Research

Inhibition of different histone acetyltransferases (HATs) uncovers transcription-dependent and -independent acetylation-mediated mechanisms in memory formation

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Acetylation of histones changes the efficiency of the transcription processes and thus contributes to the formation of long-term memory (LTM). In our comparative study, we used two inhibitors to characterize the contribution of different histone acetyltransferases (HATs) to appetitive associative learning in the honeybee. For one we applied garcinol, an inhibitor of the HATs of the p300 (EP300 binding protein)/CBP (CREB-binding protein) family, and the HATs of the PCAF (p300/CBP-associated factor) family. As comparative agent we applied C646, a specific inhibitor that selectively blocks HATS of the p300/CBP family. Immunohistochemical analysis reveals differences in histone H3 acetylation in the honeybee brain, in response to the injection of either C646 or garcinol. Behavioral assessment reveals that the two drugs cause memory impairment of different nature when injected after associative conditioning: processes disturbed by garcinol are annihilated by the established transcription blocker actinomycin D and thus seem to require transcription processes. Actions of C646 are unaltered by actinomycin D and thus seem to be independent of transcription. The outcome of our different approaches as summarized suggests that distinct HATs contribute to different acetylation-mediated processes in memory formation. We further deduce that the acetylation-mediated processes in memory formation comprise transcription-dependent and transcription-independent mechanisms.

Short-term memory relies on post-translational protein modifications. The formation of long-term memory (LTM) requires gene expression, in addition. LTM and underlying gene expression are regulated by transcription factors, the recruitment of transcriptional coactivators, and by the chromatin structure itself. Remodeling of chromatin is mediated by DNA methylation and post-translational modifications of histones. In particular, the reversible and dynamic acetylation on the amino-terminal tails of histones has been identified as critical regulator of transcriptional processes in neuronal plasticity and memory formation (Sharma 2010; Lubin et al. 2011; Zovkic et al. 2013). Acetylation is mediated by histone acetyltransferases (HATs), and is reversed by histone deacetylases (HDACs) (Selvi et al. 2010; Haggarty and Tsai 2011). Knockout of transcriptional coactivators with intrinsic HAT activity, such as CBP (CREB-binding protein), p300 (EP300 binding protein), PCAF (p300/CBP-associated factor) leads to an impairment of LTM in aversive and appetitive learning (Oliveira et al. 2007, 2011; Barrett et al. 2011). Accordingly, the knockout of HDACs results in an enhanced LTM (Guan et al. 2009; McQuown and Wood 2011). This picture is supported by studies using pharmacological tools to target different HATs and HDACs (Dekker and Haisma 2009; Bowers et al. 2010; Selvi et al. 2010). Focusing here on the HATs, which have been tested in different invertebrate and mammalian learning paradigms (Marek et al. 2011; Merschbaecher et al. 2012; Zhao et al. 2012; Maddox et al. 2013a,b), it has not been addressed how the different HATs (CBP, p300, PCAF, etc.) contribute to particular mechanisms in memory formation.

The vast majority of work deals with acetylation in LTM processes (Sharma 2010; Lubin et al. 2011; Zovkic et al. 2013). Interestingly, manipulation of acetylation-dependent processes by the HAT inhibitor garcinol also affects transient memory in honeybees (Merschbaecher et al. 2012). This transient memory has been shown to be insensitive to the transcription blocker actinomycin D (Grünbaum and Müller 1998; Wüstenberg et al. 1998). Considering this, we assume that acetylation-dependent processes can be found in transcription-dependent as well as in transcription-independent mechanisms.

Employing the two HAT inhibitors, garcinol and C646, we analyzed the role of different HATs in the formation of appetitive olfactory memory in honeybees. Garcinol is a naturally occurring product extracted from the plant Garcinia indica. It inhibits PCAF (IC50 ≈ 5 μM) representatives of the GNAT (Gcn5-related N-acetyltransferase) family and HATS of the p300/CBP family (IC50 ≈ 7 μM) (Balasubramanyam et al. 2004). The second inhibitor, C646, exclusively inhibits HATS of the p300/CBP family (IC50 < 1 μM) (Bowers et al. 2010). To identify transcription-dependent processes we applied the two inhibitors also in combination with the transcription blocker actinomycin D. Actinomycin D intercalates into DNA and thus impairs transcription (Sobell 1985). Our comparative approach demonstrates that these two HAT inhibitors differ in their effects on histone acetylation and their impact on memory formation. Moreover, coapplication...
of a general inhibitor of transcription, actinomycin D, uncovers processes mediated by the distinct HATs that differ in their sensitivity to this inhibitor.

Results

Garcinol and C646 differ in their effects on histone acetylation in the honeybee brain

HATS are highly conserved between species and the domains essential for inhibitor interaction exist in mammals and invertebrates. This is also true for the amino acids indispensable for the specific interaction between C646 and HATS of the p300/CBP family (Bowers et al. 2010), which are conserved between honeybee and mammals (Fig. 1). Different studies show that p300 and CBP preferentially acetylates lysine 18 of histone H3 (H3K18), while HATS of the PCAF family favor acetylation of H3K9 (Schlitz et al. 1999; Horwitz et al. 2008; Jin et al. 2011). To confirm the different actions of the used HAT inhibitors in honeybees, we measured the levels of H3K9ac and H3K18ac after injection of garcinol or C646. We applied C646 (10 μM) and garcinol (60 μM) at concentrations and time points that have been used previously (Bowers et al. 2010; Federman et al. 2012; Merschbaecher et al. 2012). Two hours after injection we quantified the relative amount of H3, H3K9ac, and H3K18ac in each of the brains by using the ELISA technique (Fig. 2). The H3 signals did not differ between the groups (Student’s t-test, garcinol/vehicle: t = 0.25, df = 18, P = 0.81; C646/vehicle: t = 0.61, df = 22, P = 0.55) and thus are used to normalize the H3K9ac and H3K18ac signals in each of the samples. Garcinol causes a significant decrease in the relative acetylation levels of H3K9 (Student’s t-test, H3K9ac/H3: t = 2.48, df = 18, P = 0.022) and H3K18 (Student’s t-test, H3K18ac/H3: t = 2.55, df = 18, P = 0.02). Injection of the p300/CBP-specific inhibitor C646 only decreases the level of H3K18ac (Student’s t-test, H3K18ac/H3: t = 2.8, df = 22, P = 0.04) but not that of H3K9ac/H3: t = 0.25, df = 22, P = 0.8). This is in agreement with the observations from mammalian systems and demonstrates the distinct action of garcinol and C646 on HATS and thus acetylation processes in the honeybee brain.

Garcinol and C646 do not affect HDAC activity in the honeybee brain

HATS and HDACs closely interact with each other and regulate their activities by auto-acetylation, acetylation and deacetylation (Selvi et al. 2010; Valor et al. 2013; Eom et al. 2014). The used HAT inhibitors may thus affect HDAC activity. Since this aspect has not been addressed in insects we tested for potential effects of garcinol and C646 on HDAC activity in the honeybee brain. The inhibitors were either injected into animals followed by fast brain dissection or added to brain homogenates. The HDAC activities were measured immediately (Table 1). To verify the sensitivity of the used HDAC assay we included the HDAC inhibitor TSA (Trichostatin A) as positive control in each of the experiments. As expected, TSA reduces HDAC activity significantly in all setups. This test result legitimates the employed HDAC assay. The HAT inhibitors garcinol and C646 do not affect HDAC activity in the honeybee brain (Table 1). This excludes the possibility that the changes in histone acetylation (Fig. 2) are due to indirect effects of garcinol and C646 on HDAC activities.

HAT inhibitors garcinol and C646 impair different memory phases

Before analyzing the impact of the HAT inhibitors on appetitive olfactory learning we ensured that C646 and garcinol had no effect on the processing of the sensory stimuli used for associative conditioning (odor and sucrose solution). As test we employed the responsiveness to gustatory stimuli (0–1 M sucrose solution) applied to the antennae and in sensitization and habituation tests. As shown in Table 2 neither of the tests showed an effect of injected garcinol or C646, thus excluding an interference of garcinol or C646 with sensory processing or nonassociative learning such as habituation or sensitization. Having found no

Figure 1. Alignment of C646 binding site of human p300 and honeybee homologue. Multiple sequence alignment of the acetyl transferase domain of human p300 (3BIY/A) (H. sapiens) and honeybee homologue (XP_006568997.1) (A. mellifera) was performed by ClustalO (http://www.ebi.ac.uk/Tools/msa/clustalo/). The domain of interaction including the amino acids (bold) essential for the specific C646 interaction with p300 is shown.

Figure 2. Systemic application of HAT inhibitor C646 or garcinol causes different changes in acetylation at position H3K9 and H3K18 in honeybee brain. Garcinol (A) or C646 (B) and the vehicle (DMSO) were injected into the hemolymph of honeybees. The central brains without optical lobes were dissected 2 h after injection and the levels of H3K18ac and H3K9ac were quantified and related to the levels of H3 in each of the samples. After normalization to the corresponding control ratio of a given experiment, the data were pooled and the relative mean ± SD was calculated. The numbers of animals tested are shown in brackets. Asterisks indicate significant differences (Student’s t-test (two-tailed); * P < 0.05) (details in Results).
The table below shows the HDAC activity in honeybee brain is not affected by HAT inhibitors garcinol or C646.

**Table 1. HDAC activity in honeybee brain is not affected by HAT inhibitors garcinol or C646**

| Application | Vehicle | Garcinol (60 μM) | C646 (10 μM) | TSA (10 μM) |
|-------------|---------|------------------|---------------|-------------|
| To brain homogenate | 1.00 ± 0.24 (9) | 0.93 ± 0.17 (9) | 0.97 ± 0.21 (9) | 0.08 ± 0.05 (9) |
| |  | P = 0.65 | P = 0.68 | P < 0.001 |
| Injection into hemolymph | 1.00 ± 0.14 (10) | 0.97 ± 0.22 (12) | 0.89 ± 0.19 (12) | 0.05 ± 0.01 (12) |
| |  | P = 0.78 | P = 0.16 | P < 0.01 |

HDAC activity was determined after adding the indicated drugs to brain homogenate or immediately after brain dissection if honeybees had received hemolymph injection. Final concentration in the homogenate or in the bee (as estimated by body weight) is indicated in brackets. In each of the experiments, the data were normalized with respect to the vehicle (DMSO) control group. The number of tested homogenate samples or injected bees is indicated in brackets. One-way ANOVA: df = 3, F = 60 P < 0.001; Tukey HSD test: TSA-treated group versus each of the other groups (P < 0.01). Groups treated with vehicle, garcinol, or C646 do not differ from each other.

Unwanted side effects of garcinol or C646 injection, we tested the contribution of garcinol and C646 on associative memory formation. In accordance with all previous studies that addressed the role on memory formation we injected the HAT inhibitors 30 min after associative conditioning. First we tested the effects of the HAT inhibitors on memory formation after strong associative training (three-trial conditioning). Three-trial conditioning induces a stable LTM (Grünbaum and Müller 1998; Wüstenberg et al. 1998). Garcinol (60 μM final concentration in the bee) injected 30 min after conditioning does not affect memory tested at 2 h, but drastically reduces memory tested at 1 and 2 d (χ^2 test, 1 d: c^2 = 12.9, P = 0.0002; 2 d: c^2 = 10.56, P = 0.0007) (Fig. 3A). Likewise injected C646 (10 μM final concentration in the bee) significantly suppresses memory at all measured time points after strong conditioning (χ^2 test, 2 h: c^2 = 5.24, P = 0.02; 1 d: c^2 = 10.5, P = 0.0008; 2 d: c^2 = 12.84, P = 0.0002) (Fig. 3B).

Testing the effects of the HAT inhibitors on memory induced by weak training (single-trial conditioning) reveals further differences between garcinol and C646. Garcinol-injected 30 min after single-trial conditioning only impairs memory tested at 1 d (χ^2 test, c^2 = 7.32, P = 0.006), but has no effect on memory tested 2 h (χ^2 test, c^2 = 0.1, P = 1) and 2 d after conditioning (χ^2 test, c^2 = 0.1, P = 0.7) (Fig. 4A). C646 injected after single-trial conditioning reduces memory at all measured time points after strong conditioning (χ^2 test, 2 h: c^2 = 4.85, P = 0.02; 1 d: c^2 = 4.07, P = 0.03; 2 d: c^2 = 6.31, P = 0.008) (Fig. 4B) just like it does after three conditioning trials.

Blocking transcription has different effects on memory impairment by garcinol and C646

Although different, the memory phases affected by the two HAT inhibitors have one feature in common, they are all insensitive to treatment with the transcription blocker actinomycin D (Grünbaum and Müller 1998; Wüstenberg et al. 1998). HATs can also acetylate proteins that are not implicated in transcriptional regulation (Friedmann and Marmorstein 2013). Memory impairments caused by HAT inhibitors could therefore be differently susceptible to actinomycin D treatment. In our respective experiments, we reconfirmed that injection of actinomycin D 30 min after three-trial conditioning does not affect memory tested up to 2 d (Fig. 5A) (χ^2 test, 2 h: c^2 = 0.04, P = 0.84; 1 d: c^2 = 0.03, P = 0.86; 2 d: c^2 = 0, P = 1). Bees injected with actinomycin D in a mixture with garcinol show no memory impairment (Fig. 5B) (χ^2 test, vehicle versus garcinol + Act D; 1 d: c^2 = 0.4, P = 0.53; 2 d: c^2 = 0, P = 1. Vehicle versus garcinol; 1 d: c^2 = 10.3, P = 0.001; 2 d: c^2 = 5.2, P = 0.017). Bees injected with C646 in a mixture with actinomycin D show a similar low performance as bees injected with C646 alone (Fig. 5C) (χ^2 test, vehicle versus C646; 2 h: c^2 = 9.6, P = 0.001; 1 d: c^2 = 10.7, P = 0.001; 2 d: c^2 = 10.9, P = 0.001. Vehicle versus C646 + Act D; 2 h: c^2 = 10.9, P = 0.001; 1 d: c^2 = 16.3, P = 0.0001; 2 d: c^2 = 10.9, P = 0.001).

Single-trial conditioning again results in a memory different to that caused by three-trial conditioning. In accordance with the aforementioned studies, actinomycin D injection after single-trial conditioning does not affect the memory induced (Fig. 6A) (χ^2 test, 2 h: c^2 = 0.86; 1 d: c^2 = 0, P = 1). Nonetheless, injection of actinomycin D and garcinol in a mixture leaves memory at 1 d after single-trial conditioning intact, quite in contrast to garcinol alone (χ^2 test, 1 d: vehicle versus garcinol, c^2 = 20.5, P = 0.001. Vehicle versus garcinol + Act D, c^2 = 0.45, P = 0.5) (Fig. 6B).

C646 injected in a mixture with actinomycin D leaves bees as memory-impaired (χ^2 test, 2 h: c^2 = 4.6, P = 0.02; 1 d: c^2 = 5.9, P = 0.013; 2 d: c^2 = 6.6, P = 0.007) as it C646 was injected alone (χ^2 test, 2 h: c^2 = 4.6, P = 0.023; 1 d: c^2 = 5.2, P = 0.014; 2 d: c^2 = 5.9, P = 0.01) (Fig. 6C). Thus, actinomycin D interferes with the memory impairment caused by garcinol, while the impairing action of C646 is not affected by actinomycin D in both, single- and three-trial conditioning.

**Table 2. HAT inhibitors C646 and garcinol do not affect gustatory responsiveness or nonassociative learning**

| Behavioral test | Vehicle | C646 | Garcinol |
|-----------------|---------|------|---------|
| Gustatory response score | 2.9 (35) | 2.6 (35) | 2.6 (30) |
| Sensitization (PER) | 10% (40) | 13% (40) | 8% (30) |
| Habituation criterion | 24 ± 12 (50) | 26 ± 15 (50) | 24 ± 8 (30) |

Gustatory responsiveness, sensitization, and habituation were tested 2 h after injection of garcinol, C646, or vehicle (DMSO). The number of bees tested in the different groups is indicated in brackets. The means of the gustatory response scores do not differ between the groups (Mann–Whitney test). Sensitization is presented as the percentage of animals that elicited the probe’s (PER) to an odor stimulus 15 sec after a sensitizing stimulus (1 M sucrose) to the antennae. Fisher exact test (2 x 3) shows no difference. The mean ± SD of the habituation criterion does not differ between the different groups (Student’s t-test).

**Discussion**

By comparing the effects of two different HAT inhibitors on memory formation we demonstrated that: (i) garcinol and C646 affect different memory phases and (ii) when injected in a mixture with actinomycin D, garcinol does not affect memory formation. This does not count for C646. Both findings are new facets of our understanding of memory formation. Previous studies addressed the function of either garcinol or C646 in memory formation (Marek et al. 2011; Merschbaecher et al. 2012; Zhao et al. 2012; Maddox et al. 2013a,b). In mice, injection of garcinol (Maddox et al. 2013a) or C646 (Maddox et al. 2013b) into the amygdala impairs newly acquired as well as reactivated fear memories. Both inhibitors show the same effect in independent studies. They impair memory at 24 h (LTM) but not at 3 h (STM), no matter whether...
tested following conditioning or following memory reactivation. Since both inhibitors act on p300 and CBP the authors propose a requirement of both in amygdala-dependent fear conditioning.

In our comparative approach we now show that the p300/CBP inhibitors C646 and garcinol (garcinol also inhibits HATs of the PCAF family) differ in their effects on appetitive associative memory in honeybees, suggesting different functional pathways. Unlike garcinol, C646 causes a memory impairment starting 2 h after conditioning. The transcription blocker actinomycin D cannot alter this effect, which suggests an implication of cellular processes mediated via acetylation of nonhistone proteins by C646-sensitive p300/CBP.

Studies in different species reveal convincing evidence for a fundamental role of post-translational protein acetylation in basic cellular processes by regulating stability, localization, interaction, and activity of proteins (Choudhary et al. 2009; Spange et al. 2009; Yuan and Marmorstein 2013). Members of the HAT family p300/CBP sensitive to C646 are known for their role in transcriptional regulation and histone modification. They also acetylate nonhistone substrates (Sadoul et al. 2007; Friedmann and Marmorstein 2013; Valor et al. 2013). However, investigations on the role of protein acetylation, especially on nonhistone targets, are at an early stage. Although not derived from investigations on neuronal plasticity, the list of nonhistone proteins targeted by HATs from the p300/CBP and MYST families (named for the founding members in yeast: MOZ, YBF2, SAS2 and TIP60) contains candidates such as STAT, NF-κB, steroid receptors, importin-α. They all contribute to processes underlying neuronal plasticity and learning (Nicolas et al. 2013; Srivastava et al. 2013; Snow et al. 2014; Panayotis et al. 2015) and therefore qualify as downstream targets of C646.

Comparatively dealing with garcinol we could contradict the well-documented notion that garcinol-sensitive memory is transcription independent (Gründen and Müller 1998; Wüstenberg et al. 1998). Garcinol injected in a mixture with the transcription blocker actinomycin D should, under the known assumption, be impaired as when given pure garcinol. In contrast, we show that when injected with the mixture, bees’ learning is enhanced or rescued, independent of single- or three-trial conditioning (Figs. 5, 6). This raises questions regarding the actions of actinomycin D and garcinol on transcription processes that contribute to memory formation.

Memory formation requires a balanced expression of genes that promote or inhibit synaptic plasticity (Abel et al. 1998; Walkinshaw et al. 2015). Changing the balance toward either promoting or inhibiting genes results in an improved or an impaired memory, respectively. Actinomycin D intercalates into DNA (Sobell 1985), prevents active transcription processes in general, and thus equalizes the imbalance.

In contrast to actinomycin D, the effect of garcinol and other HAT inhibitors on gene expression is more complex. Results derived from cell lines show that, although the majority of differentially expressed genes is downregulated, other genes are upregulated (Balasubramanyam et al. 2004; Gaddis et al. 2015).

Figure 3. HAT inhibitors garcinol and C646 differently affect memory phases induced by strong associative training. Thirty minutes after strong associative olfactory training (three-trial conditioning) honeybees are injected with garcinol (A), C646 (B), or vehicle (DMSO). The columns represent the percentage of animals that elicit the proboscis (PER, proboscis extension response) during acquisition and retrieval tests at 2 h, 1 and 2 d. The numbers of animals tested in each of the groups is shown in brackets. Asterisks indicate significant differences between treatments as tested for each time point separately (details in Results). Chi-square/Fisher exact test (two-tailed); (*) P < 0.05.

Figure 4. HAT inhibitors garcinol and C646 affect different memory phases after weak training. Honeybees were injected with garcinol (A), C646 (B), or the vehicle (DMSO) 30 min after weak associative training (single-trial conditioning). The columns show the percentage of animals that elicit the proboscis (PER) in the retrieval test at 2 h, 1 and 2 d after conditioning. The numbers of animals tested in each of the groups is shown in brackets. Asterisks indicate significant differences between treatments as tested for each time point separately (details in Results). Chi-square/Fisher exact test (two-tailed); (*) P < 0.05.
Thus it is feasible that memory impairment caused by garcinol results from an imbalance of genes with promoting and inhibiting functions in memory formation. The imbalance favors expression of inhibitory genes and thus leads to a reduced memory. In this scenario, coapplication of actinomycin D would especially block inhibitory genes, restore the balance, and thus improve memory again.

In summary, the results support the emerging idea that acetylation-mediated mechanisms contribute to processes of memory formation at various levels of regulation. Building a thorough understanding of the acetylation-mediated processes will require more sophisticated tools to characterize the role of the different HATs, HDACs, and their substrates. Techniques that allow one to discriminate the contribution of processes at the transcriptional level in the nucleus from processes in the cytosol, dendrites, and synapses will have to be developed.

Materials and Methods

Animals

The experiments were performed in the years 2012 (April–October) and 2013 (February–September) in Saarbrücken, Germany. Bees (Apis mellifera) were collected from at least three different hives of the university apiary to avoid hive-dependent
Behavioral analysis

All behavioral tests were performed according to Merschbaecher et al. (2012). For the gustatory responsiveness test, honeybee foragers were caught on the day of the experiment. They were mounted as described above and fed 2–3 drops of 1 M sucrose solution to just about sustain them until performance. The responsiveness to appetitive stimuli was measured by monitoring the PER after stimulating the antennae with increased sucrose concentrations (0, 30, 100, 300 mM and 1 M) with an interstimulus interval of 2 min. For each bee, the sum of the PER elicited by the five gustatory stimuli represents the gustatory response score (0–5).

For tests involving nonassociative learning (sensitization, habituation) bees were caught and mounted as described above. Sensitization was tested by measuring the PER to an odor stimulus (clove oil) that was applied 15 sec after an acute appetitive stimulus (1 M sucrose) to the antennae of hungry bees. Animals showing no reaction to the sensitizing sucrose stimulus were excluded from the experiment (<5%). For habituation, animals were repeatedly stimulated (1 sec inter-stimulus interval) with 1 M sucrose solution at one antenna only. The number of elicited PER until five consecutive PER failures is defined as habituation criterion. Animals that were not habituated after 50 stimuli were excluded from the analysis (<5%). Dishabituation was performed with a sucrose stimulus (1 M) to the contra-lateral antenna following the five consecutive failures. Animals not responding to the dishabituating stimulus were excluded from the analysis (<5%).

For associative olfactory conditioning, bees were caught the day before the experiment, mounted, and fed 2–3 drops of 1 M sucrose solution. They were then starved overnight for at least 16 h in a dark humidity chamber at 70% humidity and 20°C–25°C. An associative olfactory-conditioning trial consisted of pairing an odor stimulus (conditioned stimulus, CS; clove oil for 5 sec) with an appetitive reward stimulus (unconditioned stimulus, US; 1 M sucrose for 5 sec). Animals received either one or three successive conditioning trials with an inter-trial interval of 2 min. Memory was tested 2 h, 1 and 2 d after training. Animals not responding to the US during conditioning were excluded from the experiment (<2%). Two hours after each memory test, the animals were fed 2–3 drops of 1 M sucrose solution.

Drug application

One microliter of either of the following solutions was injected with a calibrated glass capillary into the hemolymph of the thorax: C646 (cat#SML0002, Sigma-Aldrich), 1 mM in DMSO; garcinol (cat#10566, Biomol), 6 mM in DMSO; actinomycin D (cat#A1410, Sigma Aldrich), 1.8 mM in DMSO, mixture C646/actinomycin D (1 mM/1.8 mM) in DMSO, mixture garcinol/actinomycin D (6 mM/1.8 mM) in DMSO. The corresponding control groups were injected with 1 μL DMSO only. For the administered in vivo concentration, we assumed a mean bodyweight of 100 mg/bee.

Quantification of protein acetylation

To quantify H3K18ac, H3K9ac, and H3 in each of the brain samples we used the enzyme-linked immunosorbent assay (ELISA) (Merschbaecher et al. 2012). Following the described trial sections, honeybees were chilled on ice, the heads were cut off and mounted on wax. The cuticle was opened, the central brain with the mushroom bodies dissected and homogenized in 250 μL homogenization buffer (PBS containing 0.5% BSA, 1 mM EDTA, 1 mM Na2HPO4, 1.8 mM KH2PO4 containing 1 mM NaN3 and 5 mM sodium butyrate) within 30 sec. Each sample was divided onto three separate micro titer plates (P96 Maxisorp, NUNC-IMMUNO, Langenselbold, Germany), 50 μL aliquot per plate/antigen. The samples of the different groups within one experiment were placed on the same ELISA plates (separate copy plates for each antibody).

The samples were then diluted in five consecutive steps (1:2) with homogenization buffer. After 1 h coating, the wells were blocked for 1 h with blocking buffer (PBS containing 0.5% BSA) and then incubated for 1 h at room temperature with the different primary antibodies [anti-acetyl histone H3 (K18) (1:1000) (cat#SAB4500349, Sigma Aldrich), anti-acetyl histone H3 (K9) (1:1000) (cat#SAB4500347, Sigma Aldrich), anti-histone H3 (1:5000) (cat#4499P, Cell signaling), all diluted in PBS containing 0.5% BSA]. Washing was followed by an incubation with anti-rabbit IgG alkaline phosphatase conjugated antibody (1:4000 in PBS containing 0.5% BSA) (cat#A3687, Sigma Aldrich) for 1 h at room temperature. For the color reaction phosphatase substrate solution (1 mM 4-nitrophenylphosphate disodium salt (p-NPP) in 0.1 M Tris/HCl pH 8.7, 1 mM MgCl2) was added and the conversion of the substrate was quantified with a plate reader (safire², Tecan, Crailsheim, Germany) at 405 nm using 600 nm as background.

The ELISA data were evaluated as described previously (Merschbaecher et al. 2012). The slope calculated from the optical density values of the dilution steps represents the relative amount of antigen in a given sample. The means of the relative amounts of H3 did not differ between the groups (Tables 1, 2). In each sample, the normalized H3, H3K18ac, and H3K9ac values were used to calculate the H3K18ac/H3 or H3K9ac/H3. After normalization to the corresponding control ratio of a given experiment, the mean ± SD was calculated. The two-tailed Student’s t-test was used to compare the data.

HDAC assay

The HDAC activity in the honeybee brain was determined using the HDAC Fluorimetric Assay (BML-AK500) (Enzo Life Sciences, Lorrach, Germany) and the AmpliTekTM Fluorimetric HDAC Activity Assay Kit, Green Fluorescence (AAT Bioquest, Sunnyvale US). Garcinol, C646, TSA, or DMSO were either added to fresh brain homogenates or injected into the hemolymph 30 min prior to brain dissection. To quantify the effects of the inhibitors in the homogenates, brains were dissected and homogenized in the buffer provided with the HDAC assay. After splitting each homogenate into four fractions and adding garcinol, C646, TSA, and DMSO, respectively, HDAC assays were performed as described by the suppliers. To analyze the action of the inhibitors in the brain, the HDAC activity was determined after injection of the drugs into the hemolymph. Thirty minutes after systemic injection, brains of individual bees are dissected (<30 sec) and subjected to the HDAC assays. The measurements were performed according to the suppliers’ instructions. Assay conditions were optimized in pretests and revealed that one-tenth of a honeybee brain in a single measurement is best to determine HDAC activity in the linear range.

Statistical analysis

SYSTAT10 was used for the statistical analysis. The statistical tests used and the parameters are indicated in the section results or in the figure legends. P < 0.05 is considered as significant for the comparison of two groups. If groups are compared twice, the Bonferroni correction was applied and P < 0.025 is considered as significant.

Acknowledgments

We thank Angelika Gardezi for excellent technical assistance and Dr. Susanne Meuser for invaluable discussions and comments.
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Received June 22, 2015; accepted in revised form November 18, 2015.

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Learn. Mem. 2016, 23:
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