The Role Of Cuticular Pheromones In Courtship Conditioning Of Drosophila Males

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The role of cuticular pheromones in courtship conditioning of *Drosophila* males

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Courtship conditioning is an associative learning paradigm in *Drosophila melanogaster*, wherein male courtship behavior is modified by experience with unreceptive, previously mated females. While the training experience with mated females involves multiple sensory and behavioral interactions, we hypothesized that female cuticular hydrocarbons function as a specific chemosensory conditioned stimulus in this learning paradigm. The effects of training with mated females were determined in courtship tests with either wild-type virgin females as courtship targets, or with target flies of different genotypes that express distinct cuticular hydrocarbon (CH) profiles. Results of tests with female targets that lacked the normal CH profile, and with male targets that expressed typically female CH profiles, indicated that components of this CH profile are both necessary and sufficient cues to elicit the effects of conditioning. Results with additional targets indicated that the female-specific 7,11-dienes, which induce naive males to court, are not essential components of the conditioned stimulus. Rather, the learned response was significantly correlated with the levels of 9-pentacosene (9-P), a compound found in both males and females of many *Drosophila* strains and species. Adding 9-P to target flies showed that it stimulates courting males to attempt to copulate, and confirmed its role as a component of the conditioned stimulus by demonstrating dose-dependent increases in the expression of the learned response. Thus, 9-P can contribute significantly to the conditioned suppression of male courtship toward targets that express this pheromone.

Neural mechanisms that are responsible for behavioral plasticity in *Drosophila melanogaster* have been addressed primarily in the context of a classical conditioning paradigm in which flies learn to associate electric shocks with olfactory cues (Tully and Quinn 1985). As a result of intensive investigations involving diverse experimental approaches and levels of analysis, a detailed model has evolved of the essential neural substrates and processes underlying this form of olfactory classical conditioning (Davis 1996; Roman and Davis 2001; Waddell and Quinn 2001; Dubnau et al. 2003). A more complex associative learning paradigm is experience-dependent courtship conditioning, wherein male courtship behavior is modified by experience with an unreceptive, previously mated female (Siegel and Hall 1979; Tompkins et al. 1983). In spite of the ecological relevance of this form of learning, progress in deciphering its neural mechanisms has been impeded by its complexity, specifically by the multisensory nature of the training experience with mated females (Siwicki and Ladewski 2003).

In his first encounter with a potential mate, a male fruit fly follows the target, taps her cuticle with his foretarsi, and vibrates one wing to generate a courtship song (Hall 1994; Greenspan and Ferveur 2000). If the target decreases her locomotion (Tompkins et al. 1982), the male approaches from the rear, extends his proboscis to contact her posterior abdomen, and attempts to copulate (Bastock and Manning 1955; Hall 1994). Socially naive males court not only virgin females, but also unreceptive targets like quinine or cis-vaccenyl acetate (Ackerman and Siegel 1986; Zawistowski 1988). These studies showed that pairing an aversive chemosensory stimulus with organic extracts of females can produce a conditioned suppression of male courtship toward virgins. In the present study, the specific identity of compounds that function as the conditioned stimulus during training with mated females has been addressed by manipulating the pheromonal profiles of flies used to test courtship memory.

Courting males sample the pheromones of target flies via...
olfactory sensilla on the antennae and maxillary palps and gustatory sensilla on the foretarsi and proboscis (Stocker and Gendre 1989; Stocker 1994; Bray and Amrein 2003). Long-chain hydrocarbons on the cuticle of D. melanogaster are sexually dimorphic and include substances that can either stimulate or inhibit male courtship, suggesting a role as sex pheromones (Jallon 1984; Cobb and Ferveur 1996; Ferveur 1997). Recent evidence for the aphrodisiac and anti-aphrodisiac properties of these cuticular compounds derives from genetic manipulations of the hydrocarbons of living courtship targets. Specifically, UAS-transformer (UAS-tra) in combination with various GAL4 drivers can dramatically alter the cuticular hydrocarbon (CH) profiles of both male and female flies (Ferveur and Sureau 1996; Savarit et al. 1999; Savarit and Ferveur 2002a). We have used these phenomally altered genotypes to test the hypothesis that cuticular pheromones expressed by females are the specific chemosensory cues that function as a conditioned stimulus in courtship conditioning. Moreover, by evaluating individual components of the CH profile for possible correlations with learning and memory, we have identified 9-pentacosene (9-P) as the most important chemosensory cue for eliciting a learned response from trained males. By adding pure 9-P to test flies, we discovered that it stimulates males to attempt to copulate, and that males are less responsive to this aphrodisiac after training with mated females. The results provide direct evidence for the aphrodisiac properties of 9-P and for the identity of the conditioned stimulus responsible for changes in male courtship behavior following experience with mated females.

### Table 1. Cuticular hydrocarbons (CHs) identified in hexane extracts of flies used as courtship targets

| Chain length | No. | Abbreviation | Hydrocarbon          | LIS\(^a\) | LI60\(^d\) |
|--------------|-----|--------------|----------------------|----------|----------|
|              |     |              |                      | R        | P        | R        | P        |
| 23C          | 1   | 7,11-TD      | 7,11-Tricosadiene    | 0.505    | 0.05     | 0.656    | NS       |
|              | 2   | 23Br         | 2-Methyl-dodacocane  | <0.5     | NS       | <0.5     | NS       |
|              | 3   | 9-T          | 9-Tricosene          | 0.555    | 0.03     | 0.684    | NS       |
|              | 4   | 7-T          | 7-Tricosene          | <0.5     | NS       | <0.5     | NS       |
|              | 5   | 5-T          | 5-Tricosene          | <0.5     | NS       | <0.5     | NS       |
|              | 6   | 23Lin        | n-Tricosane          | <0.5     | NS       | <0.5     | NS       |
| 25C          | 7   | 7,11-PD      | 7,11-Pentacosadiene  | <0.5     | NS       | 0.658    | NS       |
|              | 8   | 25Br         | 2-Methyl-tetracocane | <0.5     | NS       | <0.5     | NS       |
|              | 9   | 9-P          | 9-Pentacosene        | 0.653    | 0.006    | 0.823    | 0.01     |
|              | 10  | 7-P          | 7-Pentacosene        | <0.5     | NS       | <0.5     | NS       |
|              | 11  | 5-P          | 5-Pentacosene        | <0.5     | NS       | <0.5     | NS       |
|              | 12  | 25Lin        | n-Pentacosane        | <0.5     | NS       | <0.5     | NS       |
| 27C          | 13  | 9,13-HD      | 9,13-Heptacosadiene  | <0.5     | NS       | 0.546    | NS       |
|              | 14  | 7,11-HD      | 7,11-Heptacosadiene  | <0.5     | NS       | 0.632    | NS       |
|              | 15  | 27Br         | 2-Methyl-hexacosane  | <0.5     | NS       | 0.586    | NS       |
|              | 16  | 9-H          | 9-Heptacosane        | <0.5     | NS       | 0.666    | NS       |
|              | 17  | 7-H          | 7-Heptacosene        | <0.5     | NS       | 0.700    | NS       |
|              | 18  | 5-H          | 5-Heptacosene        | 0.542    | 0.03     | 0.700    | NS       |
|              | 19  | 27Lin        | n-Heptacosane        | <0.5     | NS       | <0.5     | NS       |
| 29C          | 20  | 9,13-ND      | 9,13-Nonacosadiene   | <0.5     | NS       | <0.5     | NS       |
|              | 21  | 7,11-ND      | 7,11-Nonacosadiene   | <0.5     | NS       | 0.566    | NS       |
|              | 22  | 29Br         | 2-Methyl-octacosane  | <0.5     | NS       | <0.5     | NS       |
|              | 23  | 9-N          | 9-Nonacosene         | <0.5     | NS       | <0.5     | NS       |
|              | 24  | 29Lin        | n-Nonacosane         | <0.5     | NS       | <0.5     | NS       |

\(^{a}\)Data in the four columns on the right describe correlations between Learning Indexes (LIS\(^b\) and LI60\(^{d}\)) determined in tests with targets of different genotypes and the levels of 16 unsaturated hydrocarbons in cuticular extracts of the target flies. Six of these correlations are shown as scatterplots in Figure 4. A total of 16 LIS\(^b\) values were determined independently in tests of trained and naive males with nine different target genotypes. The LI60\(^{d}\) data set comprised eight values determined independently in tests with seven different target genotypes. The indicated correlation coefficients (\(R\)) and \(P\)-values were determined by simple regressions (without correcting for multiple comparisons). NS indicates \(P > 0.05\). By using Monte Carlo simulations to evaluate the significance of correlations determined in multiple independent comparisons (see Materials and Methods), 9-P is the only hydrocarbon with which LIS\(^b\) is significantly correlated (\(P < 0.05\)).

\(^{b}\)Refers to the bar graphs in Figure 1, depicting the levels of each CH measured in extracts of individual flies of different genotypes.
stimulate male courtship, but are not necessary for courtship and mating (Antony et al. 1985; Ferveur and Sureau 1996; Savarit et al. 1999). Another aphrodisiac hydrocarbon is 7-pentacosene (7-P, bar #10), which is expressed at similar levels by CS males and females (Antony and Jallon 1982; Antony et al. 1985; Ferveur and Sureau 1996; Savarit and Ferveur 2002a). The most effective known inhibitor of male courtship is 7-tricosene (7-T, bar #4), a 23C monoene (Jallon 1984; Ferveur and Sureau 1996). This antiaphrodisiac is present at low levels in CS female extracts and is the single most abundant CH in extracts of CS males (Fig. 1A,B, bar #4). Because analyses of CS virgin and mated female extracts yielded indistinguishable profiles (data not shown), Figure 1A also represents the CH profile experienced by males during training with mated females. Experimental targets included males and females of various genotypes that expressed different combinations of the compounds in CS female extracts; their CH profiles (Fig. 1C–H) are described below in the contexts of the corresponding behavioral data.

Initial results confirmed that 1 h of training with CS mated females results in significant suppression of male courtship toward CS virgin female targets, in tests at both 5 min and 60 min after training (Fig. 2A). The role of the females’ CH profile in eliciting the males’ learned responses was tested with hs-tra virgin females, which are severely depleted of all CHs (Fig. 1C). Naive males courted hs-tra females actively, confirming that the CHs expressed by normal virgins are not required to elicit male courtship (Savarit et al. 1999). Trained males tested with hs-tra virgins displayed no significant suppression of courtship when tested either 5 min or 60 min after mated-female training (Fig. 2B). Since male courtship of these CH-depleted virgins is not affected by experience with mated females, the results suggest that the conditioned suppression of courtship observed in tests with normal virgins (Fig. 2A) depends on some component(s) of the normal female CH profile.

This interpretation was tested with courtship targets bearing a variety of distinctive CH profiles. These included male flies expressing characteristically female profiles (Fig. 1D,E), virgin females selectively depleted for unsaturated CHs (Fig. 1G), and males expressing subsets of the female CH profile (Fig. 1F,H). For example, males heterozygous for certain enhancer-trap pGAL4 inserts and UAS-tra express female CH profiles (Fig. 1D,E) but retain male morphology (Ferveur et al. 1997; Savarit and Ferveur 2002a). When these GAL4-tra males were used as test targets, they elicited active courtship from naive CS male subjects, while trained males courted them significantly less actively both 5 min and 60 min after training (Fig. 2D,E). These results show that a female CH profile on a decapitated male target is sufficient to elicit active courtship from naive males and is a less effective aphrodisiac cue after subjects have been trained with mated females. They further support the hypothesis that component(s) of the hydrocarbon blend, and not female morphological features, function as a conditioned stimulus.

Three additional target genotypes allowed for independent manipulations of different components of the female CH profile. Males of the wild Tai2 strain naturally express very high levels of monounsaturated pentacosenes 9-P and 7-P (bars #9 and 10, respectively, in Fig. 1F), but no detectable 7,11-dienes, the aphrodisiacs found specifically on CS females. Because 7-P can also act as an aphrodisiac for CS males (Antony et al. 1985; Sureau and Ferveur 1999), naive subjects court actively in the presence of decapitated Tai2 males (Fig. 2F). If 7,11-dienes were essential components of the conditioned stimulus, then trained males would be expected to court Tai2 targets as actively as naives. In fact, trained subjects displayed very strong suppression of court-
Figure 2. The effects of training with mated females on subsequent courtship of Canton-S males toward different targets. The mean Courtship Indexes (CIs) are plotted (± SEM) for groups of Naive (black bars, N = 38–54 per group) and Trained (gray bars, N = 19–37 per group) male subjects tested with decapitated targets of the indicated genotypes, either 5 min (lighter gray bars) or 60 min (dark gray bars) after training. The test targets were (A) Canton-S virgin females, (B) hs-tra virgin females, (C) desat11573 virgin females, (D) 376-GAL4/+; UAS-tra/+ males, (E) Tp2602-GAL4/+; UAS-tra/+ males, (F) Tai2 males, and (G) desat11573 x Tai2 males. Differences between the CIs of Trained groups and Naive controls with each target were evaluated by ANOVA; ** P < 0.01; *** P < 0.001.

Figure 3. The effects of training with mated females on males’ attempts to copulate with different targets during 10-min courtship assays performed 5 min after training. The percentage of males that attempted to copulate with decapitated targets of the indicated genotypes is plotted for groups of Naive (black bars, N = 19–26 per group) and Trained (gray bars, N = 20–26 per group) male subjects. Fewer Trained males than Naives attempted to copulate with (A) Canton-S virgin females (χ² = 6.059, df = 1, [*] P < 0.02) and (D) Tp2602-tra males (χ² = 6.584, df = 1, [*] P < 0.04), but the frequencies of copulation attempts with other targets were independent of subjects’ training status (by χ² tests of association). These targets were (B) hs-tra virgin females, (C) desat1 virgin females, (E) Tai2 males, and (F) desat1 x Tai2 males.
The effects of 9-pentacosene on courtship of naive and trained males

Results of this correlation analysis suggest that courtship conditioning might alter males’ responses to 9-P, such that it either acquires anti-aphrodisiac properties or becomes less potent in its aphrodisiac effects as a consequence of training. We tested this directly by adding pure, synthetic (Z)-9-pentacosene (kindly provided by L.M. Hanks [University of Illinois] and J.G. Millar [University of California–Riverside]) to test targets that naturally express female CHs.

The effects of 9-pentacosene were to increase the fraction of males attempting to copulate with the CH-depleted targets in control tests (Fig. 6B,C,D). These effects of training were less evident with the higher 9-P dose, however, because both naive and trained groups increased their copulation attempts between the 0.2-µg and 2-µg-9-P doses. Both naive and trained males also increased their copulation attempts with CS females between control and 2-µg-9-P conditions (Table 2). Clear increases in LI5 were found both between control and 0.2 µg of 9-P and between 0.2 µg and 2 µg of 9-P, when effects of training were assessed with hs-tra females, desat1 females, and immature CS males. No such effect was found in tests with CS females.

Even more dramatic effects of 9-P emerged when the specific behavioral stages of courtship were analyzed. Results of earlier experiments (see Fig. 3 above) showed that training dramatically decreased the fraction of males attempting to copulate with test targets expressing female CH profiles, while very few males in either trained or naive groups attempted to copulate with the CH-depleted hs-tra or desat1 mutant females. The essential features of these results were confirmed in control conditions (hexane only) of the 9-P experiments: naive males attempted to copulate with CS female targets at a much greater frequency than trained males (Fig. 6A), but similar numbers of trained and naive males attempted to copulate with the CH-depleted targets in control tests (Fig. 6B,C,D). The effects of adding 9-P were to increase the fraction of males attempting to copulate with CS virgin females (Fig. 6A), hs-tra virgins (Fig. 6B), and desat1 virgins (Fig. 6C), but a slight increase in copulation attempts with immature male targets was not significant (Fig. 6D). Effects of training, as indicated by a higher frequency of copulation attempts by naive males than by trained males, were increased between control and 0.2-µg-of-9-P conditions with the CH-deficient courtship targets (Fig. 6B,C,D). These effects of training were less evident with the higher 9-P dose, however, because both naive and trained groups increased their copulation attempts between the 0.2-µg and 2-µg-9-P doses. Both naive and trained males also increased their copulation attempts with CS females between control and 2-µg-9-P conditions. Overall, the results indicate that 9-P increased copulation attempts by CS males; the lower dose of 9-P elicited this effect for naive males, while trained males responded only to the higher dose.

Discussion

The goal of this study was to identify specific female cues that function as a conditioned stimulus in the courtship conditioning paradigm of D. melanogaster, wherein males learn to suppress their courtship toward virgin females as a result of unsuccessful courtship with previously mated females. The results confirm previous suggestions that the conditioned stimulus is a chemo-
sensory cue (Tomkins et al. 1983; Gailey et al. 1984; Ejima et al. 2005), and provide direct evidence that a minor cuticular hydrocarbon functions as a significant component of this conditioned stimulus. Based on the reasoning that trained males would exhibit a learned response only in courtship tests with targets that expressed a conditioned stimulus, flies with different combinations of putative female aphrodisiac cues were used as test targets, and assessed for their ability to elicit a learned response from CS males. After 1 h of experience courting a previously mated CS virgin females (Fig. 2B), Tests with farnesized males (pGAL4/+; UAS-tra/+) that expressed typically female CH profiles revealed that this profile is sufficient both to stimulate naive male courtship and to elicit conditioned suppression of courtship after experience with mated females (Fig. 2B). This raises the question of whether specific compounds or the overall female profile are responsible for the learned response. The data indicate that a target’s ability to elicit a learned response was most strongly correlated with the levels of a single monotene, 9-pentacosene (9-P). Specifically, LIs based on tests either 5 min or 60 min after training were significantly correlated with target levels of a single unsaturated hydrocarbon, 9-P (Table 1). Surprisingly, the LIs were not significantly correlated with levels of any principal compound previously known to possess aphrodisiac properties, that is, the 7,11-dienes or 7-P (Fig. 2D,E). Together, these results strongly support the hypothesis that components of the female CH profile function as the conditioned stimulus in courtship conditioning.

Table 2. Effects of 9-P on Learning Indexes (LIS’S) obtained in tests with different courtship targets

| Courtship target | Control | 0.2 µg 9-P | 2 µg 9-P |
|------------------|---------|------------|---------|
| Canton-S females | 70      | 69         |         |
| hs-tra females   | 20      | 3.4        | 23      |
| desat1 females   | 23      | 40         | 47      |
| Immature Canton-S males | 22 | 34       | 45      |

The courtship activity (mean CI) of naive males was not well correlated with the hydrocarbon profiles of the different test targets, confirming the results of prior studies showing that female CHs are sufficient, but not necessary, to stimulate naive male courtship (Antony et al. 1985; Savarit et al. 1999). In particular, naive males courted hydrocarbon-depleted hs-tra virgins and males depleted of unsaturated CHs as actively as they courted targets with typically female CH profiles. This is consistent with the view that naive males respond to a combination of aphrodisiac and anti-aphrodisiac pheromones on a decapitated target fly. In contrast, the conditioned suppression of courtship resulting from experience with mated females was closely correlated with the hydrocarbon profiles of the test targets. Results of testing with hydrocarbon-depleted female targets indicated that CHs are essential stimuli for the conditioned suppression of courtship after experience with mated females (Fig. 2B). Tests with farnesized males (pGAL4/+; UAS-tra/+) that expressed typically female CH profiles revealed that this profile is sufficient both to stimulate naive male courtship and to elicit conditioned suppression of courtship in trained male subjects (Fig. 2D,E). Together, these results strongly support the hypothesis that components of the female CH profile function as the conditioned stimulus in courtship conditioning.

This raises the question of whether specific compounds or the overall female profile are responsible for the learned response. The data indicate that a target’s ability to elicit a learned response was most strongly correlated with the levels of a single monotene, 9-pentacosene (9-P). Specifically, LIs based on tests either 5 min or 60 min after training were significantly correlated with target levels of a single unsaturated hydrocarbon, 9-P (Table 1). This raises the question of whether specific compounds or the overall female profile are responsible for the learned response. The data indicate that a target’s ability to elicit a learned response was most strongly correlated with the levels of a single monotene, 9-pentacosene (9-P). Specifically, LIs based on tests either 5 min or 60 min after training were significantly correlated with target levels of a single unsaturated hydrocarbon, 9-P (Table 1). This raises the question of whether specific compounds or the overall female profile are responsible for the learned response. The data indicate that a target’s ability to elicit a learned response was most strongly correlated with the levels of a single monotene, 9-pentacosene (9-P). Specifically, LIs based on tests either 5 min or 60 min after training were significantly correlated with target levels of a single unsaturated hydrocarbon, 9-P (Table 1). This raises the question of whether specific compounds or the overall female profile are responsible for the learned response. The data indicate that a target’s ability to elicit a learned response was most strongly correlated with the levels of a single monotene, 9-pentacosene (9-P). Specifically, LIs based on tests either 5 min or 60 min after training were significantly correlated with target levels of a single unsaturated hydrocarbon, 9-P (Table 1). This raises the question of whether specific compounds or the overall female profile are responsible for the learned response. The data indicate that a target’s ability to elicit a learned response was most strongly correlated with the levels of a single monotene, 9-pentacosene (9-P). Specifically, LIs based on tests either 5 min or 60 min after training were significantly correlated with target levels of a single unsaturated hydrocarbon, 9-P (Table 1).
This was confirmed by Ferveur and Sureau (1996), using genetic manipulations to produce live targets with distinctive CH profiles, who also found that even low levels of 7,11-ND (10–15 ng) could strongly potentiate the aphrodisiac effects of 7,11-HD. Both strategies produced evidence for the anti-aphrodisiac properties of 7-T and the aphrodisiac effects of 7-P (Antony et al. 1985; Sureau and Ferveur 1999). Prior evidence regarding the putative aphrodisiac properties of 9-P is limited to a positive correlation with the frequency of copulation attempts (Ferveur and Sureau 1996). When applied to hydrocarbon-stripped targets, 9-P failed to stimulate male courtship (Antony et al. 1985).

In the present study, one clear effect of adding exogenous, pure (Z)-9-pentacosene to courtship chambers was to increase the fraction of CS males that attempted to copulate with CS virgin females, and with CH-depleted virgins of the hs-tra and desatI genotypes (Fig. 6). This result strongly supports the hypothesis that the main aphrodisiac effect of 9-P is to increase the likelihood of copulation attempts (Ferveur and Sureau 1996; Ferveur 1997). Interestingly, 9-P did not increase the mean CI in most of these test groups: Only naive males tested with desatI females had a higher mean CI in the presence of 9-P (Fig. 5C). This selective effect on the mean CI of naive males with desatI females, but not trained males, as well as the dose-dependent effects of 9-P on the Learning Index (Table 2), clearly establish a role for 9-P as a component of the conditioned stimulus. With most targets, however, 9-P did not stimulate overall courtship activity; rather, it increased the probability for courting males to attempt copulations. Moreover, because this effect was elicited by the lower dose of 9-P for naive males, and only at the higher dose for trained males, it suggests that a specific effect of training with mated females is to reduce the responsiveness of trained males to the aphrodisiac effects of this pheromone.

These results indicate a dual role of 9-P: (1) increasing the probability that courting males will attempt to copulate, and (2) contributing to the difference between naive and trained males in overall courtship. Yet, depending on the target genotype, different doses of 9-P incurred different effects on the global amount of courtship (CI) and on the frequency of copulation attempts. For example, 9-P only increased the CI of naive males when present at 2 µg on desatI females, but both 0.2-µg and 2-µg doses tended to increase the probability to attempt copulation when combined with any of the four targets. This difference can be explained because the duration of a copulation attempt is very brief compared to the wing vibration and licking behaviors that make up most of the CI measure (Ferveur and Sureau 1996). The selective effect of 9-P on CI with desatI females suggests that its effect could be potentiated by other CHs that are present in these targets but absent in hs-tra females and immature males. At least two relatively abundant CHs fit this criterion: n-tricosane (#6; Table 1) and 2-methyl-tetradecane (#8). On the other hand, it is likely that 9-P had minimal effects on CIs with CS females because all the CHs necessary to induce high levels of courtship were already present, and the exogenous 9-P did not increase the overall sex appeal of these targets. It was also with the three CH-depleted genotypes, but not with CS females, that dose-dependent effects of 9-P were found on LIs, a measure of the relative suppression of overall courtship in trained males, whereas 0.2 µg of 9-P was sufficient to elicit the maximum effect of training on the probability to attempt copulation. It is worth noting that the 0.2-µg dose of 9-P (applied to a filter paper under the target fly) is very similar to the biological dose found on CS females (50–70 ng). Thus, the results demonstrate that CS males can detect and respond to physiologically relevant levels of this pheromone.

It is likely that 9-P is one component of a combination of CHs that function as the conditioned stimulus. This is supported by the fact that 9-P was not sufficient to elicit effects of training on courtship of CH-depleted hs-tra females (Figs. 5B, 6B). It is also supported by the modest, albeit significant, correlation between the LIs and 9-P levels in different test targets (Fig. 4A). When possible synergistic effects of other unsaturated CHs with 9-P were examined by running multiple regressions, there were no cases in which the levels of 9-P plus another CH predicted the LI-values significantly better than did 9-P alone. Thus, the identities of the putative interacting components of the conditioned stimulus remain to be determined.

The surprising result of this study is that this particular compound emerged as a conditioned stimulus. The levels of 7,11-dienes in a courtship target were not correlated with the expression of the males’ learned response (Fig. 4E,F), indicating that conditioning does not dramatically alter the potency of these highly abundant, female-specific pheromones. In contrast, 9-P is

Figure 6. The effects of exogenous 9-P on the frequency of copulation attempts of Naive (black bars) and Trained (gray bars, tested 5 min after training) males with different courtship targets, from the same experiment for which CIs are plotted in Figure 5. Possible effects of training and of 9-P were evaluated independently by χ² tests of association. The effect of 9-P (with Naive and Trained groups combined; thus brackets are positioned above each pair of bars) was to significantly increase the percent of males attempting to copulate with (A) Canton-S females (χ² = 6,988, df = 1, [**] P < 0.01), (B) hs-tra females (χ² = 11,01, df = 2, [***] P < 0.01), and (C) desatI females (χ² = 8,200, df = 2, [*] P < 0.02), but not with (D) immature Canton-S males (χ² = 2,800, df = 2, P = 0.2). Similarly, effects of training were assessed across all 9-P doses; these tests revealed a significant association between training status and attempts to copulate with (A) Canton-S females (χ² = 18,26, df = 1, P < 0.0001), (C) desatI females (χ² = 5,927, df = 1, P < 0.02), and (D) immature Canton-S males (χ² = 7,420, df = 1, P < 0.01), but not with (B) hs-tra females (χ² = 0.001, df = 1, P = 0.97).

1985).
a relatively minor component of the Canton-S CH profile, and is expressed by both mature males and females.

Having identified a role for 9-P as a chemosensory stimulus that is particularly susceptible to conditioning during courtship of mated females, the results may shed new light on the question of whether courtship conditioning ability is relevant to the behavior of males in the wild. While it is likely to be maladaptive for males to suppress their courtship toward conspecific virgin females, Gailey and colleagues (1984) proposed that courtship toward mated females might be decreased to a greater degree, thus improving male selectivity for virgins. As 9-P is not very specific (it is found in cuticular extracts of both males and females of several Drosophila species) (Jallon and David 1987; Howard et al. 2003), it is possible that experience-dependent modification of the responsiveness to this widely expressed hydrocarbon might provide a male with the ability to learn to discriminate appropriate from inappropriate courtship targets. For example, females of Drosophila affinis were reported to express much higher levels of 9-P than D. melanogaster females (Jackson 1996; Ferveur 2005). These two species are sympatric in some areas of North America, and they have been observed courting in interspecific groups gathered on food sources (McRobert and Tompkins 1986). These observations suggest that the ability of CS males to learn to be less responsive to 9-P might allow them to learn to avoid courting D. affinis females. In an environment where misdirected courtship of heterospecific females is likely to occur, there would presumably be some advantage for males that could learn to become less sensitive to particular pheromones that are more abundant in heterospecific females than in conspecifics.

D. melanogaster males also court immature males and females, which express distinctly different blends of CHs than mature flies (Pechine et al. 1988; Ejima et al. 2005). Experience conditioning immature males results in habituation to the aphrodisiac effects of specific 31C monoenes in the immature male CH blend (Gailey et al. 1982; Vaia et al. 1993). The CH profiles of immature males and females are similar to each other in that they are comprised mostly of longer 27C–37C compounds and contain only trace amounts of compounds smaller than 27C (Pechine et al. 1988; Ejima et al. 2005). It is difficult to predict how experience courting a mated female, and the corresponding decrease in aphrodisiac potency of 9-P, might affect a male’s subsequent courtship of immature flies with little or no 9-P and an abundance of other CHs with unknown behavioral effects. This difficulty is compounded by the possibility that some longer-chain CHs of immature flies may be detected and processed through some of the same sensory pathways as mature CH pheromones, a possibility that is reinforced by recent evidence that many gustatory receptor neurons express combinations of receptors (Amrein 2004; Thorne et al. 2004). At present, therefore, it is not possible to interpret the effects of mated-female training on male courtship of immature males and females (Ejima et al. 2005) in terms of specific modifications of responses to specific CHs.

The present results suggest an additional level of complexity in the functional organization of the neural systems that control and modulate male courtship. Some CHs (7,11-dienes and 7-P) stimulate naive males to court actively; and others (7-T) inhibit naive courtship, suggesting that there are at least two distinct gustatory pathways by which contact pheromones influence courtship control centers (Bray and Amrein 2003; Stockinger et al. 2005). Males repeatedly sample all of these pheromones during training with mated females, yet the experience-dependent changes in subsequent courtship behavior are strongly correlated only with targets’ levels of 9-P. This suggests that the sensorimotor circuits by which 7,11-dienes and 7-P stimulate courtship may be less modifiable than pathways activated by 9-P. It follows from this hypothesis that these pheromones may be detected and processed by distinct neural pathways, a prediction that can be tested by manipulating the genes and cells involved in pheromone detection (Xu et al. 2002; Bray and Amrein 2003; Stockinger et al. 2005).

In this study, we have provided direct support for the hypothesis that the conditioned stimulus in courtship conditioning of D. melanogaster males is a chemical, rather than behavioral or anatomical, cue provided by the female (Tompkins et al. 1983; Gailey et al. 1984). Our results indicate that components of the female cuticular hydrocarbon profile function as the conditioned stimulus in courtship conditioning. In particular, the most relevant stimulus is 9-pentacosene, an unsaturated hydrocarbon found on the cuticle of both males and females. Together with previous evidence concerning the relative potencies of various cuticular substances at eliciting male courtship, the present results suggest that naive males respond to a combination of aphrodisiacs and anti-aphrodisiacs on a target fly, while conditioned males are less responsive than naïves to the aphrodisiac effects of 9-P. Identifying the sensory pathways responsible for detection of this chemical conditioned stimulus will allow for a more definitive analysis of the neural mechanisms responsible for this form of associative learning.

Materials and Methods

Drosophila strains

All flies were reared at 25°C on cornmeal-sucrose-yeast-agar medium, and were exposed to a 12-h light/12-h dark cycle. Wild-type Canton-S (CS) and Tai2 strains and transgenic strains for pheromonal manipulations (UAS-tra 20J7 and pGALA4 enhancer trap strains) were from lines maintained in the Ferveur lab. The hsP70-GALA4.PB strain was obtained from the Bloomington Stock Center (1799).

Hydrocarbon-depleted hs-tra females were the progeny of crosses between UAS-tra 2017 males and hsP70-GALA4.PB females: hsP70-GALA4.PB/+; UAS-tra/+; UAS-tra/+ virgin females were subjected to a 1-h heat shock (37°C) 5–6 h after eclosion (Savarit et al. 1999; Savarit and Ferveur 2002b). It is thought that early toxic effects of hsP70-GALA4 induction in cells responsible for CH synthesis (i.e., oenocytes) may explain the near absence of detectable CHs in these flies 4–5 d later (Savarit and Ferveur 2002b), when they were used as test targets in courtship assays. Male targets with female CH profiles are the progeny of crossing UAS-tra 2017 males with females of different enhancer trap GAL4 strains (27C, GAL4, Tp2602-GAL4, and Tp3502-GAL4), which drive expression in oenocytes and thereby feminize the profile of CHs in the respective GAL4/+; UAS-tra/+ male progeny (Savarit and Ferveur 2002a). In desaturase1 mutant females, a pGAL4 insertion in the desaturase1 gene on Chromosome III results in a dramatic reduction of unsaturated CHs and increase in saturated compounds (Marcillac et al. 2005). males of the desaturase1<sup>−/+</sup> x Tai2 genotype were from a stock in which Chromosome II of the Tai2 strain (that changes the 7-T; 7-P ratio) was combined with Chromosome III of the desaturase1<sup>−/+</sup> strain.

Hydrocarbon extraction and gas chromatography

Cuticular hydrocarbons (CHs) of individual flies were extracted by a standard hexane wash, and levels of identified hydrocarbons were measured by gas chromatography, as previously described (Ferveur 1991; Savarit and Ferveur 2002a). Peaks were identified based on time of elution, and amounts were calculated based on comparison to internal standards. Data from 6–12 individual flies were averaged; there was little variation in CH levels among flies of the same genotype.

Behavioral assays

All flies were collected under CO₂ anesthesia within 5 h of eclosion. CS male subjects were aged for 4–5 d in individual small
glass tubes containing normal cornmeal medium. They were transferred to fresh individual tubes 15–24 h prior to experiments. CS females used for training were aged 4–6 d in groups of 5–15, then paired individually with CS males for pre-making.

Females observed to mate were stored in groups of five until being used to train subject males the following day.

Training and testing took place during the first 5 h after lights-on, at ambient temperature (23–26°C) with fluorescent room lighting. Courtship chambers were created by placing a watch glass (3.5 cm²) on a glass plate (Savarit et al. 1999). For the training phase, healthy CS males were placed in individual chambers; 5 min later, mated females were randomly added to about half the chambers. Both sham-trained males (Naïve) and males with mated females (Trained) were left in the chambers for 60 min, then the male subjects were aspirated to clean courtship chambers. After 5 or 60 min, a decapitated target fly was added to assay the male’s courtship activity.

Target flies were anesthetized with CO₂ and decapitated –20 min prior to testing. Only targets that resumed a standing position were used. Once a target was added to a testing chamber, an observer who was blind to the training status of the subject flies recorded the total duration of all courtship activity by each subject male during a 10-min testing period. The latency to initiate court and the occurrence of specific behaviors (tapping, wing vibration, licking, and attempting copulation) were also recorded.

In some experiments, synthetic (Z)-9-pentacosene (kindly provided by J.G. Millar and L.M. Hanks) was added to filter paper disks (Whatman #1) in courtship chambers in a 20-µL volume of hexane, and the hexane was allowed to evaporate before the flies were added. Control tests were performed on filter disks treated with 20 µL of hexane alone.

Data analyses
A Courtship Index (CI) was calculated for each male subject by dividing the total time courting by the duration of the testing period (600 sec). After confirming that the CI values were approximately normally distributed within each group, the CIs of naïve and trained groups with each type of target were compared by ANOVAs (using StatView 5.0.1). The effects of training and of 9-Δ dose on the fraction of males in each group that attempted to copulate with a target were evaluated by χ² tests of association (using JMP 5.0.1).

Learning Indexes (LIs) were calculated to express the effects of training on males tested at 5 min and 60 min with each target type.

Learning Index (LI) = \[ \frac{\text{mean Naïve} - \text{mean Trained} \times 100}{\text{mean Naïve}} \]

Correlations between LI values and the levels of 16 individual hydrocarbons in cuticular extracts of target genotypes were determined by simple regression analyses (using StatView 5.0.1). An adjustment for multiple comparisons was determined by running Monte Carlo simulations, in which the maximum correlation coefficient \( R_{max} \) was determined for 1000 random data sets of 16 Y-values and 16 different randomly generated X-values. The simulation indicated that maximum correlation coefficients greater than \( R_{max} = 0.647 \) occurred with \( P < 0.05 \).

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