Title:

Intrasexual cuticular hydrocarbon dimorphism in a wasp sheds light on hydrocarbon biosynthesis genes in Hymenoptera

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Abstract:

Cuticular hydrocarbons (CHCs) cover the cuticle of insects and serve as desiccation barrier and for chemical communication. While the main enzymatic steps of CHC biosynthesis are well understood, few of the underlying genes have been identified. Here we show how exploitation of intrasexual CHC dimorphism in a mason wasp, *Odynerus spinipes*, in combination with whole-genome sequencing and comparative transcriptomics facilitated identification of such genes. RNAi-mediated knockdown of twelve candidate gene orthologs in honey bees, *Apis mellifera*, confirmed nine genes impacting CHC profile composition. Most of them have predicted functions consistent with current knowledge of CHC metabolism. However, we found first-time evidence for a fatty acid amide hydrolase also influencing CHC profile composition. *In situ* hybridization experiments furthermore suggest trophocytes participating in CHC biosynthesis. Our results set the base for experimental CHC profile manipulation in Hymenoptera and imply that the evolutionary origin of CHC biosynthesis predates the arthropods' colonization of land.

Introduction

Cuticular hydrocarbons (CHCs) are long-chained molecules that cover the cuticle of most, if not all, insects¹. They function as important desiccation barrier and are thought to have fostered the transition from marine to terrestrial habitats in the lineage leading to Hexapoda². CHCs additionally serve as semiochemicals for inter- and intraspecific communication². In Hymenoptera, they play a particularly important role in establishing eusocialism by enabling kin, nest-mate and caste recognition². They have also been exploited for chemical deception and chemical mimicry². Identifying genes involved in CHC biosynthesis of Hymenoptera is of major interest, as it would open the door for CHC profile manipulation (e.g., via CRISPR-Cas9 or RNA interference) in behavioral, ecological, and socio-biological studies on Hymenoptera.

Most insects synthesize CHCs *de novo* from fatty acid precursors³ in oenocytes. These secretory cells of ectodermal origin are found in the fat body of all major insect lineages⁴. The main enzymatic steps of CHC biosynthesis are well understood and involve fatty acid synthases (FAS), fatty acid elongases (ELO), fatty acid desaturases (Desat), fatty acyl-CoA reductases (FAR), and cytochrome p450 decarbonylases (CYP4G subfamily)^{5,6,7,8,9,10} (**Figure 1**). However, the details of CHC biosynthesis and which genes encode the participating enzymes are not yet fully elucidated, particularly in the species-rich insect order Hymenoptera¹⁰. Furthermore, insights obtained from studying species of one insect lineage (e.g., Diptera) are not necessarily representative for species of a distantly-related lineage (e.g., Hymenoptera). Most genes associated with CHC biosynthesis belong to multi-gene families, and some of these exhibit remarkably high turnover rates (e.g., fatty acid desaturases¹¹). Therefore, it is highly desirable to identify CHC biosynthesis-related genes in multiple insect lineages. A promising approach to this end is transcriptomic analysis comparing gene expression between samples of the same species that differ in their CHC composition from each other.

In most species, individuals of the same sex share the same set of CHCs, which is referred to as CHC profile type¹². Intriguingly, previous research found females of the mason wasp *Odynerus spinipes* to be able to express two CHC profile types¹³. The

two chemotypes (c1 and c2) qualitatively differ in 77 CHCs from each other¹³. This CHC dimorphism offers an exceptional opportunity to identify genes involved in CHC biosynthesis, as *O. spinipes* females with different chemotypes do not appear to differ in any other trait from each other^{13,14}. Hence, they should systematically differ only in the expression of CHC biosynthesis-related genes. *O. spinipes* females additionally show age-related quantitative changes in their CHC profile composition (Moris V. et al., unpublished data), which can also be used to identify candidate genes.

In this study, we show how exploitation of the CHC dimorphism in *O. spinipes* females facilitated identification of CHC biosynthesis-related genes in Hymenoptera. Specifically, 1) we tested whether *O. spinipes* females keep a given chemotype during their adult life. 2) We sequenced, assembled, and annotated the genome of *O. spinipes* to aid accurate analysis of gene expression levels. 3) We identified candidate genes differentially expressed between *O. spinipes* females with different CHC composition. 4) We identified candidate gene orthologs in the honey bee (*Apis mellifera*) genome, a more trackable organism for in-depth molecular analyses. 5) We investigated which cell types of the worker honey bee fat body express the candidate genes. 6) We tested whether knockdown of selected candidate genes in worker honey bees via RNA interference (RNAi) impacts CHC profile composition.

Results

***Odynerus spinipes* chemotype persistency**

None of the *O. spinipes* females whose CHC profile we sampled multiple times during their adult life changed their chemotype (c1/c2): neither those kept under laboratory conditions ($N_{c1} = 10$; $N_{c2} = 17$; sampled on average three times during the first 16 days after their eclosion; **Supplementary Table 1**), nor those studied in field populations ($N_{c1} = 8$; $N_{c2} = 10$; sampled on average two times and with a time difference of on average five days; **Supplementary Table 2**). These results are consistent with the hypothesis that adult *O. spinipes* females cannot change their chemotype and justify

associating CHC chemotype differences between females with gene expression differences between these females.

Odynerus spinipes draft genome

To foster comparative transcriptomic analyses on *O. spinipes*, we sequenced the genome of this species using short read sequencing technology. The assembly of the *O. spinipes* genome resulted in 21,922 scaffolds spanning a total of 171 Mb, 89.5% of the total genome, which we estimated to be ca. 191 Mbp. The N50 value and L50 value of the draft genome assembly are 3,637 Mbp and twelve scaffolds, respectively. Exploiting whole body transcript libraries of one adult male and of two adult females (representing each chemotype) to foster annotation of nuclear-encoded protein-coding genes (**Supplementary Table 3**), we inferred a total of 16,677 gene models.

Candidate genes in *Odynerus spinipes*

We first identified statistically significant differentially expressed genes in transcriptomes of females with different chemotype in two batches of adult females that differed in sampling time (12–38 [batch 1] and 48–62 [batch 2] hours after eclosion) (**Supplementary Tables 4–6**). Considering only genes that were consistently differentially expressed in the two batches, we retained five candidate genes (**Table 1; Supplementary Tables 7 and 8**). We next searched for genes that were significantly differentially expressed in the two above batches of adult female wasps that differed in their age from each other, ignoring the wasps' chemotype. However, we only considered genes that belonged to four gene families known to be involved in CHC biosynthesis (i.e., FAS, ELO, Desat, FAR). This procedure resulted in the identification of eleven candidate genes (**Table 1; Supplementary Table 9**).

Combining the results from both approaches and accounting for redundancy (g7616 was identified with both search strategies), we identified a total of 15 candidate genes affecting CHC biosynthesis in *O. spinipes* (**Table 1**). Based on sequence homology, the candidate genes encode one fatty acid synthase, six fatty acid elongases, two fatty

acid desaturases, three fatty acyl-CoA reductases, one fatty acid amine hydrolase, one parathyroid hormone-related peptide receptor-like protein, and one a rhythmically expressed gene 5 protein.

Honey bee candidate gene orthologs

We applied a phylogenetic approach to identify orthologs of twelve candidate genes belonging to multi-gene families (i.e., FAS, ELO, Desat, FAR) in the honey bee genome. The majority (ten) of the *O. spinipes* candidate genes share a 1:1 ortholog relationship to honeybee genes (**Figures 2 and 3; Supplementary Figures 1 and 2; Table 1**). However, the fatty acid desaturase candidate gene (g14712) of *O. spinipes* was found to have three co-orthologs in *A. mellifera*. We decided to consider all of them for further analyses (**Table 1; Figure 3**). We found one fatty acid desaturase candidate gene (g14708) to not have an ortholog in *A. mellifera*. We decided to include the phylogenetically most closely related *A. mellifera* gene for further experimental assessment (**Table 1; Figure 3**).

We applied the best reciprocal hit criterion to identify orthologs of the remaining three candidate genes (g283, g2290, g3059) in the honey bee genome (**Table 1**). In all instances except one we did find 1:1 orthologs. The exception was a gene predicted to encode a fatty acid amide hydrolase (FAAH) (g2290) that we found to be represented by three co-orthologs in the honey bee genome. Because this gene family had previously not been implicated with CHC biosynthesis, we considered only one ortholog (GB53695), that with the highest amino acid sequence similarity to g2290, in subsequent experiments.

We extended our list of candidate genes by seven genes (**Table 2**). Two of them are *A. mellifera* orthologs of genes shown to be involved in the CHC biosynthesis in the fruit fly *Drosophila melanogaster* (i.e., the honey bee ortholog of the two fatty acid synthases FASN1/FASN2, **Supplementary Figure 1**, and the honey bee ortholog of the three desaturases Desat 1, 2, and F, **Figure 3**). The remaining five genes encode fatty acid elongases that belong to clades phylogenetically closely related to genes

known to be involved in CHC biosynthesis (**Figure 2**). The total number of candidate genes selected for further investigation in *A. mellifera* was thus 24 (**Tables 1** and **2**).

Candidate gene expression sites

In situ hybridization (ISH) experiments on fat body tissues from honey bee metasomas revealed 18 of the 24 candidate genes to be expressed in oenocytes: a fatty acid synthase (GB52590), eight fatty acid elongases (GB46038, GB54396, GB54397, GB51250, GB51247, GB54399, GB54404, GB55040), four fatty acid desaturases (GB40659, GB42218, GB48195, GB51236), three fatty acyl-CoA reductases (GB49380, GB50627, GB52087), one fatty acid amide hydrolase (GB53695), and one parathyroid hormone-related peptide receptor-like protein (GB52820) (**Figure 4; Tables 1** and **2**; negative controls performed with sense probes are shown in **Supplementary Figure 3**). Six of them (three fatty acid elongases, one fatty acid desaturase, one fatty acyl-CoA reductase, one fatty acid amide hydrolase) were found to be co-expressed in trophocytes, a fat body cell type different from oenocytes (**Figure 4; Tables 1** and **2**). We found two candidate genes, one fatty acid desaturase (GB51238) and one fatty acid elongase (GB54401), to be expressed in trophocytes only (**Figure 4; Table 1**). Three candidate genes, predicted to encode a fatty acid synthase (GB53412), a fatty acid elongase (GB40681), and a parathyroid hormone-related peptide receptor-like protein (GB52820), respectively, were found to be expressed in cells located between the cuticle and the fat body layer (GB52820 was found also slightly expressed in oenocytes and trophocytes). We refer to these cells as hexagonal cells (**Figure 4; Tables 1** and **2**). *In situ* hybridization experiments on one candidate gene, predicted to encode a rhythmically expressed protein (GB44756), were inconclusive (marked by an asterisk in **Figure 4**), because the staining intensity of the treatment group did not differ from that of the negative controls (**Supplementary Figure 3**). We were unable to study cell-specific expression of one candidate gene (GB54302) encoding a fatty acid elongase, as its probe synthesis failed.

Candidate gene knockdown experiments

We assessed twelve candidate genes (selection criteria given in **Supplementary Methods**) for influencing CHC profile composition by conducting RNAi-mediated knockdown experiments on worker honey bees (**Figures 5** and **6**). Knockdown of nine candidate genes resulted in statistically significant changes of the CHC profile composition of the treated bees relative to control experiments in which we injected double-stranded RNA (dsRNA) of green fluorescent protein (GFP) (powered partial least squares discriminant analysis [PPLS-DA]: $p \leq 0.05$ after Benjamini-Hochberg correction for multiple testing). The nine genes were: four fatty acid elongases (GB51250, GB51274, GB54397, GB54404) (**Figure 5**), four fatty acid desaturases (GB40659, GB42218, GB48195, GB51238) (**Figure 6**), and one fatty acid amide hydrolase (GB53695) (**Figure 5**). CHC profile compositional changes associated with the knockdown of fatty acid desaturase GB51236 were statistically insignificant after Benjamini-Hochberg correction for multiple testing (PPLS-DA, $p = 0.087$). RNAi-mediated knockdown of the remaining two genes, a fatty-acid synthase (GB52590) and a fatty-acyl-CoA reductase (GB52087), caused a premature death of the treated bees within the first two days after dsRNA injection.

While we found RNAi of the nine genes outlined above to generally result in a strong phenotypic effect 2–5 days after dsRNA injection, we did not detect statistically significant differences in expression levels of most target genes at the same point in time ($p = 0.126$ – 0.607 ; Welch's t-tests; **Figures 5** and **6**). The exceptions are three fatty acid desaturases (GB42218, GB48195, GB51238) that showed significantly lower expression levels in treated bees relative to control bees after Benjamini-Hochberg correction for multiple testing (GB42218: Wilcoxon signed-rank test, $W = 122$, $p = 0.002$; GB48195: Welch's t-test, $t_{(10.85)} = 4.01$, $p = 0.010$; GB51238: Welch's t-test, $t_{(9.67)} = 3.21$, $p = 0.033$; **Figure 6**). Note that we were unable to measure the expression levels of GB54397 (encoding a fatty acid elongase) and of GB40659 (encoding a fatty acid desaturase), because the qRT-PCR oligonucleotide primers did not efficiently amplify a unique stretch of the genes' cDNA.

PPLS-DA of the CHC profiles provided information on the compounds that dominantly discriminate the RNAi treatment group and the control group. Knockdown

of fatty acid elongases caused major CHC profile perturbations in several compound classes (**Supplementary Figure 4**), but it most notably impacted CHC chain lengths (**Figure 5**). Specifically, we found the chain lengths of CHCs positively correlating more than 50% along the first principal component of the PPLS-DA (**Figure 5**) with a fatty acid elongase dsRNA treatment (GB51247, GB51250, GB54397, GB54404) to be significantly lower than CHC chain lengths positively correlating with a control dsRNA treatment after correcting for multiple testing (Holm-Bonferroni) (GB51247: Welch's t-test, $t_{(12.04)} = 6.67$, $p < 0.001$; GB51250: Welch's t-test, $t_{(15.98)} = 6.18$, $p < 0.001$; GB54397: Welch's t-test, $t_{(27.76)} = 3.54$, $p = 0.003$; GB54404: Welch's t-test, $t_{(24.93)} = 2.58$, $p = 0.016$) (**Figure 7**). We found the opposite pattern when interfering with the expression of GB53695 (encoding a fatty acid amide hydrolase): the CHC profiles of the treatment bees contained CHCs with significantly longer chain lengths than the CHC profiles of the control bees (Welch's t-test, $t_{(27.71)} = -5.92$, $p < 0.001$; **Figures 5** and **7**; **Supplementary Figure 4**).

Knockdown of fatty acid desaturases also caused major CHC profile perturbations and impacted the relative abundance of several compound classes (i.e., alkanes, alkenes, and methyl-branched alkanes) (**Figure 6**; **Supplementary Figure 5**). Depending on what gene we targeted with dsRNA, we found statistically significant differences in the relative abundance of alkenes with specific double bond positions between treated bees and control bees. Specifically, we found the relative abundance of alkenes with a double bond at even positions (8,10) to be significantly smaller in bees treated with dsRNA of GB48195 (Wilcoxon signed-rank test, $W = 87$, $p = 0.022$) and of GB51238 (Welch's t-test, $t_{(11.94)} = 5.31$, $p = 0.002$) than in control bees after correcting for multiple testing (Holm-Bonferroni) (**Figure 8**). We did not study the impact of target gene knockdown on alkadienes (with two double bonds), since the relative abundance of alkadienes was close to the detection limit of the used GC-MS instruments.

Discussion

CHC biosynthesis-related genes in Hymenoptera

Studying intrasexual CHC profile differences in the mason wasp *O. spinipes* provided promising candidate genes that modulate CHC profile composition in Hymenoptera. Knockdown experiments on twelve selected candidate genes in *A. mellifera* resulted in statistically significant CHC profile changes in nine of the experiments. In the light of these exciting results, the remaining candidate genes not further experimentally assessed in the present investigation are highly promising targets for future reverse genetic experiments.

Fatty acid elongases

Our study provided strong evidence for four fatty acid elongases (GB51247, GB51250, GB54397, GB54404) modulating CHC profile composition in Hymenoptera (**Figures 5** and **7**). The fact that we did not measure a statistically significant decrease in gene expression levels at the point in time when we sampled the bees' CHC profiles (**Figure 5**) could be due to a combination of two factors: 1) lag-time between gene expression level changes and CHC profile changes, and 2) short-term duration of the knockdown effect on the target genes' expression levels. Because the time points for sampling dsRNA-treated bees had been chosen based on when phenotypic changes became visible in a subset of samples (typically 2–5 days after dsRNA injection; **Supplementary Table 10**), they were possibly too late to still detect transient gene expression level changes. We consider the alternative explanation, namely that RNAi treatment did not have an effect on the target genes' expression levels, as unlikely, because of the observed phenotypic changes were in most instances consistent with the predicted function of the target genes.

Considering the gene tree, at least four major clades of fatty acid elongases seem to be involved in CHC biosynthesis in Hymenoptera: ELO B1, ELO B2, ELO B3, and ELO C7 (**Figure 2**). An additional clade (ELO C3; **Figure 2**) is known to be involved in CHC biosynthesis in *D. melanogaster*^{15,16}. The wide distribution of known CHC biosynthesis-related genes and of candidate genes shown in this study to be expressed in oenocytes or trophocytes (see below) (**Figure 2**) within this gene family

makes genes belonging to the clades ELO C1–7 promising targets for future knockdown experiments. In contrast, functional involvement of genes of clade ELO A in CHC biosynthesis currently appears unlikely, as the only gene of this clade functionally studied so far (i.e., *baldspot* of *D. melanogaster*) participates in spermatogenesis¹⁷.

Fatty acid elongases have been shown to catalyze the elongation of long fatty acyl-CoAs^{2,10,18}. Consistent with this idea, we found knockdown of fatty acid elongase genes to result in a decrease in the relative abundance of CHCs with chain lengths > 27 carbon atoms (**Figures 5 and 7**). Yet, we did not find any additional differences between the knockdown phenotypes. Lack of distinct phenotypes could be due to off-target effects (i.e., knockdown of a gene coding for one fatty acid elongase simultaneously decreases the expression of another gene coding for a fatty acid elongase). However, except for two target genes, this scenario seems unlikely, as the nucleotide sequences of the synthesized dsRNAs (provided on zenodo: 10.5281/zenodo.5552394) differed substantially from each other. The exception is GB51250 and GB54404, which differed only in 43% of their nucleotides from each other. An alternative explanation for the lack of distinct phenotypic effects could be that the targeted fatty acid elongases exhibit, in stark contrast to those of mammals¹⁸, little substrate specificity. Finally, manipulation of the elongation steps could cause a shift of substrate concentrations along the entire CHC biosynthetic pathway, making it difficult to detect substrate specificity. Additional tools have to be used in future investigations to further interrogate the specific function and the substrate specificity of the enzymes encoded by the here identified genes (e.g., heterologous expression, as done by Pei et al.¹⁹, or *in vivo* synthesis using labeled precursors). The results of our study provide the foundation for such investigations.

Fatty acid desaturases

We identified four fatty acid desaturase genes (GB40659, GB42218, GB48195, GB51238) that modulate CHC composition in worker honey bees. Previous research on *D. melanogaster* had led to the identification of three closely related fatty acid

desaturases (Desat1, 2, and F) for being involved in CHC biosynthesis and introducing double bonds in fatty acyl-CoA^{20,21,22,23}. All three desaturases belong to clade Desat A1 (**Figure 3**). In the honey bee, clade Desat A1 is represented only by a single gene (GB48195), which our knockdown experiments confirmed to be also involved in CHC biosynthesis in Hymenoptera. Intriguingly, the three additional fatty acid desaturases identified in our study to influence CHC profile composition in worker honey bee belong to two clades previously not associated with CHC biosynthesis (**Figure 3**): Desat A2 (GB42218, GB51238) and Desat B (GB40659). This demonstrates that fatty acid desaturases from very distantly related clades participate in CHC metabolism (i.e., Desat A1, Desat A2, Desat B; possibly also Desat D, which contains the *O. spinipes* candidate gene g14708, but which lacks an ortholog in *A. mellifera*).

Knockdown of the above-outlined four fatty acid desaturases had a strong effect in worker honey bees on both target gene expression (except GB40659) and on CHC profile composition (**Figure 6**). Moreover, we showed that knockdown of GB48195 and of GB51238 resulted in a decrease of alkenes, consistent with the idea of the encoded proteins introducing double bonds in fatty acyl-CoA. While all desaturases in *D. melanogaster* known to be involved in CHC biosynthesis (Desat1, 2, and F) show $\Delta 9$ -specificity and thus facilitate synthesis of alkenes with double bonds at uneven positions^{20,22}, GB48195 and GB51238 are the first fatty acid desaturases identified to be involved in the production of alkenes with double bonds at even positions (8 and 10) (**Figure 8**). Knockdown of the remaining two fatty acid desaturases (GB40659, GB42218) led to broad and complex compositional changes involving multiple CHCs. As discussed above, lack of distinct phenotypes could be due to off-target effects. However, this scenario seems also in this case unlikely, as the nucleotide sequences of the synthesized dsRNAs (provided on zenodo: 10.5281/zenodo.5552394) differed between all genes except two substantially from each other. The exception is GB42218 and GB51238, whose dsRNAs differed only in 29% of their nucleotides from each other. We consider off-target effects in this specific instance to unlikely have significantly impacted the results, however, as the nucleotide sequence used for antisense riboprobe (ISH) and the dsRNA probe (RNAi) caused different staining in the ISH experiments and different phenotypes in the knockdown experiments,

respectively.

Fatty-acid amide hydrolase

We identified a fatty-acid amide hydrolase (GB53695) influencing CHC profile composition in worker honey bees. This result is remarkable, as fatty-acid amide hydrolases had not been associated with CHC biosynthesis before. Knockdown of GB53695, which we found to be expressed in oenocytes, resulted in a significant increase of the relative abundance of CHCs with chain lengths > 27 carbon atoms (**Figure 7**). However, we did not detect statistically significant gene expression changes (**Figure 5**). Fatty-acid amide hydrolases are characterized by their amidase signature and are — in mammals — known to degrade fatty acid amides of endogenous lipids into fatty acids by removing the amine group²⁴. How the functions of fatty-acid amide hydrolases relate to the modulation of CHC profiles in Hymenoptera remains unclear at this point. We deem it possible that the studied fatty-acid amide hydrolase impacts the availability of CHC precursors, by, for instance, releasing short-chain fatty acids that are then possibly reduced to aldehydes, which could be transformed to CHCs by a cytochrome p450 decarbonylase (**Figure 1**).

Fatty acid synthases and fatty acyl-CoA reductases

Knockdown of candidate genes encoding a fatty acid synthase (GB52590) and a fatty acyl-CoA reductase (GB52087) resulted in a premature death of the treated bees. A similar observation was made by other researchers when targeting genes of these two families in various insects (e.g., ^{25,26,27,28,29}). We hypothesize that interfering with the expression of genes of the two gene families reduced — due to their central role in CHC biosynthesis — the overall availability of CHCs to an extent that it caused desiccation-related death of the treated bees^{26,28,29}. Knockdown of fatty acid synthase could additionally have reduced the synthesis of other molecules fundamentally important for survival (e.g., cuticular lipids²⁶). Despite the fact that our RNAi experiments on GB52590 and GB52087 remained inconclusive, alternative approaches should consider them highly promising candidate genes, as multiple lines

of evidence strongly suggest both genes to be involved in CHC biosynthesis: a) ortholog relationship to *D. melanogaster* genes known to be involved in CHC biosynthesis (i.e., GB52590, ortholog of FASN3²⁸ fatty acid synthase which belongs to clade FAS B2; **Supplementary Figure 1**; GB52087, ortholog of the three fatty acyl-CoA reductases CG10097, CG13091, and CG17560¹⁶, all of which belong to clade FAR F; **Supplementary Figure 2**), b) expression in honey bee oenocytes (**Figure 4**), c) statistically significant expression differences of their *O. spinipes* orthologs (i.e., g3158, ortholog of GB52590; g1571, ortholog of GB52087) between 12–38 hours-old *O. spinipes* females with different chemotype (**Supplementary Tables 5 and 7**) and between females differing in age from each other (**Table 1**; **Supplementary Table 9**). Two other FAR-encoding candidate genes, GB50627 (clade FAR F) and GB49380 (clade FAR C) also seem promising candidate genes, as they are both expressed in honey bee oenocytes (**Figure 4**) and their *O. spinipes* orthologs (g2413 and g7842) were found to be differentially expressed between females differing in age from each other (**Table 1**; **Supplementary Table 9**).

Evolutionary roots of CHC biosynthesis

The phylogenetic distribution of genes involved in CHC biosynthesis across Euarthropoda (i.e., Chelicerata, Myriapoda, “Crustacea”, and Hexapoda) has implications for our understanding of the evolution of CHC biosynthesis. All major gene families (**Figure 1**) and most subclades within them harboring genes known to be involved in CHC biosynthesis (FAS A, ELO C, ELO B, Desat A1, and potentially FAR C) are present in the extant major lineages of Euarthropoda. CHCs and oenocytes have also been reported from all major Euarthropoda lineages^{30,31,32,33,34,35,36,37}. We therefore consider it reasonable to hypothesize that the capacity of CHC biosynthesis evolved before the divergence of the major lineages of Euarthropoda and, consequently, before the colonization of land. The only gene whose phylogenetic distribution is currently not fully consistent with this hypothesis is the cytochrome p450 decarbonylase (CYP4G gene family) (**Figure 1**) that catalyzes the final step of CHC biosynthesis^{8,38} and which has — so far — only been found in Hexapoda^{38,39}. As the

cytochrome p450 gene family is known for its high turnover rate³⁸, we do not put too much weight in this single exception.

Possible importance of trophocytes for CHC biosynthesis

We found almost all genes whose knockdown resulted in a CHC profile change to be expressed in oenocytes — consistent with the common view that CHC metabolism occurs in oenocytes (e.g.,^{8,40,41,42}). A major exception was GB51238, encoding a fatty acid desaturase. We found this gene exclusively and highly expressed in trophocytes (**Figure 4**), another cell type of the insects' fat body. While we cannot exclude the possibility that the gene is expressed at a very low level in oenocytes, we deem the result noteworthy, as other researches previously already suggested fat body cells other than oenocytes to be possibly involved in CHC biosynthesis^{28,43}. Given the possible implications of CHC biosynthesis involving additional types of fat body cells, we recommend future studies to not exclusively focus on oenocytes when studying CHC metabolism.

Molecular basis of the CHC profile dimorphism in *O. spinipes* females

The knockdown experiments on *A. mellifera* helped better understanding the molecular basis for chemotype differences in *O. spinipes* females. *O. spinipes* females with chemotype 2 differ from those with chemotype 1 primarily by having alkenes with double bonds at even chain positions (8, 10, 12, and 14; chemotype 1 females have them at uneven chain positions) and by featuring larger quantities of alkenes with chain lengths > 25 carbon atoms (**Supplementary Figure 6**). Intriguingly, knockdown of one (GB51238) of the three co-orthologs of fatty acid desaturase g14712 caused a decrease in the relative abundance of alkenes with double bonds at even chain positions in worker honey bees. The phenotypic change is thus fully consistent with the idea that higher expression of the gene's ortholog (g14712) in *O. spinipes* females with chemotype 2 gives rise to a higher abundance of alkenes with double bonds at even chain positions. Likewise, knockdown of the fatty acid elongase (g7616) ortholog

in worker honey bees (GB51247) caused a decrease of the relative abundance of CHCs with chain lengths larger than 27 carbon atoms. The result is in line with the idea that higher expression of the gene's ortholog (g7616) in *O. spinipes* females with chemotype 2 gives rise to a higher abundance of alkenes with chain lengths > 25 carbon atoms. It is remarkable that, considering the large number of qualitative and quantitative differences between the two *O. spinipes* CHC chemotypes¹³, most of the differences seem to be due to the action of only two enzymes in the CHC biosynthesis pathway.

While comparative analysis of gene expression levels between *O. spinipes* females with different chemotypes led to the identification of additional candidate genes than the above-discussed fatty acid elongase and fatty acid desaturase (**Table 1**), their role in shaping the two distinct phenotypes remains unclear. For example, we found a fatty acid amid hydrolase (g2290) being more expressed in females with chemotype 2 than in females with chemotype 1. However, knockdown of one of the three co-orthologs (GB53695) in the worker honey bees caused a decrease of short chain CHCs, contrary to what we would have predicted from the CHC profile characteristics of *O. spinipes* females with chemotype 2. The predicted functions of other candidate genes (e.g., rhythmically expressed gene 5 protein [g283]; parathyroid hormone or parathyroid hormone-related peptide receptor-like protein [g3059]) also provide no clue on how they could contribute to the observed CHC chemotype differences.

Our study demonstrates how comparative transcriptomic analyses on a non-model organism featuring unique characteristics can deliver valuable information that research on established model organisms may not be able to provide. It furthermore enables future experimental manipulation of CHC profiles in Hymenoptera and subsequent functional investigations on the activity and substrate specificity of the identified genes.

Methods and Material

Odynerus spinipes chemotype persistency analysis

Lab experiments

We collected prepupae of *O. spinipes* females and males from trap nests (details in **Supplementary Methods**). Each female was placed in a separate observation cage (30 cm x 30 cm x 30 cm, Bioform, Nürnberg, Germany) and provided with a conspecific male and absorbent paper soaked with honey and water. The presence of males was meant to simulate field conditions for the females. The cages were kept in a climate chamber (70% humidity, 23 °C during day, 18 °C during night, and with a 12/12 h day/night cycle). We sampled the cuticular hydrocarbons (CHCs) of 23 females with the aid of Solid Phase Micro-Extraction (SPME) fibers (see below) at two days, seven/eight days, and 14 days (or immediately after they died) after they had eclosed (**Supplementary Table 1**).

Field experiments

We sampled the CHCs of 18 females two to four times during their adult life in the field (**Supplementary Table 2**). Females were caught at their nest entrance, color-coded with water resistant paint markers (Edding 780, 0.8 mm, Ahrensburg, Germany), CHC-sampled with the aid of SPME fibers, and released the next morning at their nest site. The sampling time coincided with the start of nest building activity.

Cuticular hydrocarbon extraction

CHCs from living wasps were sampled by stroking a SPME fiber (Supelco, coating: polydimethylsiloxane, 100 μ m, Sigma-Aldrich, Bellefonte, PA, USA) for 2 min. (field populations) or 4 min. (lab population) over the cuticle of a female's metasoma. Wasps processed in 2016 were anesthetized by exposing them for 1 min. to CO₂, those

processed in 2017 by cooling them down for 3 min. at -20 °C (because we suspected CO₂ treatment to cause amnesia in the treated wasp, as wasps treated this way in 2016 rarely returned to their nest; see ⁴⁴).

Gas chromatography and mass spectrometry

All CHC extracts were either analyzed with a gas chromatograph coupled to a mass spectrometer (GC-MS) or with a gas chromatograph-flame ionization detector equipped with a SLB-5 MS non-polar capillary (GC-FID) applying the same temperature profile (details in **Supplementary Methods**). The GC-MS data were recorded and quantified with the software HP Enhanced ChemStation G1701AA (version A.03.00; Hewlett Packard, Palo Alto, California, USA). The GC-FID data were recorded and quantified with the software GCSolution (Shimadzu, Wommel, Belgium). The chemotype of the female wasps was inferred from the presence/absence of chemotype-specific alkenes^{13,14} with diagnostic Kovats indices⁴⁵.

***Odynerus spinipes* draft genome**

NGS library preparation and sequencing

We sequenced the nuclear genome at a base-coverage depth of ca. 86x using Illumina HiSeq2000 sequencing technology and three different types of NGS libraries: 250 bp paired-end [10.58 Gbp] (55x), 800 bp paired-end [2.45 Gbp] (13x), and 3 kbp mate-pair [3.4 Gbp] (18x), generated from a single pool of DNA extracted from males and females (details in **Supplementary Methods**). To foster gene annotation, we sequenced whole-body transcriptomes of one adult male and of two adult females (one of each chemotype). DNA and RNA extraction, library preparation, and Illumina DNA sequencing were done by BGI Tech Solutions (Hongkong).

Genome assembly and annotation

All raw reads of genomic DNA were processed with trimmomatic⁴⁶ version 0.33 (details in **Supplementary Methods**). The processed reads were then used to assemble the genome with Platanus⁴⁷ version 1.2.4 using all three libraries in each of the three assembly steps (contig construction, scaffolding, and gap closing) and applying the software's default parameters. The protein-coding gene repertoire was annotated with the BRAKER *ab initio* gene prediction pipeline⁴⁸ version 2.1 and providing the RNA-seq data of the three above-mentioned adult wasps as intrinsic evidence (details in **Supplementary Methods**).

Comparative gene expression analysis

Odynerus spinipes samples

We sequenced metasoma transcriptomes of twelve *O. spinipes* females raised under laboratory conditions from field-collected prepupae (**Supplementary Table 4**): six of them (three of each chemotype) were sampled 12–38 h after they had eclosed, the other six (three of each chemotype) were sampled 48–62 h after they had eclosed. The prepupae were collected from three field sites (**Supplementary Table 4**).

Prior to RNA fixation, the eclosed females were kept separate from each other at 21 °C in polystyrene tubes (53 mm x 100 mm, Bioform, Nuremberg, Germany) containing moistened cotton and experienced a 12/12 h day/night cycle. All samples were euthanized at the same time of the day (1 pm) to minimize a potential impact of expression differences by rhythmically expressed genes (note that expression of *Desat1* follows such a diurnal rhythm in *D. melanogaster*⁴⁹). Wasps were euthanized by submersing them for 60–120 sec. in hexane (*n*-hexane for gas chromatography, SupraSolv, Merck, Germany). The CHC extracts were used to infer the wasps' chemotypes as described above. Each wasp's metasoma was ground with a sterile pestle in 1 mL RNAlater (QIAGEN, Hilden, Germany) and kept at -80 °C before further processing it.

Transcriptome sequencing

RNA extraction, NGS library preparation, and 150 bp paired-end nucleotide sequencing with an Illumina NextSeq 500 (Illumina, San Diego, USA) were done by StarSEQ (Mainz, Germany) using standard protocols. We collected on average 20–44 million reads per sample.

Gene expression analysis

The RNA-seq data were analyzed with software integrated in the Galaxy Europe platform (<https://usegalaxy.eu>). We quality-trimmed the raw reads and removed NGS library adapters with Trim Galore version 0.4.3.0 (http://www.bioinformatics.babraham.ac.uk/projects/trim_galore/). The quality of the raw reads was assessed with FastQC version 0.67 (<http://www.bioinformatics.babraham.ac.uk/projects/fastqc/>). The gene model file (GTF) of the *O. spinipes* genome was used to model splice junctions with the software STAR⁵⁰ version 2.5.2b0 via mapping the transcript raw reads onto the genome assembly. We inferred a matrix of normalized read counts per gene, considering the genes in the above GTF file, with featureCounts⁵¹ version 1.5.3.

Because our initial experimental design included only one batch of six transcriptomes (three of each chemotype) (**Supplementary Table 4**), we applied a two-step procedure for identifying genes differentially expressed between females with different chemotypes: we first searched with DESeq2⁵² version 2.11.39 for differentially expressed genes in each of the two batches, using a False discovery rate (FDR) of 0.05 as cutoff. We then reciprocally assessed genes that this analysis indicated to be differentially expressed between the two chemotypes in the other batch, this time applying a more conservative Holm-Bonferroni correction for multiple testing. Only genes that were significantly differentially expressed in both analyses were kept as candidate genes. We assessed with edgeR⁵³ version 3.28.0 in the R statistical software version 3.4.1 (<http://www.R-project.org>) whether the five genes identified with the above-described procedure were also found differentially expressed in the two batches ($p\text{-value} < 0.05$ after Holm-Bonferroni correction for five tests). We used

DESeq2 as well as edgeR to assess statistical significance of gene expression differences between *O. spinipes* females of different age classes, ignoring the wasps' chemotype. We considered genes, judged to differ in their expression by either of the two programs, as candidates if they encoded fatty acid synthases (FAS), fatty acid elongases (ELO), fatty acid desaturases (DESAT), or fatty acyl-CoA reductases (FAR).

Identification of candidate gene orthologs in the honey bee genome

Honey bee orthologs of *O. spinipes* candidate genes belonging to the gene families Desat, ELO, FAR, and FAS were identified by their phylogenetic position in the corresponding gene trees. To this end, we identified the proteins of the four gene families by searching for proteins with the protein family-characteristic protein domains (see below) available as HMM models from PFAM database⁵⁴ (PFAM-A, version 28). The HMM models were searched with the aid of hmmer⁵⁵ version 3.1b1 against the protein sets of 37 Euarthropoda, including *O. spinipes*, with sequenced genomes (details in **Supplementary Methods**). To identify fatty acid synthases, we searched for proteins containing the PFAM domains ketoacyl-synt KS (Pfam family accession PF00109), acyl-transferase-1 (PF00698), and PS-DH (PF14765) (e-value < 1e-5). To identify fatty acid elongases, we searched for proteins containing the ELO PFAM domain PF01151 (e-value < 1e-30), and consisting at least of 200 amino acids. To identify fatty acid desaturases, we searched for proteins containing the PFAM domain PF00487 (e-value < 1e-20). Note that we did not consider sphingolipid desaturases, as they show little amino acid sequence similarity to other desaturases (see ¹¹). To identify fatty acyl-CoA reductases, we searched for the simultaneous presence of the two PFAM domains PF07993 (e-value < 1e-30) and PF03015 (e-value < 1e-20). The amino acid sequences of a given gene family were aligned with MAFFT⁵⁶ version 7.123 and phylogenetically analyzed with the software IQ-TREE⁵⁷ version 1.6 (details in **Supplementary Methods**).

Honey bee orthologs of the remaining *O. spinipes* candidate genes and which did not belong to the above gene families were identified by applying the best reciprocal hit criterion and searching the protein repertoires of the two species with the blastp

program of the BLAST+ software suite^{58,59}.

***In situ* hybridization**

Synthesis of sense and anti-sense riboprobes

RNA probes were synthesized according to standard protocols⁶⁰ from cloned partial mRNA sequences of all candidate genes (**Tables 1** and **2**) using DIG and FLU RNA Labeling Kits (Roche, Basel, Switzerland). Strand cDNA was synthesized from isolated total RNA (1 µg, DNase-treated) of fat body tissues isolated from *A. mellifera* using the Quantitect kit (QIAGEN GmbH, Hilden, Germany). The resulting cDNA library was used to amplify target regions via polymerase chain reactions (PCRs) (Mastercycle DNA Engine Thermal cycler PCR, Eppendorf AG, Hamburg, Germany) using the Phusion high fidelity kit (New England Biolabs, Ipswich, MA, USA). Amplicons were cloned with the NEB PCR Cloning kit (New England Biolabs GmbH, Ipswich, MA, USA), and the plasmids were extracted using the Gen Elute Plasmid Miniprep kit (Sigma-Aldrich, Steinheim, Germany) and were sent to Genewiz (Genewiz, Leipzig, Germany) for nucleotide sequencing. The cloned gene fragments (387–774 bp long) were used to synthesize anti-sense and sense RNA probes using T7 (Roche, Mannheim, Germany) or SP6 (Roche, Mannheim, Germany) RNA polymerases. Sense probes served as negative control to each anti-sense probe. Oligonucleotide primers used for probe cloning were designed with the online software Primer-BLAST⁶¹ and synthesized by Metabion (Planegg, Germany) or Sigma-Aldrich (Steinheim am Albuch, Germany) (**Supplementary Table 11**).

Whole mount RNA in situ hybridization

In situ hybridization was done on worker honey bees collected from a single hive at Würzburg University. All bees eclosed 4–8 days prior to their processing, were reared from prepupae in climate chambers at 28 °C, and were collectively kept in observation cages at 32 °C with 30% sugar solution. Metasoma tergites and sternites were

dissected from living bees (placed on ice for 5 min) in 4% paraformaldehyde (PFA) (Sigma-Aldrich, Steinheim am Albuch, Germany; dilution in Phosphate buffered saline [PBS]). The dissected segments were fixed overnight in a solution consisting of 2 mL heptane (Carl Roth, Karlsruhe, Germany) and 2 mL of 4% PFA on a rotator at room temperature. The fixation was finalized following the protocol given by Dearden et al.⁶². Whole-mount RNA *in situ* hybridization was done as described by Dearden et al.⁶³, with minor modifications outlined in **Supplementary Methods**. All mounting media included DAPI (0.5 μ g/mL; Thermofisher Scientific, Waltham, Massachusetts, USA) to achieve nuclear counterstaining. High resolution images of stained tissues mounted on glass slides were obtained with a Zeiss Axio Imager A1 fluorescence microscope, a Zeiss AxioCam MRc, DAPI fluorescence filters, a Zeiss HXP120 UV light unit, and the Zeiss Axiovision software (Carl Zeiss Jena GmbH, Jena, Germany). Overlay of brightfield images and DAPI fluorescence was done with ImageJ/Fiji (<https://fiji.sc/>). Trophocytes were distinguished from oenocytes by their polyploid or polygonal cells (oenocytes are round) with a large irregular nucleus (the nucleus of oenocytes is round or oval)⁶⁴.

RNAi-mediated knockdown experiments

dsRNA synthesis

Double-stranded RNA (dsRNA) was synthesized in vitro from 387–774-bp-long inserts — previously generated for RNA probe synthesis — using MEGAscript RNAi kit (Ambion, Texas, USA) (details in **Supplementary Methods**). T7 promotor was added to the oligonucleotide primers (listed in **Supplementary Table 11**) used to amplify the plasmids. A plasmid with a DNA fragment encoding green florescent protein (GFP), kindly provided by Anne-Kathrin Rohlfing (nucleotide sequence on zenodo: 10.5281/zenodo.5552394), was used to synthesize dsRNA that served as negative control. We verified that the GFP nucleotide sequence had no significant match in the honey bee genome⁶⁵ by searching it with tools of the BLAST+ software suite^{58,59} against the honey bee genome (GCF_000002195.4_Amel_4.5). The identity of the

amplicons was verified by nucleotide-sequencing them. All dsRNA probes were diluted in MOPS Ringer solution (270 mM NaCl, 3.2 mM KCl, 1.2 mM CaCl₂, 10 mM MgCl₂, 10 mM γ-(N-Morpholino) propane sulfonic acid (MOPS), pH 7.4) to reach a final concentration of 5 μg/μL.

RNAi

Knockdown experiments were done on worker honey bees reared as described above and chilled prior to the dsRNA injection. All bees of a given experiment came from the same hive, eclosed 6–7 days prior to dsRNA treatment and belonged to the same caste (workers, nurse bee). We injected between the 2nd and 3rd tergite of each bee's the metasoma 1 μL dsRNA (5 μg) with a Hamilton microliter syringe (MICROLITER Series 7000 Tip type 2, 7105 KH, Carl Roth, Karlsruhe, Germany). Per experiment, we treated 6–12 bees with dsRNA of the target gene of interest and 6–13 bees with dsRNA of GFP, which served as control (**Supplementary Table S11**). Bees of different experiments were kept in separated observation cages and were fed with 30% sugar water. Note that we were forced to use the same control group when assessing knockdown of GB51250 and GB48195 (which have been carried out at the same time and with bees from the same hive), as we lacked bees to make an extra control group at that specific time.

Analysis of CHC profile changes

After dsRNA injection, we daily sampled the CHCs of six bees (three of the treatment group and three of the control group) from a given RNAi experiment with SPME fibers, probing each bee's metasoma. As soon as notable CHC profile differences between treatment bees and control bees were detected (sampling times given in **Supplementary Table 10**), all samples from the corresponding RNAi experiment were euthanized by immersing them for 2 min. in hexane (n-hexane for gas chromatography, SupraSolv, Merck, Germany) and were subsequently frozen in liquid nitrogen. The bees' metasoma were subsequently stored at -80 °C before further processing them.

CHC extracts were analyzed with a GC-MS (details in **Supplementary Methods**). CHC abundance values were normalized by dividing them by the total abundance of CHCs identified in a given CHC extract (available on zenodo: 10.5281/zenodo.5552394). The normalized data were analyzed in Rstudio (R version 4.0.3), using functionalities from the following packages: RVAideMemoire version 0.9-80 (<https://CRAN.R-project.org/package=RVAideMemoire>), vegan version 2.5-7, Hotelling version 1.0-7, pls version 2.7-3, FactoMineR⁶⁶ version 2.4, and Factoextra version 1.0.6 (<https://CRAN.R-project.org/package=factoextra>). The statistical significance of CHC profile differences between groups was assessed with a powered partial least squares discriminant analysis and applying pairwise permutation tests^{67,68}. Differences in chain lengths and the relative abundance of specific alkenes were assessed with Welch's t-test or with the Wilcoxon signed-rank test, depending on whether or not the data appeared normally distributed (assessed with the Shapiro-Wilk test), using the stats package version 3.6.2 in Rstudio. *P*-values were corrected with the Holm-Bonferroni method for multiple testing. Boxplots were drawn with the ggplot2 package (<https://ggplot2.tidyverse.org>) in Rstudio.

Analysis of gene expression changes

We extracted the total RNA from the metasoma of each bee treated with dsRNA using RNeasy Plus Mini kit (QIAGEN, Hilden, Germany) and quantified the relative expression of the target genes via real-time quantitative PCR (RT-qPCR) using the QuantiTect SYBR Green RT-PCR kit (QIAGEN, Hilden, Germany) and a StepOnePlus Real-Time PCR System (Applied Biosystems-Life technologies, ThermoFisher Scientific, Waltham, Massachusetts, USA). We quantified expression levels in at least five samples per group (i.e., treatment and control) and in three technical replicates per sample. RT-qPCR oligonucleotide primers used to assess the target genes' expression levels are listed in **Supplementary Table 12** and were designed with Primer-BLAST⁶¹. If possible, oligonucleotides were designed so that the resulting amplicon was 150–250 bp in length, that the oligonucleotides' melting temperature was between 57 and 63 °C, that the oligonucleotides would span exon-exon junctions, that

the amplified region was localized at the 5' to the dsRNA target region⁶⁹, and that the amplified region was not overlapping with the dsRNA target region. We used GB49975 (predicted ribosomal protein L1) as reference gene (**Supplementary Table 11**), because its *O. spinipes* ortholog was one of the genes that exhibited the least expression differences between females with different chemotype. All oligonucleotide primer pairs exhibited an amplification efficiency of more than 95%, and each amplicon showed a single melting temperature maximum. The expression data were analyzed following the protocol given by Pfaffl, 2001⁷⁰. Gene expression levels were statistically assessed with the Welch's t-test and with the Wilcoxon signed-rank test, as described above. *P*-values were corrected with the Benjamini-Hochberg method for multiple testing.

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Author contribution table

O.N., T.S., and V.C.M. conceived and designed the project. A.D., B.M., J.W., M.P., O.N., and S.M. sequenced, assembled, and annotated the *Odynerus spinipes*

genome. J.P.O. and V.C.M. performed the comparative transcriptomic analyses. L.P. and V.C.M. inferred the gene trees. D.L. and V.C.M. designed the *in situ* hybridization experiments. M.T., O.N., R.S., T.S., and V.C.M. contributed to the design of the knockdown experiments. V.C.M. conducted the chemotype persistency, the *in situ* hybridization, the knockdown experiments, and all chemical and statistical analyses. D.L., O.N., R.S., T.S. provided instruments and reagents. O.N. and V.C.M. wrote the manuscript, with contributions from T.S. All authors provided feedback on the manuscript.

Competing Interests statement

The authors declare no competing financial interests.

Tables

Table 1. Cuticular hydrocarbon biosynthesis candidate genes in the mason wasp *Odynerus spinipes*. The table informs about 1) between which groups differential expression differences (DGE) were detected, 2) orthologous/homologous genes in the honey bee (*Apis mellifera*) genome, 3) the genes' predicted functions, and 4) the expression of the candidate genes in bee fat body cells.

Wasp gene ID	DGE between	Honey bee gene ID	Predicted function	Cell type
g3158	age classes	GB52590	fatty acid synthase	oenocytes (weak)
g6537	age classes	GB46038	fatty acid elongase	oenocytes trophocytes +
g7609	age classes	GB54399	fatty acid elongase	oenocytes (weak)
g7610	age classes	GB54397	fatty acid elongase	oenocytes
g7616	age classes / chemotypes	GB51247	fatty acid elongase	oenocytes
g7620	age classes	GB54401	fatty acid elongase	trophocytes
g7621	age classes	GB54302	fatty acid elongase	NA
g14708	age classes	NA ¹	fatty acid desaturase	oenocytes
g14712	chemotypes	GB42218 GB51236 GB51238	fatty acid desaturase	oenocytes oenocytes trophocytes
g2413	age classes	GB50627	fatty acyl-CoA reductase	oenocytes
g1571	age classes	GB52087	fatty acyl-CoA reductase	oenocytes trophocytes +
g7842	age classes	GB49380	fatty acyl-CoA reductase	oenocytes
g283	chemotypes	GB44756	rhythmically expressed gene 5 protein	NA
g2290	chemotypes	GB53695 ²	fatty acid amine hydrolase	oenocytes trophocytes (weak) +
g3059	chemotypes	GB52820	parathyroid hormone-related peptide receptor-like protein	oenocytes (weak) + trophocytes (weak) + hexagonal cells [strong]

¹ Since we found no 1:1 ortholog in the honey bee genome, we assessed the phylogenetically most closely related homolog, GB40659, for being involved in CHC biosynthesis.

² One of three co-orthologs in the honey bee, the other two being GB48850 and GB47832 (not studied by us).

Table 2. Additional cuticular hydrocarbon biosynthesis candidate genes studied by us in the honey bee (*Apis mellifera*). Given are 1) the fly and 2) the honey bee gene IDs, 3) the genes' predicted functions, and 4) the expression of the candidate genes in honey bee fat body cells.

Fly gene ID	Honey bee gene ID	Predicted function	Cell type
FASN1/2	GB53412	fatty acid synthase	hexagonal cells
CG30008	GB55040	fatty acid elongase	oenocytes + trophocytes
CG30008	GB40681	fatty acid elongase	hexagonal cells
CG5326	GB54396	fatty acid elongase	oenocytes + trophocytes
CG31522	GB51250	fatty acid elongase	oenocytes
CG2781	GB54404	fatty acid elongase	oenocytes
Desat1, 2, and F	GB48195	fatty acid desaturase	oenocytes + trophocytes

List of figures

Figure 1. Schematic overview of the main steps of cuticular hydrocarbon biosynthesis. The pathway starts with a fatty acid synthase (FAS) that binds acetyl-coenzyme A (CoA) with malonyl-CoA units. The resulting acyl-CoA (with an even number of carbon atoms) is repeatedly elongated by a FAS which binds acyl-CoA with malonyl-CoA. During each elongation step, the chain length of the fatty acyl-CoA increases by two carbon atoms. Fatty acyl-CoA chains with an uneven number of carbon atoms are thought to arise from substituting malonyl-CoA with propionyl-CoA during the elongation step. The consecutive elongation of the fatty acyl-CoA chain by FAS results in long chain fatty acyl-CoAs. Additional elongation of these fatty acyl-CoA chains to very-long fatty acyl-CoA requires the action of fatty acid elongases that add additional malonyl-CoA molecules to the long chain fatty acyl-CoA. These are reduced into aldehydes by fatty acyl-CoA reductases. Before the latter step, double bonds can be inserted in the (very-)long chain fatty acyl-CoAs by fatty acid desaturases, resulting in the biosynthesis of unsaturated hydrocarbons with a different number of double bonds (alkenes, alkadienes). The biosynthesis of methyl-branched alkanes is thought to depend on the action of a specialized fatty acid synthase that incorporates methylmalonyl-CoA instead of malonyl-CoA into the chain at specific positions during the chain elongation. The last step of CHC biosynthesis is the oxidative decarbonylation of aldehydes leading to hydrocarbons (with the loss of one carbon) by a cytochrome p450. Based on ^{2,5,6,7,8,9}.

Figure 2. Gene tree of fatty acid elongases from 37 Euarthropoda showing copy numbers and involvement of genes in CHC biosynthesis. Branch support was calculated from ultrafast bootstrap (UFBoot2) replicates (in %) and Shimodaira-Hasegawa approximate likelihood ratio tests ([SH]-aLRT in %).

Figure 3. Gene tree of fatty acid desaturases from 37 Euarthropoda. Color coding and symbols are as in the legend of **Figure 2**.

Figure 4. Photomicrographs of worker honey bee (*Apis mellifera*) fat body cells stained with DAPI (in light blue) and RNA *in situ* hybridizations (dark blue/purple) visualizing the expression of fatty acid synthases (FAS), fatty acid elongases (ELO), fatty acid desaturases (Desat), fatty acid reductases (FAR), and additional candidate gene transcripts (others). Three fat body cell types are indicated: hexagonal cells (h), oenocytes (o), and trophocytes (t). Two separate photomicrographs visualize the expression of GB52820. DAPI was used to counterstain nuclei. Staining in the photomicrograph marked with an asterisk (*) did not significantly differ from that of the corresponding control (**Supplementary Figure 3**). Scale bar: 50 μ m.

Figure 5. Effects of RNAi-mediated knockdown of four fatty acid elongases (GB51247, GB51250, GB54397, GB54404) and of one fatty acid amide hydrolase (GB53695) in worker honey bees (*Apis mellifera*) 2–5 days after dsRNA treatment. The first two columns show score plots and correlation circle plots from a powered partial least squares discriminant analysis (PPLS-DA) of the cuticular hydrocarbon (CHC) profile data of target gene-treated and of control bees. *P*-values indicate the statistical probability of group differences representing random variation after Benjamini-Hochberg correction for multiple ($N = 10$) testing. The correlation circle plots indicate how many CHCs of a given chain length class (i.e., ≤ 27 [red] or > 27 [blue] carbon atoms) correlate with the first two principal components. Box plots in the third column show gene expression levels ($\log_2$ -fold change) of the target genes in target gene-treated and in control bees. None of the expression levels differed statistically significantly between target gene-treated and control bees after applying Benjamini-Hochberg correction for multiple testing ($N = 9$; Welch's t-test and Wilcoxon signed-rank test). Additional information is shown in **Supplementary Figure 4**.

Figure 6. Effects of RNAi-mediated knockdown of four fatty acid desaturases (GB40659, GB48195, GB42218, GB51238) in worker honey bees (*Apis mellifera*) 2–5 days after dsRNA treatment. The first two columns show score plots and correlation circle plots from a powered partial least squares discriminant analysis (PPLS-DA) of cuticular hydrocarbon (CHC) profile data of target gene-treated and of control bees. *P*-values indicate the statistical probability of group differences representing random variation after Benjamini-Hochberg correction for multiple ($N = 10$) testing. The correlation circle plots indicate how many CHCs of a given compound class (i.e., alkanes [green], alkenes/alkadienes [blue], methyl-branched alkanes [yellow]) correlate with the first two principal components. Box plots in the third column show gene expression levels ($\log_2$ -fold change) of the target genes in target gene-treated and in control bees. Asterisks indicate the statistical significance of expression differences between target gene-treated and control bees after applying Benjamini-Hochberg correction for multiple testing ($N = 9$; Welch's t-test/Wilcoxon signed-rank test; $p \leq 0.05$ [*], $p \leq 0.01$ [**]). Additional information is shown in **Supplementary Figure 5**.

Figure 7. Chain lengths of cuticular hydrocarbons (CHCs) positively associated in a powered partial least squares discriminant analysis along the first principal component with bees treated with dsRNA targeting fatty acid elongases (GB51247, GB51250, GB54397, GB54404; blue) or a fatty acid hydrolase (GB53695; orange) and with bees treated with dsRNA targeting GFP (control; grey). Asterisks indicate statistical significance of abundance differences between the two treatment groups after Holm-Bonferroni correction for multiple ($N = 5$) testing (Welch's t-test: $p \leq 0.05$ [*], $p \leq 0.01$ [**], $p \leq 0.001$ [***]).

Figure 8. Relative abundance of alkenes with double bonds at even and at uneven positions in bees treated with dsRNA of fatty acid desaturases (pink) or dsRNA of GFP (control; grey). Asterisks indicate statistically significant abundance differences between the two groups after Holm-Bonferroni correction for multiple (N = 10) testing (Welch's t-test/Wilcoxon signed-rank test: $p \leq 0.05$ [*], $p \leq 0.01$ [**], $p \leq 0.001$ [***]).

Supplementary Files (on Zenodo: 10.5281/zenodo.5552394)

- *Odynerus spinipes* draft genome assembly (ODSP_platanus.1.2.4.n0.s20.fa)
- Soft-masked version of the draft genome (ODSP_platanus.1.2.4.n0.s20.fasta.softmasked)
- Gene models of predicted protein-coding genes (ODSP_platanus.1.2.4.n0.s20_BRAKER1.9_annot.gff)
- mRNA file of predicted gene protein-coding genes (mRNA_ODSP_platanus.1.2.4.n0.s20.fa)
- Protein file of predicted gene protein-coding genes (ODSP_BRAKER_augustus_aa.fa)
- Deseq2 results table (age class comparison) (AGE_DESEQ2_sign_results_ordered.xlsx)
- EdgeR results table (age class comparison) (AGE_EdgeR_lrt_sign_results_ordered.xlsx)
- qRT-PCR raw values (Expression_knock_downs_all_Apis_ids.xlsx)
- Cuticular hydrocarbon compound abundances in *Apis mellifera* samples treated with dsRNA (CHCs_abundance_knock_downs_Apis_ids.xlsx)
- Nucleotide sequence encoding green fluorescent protein (GFP) (Primer_GFP_sequence_used_Negative_control_RNAi.docx)
- Nucleotide sequence of candidate genes used for probe and dsRNA synthesis (Sequences_ISH_dsRNA_Apis_ids.fa)