. Huang, A. M. Sanchez-Moreiras, C. Abel, R. Sohrabi, S. Lee, J. Gershenzon and D. Tholl, *New Phytol.*, 2012, **193**, 997–1008.
- 120 T. van der Niet, A. Jürgens and S. D. Johnson, *Plant Biol.*, 2015, **17**, 226–237.
- 121 M. G. Balducci, D. J. Martins and S. D. Johnson, *Plant Syst. Evol.*, 2019, **305**, 765–775.
- 122 H. Braunschmid and S. Dötterl, *Front. Plant Sci.*, 2020, **11**, 584081.
- 123 K. Gross, M. Sun and F. P. Schiestl, *PLoS One*, 2016, **11**, e0147975.
- 124 C. Micheneau, J. Fournel, B. H. Warren, S. Hugel, A. Gauvin-Bialecki, T. Pailler, D. Strasberg and M. W. Chase, *Ann. Bot.*, 2010, **105**, 355–364.
- 125 K. E. Steiner, R. Kaiser and S. Dötterl, *Am. J. Bot.*, 2011, **98**, 1663–1679.
- 126 A. Felemban, J. Braguy, M. D. Zurbriggen and S. Al-Babili, *Front. Plant Sci.*, 2019, **10**, 1168.
- 127 L. Aloum, E. Alefishat, A. Adem and G. Petroianu, *Molecules*, 2020, **25**, 5822.
- 128 N. H. Williams and C. H. Dodson, *Evolution*, 1972, **26**, 84–95.
- 129 H. Braunschmid, R. Guilhot and S. Dötterl, *Diversity*, 2021, **13**, 1–15.
- 130 L. Dormont, R. Delle-Vedove, J.-M. Bessière and B. Schatz, *Phytochemistry*, 2014, **100**, 51–59.
- 131 J. Jersáková, J. Spaethe, M. Streinzer, J. Neumayer, H. Paulus, S. Dötterl and S. D. Johnson, *Bot. J. Linn. Soc.*, 2016, **180**, 269–294.
- 132 T. Van der Niet, A. Jürgens and S. D. Johnson, *S. Afr. J. Bot.*, 2010, **76**, 726–738.
- 133 K. Brandt, I. C. Machado, D. M. do Amaral Ferraz Navarro, S. Dötterl, M. Ayasse and P. Milet-Pinheiro, *AoB Plants*, 2020, **12**, plaa030.
- 134 A. Flach, R. C. Dondon, R. B. Singer, S. Koehler, M. D. C. E. Amaral and A. J. Marsaioli, *J. Chem. Ecol.*, 2004, **30**, 1045–1056.
- 135 M. C. Hetherington-Rauth and S. R. Ramírez, *Ann. Bot.*, 2016, **118**, 135–148.
- 136 S. D. Johnson, *Plant Syst. Evol.*, 2005, **251**, 153–160.
- 137 G. C. Kite and G. A. Salazar, *Rev. Mex. Biodivers.*, 2008, **79**, 153–157.
- 138 J. Li, G.-F. Zhu and Z.-H. Wang, *J. Essent. Oil-Bear. Plants*, 2017, **20**, 385–394.
- 139 A. Tava, R. Cecotti and M. Confalonieri, *J. Essent. Oil Res.*, 2012, **24**, 39–44.
- 140 T. van der Niet, R. J. Cozien, B. Bytebier and S. D. Johnson, *Plant Syst. Evol.*, 2017, **303**, 387–401.
- 141 C. C. Salzmänn, S. Cozzolino and F. P. Schiestl, *Plant Biol.*, 2007, **9**, 720–729.
- 142 S. D. Johnson, M. G. Balducci and A. Shuttleworth, *Biol. J. Linn. Soc.*, 2020, **129**, 213–226.
- 143 F. K. Huber, R. Kaiser, W. Sauter and F. P. Schiestl, *Oecologia*, 2005, **142**, 564–575.
- 144 C. E. P. Nunes, M. F. G. V. Peñafior, J. M. S. Bento, M. J. Salvador and M. Sazima, *Oecologia*, 2016, **182**, 933–946.
- 145 K. H. Tan and R. Nishida, *J. Insect Sci.*, 2012, **12**, 56.
- 146 R. Steen, H. R. Norli and G. Thöming, *Arthropod-Plant Interact.*, 2019, **13**, 581–592.
- 147 R. Nishida, K.-H. Tan, S.-L. Wee, A. K.-W. Hee and Y.-C. Toong, *Biochem. Syst. Ecol.*, 2004, **32**, 245–252.
- 148 R. Nishida, O. Iwahashi and K. H. Tan, *J. Chem. Ecol.*, 1993, **19**, 713–722.
- 149 T. Katte, K. H. Tan, Z.-H. Su, H. Ono and R. Nishida, *Appl. Entomol. Zool.*, 2020, **55**, 55–64.
- 150 T. Keng-Hong and R. Nishida, *J. Chem. Ecol.*, 2005, **31**, 497–507.
- 151 M. Nakahira, H. Ono, S. L. Wee, K. H. Tan and R. Nishida, *Biochem. Syst. Ecol.*, 2018, **81**, 86–95.
- 152 K.-H. Tan, J. J. Vermeulen, T. Katte, H. Ono and R. Nishida, *Arthropod-Plant Interact.*, 2021, **15**, 447–455.
- 153 K. H. Tan and R. Nishida, *Biochem. Syst. Ecol.*, 2007, **35**, 334–341.
- 154 K.-H. Tan and R. Nishida, *J. Chem. Ecol.*, 2000, **26**, 533–546.
- 155 S. D. Johnson and N. Hobbhahn, *S. Afr. J. Bot.*, 2010, **76**, 739–748.
- 156 P. Milet-Pinheiro, J. B. F. Silva, D. M. A. F. Navarro, I. C. S. Machado and G. Gerlach, *Plant Species Biol.*, 2018, **33**, 158–163.
- 157 T. Van Der Niet, D. M. Hansen and S. D. Johnson, *Ann. Bot.*, 2011, **107**, 981–992.
- 158 R. Delle-Vedove, N. Juillet, J.-M. Bessire, C. Grison, N. Barthes, T. Pailler, L. Dormont and B. Schatz, *Phytochemistry*, 2011, **72**, 735–742.
- 159 L. J. Nielsen and B. L. Møller, *Phytochemistry*, 2015, **120**, 3–18.
- 160 A. Jürgens, U. Glück, G. Aas and S. Dötterl, *Bot. J. Linn. Soc.*, 2014, **175**, 624–640.
- 161 F. Mas, A. Harper, R. Horner, T. Welsh, P. Jaksons and D. M. Suckling, *J. Sci. Food Agric.*, 2018, **98**, 4445–4453.
- 162 S. Dötterl, A. Jürgens, K. Seifert, T. Laube, B. Weißbecker and S. Schütz, *New Phytol.*, 2006, **169**, 707–718.
- 163 F. B. Mortzfeld, C. Hashem, K. Vranková, M. Winkler and F. Rudroff, *Biotechnol. J.*, 2020, **15**, 2000064.
- 164 B. Bohman, R. D. Phillips, M. H. M. Menz, B. W. Berntsson, G. R. Flematti, R. A. Barrow, K. W. Dixon and R. Peakall, *New Phytol.*, 2014, **203**, 939–952.
- 165 B. Bohman, L. Jeffares, G. Flematti, R. D. Phillips, K. W. Dixon, R. Peakall and R. A. Barrow, *Org. Lett.*, 2012, **14**, 2576–2578.



- 166 B. Bohman, L. Jeffares, G. Flematti, L. T. Byrne, B. W. Skelton, R. D. Phillips, K. W. Dixon, R. Peakall and R. A. Barrow, *J. Nat. Prod.*, 2012, **75**, 1589–1594.
- 167 M. Iranshahi, *J. Essent. Oil Res.*, 2012, **24**, 393–434.
- 168 B. Bohman, R. D. Phillips, G. R. Flematti, R. A. Barrow and R. Peakall, *Angew. Chem., Int. Ed.*, 2017, **56**, 8455–8458.
- 169 B. Bohman, R. D. Phillips, G. R. Flematti and R. Peakall, *Fitoterapia*, 2018, **126**, 78–82.
- 170 A. M. Weinstein, B. Bohman, G. R. Flematti and R. D. Phillips, *Plants*, 2022, **11**, 260.
- 171 D. D. Gervasi and F. P. Schiestl, *Nat. Commun.*, 2017, **8**, 1–8.
- 172 P. Zito, S. Rosselli, M. Bruno, A. Maggio and M. Sajeve, *Plant Signaling Behav.*, 2019, **14**, 1552056.
- 173 N. Joffard, B. Buatois and B. Schatz, *Sci. Nat.*, 2016, **103**, 77.
- 174 N. P. Ferreira, L. U. R. Chiavelli, C. R. Savaris, S. M. Oliveira, D. L. Lucca, M. A. Milaneze-Gutierrez, R. T. Faria and A. M. Pomini, *Biochem. Syst. Ecol.*, 2019, **86**, 103918.
- 175 B. Bohman, A.-K. Borg-Karlson and R. Peakall, in *Biology of Plant Volatiles*, ed. E. Pichersky and N. Dudareva, CRC Press, 2020, pp. 39–56.
- 176 L. Jirovetz, J. E. Gonzalez, G. Silvera, A. Nikiforov and A. Woidich, *J. Essent. Oil Res.*, 1992, **4**, 435–438.
- 177 H. R. Buser, H. Arn, P. Guerin and S. Rauscher, *Anal. Chem.*, 1983, **55**, 818–822.
- 178 E. Dunkelblum, S. H. Tan and P. J. Silk, *J. Chem. Ecol.*, 1985, **11**, 265–277.
- 179 T. W. Mitchell, H. Pham, M. C. Thomas and S. J. Blanksby, *J. Chromatogr. B*, 2009, **877**, 2722–2735.
- 180 L. Xu, H. Guan, L. Liu, S. Mao, J. Feng, Z. Su and L. Liu, *J. Chromatogr. A*, 2022, **1672**, 463009.
- 181 N. Shimizu, N. Mori and Y. Kuwahara, *Biosci., Biotechnol., Biochem.*, 1999, **63**, 1478–1480.
- 182 B. Bohman, A. Kartton, R. C. M. Dixon, R. A. Barrow and R. Peakall, *J. Chem. Ecol.*, 2016, **42**, 17–23.
- 183 P. A. Aronov and B. D. Hammock, *Mass Spectrom. Rev.*, 2007, **26**, 51–78.
- 184 R. Spicer, R. M. Salek, P. Moreno, D. Cañueto and C. Steinbeck, *Metabolomics*, 2017, **13**, 106.
- 185 S. H. Kang and T. Kitahara, *Nat. Prod. Lett.*, 1996, **9**, 13–20.
- 186 T. Kitahara, K. S. Hyun and M. Santelli, *Nat. Prod. Lett.*, 1994, **5**, 157–164.
- 187 T. Kitahara, M. Inoue, S. Tamogami and R. Kaiser, *Tetrahedron*, 1996, **52**, 1487–1492.
- 188 Y. Masuzawa, S. Tamogami and T. Kitahara, *Nat. Prod. Lett.*, 1999, **13**, 239–246.
- 189 J. Poldy, R. Peakall and R. A. Barrow, *Org. Biomol. Chem.*, 2009, **7**, 4296–4300.
- 190 B. Bohman, B. Berntsson, R. C. M. Dixon, C. D. Stewart and R. A. Barrow, *Org. Lett.*, 2014, **16**, 2787–2789.
- 191 T. Eltz, *J. Chem. Ecol.*, 2006, **32**, 907–915.
- 192 D. Goulson, J. W. Chapman and W. O. H. Hughes, *J. Insect Behav.*, 2001, **14**, 669–678.
- 193 R. D. Phillips, N. Reiter and R. Peakall, *Ann. Bot.*, 2020, **126**, 345–362.

