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Plant essential oils and formamidines as insecticides/acaricides: what are the molecular targets?

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Abstract – The parasitic mite *Varroa destructor* is the main cause of the severe reduction in beekeeping during the last few decades. Therefore, efforts have been made to develop chemical treatments against the parasite. In the past, synthetic products were preferentially used to combat *Varroa* mites. Nowadays, mainly plant essential oils and organic acids are applied because they are safer and impose less unfavorable effects on the environment. Essential oils contain mixtures of mostly volatile and odorous terpenoid constituents. The molecular targets of these substances are tyramine and/or octopamine receptors that control and modulate vital functions ranging from metabolism to behavior. Disturbing the native function of these receptors in the mite results in deleterious effects in this parasite. This overview considers not only tyramine and octopamine receptors but also other potential targets of essential oils including ionotropic GABA_A receptors, TRP type ion channels, and acetylcholinesterase.

GABA / G protein-coupled receptor / octopamine / thymol / tyramine

1. PLANT ESSENTIAL OILS— AN INTRODUCTION

The availability of environmentally safe and efficient chemicals against arthropod pests is an important aspect for veterinary and crop industries. At present, chemical compounds that rapidly deliver their insecticidal effect are preferentially used to achieve a timely reduction of pests and ectoparasites on their respective hosts. This condition is fulfilled by compounds that modify the activity of voltage-gated and/or ligand-gated ion channels in the central nervous system (CNS) and by compounds that impair the activity of neuronal enzymes such

as acetylcholinesterase (AChE) or that manipulate the overall metabolic status of the parasite (for a review, see Isman 2006). Substances such as synthetic pyrethroids inhibit voltage-gated sodium channels that are important for the firing of neuronal action potentials. Phenylpyrazolines or polychloroalkanes (e.g., lindane) block the activity of GABA_A channels, important members of the ligand-gated ion channel family involved in inhibitory signal generation in neurons. Organophosphates inhibit the enzymatic activity of AChE, an enzyme controlling the concentration of the excitatory neurotransmitter acetylcholine (ACh) in the synaptic cleft. Isoflavonoids such as rotenone poison mitochondria and thus cause the breakdown of cellular energy production. Metabotropic (G protein-coupled) octopamine receptors are the target of formamidine pesticides such as amitraz which manipulate

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cellular signal transduction processes. Despite the efficient and wide use of these substances, they seem to harbor certain disadvantages. (1) Parasites can develop resistance to the chemicals, probably because the generation times of the arthropod pests are short. (2) Cross-reactivity of such substances might also be significant in beneficial insects such as the honeybee. In addition, these substances might cause undesired allergic or toxic effects in humans. For such reasons, there is a continual demand for compounds that, on the one hand, efficiently reduce parasites or pests and, on the other hand, are safe for the host, the environment, and non-target species, including man, that come into contact with such insecticides/acaricides. Over the years, plant essential oils have been considered as an attractive alternative that might fulfill these criteria.

Plant essential oils are complex mixtures of odorous substances obtained from botanical raw materials by water vapor extraction, dry distillation, or mechanical treatment without heating (Isman 2006; Vigan 2010). Because of their odorous character, plant essential oils are widely used as fragrances and flavorings in perfume and food industries. To identify and to further characterize the individual components of essential oils, either separation by gas chromatography or HPLC has to be performed. These analyses have identified monoterpenes, sesquiterpenes, and related aromatic compounds as the main constituents of plant essential oils. Terpenoids are derived from five-carbon isoprene units. Coupling of two isoprene units leads to ten-carbon structures known as monoterpenoids, whereas coupling of three isoprene units leads to sesquiterpenoids (Table I). Terpenoids can be linear (acyclic) or contain cyclic structures (Table I). Further biochemical modifications by endogenous plant enzymes can cause oxidation, rearrangement, or additional cyclization (Table I) that increases the variability of essential oil constituents (Chappell 1995; Holstein and Hohl 2004). At approximately 90%, the mono- and sesquiterpenoids outnumber the other constituents of plant essential oils. Nevertheless, some aromatic compounds are worth mentioning (Table I).

Similar to their heterogeneous composition, plant essential oils display a broad spectrum of biological activity. They can be toxic for insects and microbes, but can also act as insect antifeedants and repellents (Anthony et al. 2005; Isman 2006; Nerio et al. 2010; Vigan 2010). The toxic effect of essential oils is most likely mediated by neurological mechanisms. Indeed, plant essential oils have been suggested to exert their bioactivity by interacting with various molecular targets including tyramine and octopamine receptors (Enan 2001, 2005a, b; Price and Berry 2006), ionotropic GABA receptors (Priestley et al. 2003; Tong and Coats 2010), and AChE (Grundy and Still 1985; Ryan and Byrne 1988; Keane and Ryan 1999). As previously mentioned, octopamine receptors have also been identified as a target site of the formamidine class of insecticides (Evans and Gee 1980; Gole et al. 1983; Downer et al. 1985; Dudai et al. 1987; Chen et al. 2007).

In mammalian tissues, *p*-tyramine, β -phenylethylamine, tryptamine, and octopamine are present at very low (nanomolar) concentrations and are therefore referred to as “trace amines.” Trace amine-associated receptors (TAARs) have recently been discovered in mammals (for a review, see Zucchi et al. 2006). It is worth emphasizing, however, that thus far, only two members of the TAAR family have been shown to be responsive to trace amines. In addition, mammalian TAARs are not closely related to arthropod tyramine and octopamine receptors. Thus, it appears that in mammals, TAARs evolved independently from the arthropod tyramine and octopamine receptors and acquired the ability to interact with different amines, including the decarboxylated thyroid hormone derivatives known as thyronamines, several volatile amines, and possibly other as yet unidentified endogenous compounds (for a review, see Zucchi et al. 2006). Therefore, arthropod tyramine and octopamine receptors might emerge as promising targets for insecticides/acaricides (e.g., plant essential oil components) with no or low toxicity in vertebrates.

Table I. Naturally occurring essential oil components and synthetic substances mentioned in the text.

Compound	Molecular formula	Substance class	Sources (examples)
Linalool [licareol=(<i>R</i>)-(-)-linalool; coriandrol=(<i>S</i>)-(+)-linalool]	C ₁₀ H ₁₈ O	Monoterpenoid alcohol, linear	<i>Coriandrum sativum</i> (coriander) <i>Ocimum basilicum</i> (basil) <i>Citrus × sinensis</i> (sweet orange)
Citral [geranial=citral A; neral=citral B]	C ₁₀ H ₁₆ O	Monoterpenoid aldehyde, linear	<i>Backhousia citriodora</i> (lemon myrtle) <i>Cymbopogon</i> spp. (lemongrass) <i>Leptospermum liversidgei</i> (lemon tea tree)
Menthol	C ₁₀ H ₂₀ O	Monoterpenoid alcohol, monocyclic, unsaturated	<i>Mentha × piperita</i> (peppermint)
α-Terpineol	C ₁₀ H ₁₈ O	Monoterpenoid alcohol, monocyclic, unsaturated	<i>Melaleuca leucadendra</i> (cajuput oil) <i>Pinus</i> spp. (pine oil) <i>Citrus × aurantium</i> (petitgrain oil)
Carveol	C ₁₀ H ₁₆ O	Monoterpenoid alcohol, monocyclic, unsaturated	<i>cis</i> -(-)-Carveol: <i>Mentha spicata</i> (spearmint)
Pulegone	C ₁₀ H ₁₆ O	Monoterpenoid ketone, monocyclic	<i>Nepeta cataria</i> (catnip) <i>Mentha × piperita</i> (peppermint) <i>Mentha pulegium</i> (pennyroyal)
<i>R</i> -(-)-Carvone, <i>S</i> -(+)-carvone	C ₁₀ H ₁₄ O	Monoterpenoid ketone, monocyclic, unsaturated	<i>S</i> -(+)-carvone <i>Carum carvi</i> (caraway) <i>Anethum graveolens</i> (dill) <i>R</i> -(-)-carvone <i>Mentha spicata</i> (spearmint)
α-Terpinene, γ-terpinene	C ₁₀ H ₁₆	Monoterpenoid, monocyclic, unsaturated	<i>Melaleuca alternifolia</i> (tea tree oil) <i>Citrus × limon</i> (bergamot oil)
<i>p</i> -Cymene	C ₁₀ H ₁₄	Monoterpenoid, monocyclic, aromatic	<i>Cuminum cyminum</i> (cumin) <i>Chenopodium ambrosioides</i> (epazote) <i>Peumus boldus</i> (boldo)
Carvacrol	C ₁₀ H ₁₄ O	Monoterpenoid phenol	<i>Origanum vulgare</i> (oregano) <i>Thymus</i> spp. (thyme) <i>Lepidium</i> spp. (peppermint)
Thymol	C ₁₀ H ₁₄ O	Monoterpenoid phenolic derivative of cymene	<i>Thymus</i> spp. (thyme) <i>Trachyspermum copticum</i> (ajwain) <i>Origanum vulgare</i> (oregano)
Anethole (<i>para</i> -methoxyphenylpropene,	C ₁₀ H ₁₂ O	Monoterpenoid ether, aromatic, unsaturated	<i>Pimpinella anisum</i> (anise) <i>Foeniculum vulgare</i> (fennel)

Table 1 continued.

Compound	Molecular formula	Substance class	Sources (examples)
<i>p</i> -propenylanisole, isoeustragole			<i>Syzygium anisatum</i> (anise myrtle)
Eugenol	C ₁₀ H ₁₂ O ₂	Monoterpenoid, methoxy-phenol, unsaturated	<i>Syzygium aromaticum</i> (clove oil) <i>Pimenta dioica</i> (allspice) <i>Myristica fragrans</i> (nutmeg)
(+)-Camphor, (-)-camphor	C ₁₀ H ₁₆ O	Monoterpenoid ketone, bicyclic	(+)-Camphor <i>Cinnamomum camphora</i> (camphor laurel) <i>Rosmarinus officinalis</i> (rosemary) (-)-Camphor <i>Tanacetum parthenium</i> (feverfew) <i>Matricaria recutita</i> (German chamomile)
α-Pinene; β-pinene	C ₁₀ H ₁₆	Monoterpenoid, bicyclic, unsaturated	<i>Pinus</i> spp. (pine resin) <i>Rosmarinus officinalis</i> (rosemary)
(+)-Borneol	C ₁₀ H ₁₈ O	Monoterpenoid ketone, bicyclic	<i>Cinnamomum camphora</i> (camphor oil) <i>Coriandrum sativum</i> (coriander) <i>Tanacetum vulgare</i> (tansy)
Bornyl acetate	C ₁₂ H ₂₀ O ₂	Acetic acid ester of borneol	<i>Abies</i> spp. (fir) <i>Pinus</i> spp. (pine) <i>Picea</i> spp. (spruce)
Eucalyptol (1,8-cineol, 1,8-cineole, cineol, cineole)	C ₁₀ H ₁₈ O	Monoterpenoid, oxa-bicyclic	<i>Eucalyptus</i> spp. (eucalyptus oil) <i>Cinnamomum camphora</i> (camphor laurel) <i>Laurus nobilis</i> (bay leaf)
Farnesol	C ₁₅ H ₂₆ O	Sesquiterpene, linear	<i>Cymbopogon</i> spp. (lemongrass, citronella oil) <i>Citrus × aurantium</i> (bitter orange, neroli oil) <i>Polianthes tuberosa</i> (tuberose)
Cinnamic alcohol	C ₉ H ₁₀ O	Aromatic, unsaturated alcohol	<i>Liquidambar orientalis</i> (storax) <i>Myroxylon balsamum</i> (balsam Peru) <i>Cinnamomum</i> spp. (cinnamon)
Cinnamaldehyde	C ₉ H ₈ O	Aromatic, unsaturated aldehyde	<i>Cinnamomum</i> spp. (cinnamon)
Capsaicin	C ₁₈ H ₂₇ NO ₃	Capsaicinoid, 8-methyl-	<i>Capsicum</i> spp. (chili pepper)

Table 1 continued.

Compound	Molecular formula	Substance class	Sources (examples)
Chlordimeform	C ₁₀ H ₁₃ ClN ₂	<i>N</i> -vanilyl-6-nonenamide, aromatic Amidine, organochloride	Synthetic
Demethylchlordimeform	C ₉ H ₁₁ ClN ₂	Amidine, organochloride, aromatic	Synthetic
Amitraz	C ₁₉ H ₂₃ N ₃	Amidine, triazapentadiene, aromatic	Synthetic

For each substance, the molecular formula, characteristic structural properties, and a selection of natural sources synthesizing the compound are given

2. USE OF PLANT ESSENTIAL OILS FOR *VARROA* TREATMENT OF HONEYBEE COLONIES

The honeybee, *Apis mellifera*, pollinates about 80% of crop and wild plants. This makes it the fourth most important farm animal in Germany (after cattle, pigs, and poultry). Hence, it is alarming that the honeybee stocking density has rapidly declined over the last few decades. One reason for this dramatic reduction in beekeeping is the parasitic mite *Varroa destructor*, which was introduced into Germany in the 1970s. *V. destructor* can only reproduce in a honeybee colony, precisely in sealed brood cells. It attaches to the body of the larva, pupa, or adult bee and weakens its host by sucking hemolymph (for reviews, see Rademacher 1990; Sammataro et al. 2000; Rosenkranz et al. 2010). The mite acts as a vector and spreads RNA viruses such as deformed wing virus to the bee (for reviews, see Genersch and Aubert 2010; Le Conte et al. 2010). A significant mite infestation will lead to the death of the honeybee colony, usually during the summer or winter months. *V. destructor* is the parasite with the most pronounced economic impact on the beekeeping industry. It is a major contributing factor to colony collapse disorder (CCD) and to non-CCD winter losses, which constitute severe problems in beekeeping worldwide (Vanengelsdorp et al. 2009; Guzman-Novoa et al. 2010; for a review, see Le Conte et al. 2010). Honeybees in CCD colonies are characterized by higher pathogen loads and are

co-infected with a greater number of pathogens compared with control populations, suggesting an interaction between pathogens and other stress factors in CCD and a possible legacy effect of *Varroa* parasitism (Vanengelsdorp et al. 2009).

Numerous efforts have been made to develop chemical treatments against *V. destructor* (Mutinelli and Rademacher 2003; Rosenkranz et al. 2010). Nowadays, products based on organic acids (e.g., formic acid and oxalic acid; Rademacher and Imdorf 2004; Rademacher and Harz 2006; Calderone 2010), essential oils (e.g., thymol; Imdorf et al. 1995, 1999; Rademacher and Radtke 2001; Floris et al. 2004), pyrethroids (e.g., fluvalinate and flumethrin), organophosphates (e.g., coumaphos; Milani and Lob 1998), and formamidines (e.g., amitraz; Floris et al. 2001) are used to treat infected colonies. Essential oils (and formamidines) most probably exert their therapeutic effects by binding to tyramine and/or octopamine receptors. Amongst others, thymol (e.g., Apiguard and Thymovar) and combinations of thymol and other essential oils such as eucalyptol, camphor, and menthol (e.g., Api Life VAR) are also used for *Varroa* treatment (Imdorf et al. 1999) and have been approved as veterinary drugs in many European countries. Calderone and Spivak (1995) found that a blend of natural products similar to Api Life VAR provided more than 98.5% control of *V. destructor*. In a recent study, Ghasemi et al. (2011) determined the fumigant toxicity of essential oils taken from *Thymus kotschyanus*,

Ferula assa-foetida, and *Eucalyptus camaldulensis* against *V. destructor* and *A. mellifera*. Interestingly, whereas the essential oil of *T. kotschyanus* is the most potent fumigant for *V. destructor*, it has the lowest insecticidal activity against *A. mellifera* (Ghasemi et al. 2011). Noteworthy is that menthol is used to treat hives infested with the tracheal mite, *Acarapis woodi*, which is an internal parasite of honeybees. On the one hand, essential oils have not yet led to resistant mite populations, but on the other hand, they are not consistently highly active in reducing mite populations in all situations (Le Conte et al. 2010). In addition, treatments using thymol as their active substance are suspected to have adverse effects on honeybee colonies. Interestingly, honeybee responses to Apiguard treatment change with honeybee age (Mondet et al. 2011). While 2-day-old bees respond neutrally to Apiguard, foragers appear to be repelled by Apiguard, but can become habituated to the treatment (Mondet et al. 2011).

3. OCTOPAMINE AND TYRAMINE RECEPTORS AS MOLECULAR TARGETS OF PLANT ESSENTIAL OILS AND FORMAMIDINE PESTICIDES

Biogenic amines are small organic compounds that act as neurotransmitters, neuromodulators, and/or neurohormones in vertebrates and in invertebrates. They represent an important group of messenger substances and mediate their diverse effects by binding to membrane receptors that belong to the large gene family coding for G protein-coupled receptors. In arthropods, the group of biogenic amine messengers consists of five members: dopamine, tyramine, octopamine, serotonin, and histamine. Receptor activation leads to changes in the concentration of intracellular second messengers (e.g., cAMP or $\text{InsP}_3/\text{Ca}^{2+}$). For each amine, multiple receptor subtypes exist that couple to various intracellular signaling pathways in a receptor-specific manner (for reviews, see Blenau and Baumann 2001, 2003; Evans and Maqueira 2005; Scheiner et al. 2006; Hauser et al. 2006;

Verlinden et al. 2010a; Blenau and Thamm 2011). In the honeybee, three dopamine receptors (AmDOP1: Blenau et al. 1998; Mustard et al. 2003; AmDOP2: Humphries et al. 2003; Mustard et al. 2003; AmDOP3: Beggs et al. 2005), one tyramine receptor (AmTYR1: Blenau et al. 2000; Mustard et al. 2005), one octopamine receptor (AmOA1: Grohmann et al. 2003), and two serotonin receptors (Am5-HT_{1A}: Thamm et al. 2010; Am5-HT₇: Schlenstedt et al. 2006) have been characterized. Aminergic receptors have also been characterized in locusts (*Locusta migratoria*: Vanden Broeck et al. 1995; *Schistocerca gregaria*: Verlinden et al. 2010b) and the American cockroach (*Periplaneta americana*; Bischof and Enan 2004; Rotte et al. 2009; Troppmann et al. 2010). These hemimetabolous insects not only are pest insects but also serve as model organisms for basic research in neurobiology and pharmacology. The successful completion of various insect genome sequencing projects has led to the annotation of additional receptor genes by bioinformatics (*Drosophila melanogaster*: Brody and Cravchik 2000; *Anopheles gambiae*: Hill et al. 2002; *A. mellifera*: Hauser et al. 2006; *Tribolium castaneum*: Hauser et al. 2008). Unfortunately, knowledge of aminergic receptors in mites remains limited. So far, only one tyramine receptor (Baxter and Barker 1999) and one serotonin receptor (Chen et al. 2004) of the cattle tick, *Boophilus microplus*, and two dopamine receptors (Meyer et al. 2011) of the blacklegged tick, *Ixodes scapularis*, have been molecularly identified. To date, no molecular data on aminergic receptors are available for *V. destructor*.

The neurotoxic activity of essential oils and/or their purified constituents (eugenol, α -terpineol, cinnamic alcohol) against various insect species is probably attributable to binding to tyramine and octopamine receptors (Enan 2001). Octopamine and essential oil constituents (10 nM) have been found to cause a significant increase in intracellular cAMP concentration ($[\text{cAMP}]_i$) in the cotton bollworm, *Helicoverpa armigera* (Kostyukovsky et al. 2002). The octopamine receptor antagonist phentolamine effectively inhibits the increase in $[\text{cAMP}]_i$

induced by essential oil treatment. These results support the hypothesis that native octopamine receptors are targets of essential oil constituents (Kostyukovsky et al. 2002).

This hypothesis has subsequently been supported by evidence from cloned and heterologously expressed receptors. Enan (2005a) has shown that the toxicity rank order of *p*-cymene, thymol, carvacrol, α -terpineol, and *R*(-)-carvone for *D. melanogaster* correlates with their binding affinity to a heterologously expressed tyramine receptor (TyrR). The main route by which this receptor acts causes a reduction in $[cAMP]_i$. When heterologously expressed in *D. melanogaster* S2 cells, the activation of the tyramine receptor also causes an increase in the intracellular Ca^{2+} concentration (Enan 2005a). Interestingly, two compounds, i.e., thymol and carvacrol, are not toxic for a tyramine receptor mutant (TyrR^{neo30}; Enan 2005a). In this mutant, the insertion of a P element abolishes the expression of a functional receptor protein (Cooley et al. 1988). Together, these results suggest that thymol and carvacrol mediate their toxic effects via tyramine receptors. Notably, the specific binding of essential oil components (eugenol, cinnamic alcohol, *trans*-anethole) to an octopamine receptor from *D. melanogaster* and its ortholog from the cockroach, *P. americana* (Pa o_{a1}), has also been reported (Enan 2005b).

As early as the middle of the 1980s, the formamidines demethylchlordimeform (DMCD), BTS-27271, and amitraz were shown to mimic the action of octopamine in elevating adenylyl cyclase activity in the nervous tissue of *P. americana* (Gole et al. 1983; Downer et al. 1985). In *D. melanogaster*, however, DMCD and chlordimeform inhibit octopamine-stimulated adenylyl cyclase activity, whereas amitraz activates the enzyme (Dudai et al. 1987). Both octopamine and chlordimeform potentiate contractions of the locust slow extensor tibiae muscle (Evans and Gee 1980). Furthermore, chlordimeform and DMCD elicit light production in the firefly (*Photinus pyralis*) lantern, which is controlled by octopaminergic neurons (Hollingworth and Murdock 1980). All these results argue in favor

of formamidines interacting with octopamine receptors. Interestingly, some formamidines show an even higher affinity for octopamine receptors in the CNS of insects than the natural ligand octopamine itself (Hiripi et al. 1994; Roeder 1995). The interaction of formamidines with octopamine receptors results in hyperexcitability, abnormal behavior, paralysis, and death in insects (Dudai et al. 1987), and also in the detachment and mortality of parasitic acarines (Stone et al. 1974). Chlordimeform HCl (K-79) has been tested for the treatment of varroosis and has been shown to be highly toxic for *V. destructor*, but only slightly toxic for bees (Rademacher 1981; Wachendörfer et al. 1981). In veterinary medicine, amitraz is used to control ectoparasitic mites and insects (Hollingworth 1976). In some European countries, amitraz is applied in the form of special plastic strips that slowly release the agent to control *V. destructor* (Floris et al. 2001). In other countries, amitraz is applied as an aerosol or a spray. In this form, amitraz is highly unstable and becomes rapidly degraded into various partly poisonous metabolites. For this reason, the compound is not licensed for the treatment of honeybee colonies in Germany. Another side effect of amitraz treatment is that mites can become resistant to the chemical (Elzen et al. 2000; for a review, see Jonsson and Hope 2007). Notably, amino acid substitutions have been identified in the receptor sequence coding for a putative octopamine/tyramine receptor in the cattle tick, *B. microplus*, in two amitraz-resistant strains (Chen et al. 2007). These substitutions are absent in all amitraz-susceptible strains. The discovery of such mutations in amitraz-resistant ticks provides evidence for these amino acid residues being part of the pesticide-binding site of the wild-type receptor.

4. GABA_A RECEPTOR ION CHANNELS AS MOLECULAR TARGETS OF PLANT ESSENTIAL OILS

Fast neuronal signaling by the neurotransmitter γ -aminobutyric acid (GABA) is mediated by ionotropic GABA_A receptors. These are pentameric complexes with an integral GABA-gated

anion channel (for a review, see Buckingham et al. 2005). Insect GABA_A receptors are targets for several insecticides, such as dieldrin, lindane, BIDN, and fipronil, all of which act as antagonists (for a review, see Bloomquist 1996). The monoterpenoid thymol, applied in concentrations between 1 and 100 μM , potentiates the action of GABA at recombinant human GABA_A receptors with various subunit compositions and also at an insect (*D. melanogaster*) ionotropic GABA receptor (Priestley et al. 2003). The enhanced response to GABA is likely to be the result of a positive allosteric action of thymol (Priestley et al. 2003). The direct agonistic action of thymol has been observed at both mammalian and insect recombinant receptors between 100 and 500 μM (Priestley et al. 2003). Thymol does not appear to compete with other GABAergic ligands. Thus, the effect of thymol is possibly mediated by a binding site that is not yet characterized and that could represent a new avenue in insecticide research (Priestley et al. 2003). Recently, Tong and Coats (2010) evaluated the pharmacological action of five monoterpenoids (α -terpineol, carvacrol, linalool, pulegone, and thymol) on native insect GABA_A receptors from houseflies and American cockroaches using radiotracer methods. The binding of [³H]*t*-butylbicycloorthobenzoate ([³H]-TBOB) to membrane preparations of housefly heads was enhanced by carvacrol, pulegone, and thymol with EC₅₀ values of 48 μM , 432 μM , and 6 mM, respectively. Moreover, these three monoterpenoids at concentrations of 500 μM and 1 mM also significantly increased the GABA-induced ³⁶Cl⁻ uptake in membrane microsacs prepared from American cockroach ventral nerve cords. The authors concluded that carvacrol, pulegone, and thymol are all positive allosteric modulators at insect GABA_A receptors (Tong and Coats 2010). In contrast, α -terpineol and linalool showed little or no effect on both [³H]-TBOB binding and ³⁶Cl⁻ uptake assays (Tong and Coats 2010). In the blowfly, *Phaenicia sericata*, thymol reduced the frequency of flight muscle impulses and, consequently, inhibited wing beat frequency (Waliwitiya et al. 2010). The similarity of thymol and GABA actions suggests that thymol acts

centrally by mimicking or facilitating the action of GABA (Waliwitiya et al. 2010).

A closer look at the above data reveals that micromolar concentrations are needed for all described effects of essential oil components on GABA_A receptor channels. This makes the hypothesis of GABA_A receptors being responsible for the insecticidal activity of essential oils rather questionable.

5. ION CHANNELS OF THE TRP FAMILY ARE INHIBITED/ACTIVATED BY MONOTERPENOIDS

Transient receptor potential (TRP) channels are essential components of signaling cascades in sensory cells specialized to detect painful stimuli, such as heat, cold, and pressure, and chemicals, such as acids or irritants. The seven subfamilies of TRP channels are clustered into two groups, with five members in group 1, i.e., TRPC (“C” for canonical), TRPV (“V” for vanilloid), TRPM (“M” for melastatin), TRPN, and TRPA. Two subfamilies constitute group 2, i.e., TRPP (“P” for polycystic) and TRPML (“ML” for mucolipin; for a review, see Venkatachalam and Montell 2007). The *D. melanogaster* TRPC channels, TRP and TRP-like (TRPL), are the founding members of the TRP superfamily. In *D. melanogaster*, TRP channels are specifically expressed in photoreceptor cells of the complex eye. Upon illumination, TRP channels are activated and cause the depolarization of the photoreceptor cell (for a review, see Raghu and Hardie 2009). The thermoTRPs, which include TRPV1-4, TRPM8, and TRPA1, constitute a subgroup of TRP channels that are activated by either hot or cold temperatures and also by natural compounds evoking “hot” (e.g., capsaicin) or “cold” (e.g., menthol) sensations, respectively (Macpherson et al. 2006; Xu et al. 2006; Vogt-Eisele et al. 2007; Lee et al. 2008).

Recently, Parnas et al. (2009) have demonstrated, by patch clamp whole cell recording from *D. melanogaster* S2 and *D. melanogaster* photoreceptor cells, that carvacrol (500 μM), a known activator of the mammalian ther-

moTRPs, TRPV3, and TRPA1, acts as an inhibitor of native TRP channels and of both native and heterologously expressed TRPL channels. Furthermore, thymol (1 mM), eugenol (3 mM), cinnamaldehyde ($IC_{50}=1.6$ mM), menthol ($IC_{50}=1.81$ mM), and carveol ($IC_{50}=4.22$ mM), which are activators of mammalian TRPV3, have also been found to inhibit the TRPL channel (Parnas et al. 2009). Interestingly, high concentrations of borneol (5 mM) and camphor (10 mM) activate the TRPL channel (Parnas et al. 2009). Whether the interaction of millimolar concentrations of plant essential oil components with ion channels of the TRP family correlates with their insecticidal/acaricidal properties needs further validation.

6. INHIBITION OF ACETYLCHOLINESTERASE ACTIVITY BY MONOTERPENOIDS

ACh is an important neurotransmitter in both the CNS and the peripheral nervous system of many organisms including arthropods. Once released, ACh has a short half-life because of the enzymatic activity of AChE, which hydrolyzes ACh to choline and acetate in the synaptic cleft. Thus, AChE inhibitors (e.g., organophosphates or carbamates such as physostigmine or neostigmine) can be used to increase the ACh concentration and to prolong its functional activity in the nervous system.

As early as 1985, however, Grundy and Still demonstrated that (+)-pulegone and related monoterpenoids inhibit both housefly and Madagascar roach (*Gromphadorhina portentosa*) AChE activity in vitro (Grundy and Still 1985). In a pioneering study on “plant–insect co-evolution and inhibition of acetylcholinesterase,” monoterpenoids (i.e., eucalyptol, pulegone, linalool, citral, and (–)-bornyl acetate) were convincingly shown to inhibit AChE purified from the electric eel (Ryan and Byrne 1988). Similar to the results obtained for AChE from electric eel, the five monoterpenoids were found to be reversible competitive inhibitors of AChE purified from the brain of the waxmoth, *Galleria mellonella* (Keane and Ryan 1999).

The K_i values were: eucalyptol, 220 nM; pulegone, 32 μ M; linalool, 240 μ M; citral, 330 μ M; (–)-bornyl acetate, 510 μ M (Keane and Ryan 1999). In addition, eucalyptol inhibits AChE activity in homogenates obtained from head louse with an IC_{50} of 77 mM (Picollo et al. 2008). Leaf oils of *E. camaldulensis* and their constituents (e.g., eucalyptol, *p*-cymene, γ -terpinene) act as both fumigants and contact toxicants on termites, *Coptotermes formosanus*; this is most likely attributable to the inhibition of AChE activity (Siramon et al. 2009). β -Pinene ($K_i=2.8$ μ M) and menthol ($K_i=48$ μ M) are potent inhibitors of AChE from the rice weevil, *Sitophilus oryzae*, one of the main stored grain pests (Lee et al. 2001).

In conclusion, many terpenoids might act as AChE inhibitors, both in arthropods and in vertebrates (for a review, see Houghton et al. 2006), but at relatively high concentrations (see above; Kostyukovsky et al. 2002). Whether terpenoid toxicity is correlated with their ability to inhibit AChE activity needs further experimental evidence (Grundy and Still 1985; Lee et al. 2001).

7. PLANT ESSENTIAL OILS AS INSECTICIDES/ACARICIDES—AN OUTLOOK

Monoterpenoids and related aromatic compounds show a broad spectrum of actions in insects and mites, with tyramine and octopamine receptors being likely candidates for their molecular targets. In addition to affecting tyramine and/or octopamine receptors, some of the compounds show other modes of action. Thymol, for example, affects insect GABA_A receptors (Priestley et al. 2003; Tong and Coats 2010) and *D. melanogaster* TRP and TRPL channels (Parnas et al. 2009).

So far, nothing is known about the aminergic receptors of *V. destructor*. This is surprising as substances belonging to two different chemical classes used for *Varroa* treatment exert their actions by binding to aminergic receptors. To understand the mode of action of essential oils and formamidines, the molecular signature(s) of

tyramine and octopamine receptors expressed in *V. destructor* and other arthropod pests therefore need to be unraveled.

A major concern about compounds currently used for *Varroa* treatment is their (potential) toxicity for the bees. Hoppe (1990) examined 55 essential oils for mite and bee toxicity. After 72 h, 24 essential oils produced a mite mortality of more than 90%. However, of these 24 oils, only nine resulted in bee mortality below 10%! Consequently, when applying these compounds, special care has to be taken to use concentrations that are toxic for mites but that have no or only minor toxicity for bees. For example, concentrations of 5–15 µg thymol, 50–150 µg camphor, or 20–60 µg menthol per liter of air killed nearly 100% of the mites without a noticeable loss of bees (Imdorf et al. 1999).

By likewise testing compounds that are active on tyramine and/or octopamine receptors of *V. destructor* and orthologous receptors of the bee, dose–response curves for receptors expressed in both species can be determined. In this way, compounds or combinations of compounds and the relevant concentrations that are highly toxic for mites but non-toxic for bees can be identified with justifiable costs.

During the last few years, protein structure-based drug design has become a valuable and integral part of the discovery of active agents. This strategy is also applicable for arthropod tyramine and octopamine receptors once their primary structure is known. Huang et al. (2007) performed homology modeling for an octopamine receptor of the silkworm, *Bombyx mori* (BmOAR1). Subsequently, the octopamine molecule was manually docked into the potential binding site of the BmOAR1 receptor according to mutagenesis data (Huang et al. 2007). In order to understand the interaction of the cockroach Pa oa₁ receptor (Bischof and Enan 2004) with its ligands, a three-dimensional model of Pa oa₁ was computed and then its agonist binding site was identified, followed by docking studies (Hirashima and Huang 2008). These molecular docking studies have shown that ligands with certain substituents possibly act as potential Pa oa₁ agonists (Hirashima and

Huang 2008). Similar receptor models should be established for the tyramine and octopamine receptors of arthropod parasites (including *V. destructor*) and pests. These models can then be used to design new pharmaceutical leads for use as potent and relatively specific agonists/antagonists for tyramine and octopamine receptors, thereby acting as potential pest control agents.

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Huiles essentielles de plantes et formamidines, comme insecticides/acaricides: quelles sont leurs cibles moléculaires?

GABA / récepteur couplé à une protéine G / octopamine / thymol / tyramine

Ätherische Öle und Formamidine als Insektizide/Akarizide. Welches sind die molekularen Wirkorte?

GABA / G Protein-gekoppelter Rezeptor / Octopamin / Thymol / Tyramin

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