The Role of Carcinine in Signaling at the Drosophila Photoreceptor Synapse

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The Drosophila melanogaster photoreceptor cell has long served as a model system for researchers focusing on how animal sensory neurons receive information from their surroundings and translate this information into chemical and electrical messages. Electroretinograph (ERG) analysis of Drosophila mutants has helped to elucidate some of the genes involved in the visual transduction pathway downstream of the photoreceptor cell, and it is now clear that photoreceptor cell signaling is dependent upon the proper release and recycling of the neurotransmitter histamine. While the neurotransmitter transporters responsible for clearing histamine, and its metabolite carcinine, from the synaptic cleft have remained unknown, a strong candidate for a transporter of either substrate is the uncharacterized inebriated protein. The inebriated gene (ine) encodes a putative neurotransmitter transporter that has been localized to photoreceptor cells in Drosophila and mutations in ine result in an abnormal ERG phenotype in Drosophila. Loss-of-function mutations in ebony, a gene required for the synthesis of carcinine in Drosophila, suppress components of the mutant ine ERG phenotype, while loss-of-function mutations in tan, a gene necessary for the hydrolysis of carcinine in Drosophila, have no effect on the ERG phenotype in ine mutants. We also show that by feeding wild-type flies carcinine, we can duplicate components of mutant ine ERGs. Finally, we demonstrate that treatment with H₃ receptor agonists or inverse agonists rescue several components of the mutant ine ERG phenotype. Here, we provide pharmacological and genetic epistatic evidence that ine encodes a carcinine neurotransmitter transporter. We also speculate that the oscillations observed in mutant ine ERG traces are the result of the aberrant activity of a putative H₃ receptor.

Introduction

An exceedingly complex regulation is involved in the synthesis, release, activity, and recycling or degradation of neurotransmitter in the nervous system of animal species. This regulation may involve neurotransmitter degradation within the synapse [1], reuptake of neurotransmitter by presynaptic neurons [2], recycling of neurotransmitter by neighboring cells [3], and/or activation of receptors that trigger positive/negative feedback loops resulting in an increase/decrease of neurotransmitter release in presynaptic neurons [4]. Many of the mechanisms and machinery components associated with this regulation have been well conserved across species, ranging from Caenorhabditis elegans to humans [5]. D. melanogaster is often utilized when studying neurotransmitter dynamics because of its malleable genetics, strong phenotypes in the presence of neurotransmission defects, and its sensitivity to numerous neuropharmacological compounds that have been shown to exert similar effects in humans (for review, see [6]).

The major neurotransmitter released by photoreceptor cells in Drosophila is histamine [7], which is biosynthesized from histidine by the enzyme histidine decarboxylase (Hdc) found within the photoreceptor cell [8]. Upon excitation by light, the photoreceptor cell depolarizes and releases histamine, which then binds to a postsynaptic histamine-gated chloride channel, resulting in the hyperpolarization of the postsynaptic neuron [9,10]. Each synaptic cartridge is surrounded by three glial cells that invaginate into the photoreceptor terminals by means of fingerlike projections known as capitate structures [11]. While the exact site of histamine reuptake is currently unknown, the histamine remaining in the synaptic cleft is thought to be rapidly taken up by glial cells, possibly at the site of these capitate structures [12]. In glial cells this histamine is converted by the enzyme N-β-alanyl dopamine synthase, encoded by the ebony gene in Drosophila, into β-alanyl-histamine, also known as carcinine [3,13]. This carcinine is then released by the glial cell as an “inactive” conjugate of histamine, again possibly at the site of capitate structures, where it is then taken up by the presynaptic neuron. Once in the photoreceptor cell, the carcinine is converted by the enzyme N-β-alanyl dopamine hydrolase, encoded by the tan gene, back into the original neurotransmitter histamine [13,14]. It is proposed that the combination of histamine biosynthesis through histidine decarboxylase and the recycling of histamine by ebony and tan enzymes defines the total pool of histamine available at the photoreceptor cell synapse. The transporters responsible for histamine uptake by glial...
Author Summary

During signaling in the nervous system, individual nerve cells transfer information to one another by a complex process called synaptic transmission. This communication involves the release of a specific neurotransmitter into the synaptic cleft, which then triggers signaling in the downstream neuron by binding to and activating specific cell surface receptors. In order to terminate the neuronal signal, the neurotransmitter must be rapidly removed from the synaptic cleft. This is done by two mechanisms: the neurotransmitter can be degraded or modified, or the transmitter can be taken up by the presynaptic neuron and packaged into vesicles for reuse. In the compound eye of the fruitfly D. melanogaster, the photoreceptor cell responds to light and releases histamine into the synaptic cleft. This signal is terminated by the removal of histamine from the synapse and the enzymatic conversion of histamine to carcinine. We have shown that it is not sufficient just to modify the histamine neurotransmitter, but it is also important to remove carcinine from the photoreceptor synapse. The failure to adequately remove carcinine results in defects in the visual transduction process. Moreover, the work suggests that carcinine itself modulates vision by regulating histamine release into the synapse.

Carcinine Signaling at the Photoreceptor Synapse

Center. HdcP218 and ort2 fly mutants were obtained from W. Pak (Purdue University) while the UAS-ine-RB and ine2 transgenic fly lines were obtained from M. Stern (Rice University). All stocks were maintained in constant darkness at room temperature. Flies carrying two or three mutations/transgenes were generated by standard genetic methodologies. All wild-type flies were of the w1118 background.

Electroretinography

Flies were anesthetized by exposure to carbon dioxide and immobilized within a rotating disc using a drop of molten myristic acid (Akros). To record voltage changes within the eye, an electrode filled with signa gel (Parker Labs) was placed on the surface of the eye, while a second gel-filled electrode was gently inserted into the thorax. For light treatments a halogen lamp controlled by a Model T132 Uniblitz shutter was used. All light treatments, unless otherwise stated, were performed using a 580 nm filter and were 4 s in duration. To reduce the effects that exogenous sources of histamine might have on ineHdcP218 flies during ERG analysis, these animals were starved for 24 h before testing. To induce depolarization spikes in ort2 flies, two 4-s pulses of 480 nm light, followed by two 4-s pulses of 580 nm of light, were delivered to the flies, and a trace was taken during the second 580 nm pulse for analysis. Voltage changes were amplified using a DAM50 amplifier (World Precision Instruments) recorded using Powerlab 4/30 (AD Instruments, Colorado Springs, CO) and analyzed using Chart 5 software (AD Instruments). Oscillation frequency was determined by counting and averaging the number of repolarization spikes observed within 0.2 s of light exposure in either ine2 or carcinine-treated fly ERG recordings.

Drug Treatment

Thioperamide, immepip, and histamine were obtained from Sigma and carcinine was obtained from Peninsula Laboratories. All compounds were reconstituted in sterilized water for long-term storage. Flies were treated overnight in vials containing Kimwipes soaked with 200 µl of 1% sucrose solution with or without drug compound. Histamine was delivered to flies at a concentration of 10% [22]. Thioperamide and immepip were used at 0.5% and carcinine at either 5% or 10%. Flies were starved for 24 h before drug treatment.

Reverse transcriptase PCR

RNA was purified from either embryos or adult fly heads of the w1118 background by employing an RNeasy Mini Kit (Qiagen Sciences). cDNA was generated from purified RNA by utilizing MMLV-Reverse Transcriptase (Fisher Scientific). Forward primers specific to either the ine-RA or ine-RB transcripts and a reverse primer common to both transcripts were obtained from Integrated DNA Technologies. The ine-RA forward primer was ATCGATGGCACACTTCCGGATTACA, the ine-RB forward primer was ATCGATGGCACACTTGAGTTCG, and the reverse primer used to generate PCR product from both transcripts was TATCTGTCGACGCAAGATTTCA. Products were generated and amplified by means of PCR using Taq polymerase, buffers obtained from Invitrogen, and a TECHNE, TC-312 thermocycler (Barto-world Scientific) using the following parameters for 35 cycles (94 °C for 30 s, 55 °C for 30 s, and 72 °C for 90 s). PCR products were separated in a 1% agarose gel and then stained using ethidium bromide.

Materials and Methods

Fly Stocks

The repo-GAL4, “long”-GMR-GAL4, w1118, tan1, tan2, e1, e11, ort2/+ fly lines were all obtained from Bloomingston Stock...
Results

Glial- or Photoreceptor-Specific Expression of Inebriated Protein Rescues ERG Oscillation Phenotype

An ERG recording from a wild-type fly (Figure 1A) contains a receptor component, or the depolarization response upon exposure to light, and on and off transient spikes that indicate the response downstream of the photoreceptor cell (arrows, Figure 1A). The ERG of ine mutants contains an intact receptor component but with the addition of an initial depolarization spike (unfilled arrowhead, Figure 1B) and prominent oscillations superimposed on the depolarization response (Figure 1B). These oscillations have a wide range of frequencies from 40–90 Hz. These mutants also possess reduced on and off transients (arrows, Figure 1B), indicating impaired photoreceptor synaptic transmission. Finally, these ine mutant ERGs often display a hyperpolarization following a light response (arrowhead, Figure 1B). We observed all of these previously described ERG phenotypes when using either ine or ine allele flies. The ine allele is the result of a deletion of the majority of the ine open reading frame common to both ine-RA and ine-RB [16], while the mutation associated with ine was identified as being a nonsense mutation in codon 125 of the ine gene and is believed to only affect the ine-RA-encoded isoform [23]. Because the ine allele is associated with reduced viability, and because we could discern no observable difference between the ERG traces of ine and ine flies, we made use solely of the ine fly line for all of our experiments and genetic crosses.

Intracellular voltage recording experiments suggest that the oscillations observed in ine mutants originate in the photoreceptor cell, and that they are not the result of synaptic feedback [24]. However, if ine does encode a neurotransmitter transporter, its expression and localization are not necessarily restricted to photoreceptor cells, as neurotransmitter transporters often function from neighboring glial cells. Indeed, a previous study demonstrated that expression of ine in either neurons or glial cells was sufficient to rescue several mutant ine-associated defects at the neuromuscular junction [23]. In order to confirm that the inebriated protein is needed at the photoreceptor cell synapse, we tested whether the ine mutant phenotype could be rescued by expressing the ineRB transcript in photoreceptor and glial cells. The UAS-ine-RB transgenic fly line [23,25] contains the ine-RB cDNA under the control of the upstream activator sequence of the yeast Gal4 transcription factor. These UAS-ine-RB flies will only express Ine-P2 when crossed with a second line of flies expressing Gal4. The Gal4 lines utilized were “long”-GMR-GALA, which expresses Gal4 protein specifically in photoreceptor cells [26], and repo-GALA, which expresses Gal4 in glial cells [27]. A strong rescue of the ine ERG phenotype was observed when ine-RB was expressed in either photoreceptor or glial cells (Figure 1C and ID) compared to non-rescued ine or wild-type control (w1118) ERGs (Figure 1A and 1B). As expected, the UAS-ine-RB transgene failed to rescue the ine phenotype if neither GALA transgene was present (unpublished data). The oscillations observed in ERG traces from these transgenic rescued

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**Figure 1. Rescue of ine2-Associated Oscillations with Expression of INE in Both Glial and Photoreceptor Cells**

ERG recordings from (A) wild type, (B) ine2, (C) ine2 expressing the UAS-ineRB cDNA in photoreceptors, and (D) ine2 expressing the UAS-ineRB cDNA in glial cells. Arrows indicate on and off transients present in (A) wild-type but not (B) ine2 mutant recordings. A sharp depolarization response (unfilled arrowhead), a hyperpolarization (filled arrowhead) response and oscillations are present in (B) ine2 ERG recordings. Note that ine expression in either photoreceptor or glial cells results in a rescue of ine2-associated oscillations, on and off transients (arrows), and the hyperpolarization response. (E) Measurement of ine-RA and ine-RB mRNA in wild-type embryos or adult heads. 1 = ine-RA product from adults, 2 = ine-RA product from embryos, 3 = ine-RB product from adult heads, and 4 = ine-RB product from embryos. The ine-RA PCR product is 336 bp, while the ine-RB product is 338 bp. Note that there is very little ine-RB mRNA found in adult heads.

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Histamine Synthesis, But Not Postsynaptic Histamine Signaling, Is Required for the *ine* Mutant ERG Phenotype

(A) Simplistic diagram showing the synthesis and activity of histamine in the *Drosophila* eye. HD, histidine; Hdc, histidine decarboxylase; HA, histamine; Ort, histamine-gated chloride channel; PC, photoreceptor cell; LC, postsynaptic laminar cell. Histamine is generated by histidine decarboxylase from histidine in the photoreceptor cell. It is then released and acts upon a histamine-gated chloride channel on the postsynaptic laminar cell to trigger downstream signaling in the eye. ERG recordings from (B) histidine in the photoreceptor cell. It is then released and acts upon a histamine-gated chloride channel on the postsynaptic laminar cell to trigger stronger and more consistent rescue of the *ine* mutant phenotype, as the *ine-RB* transcript was previously thought to remain intact in *ine* mutant flies [25]. These results suggest that *ine-RB* is normally expressed at only low levels compared to *ine-RA*, and that overexpression of *ine-RB* is sufficient in compensating for the loss of *ine-RA* associated with *ine* mutants. Reverse transcriptase PCR experiments confirm these suspicions; *ine-RB* is expressed at low levels in adult wild-type heads compared to robust expression of *ine-RB* in the developing embryo (Figure 1E). The *ine-RA* transcript was found at high levels in both wild-type embryos and adult heads (Figure 1E).

Figure 2. Histamine Synthesis, But Not Postsynaptic Histamine Signaling, Is Required for the *ine* Mutant ERG Phenotype

Histamine Is Required for the *ine* Mutant ERG Phenotype

The ability to rescue the *ine* ERG response by expressing inebriated protein in photoreceptor and glial cells suggests that inebriated functions primarily at the site of the photoreceptor cell in the eye. Histamine is believed to be the predominant neurotransmitter that signals between photoreceptor cells and second order laminar neurons in the optic lobe [28], and it is possible that inebriated serves as a histamine transporter. However, previous studies showed that when inebriated protein from the tobacco hornworm *Manduca sexta*, which has significant homology to the inebriated protein from *Drosophila*, was expressed in *Xenopus laevis* eggs, it was unable to transport histamine across the cell membrane [17]. However, these authors do propose that a second unknown protein may be required to assist inebriated in proper neurotransmitter transport function, or that inebriated may possess different substrates in *Manduca* compared to *Drosophila*. Thus, histamine could still be the substrate of inebriated in *Drosophila*. Histamine is generated by the activity of histidine decarboxylase, encoded by the *Hdc* gene in *Drosophila* (Figure 2A). Mutations in the *Hdc* gene, such as in the case of the *Hdc* allele, result in flies possessing disrupted photoreceptor synaptic transmission, as demonstrated by the lack of on and off transients in their ERGs ([8], and Figure 2C). Approximately 80% of flies that were homozygous for both the *Hdc* and *ine* alleles displayed ERGs with no oscillations (Figure 2D) when compared to *ine* controls (Figure 2B). There was a small percentage (~20%) of *Hdc* *ine* flies that displayed weak or delayed oscillations. However, the ERGs from these flies that displayed this weak rescue also possessed on and off transients, suggesting that the *Hdc* allele was either not fully penetrant in these double mutants, or that their food provided an outside source of histamine. This is not surprising, as *Drosophila* photoreceptors are known to regain some function from exogenous histamine taken up at minute
levels in their food [22]. The ERGs from Hdc

\[P218 \text{ ine}^2\] also often lack the hyperpolarization response characteristic of \text{ine}^2 flies (Figure 2D). These findings suggest that histamine production or signaling plays a strong role in the oscillation and hyperpolarization phenotype observed in \text{ine}^2 traces.

The postsynaptic receptor for histamine in Drosophila is a histamine-gated chloride channel (Figure 2A), and a subunit of this channel is encoded by the ora transientless (ort) gene [10]. The ort\(^2\) and ort\(^\text{Pbac}\) alleles both result in reduced activity of this histamine receptor in Drosophila, as shown by the lack of on and off transients in their ERG traces ([10] and Figure 2E). If \text{ine} encodes a histamine neurotransmitter transporter, then reduced function of this protein may result in an excess of histamine in the synaptic cleft, and this excess of neurotransmitter may be acting upon this postsynaptic histamine receptor to somehow generate the observed oscillations. If this were the case, then \text{ine}^2 ort double mutants should have reduced oscillations. However, neither \text{ine}^2 ort\(^3\) (Figure 2F) nor the \text{ine}^2 ort\(^\text{Pbac}\) (unpublished data) double mutants exhibited rescue of the oscillation or hyperpolarization components of the \text{ine} ERG recordings, indicating that the oscillations do not arise from histamine signaling in downstream neurons. Moreover, since ort mutations block signaling in laminar neurons, these data are consistent with the oscillations being generated in the photoreceptor cells. Surprisingly, the ort\(^2\) allele, which is the result of a frameshift mutation and therefore likely serves as a null allele for this gene [10], often displays strong depolarization spikes of its own in the receptor component of its ERG trace (Figure 2G). Therefore, mutations in Hdc, which block the formation of histamine, rescue \text{ine}^2 whereas mutations in ort, which still allow for the synthesis of histamine, fail to rescue.

**Ebony Activity Is Required for the \text{ine} Mutant ERG Phenotype**

The abolition of the \text{ine} mutant ERG phenotype upon the introduction of Hdc, but not ort, mutations, suggests that histamine is involved in generating the \text{ine}^2 ERG phenotype, but that histamine’s downstream signaling in the optic lobe is not. The recycling pathway of histamine in the eye has been well elucidated [13,14,29,30]. It has been shown that following release into the synaptic cleft, histamine is rapidly taken up by neighboring glial cells and is converted by the \(\beta\)-alanyl-dopamine synthase, encoded by the gene ebony, into \(\beta\)-alanyl-histamine, also known as carcinine (Figure 3A). This carcinine is then transported into the presynaptic photoreceptor cell and is converted back into histamine by \(\beta\)-alanyl-dopamine-hydrolase, encoded by the gene tan, for use as a recycled source of neurotransmitter (Figure 3A). Both tan and ebony mutations in Drosophila are associated with significant reductions in size of the on and off transients in ERG traces (Figure 3C and 3E), due to the loss of this recycled pool of histamine in the eye. Introduction of either tan\(^2\) (unpublished data) or tan\(^1\) mutations into an \text{ine}^2 background failed to have any effect in reducing the size of the oscillations or hyperpolarization response when compared to \text{ine}^2 mutants alone (Compare Figure 3B with 3D). However, \text{ine}^2 ebony\(^1\) (unpublished data) or \text{ine}^2 ebony\(^11\) double mutants displayed complete rescue of the oscillation phenotype in all flies tested (Compare Figure 3B with 3F). These data, combined with the fact that histamine synthesis is necessary for the presentation of a mutant ERG phenotype in \text{ine}^2 flies, provide genetic evidence that carcinine is involved in generating \text{ine}^2-associated oscillations.

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**Figure 3. Ebony, But Not Tan, Activity Is Required for the \text{ine} Mutant ERG Phenotype**

(A) Simplistic diagram showing the recycling of histamine in the Drosophila eye. HA, histamine; CA, carcinine; PC, photoreceptor cell; LC, postsynaptic laminar cell, and GC, glial cell. Histamine is taken up from the synaptic cleft by glial cells where it is converted by ebony into carcinine. Carcinine is then shipped to photoreceptor cells where it is converted back to histamine by the enzyme tan. ERG recordings from (B) \text{ine}^2, (C) tan, (D) tan\(^1\) \text{ine}^2, (E) ebony\(^1\), and (F) \text{ine}^2 ebony\(^11\) flies. Note that \text{ine} oscillations are rescued by mutations in ebony but not in tan.

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Figure 4. Carcinine Treatment Induces an ine^2-Like ERG Phenotype in Wild-Type and ebony^11 Flies

ERG recordings from untreated (A) wild-type and (D) ebony^11 flies and 5% carcinine-treated (B, C) wild-type and (E, F) ebony^11 flies. Note that while untreated wild-type and ebony^11 ERG recordings lack oscillations and a hyperpolarization response in response to light, ERG recordings from wild-type flies treated with 5% carcinine overnight display weak oscillations (arrows) in ~35% of tested animals and ERG recordings from ebony^11 flies treated with 5% carcinine exhibit depolarization spikes in response to light (unfilled arrowhead), as well as weak oscillations (arrow) and a hyper-repolarization upon the termination of light (arrowhead).

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Treatment of Wild-type and ebony Flies with Carcinine Results in ERG Abnormalities

If ine encodes a carcinine neurotransmitter transporter, as the genetic evidence above suggests, than a potential cause of the aberrant ERG phenotypes seen in ine mutants could be the buildup of carcinine within the photoreceptor synaptic cleft. In order to test whether or not carcinine is able to induce an ine^2-like ERG phenotype in wild-type animals, w^1118 flies were treated with 5% carcinine overnight and then subjected to ERG analysis. Approximately 35% of the w^1118 flies treated with carcinine displayed occasional weak oscillations or brief depolarization/repolarization spikes in the photoreceptor response of their ERG traces (Figure 4B and 4C, compare to Figure 4A). While these spikes exhibit no consistent frequency, unlike the oscillations seen in ine^2 recordings, the carcinine-induced ERG disturbances were never observed in untreated starved flies. If carcinine was delivered to ebony^11 flies, which lack the ability to synthesize carcinine from histamine, they surprisingly displayed an abnormal ERG trace. All ebony^11 flies treated with carcinine manifested phenotypes reminiscent of those seen in ine^2 ERG traces, including sharp depolarization spikes in response to a light response, weak oscillations, and a hyperpolarization peak upon the termination of light exposure (Figure 4E and 4F, compare to Figure 4D). The oscillations observed in carcinine-treated ebony^11 mutants, while only appearing briefly during the initiation of light exposure, were seen at a similar frequency as those found in ine^2 ERG recordings (63 spikes/s). As discussed below, a possible mechanism underlying these carcinine-induced ERG disturbances may involve the sensitization of a putative histamine/carcinine receptor.

Pharmacological Rescue of the ine^2 Mutant ERG Phenotype

If inebriated does serve as a carcinine transporter, and if carcinine is indeed building up within the synaptic cleft in ine^2 mutants, then this uncleared carcinine appears to somehow be acting on some synaptic receptor to initiate this aberrant oscillation phenotype. The ine^2/or^t experiments suggest that this receptor is not the post-synaptic histamine-gated chloride channel. In mammals and various other vertebrate systems, presynaptic histaminergic neurons often contain their own histamine receptors, known as H 3 receptors. The H 3 receptor is a G-protein–coupled receptor that was first identified in 1983 by Arrang et al. [4] and is now known to act as a presynaptic autoreceptor that inhibits histamine release from histaminergic neurons in the brain (for review, see [31]). Thus, H 3 receptors serve to negatively regulate histamine release and synthesis in the presence of high histamine levels in the synaptic cleft. While no H 3 receptor has been identified yet in Drosophila, there are several candidate genes that may encode such a putative receptor. There are numerous well-characterized pharmacological compounds that act as agonists, antagonists, or inverse agonists of the H 3 receptor in vivo in mammals, and recently carcinine was identified as being an inverse agonist of this receptor in mice [32]. It was shown that, rather than reduce histamine release, as occurs in the case of histamine binding to an H 3 receptor, carcinine had the opposite effect and induced both histamine synthesis and release from presynaptic histaminergic neurons in vivo.

A possible scenario to explain the oscillations seen in ine^2 ERGs is that histamine and uncleared carcinine are competing for binding to putative H 3 receptors, resulting in opposing signaling cascade responses in the photoreceptor cell. If this is the case, disrupting this balance of histamine and carcinine binding to the putative H 3 receptor in ine^2 fly eyes should result in a rescue of ERG oscillations. Indeed, treatment of ine^2 flies with 10% carcinine resulted in a rescue of oscillations in 35% of flies (unpublished data), and treatment of ine^2 flies with 0.5% thioperamide, another...
well-characterized and potent inverse agonist of the H₃ receptor in mammals, resulted in the consistent and complete ablation of oscillations in ERG traces in all flies tested (Compare Figure 5A with 5C). In addition, treatment of ort⁵ flies with 0.5% thioperamide resulted in a loss of ort⁵-associated depolarization spikes (unpublished data). Surprisingly, treatment of wild-type control flies with 0.5% thioperamide resulted in the loss of on and off transients in their ERGs (compare Figure 5B with 5D).

It should also be possible to disrupt the hypothetical balance of histamine and carcinine binding to a photoreceptor cell-specific H₃ receptor in ine⁴ by introducing an H₃ receptor agonist, such as histamine itself. Indeed, treatment of ort¹ flies with 0.5% thioperamide resulted in a loss of ort¹-associated depolarization spikes (unpublished data). Surprisingly, treatment of wild-type control flies with 0.5% thioperamide resulted in the loss of on and off transients in their ERGs (compare Figure 5B with 5D).

Discussion

Our findings indicate that the presumed neurotransmitter transporter encoded by the ine gene in Drosophila transports the histamine metabolite carcinine. We show using genetic epistasis that the oscillations observed in mutant ine ERGs require histidine decarboxylase activity and the carcinine-synthesizing enzyme ebony, but not the carcinine-hydrolyzing enzyme tan. We also reveal that treating wild-type flies with carcinine can phenocopy components of the mutant ine ERG phenotype. Finally, by rescuing the ine²-associated phenotype with drugs that target the mammalian H₃ receptor, we provide pharmacological evidence for the presence of a putative H₃ receptor in Drosophila that may be responsible for the ERG oscillations observed in flies carrying mutations in the ine gene.

The Inebriated Protein Functions in Photoreceptor and Glial Cells

Previous studies involving intracellular voltage recordings of ine mutants led the authors to conclude that the oscillations observed in ine mutant ERGs were the result of a defect occurring within the photoreceptor cell [24]. We were able to support these conclusions by expressing ine specifically in photoreceptor cells and demonstrating a rescue of the ine²-associated oscillations. Neurotransmitter transporters are often able to function from either the presynaptic neuron or from neighboring glial cells, as shown at the neuromuscular junction in ine mutants [23]. We found that glial cell-specific expression of the ine gene in ine² flies resulted in a complete rescue of the ine mutant ERG phenotype. It was somewhat unexpected that ine expression in glial cells rescued the ine² phenotypes, as glial cells have been shown to lack tan protein and thus would be unable to convert carcinine back to a recycled pool of histamine [30]. However, it is possible that glial cells do express trace amounts of the enzyme tan to hydrolyze carcinine and generate a renewable source of histamine for photoreceptor synthesis.
cells, and it is also possible that the inebriated protein is expressed in a non-autonomous manner and can be transported from glial cells to photoreceptors in the fly eye.

The finding that an ERG recording can exhibit oscillations is somewhat surprising. An ERG does not record the electrical response of a single photoreceptor, but rather is a collective measure of the retinal photoreponse. Thus, if the mutant ine2-associated ERG defects are indeed localized to the photoreceptor synapse, as our data and that of previous labs suggest, then one would expect that different photoreceptors would be excited/inhibited at different timepoints, ultimately resulting in the oscillations simply canceling themselves out. The fact that oscillations are indeed observed, and appear to be due to a defect occurring at the photoreceptor synapse, implies the existence of an uncharacterized and complex synchronization of photoreceptor cell de-repolarization.

A Postsynaptic Buildup of Carcinine in the Drosophila Eye Causes the ine2-associated Mutant ERG Phenotype

The lack of rescue of ine2-associated oscillations in flies carrying additional mutations in the postsynaptic histamine receptor gene ort, the finding that mutant ine oscillations were detected within single photoreceptor cells [24], and our observations that the mutant ine phenotype can be rescued when ine is expressed in photoreceptors, all combine to strongly suggest that the oscillation phenotype is likely a result of a defect occurring within the photoreceptor itself. In addition, by crossing ine2 animals with HdcP218 flies, we demonstrated that the ine2-associated oscillations are dependent upon histamine synthesis. All of these results indicate that histamine is somehow contributing to the aberrant ERG witnessed in ine2 flies, and that histamine appears to be acting on the presynaptic photoreceptor cell to induce this oscillation phenotype. Further epistatic analyses also revealed that ebony, but not tan, activity is required for the generation of oscillations in ine2 ERGs (Figure 6A). These genetic experiments are consistent with ine encoding either a carcinine importer found in the photoreceptor cell or a carcinine exporter found in glial cells. The homology of inebriated with other known Na+/Cl− neurotransmitter transporters (which import neurotransmitter into cells) [16] suggests that inebriated protein is transporting carcinine into the photoreceptor, and not out of glial cells.

While ebony is known to act on multiple substrates, such as dopamine to generate β-alanyl-dopamine [13], the requirement of histamine synthesis for the maintenance of ine2-associated oscillations suggests that it is β-alanyl-histamine, or carcinine, that is somehow responsible for the oscillations observed in ine2 ERGs. It should be noted, however, that ebony
mutations were not sufficient in rescuing the hyperpolarization response observed in mutant ine ERG traces (Figure 3F). The origins of this hyperpolarization response are still unclear and further research will be required to elucidate its exact meaning. In tan mutants, one would predict that there would be a buildup of carcinine. However, this buildup does not give rise to an ERG recording similar to that of ine. This is due most likely to the presence of functional inebriated protein in tan mutant flies, which should effectively clear the carcinine from the synaptic cleft for degradation within the photoreceptor cell.

By treating wild-type and ebony11 flies with carcinine and subsequently inducing components of the ine2-ERG phenotype, we provide further evidence suggesting that the sharp depolarization spike, the oscillations, and the hyperpolarization response all seen in ine2-ERGs are due to a buildup of carcinine within the photoreceptor synaptic cleft. While the oscillations observed in carcinine-treated wild-type flies do not mimic exactly the oscillations seen in ine2 ERG recordings, it is presumably difficult to replicate the carcinine and histamine balance occurring in the eyes of ine2 animals. Indeed, treatment of wild-type flies with higher (10%) or lower (1%) concentrations of carcinine were less effective in inducing the oscillations than the described 5% carcinine dose (unpublished data).

It is possible that carcinine is being degraded or modified by the fly before the compound is able to exert its effects at the photoreceptor cell. In order to eliminate the activity of one enzyme known to be involved in carcinine metabolism, tan1 flies were treated with 5% carcinine overnight. Surprisingly, none of the tan1 flies treated with carcinine showed an aberrant ERG phenotype (unpublished data). It was surprising that carcinine treatment had a strong effect in flies of the ebony11, but not the tan1, background. While the results of these tan1 and ebony11 carcinine-treatment experiments are unexpected, one possible explanation may involve the regulation of carcinine clearance/degradation. The tan1 flies presumably suffer from a perpetual excess of carcinine even before exogenous carcinine treatment, and these flies, in order to reduce their sensitivity to this compound, may consequently decrease the levels of a putative carcinine receptor, increase their rate of carcinine degradation, or increase the levels of inebriated protein for carcinine clearance. However, ebony11 flies are relatively “naive” to the effects of carcinine, as their ability to synthesize this compound has been greatly diminished, and as a result these animals may have an increased level of the supposed carcinine receptor, a decrease in inebriated receptor levels or a decrease in carcinine degradation, ultimately making them more sensitive to the effects of carcinine treatment.

It remains to be seen whether or not all of the mutant ine-associated phenotypes, including increased neuronal excitability [19,23] and increased sensitivity to osmotic stress [25], are due to the inability of these flies to transport carcinine. It is possible that the inebriated protein transports other compounds that perhaps share the common feature of β-alanine conjugation. This might help explain why none of the more common neurotransmitters were taken up by ine-transfected Xenopus oocytes [17]. In order to assist in confirming that inebriated is indeed a carcinine neurotransmitter transporter, in vitro experiments, such as neurotransmitter uptake assays, will need to be performed. In addition, the ability of inebriated protein to take up other β-alanyl-neurotransmitters/osmolytes also should be examined.

What are the ine2-Associated Oscillations?

The oscillations present within the photoreceptor response of ine2 ERGs appear as sharp depolarization/repolarization spikes, and this oscillation phenotype is dependent upon both histamine synthesis and ebony activity (Figure 6A), and is sensitive to drugs that target mammalian H3 receptors. It is perplexing that the synthesis of a single metabolite, carcinine, could be responsible for both the depolarization and repolarization spikes observed within ine mutant ERGs. We speculate that these oscillations are the result of aberrant signaling involving both carcinine and histamine at a putative H3 receptor in Drosophila (Figure 6B and 6C). H3 receptors are an unusual example of the G-protein coupled receptor family, in that they have partial constitutive activity, resulting in a constant small percentage of stimulated G-proteins [33] that trigger a reduction of histamine synthesis and release [34] as well as a decrease in extracellular calcium inflow [35,36,37]. The presence of an H3 receptor agonist, such as histamine, causes an increase in activity of the associated G-protein and therefore a stronger inhibition of both histamine release and calcium inflow. Thus, synaptic histamine serves as a negative regulator for its own release and induces a slight repolarization of a stimulated presynaptic histaminergic neuron by inhibiting presynaptic calcium channels. An H3 receptor inverse agonist is believed to act by blocking the constitutive activity of the H3 receptor, resulting in the liberation from a histamine release checkpoint as well as the release of restrictions on calcium inflow [38]. Recently, it has been shown that carcinine has the ability to act as an inverse agonist of presynaptic H3 receptors in mice [32]. While significant further research is required to confirm this hypothesis, we surmise that histamine and carcinine are exerting opposing effects on the polarization state of the histaminergic photoreceptor cell by activating or inhibiting presynaptic calcium channels via a putative Drosophila H3 receptor. While a recent search of the Drosophila genome did not uncover any direct homologs to vertebrate metabotropic histamine receptors [39], the CG7918 gene was listed as a possible candidate for encoding such a receptor, and this gene bears strong homology to genes encoding H3 receptors in mammals. In addition, the ine2-associated oscillations display sensitivity to mammalian H3 receptor agonists and inverse agonists, strengthening the possibility that an H3 receptor does exist in Drosophila. It is still unclear what the origins of the thioperamide-sensitive depolarization spikes are that are observed in ort2 ERGs. The presence of these thioperamide-sensitive spikes in ort2 ERG recordings implies the requirement of some postsynaptic retrograde signal for ERG stability, and this ort-dependent signal may be involved in the sensitization of the putative H3 receptor.

It was unexpected that thioperamide treatment of wild-type flies resulted in the loss of on and off transients within their ERG traces. It is possible that histamine release was so extreme in the presence of the potent thioperamide that histamine levels were nearly depleted in the eye, resulting in the disruption of downstream signaling events. Indeed, treatment of mice with high concentrations of carcinine, which acts as an inverse agonist of H3 receptors similar to thioperamide, was shown to result in significantly lower
confirm or dismiss the presence of a histamine/carcinine-sensitive H3 receptor in Drosophila photoreceptor cells.

Supporting Information

Acknowledgments

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