The Ih Channel Gene Promotes Synaptic Transmission and Coordinated Movement in Drosophila melanogaster

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Hyperpolarization-activated cyclic nucleotide-gated “HCN” channels, which underlie the hyperpolarization-activated current (Iₕ), have been proposed to play diverse roles in neurons. The presynaptic HCN channel is thought to both promote and inhibit neurotransmitter release from synapses, depending upon its interactions with other presynaptic ion channels. In larvae of Drosophila melanogaster, inhibition of the presynaptic HCN channel by the drug ZD7288 reduces the enhancement of neurotransmitter release at motor terminals by serotonin but this drug has no effect on basal neurotransmitter release, implying that the channel does not contribute to firing under basal conditions. Here, we show that genetic disruption of the sole HCN gene (Ih) reduces the amplitude of the evoked response at the neuromuscular junction (NMJ) of third instar larvae by decreasing the number of released vesicles. The anatomy of the (NMJ) is not notably affected by disruption of the Ih gene. We propose that the presynaptic HCN channel is active under basal conditions and promotes neurotransmission at larval motor terminals. Finally, we demonstrate that Ih partial loss-of-function mutant adult flies have impaired locomotion, and, thus, we hypothesize that the presynaptic HCN channel at the (NMJ) may contribute to coordinated movement.

Keywords: Ih gene, HCN channel, neuromuscular junction, presynaptic mechanisms, Drosophila melanogaster

INTRODUCTION

Hyperpolarization-activated Cyclic Nucleotide-gated (HCN) channels underlie the hyperpolarization-activated current, designated Iₕ (hyperpolarization-activated current) or I₇ (funny current). HCN channels have a primary structure that is similar to voltage-gated potassium channels (Gauss et al., 1998; Ludwig et al., 1998; Santoro et al., 1998); they are tetrameric and contain individual subunits which are multi-membrane spanning and possess intracellular N- and C-terminals (Lee and Mackinnon, 2017). There are four HCN genes in mammals (HCN1-HCN4), which are expressed at high level in the heart and nervous system (Accili et al., 2002; Robinson and Siegelbaum, 2003; Biel et al., 2009). HCN channels are activated by membrane hyperpolarization, and their opening is facilitated by direct binding of cAMP to a cyclic nucleotide binding site in the C-terminus (Difrancesco, 1981a; Difrancesco and Tortora, 1991; Wainger et al., 2001; Zagotta et al., 2003; Chow et al., 2012); the rate of opening and closing, and the effect of cAMP, varies
between the isoforms, which are coded by different genes (Altomare et al., 2001; Robinson and Siegelbaum, 2003). In the mammalian heart, HCN channels are especially enriched in the sinoatrial node, where their unique activation properties and sensitivity to cAMP have been proposed to promote pacemaker activity and proper electrical conduction (Difrancesco, 1993; Baruscotti et al., 2011). HCN channels and the currents they produce have also been identified in a wide range of vertebrate and invertebrate neurons where they have varying patterns of cellular localization and are thought to contribute to presynaptic and postsynaptic functions (Pape, 1996; Robinson and Siegelbaum, 2003; Biel et al., 2009).

Hyperpolarization-activated Cyclic Nucleotide-gated (HCN) channels have mixed cation permeability, they open and close with time constants that range from hundreds of milliseconds to seconds, depending on membrane voltage and the isoform involved, and mammalian isoforms do not inactivate (Difrancesco, 1981b; Pape, 1996; Robinson and Siegelbaum, 2003; Biel et al., 2009). The opening of HCN channels by hyperpolarization leads to depolarization of the resting membrane potential to \(-25\) mV, which is the reversal potential for the channels in physiological solutions. Curiously, presynaptic HCN channels have been proposed to both promote and inhibit neurotransmitter release (Biel et al., 2009; Tomlinson et al., 2010; Trevillion et al., 2010; Howells et al., 2012; Huang and Trussell, 2014; Lorenz and Jones, 2014; Shah, 2014), which may depend upon the context provided by the complement of other ion channels present in particular axons or presynaptic terminals. For example, in mice, genetic deletion of the HCN1 gene or pharmacological inhibition of I\(_h\) by the drug ZD7288 have been shown to enhance the frequency of miniature excitatory postsynaptic potentials on EC layer III pyramids by promoting the entry of calcium through T-type calcium channels; this suggests that I\(_h\) normally limits calcium entry into the synapse and reduces neurotransmission (Huang et al., 2011). Using a combination of genetic approaches, the presynaptic HCN channel has also been shown to reduce glutamate release from amacrine cells in flies by limiting calcium channel activity (Hu et al., 2015). Conversely, pharmacological inhibition of the presynaptic HCN channel by ZD7288 has been shown to reduce the stimulation of neurotransmitter release by serotonin in the neuromuscular junction (NMJ) of Drosophila melanogaster larvae and crayfish; notably, however, ZD7288 has no apparent effect on basal release (Beaumont and Zucker, 2000; Cheung et al., 2006).

Based on these pharmacological data in D. melanogaster, it appears that I\(_h\) does not impact basal neurotransmission. However, many of the studies that examine the presynaptic HCN channel use I\(_h\) inhibitors which are not specific. Perhaps the most commonly used I\(_h\) inhibitor is ZD7288, which requires a relatively long incubation and, when used at lower concentrations (<1 \(\mu\)M), does not inhibit I\(_h\) completely (Bosmith et al., 1993; Qu et al., 2008). Thus, ZD7288 is often used at concentrations which are known to affect other ion channels.

Few studies have exploited genetic strategies to either corroborate the actions of ZD7288 on neuronal HCN channels and I\(_h\) (Huang et al., 2011) or to avoid its use altogether (Hu et al., 2015). To this end, we examined the function of the single HCN gene in D. melanogaster, I\(_h\). We found that I\(_h\) null embryos are unable to hatch, whereas I\(_h\) hypomorphs are viable but as adults exhibit long periods of inactivity, impaired climbing and decreased lifespan. To obtain direct evidence for a contribution of I\(_h\) to presynaptic signaling, we performed electrophysiological recordings at the (NMJ) of late third instar larvae. We found that the evoked potentials and quantal content were reduced in I\(_h\) hypomorphs. We propose that presynaptic I\(_h\) promotes basal neurotransmitter release at the presynaptic motor terminal as well as serotonin-induced neurotransmitter release which was shown previously (Beaumont and Zucker, 2000; Cheung et al., 2006). Finally, we found that I\(_h\) hypomorphic adult flies display a notable lack of coordinated movement. Promotion of neurotransmission at motor terminals by the presynaptic HCN channel may explain the movement phenotypes of I\(_h\) mutant flies.

**METHODS**

**Genetics**

The P[Gsv2] GS50880 transposable element insertion in the I\(_h\) gene (CG8585) was crossed to a \(\Delta2-3\) transposase line to generate a series of precise and imprecise genomic excision events. Candidate alleles were screened using PCR and complementation analysis with an I\(_h\) deficiency allele, Df (2R)Exel1[17131]. A precise excision line with the smallest footprint (32 bp) was used as a control allele. Although three start codons remain in the excised P[GSV2] footprint, it was not predicted to disrupt any of the coding sequences of the gene, and I\(_h\) cDNA immediately downstream of the insertion site was not reduced in flies containing this footprint (Figure 1C). Importantly, precise excision I\(_h\)^PE alleles exhibited wild type gene activity in all molecular and behavioral tests. Imprecise excision events were observed in two lines, which did not produce homoyzygotic animals and failed to complement a deficiency through I\(_h\). Reverse-transcriptase PCR was used to test transcript levels in different alleles; confirming transcript loss in imprecise excision I\(_h\)^H (null) and wild type levels in precise excision I\(_h\)^PE (revertant control). Candidate alleles, including precise and imprecise alleles, as well as the original insertion line(s) were out-crossed five times to an isogenic w\(^{\text{cr}}\) strain to eliminate potential second site mutations and improve their utility in behavioral assays. Mutants were kept over CyO, Act-GFP TM3, Ser, Act-GFP or CyO, twiGAL4, UAS-2xEGFP or TM3, Sb, Ser, twiGAL4, UAS-2xEGFP. Isogenic w\(^{1118}\) was used as the control genotype. Flies were maintained at 25\(^\circ\)C, 70% humidity on standard cornmeal food.

**Molecular Biology**

Genomic DNA was extracted from Stage 17 embryos, late third instar larvae, or freshly-eclosed adult flies using DNAzol (Invitrogen) for screening through the entire I\(_h\) locus using PCR. For reverse-transcriptase PCR reactions, total RNA was extracted using Qiazol (Qiagen), followed by standard chloroform precipitation and purification with the Zymogen RNA clean-up kit. Total RNA was reverse transcribed with the iScript cDNA synthesis kit (BioRad). The
FIGURE 1 | Genetic excision of the fly HCN channel. (A) A cartoon showing the predicted topology of a full-length HCN channel subunit in Drosophila melanogaster. Functional channels consist of four subunits (inset) and open in response to membrane hyperpolarization. All Ih alleles used in this study are listed on the lower left. The GS transposable element (Flybase P[GSY2] GS50880) was used to generate precise and imprecise excisions. Cartoon adapted from Biel et al. (2009). (B) A genomic map of the Ih locus on the second chromosome showing exon structure of Ih transcripts. The locations of transposable element insertions are intronic and are indicated with dotted lines. Map adapted from Flybase. (C) A gel showing fragments produced by genomic PCR (left) and reverse-transcriptase PCR (right), after genetic disruption of Ih in E17 embryos as indicated. All lines were out-crossed five times to an isogenic w- wild type line. Primers correspond to the region of exons 4–6 in Ih mRNA, immediately downstream of the intronic P[GSY2] insertion site. (D) Confocal images of embryos comparing in situ hybridization of Ih mRNA expression in Ih+/CyO, Twi-GFP heterozygotes (left) with that of ih+/ih null embryos (right). Digoxigenin-UTP-labeled RNA probes to the Ih gene were generated using the same primers as in (C). Note the absence of Ih expression in ih+/ih null embryos. Fluorescence on the right arises from autofluorescence of the larval gut. Scale bars = 50 µm.
following primers were utilized to determine the presence or absence of exons 4–6: 5′-GCTGCTCCATTGCTCGGTTG, 3′-GTTCAGCGTTGTCTTGTTGC. This region translates to the first three transmembrane domains of the wild type channel.

**In situ Hybridization and Imaging**

Digoxigenin-UTP-labeled RNA probes to the **Ih** gene were generated using the Roche DIG labeling kit. Primers used were the same as described above. For each allele, timed grape plate collections were used to collect age-matched 12–15 h embryos, which were immediately fixed in 4% PBS-buffered formaldehyde. Probe hybridization was carried out as described in Weiszmann et al. (2009). Briefly, embryos were incubated overnight at 55°C in a solution containing DIG-labeled probe in 50% formamide, 3X saline-sodium citrate buffer and 0.01% Tween-20. Embryos were washed extensively, blocked with 5% donkey serum in 0.1% Triton-X-100, followed by overnight incubation with mouse **α**-DIG-POD (Fab fragments from polyclonal anti-digoxigenin antibodies, conjugated to horseradish peroxidase used at a dilution of 1:2000), followed by 30 min developing with a tyramide signal amplification (TSA) developing agent. Embryos were then washed and incubated overnight with a chicken **α**-GFPl antibody (1:1000). The following day, embryos were incubated for 1 h with **α**-chicken-Alexa 649 2° antibody (1:500), washed and mounted on slides with VectaShield.

For imaging of the (NMJ) at muscles 4 and 6/7, third instar larval files were pinned out on Syligard and fixed with 4% PBS-buffered formaldehyde. Fixed files were blocked for 2 h with 5% donkey serum in 0.1% Triton-X-100 and incubated for 2 h with **α**-DIG-POD (Fab fragments from polyclonal anti-digoxigenin antibodies, conjugated to horseradish peroxidise used at a dilution of 1:2000), followed by 30 min developing with a tyramide signal amplification (TSA) developing agent. Embryos were then washed and incubated overnight with a chicken **α**-GFPl antibody (1:1000). The following day, embryos were incubated for 1 h with **α**-chicken-Alexa 649 2° antibody (1:500), washed and mounted on slides with VectaShield.

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**Behavior**

For embryo hatching assays, timed embryo collections were used to collect age-matched Stage 17 embryos overnight. For each genotype, 20–50 dechorionated embryos were isolated on one side of a fresh grape agar plate with a small amount of yeast paste on the opposite side. After 6 h, the numbers of hatched and un-hatched embryos were counted. Plates were re-examined after 24 h, and in all cases embryos that did not hatch in the first 6 h remained un-hatched. A minimum of 100 embryos were examined for each genotype. The genotypes were compared using Fisher’s exact test.

For climbing assays, age-matched, and freshly eclosed male adult flies from each genotype were collected in groups of ten. Each vial was assayed for climbing at days 1, 5, 9, 13, 17, and 21 post-eclosion. Climbing index was calculated as the average percentage of flies to reach a 4 cm line in 8 s after they were gently tapped to the bottom of a clean, empty vial. For each genotype, 4–6 sets of 10 males were measured four times and averaged, first among trials and then among sets. One-way ANOVA with Tukey’s multiple comparisons test was performed among all genotypes for each day. For lifespan assays, fifty age-matched adults were isolated by sex in groups of ten. The number of dead flies was recorded each day until no live flies remained. Fresh food was provided daily.

**Electrophysiology**

Sharp electrode recordings were taken from muscle 6 in abdominal segments 2 or 3 of third instar larvae, as previously described (Jan and Jan, 1976; Frank et al., 2006, 2009; Brusich et al., 2015). For recording, sharp electrodes were pulled using a P-97 micropette puller (Sutter Instruments) to an approximate resistance of 15 MΩ and filled with 3 M KCl. Larvae were dissected in a modified HL3 saline containing: NaCl (70 mM), KCl (5 mM), MgCl2 (10 mM), NaHCO3 (10 mM), sucrose (115 mM = 3.9%), trehalose (4.2 mM = 0.16%), HEPES (5.0 mM = 0.12%), and CaCl2 (0.5 mM). Only muscles with a resting potential of −60 mV or lower and an input resistance of 5 MΩ or greater were accepted for recording. Recordings were made using Axopatch 200B (in bridge mode) or Axoclamp 900A amplifiers (Molecular Devices, Sunnyvale, CA), digitized using a Digidata 1440A data acquisition system (Molecular Devices), and recorded with pCLAMP 10 acquisition software (Molecular Devices). For presynaptic nerve stimulation, a Master-8 pulse stimulator (A.M.P. Instruments, Jerusalem, Israel) and an ISO-Flex isolation unit (A.M.P. Instruments) were utilized to deliver 1 ms suprathreshold stimuli to the appropriate segmental nerve through a fire-polished glass electrode filled with recording saline. For evoked events, stimuli were applied at a frequency of 1 Hz. For evoked trains of stimuli (used to analyze paired-pulse responses), a frequency of 10 Hz was used.

For each NMJ, ~100–200 spontaneous miniature excitatory postsynaptic potential (mEPSP) events were recorded. Each spontaneous event was analyzed by hand. From this analysis, an average mEPSP size for each muscle was calculated, as was an average mEPSP frequency for each muscle. To generate mEPSP cumulative size distribution histograms for each genotype, the first 75–100 mEPSPs per muscle were chosen and added to the analysis. As a result, a minimum of 800 separate spontaneous events were analyzed per genotype.

Thirty evoked excitatory postsynaptic potentials (EPSP) per muscle were recorded and averaged. Quantal content (QC) was calculated for every muscle by dividing the average EPSP/average mEPSP. Since this EPSP/mEPSP QC estimate is accurate only for sufficiently small EPSPs, QC was then corrected for non-linear summation (NLS QC) as previously described (Martin, 1955). To correct QC for non-linear summation, the calculated EPSP/mEPSP quotient for each muscle was multiplied by a correction factor, derived from (Martin, 1955): (1–EPSP/Vo)−1. Vo is defined as the maximum electromotive force and was determined for each NMJ based on the muscle resting potential.

Electrophysiological data for each genotype were reported as mean ± SEM, and analyzed for statistical significance using ANOVA with a Tukey’s post-hoc test to account for multiple comparisons (GraphPad Prism). For the mEPSP cumulative probability histogram, data were analyzed for statistical significance using a Kruskal-Wallis test with a Dunn’s multiple comparisons post-hoc test (GraphPad Prism).
RESULTS

Imprecise Excision of the Single HCN Gene, Ih, Is Homozygous Lethal

The HCN gene in D. melanogaster, Ih, produces a channel that is similar in structure and function to the four mammalian HCN isoforms and other invertebrate HCN isoforms (Gauss et al., 1998; Ludwig et al., 1998; Santoro et al., 1998; Ishii et al., 1999; Marx et al., 1999; Gisselmann et al., 2005; Jackson et al., 2012). Despite their distinctive properties, HCN isoforms have significant sequence homology with channels in the voltage-gated potassium channel superfamily, for which the crystal structure has been solved (Doyle et al., 1998; Lee et al., 2005; Long et al., 2005). By extrapolation, HCN subunits are predicted to have six transmembrane helices (S1–S6) with a cytosolic N and C-terminus. The subunits combine as tetramers to form the ion conducting channel (Figure 1A). The C-terminus contains a cyclic nucleotide-binding domain, which is connected to the sixth transmembrane domain, which forms the pore, by a region known as the C-linker (Wainger et al., 2001; Robinson and Siegelbaum, 2003; Zagotta et al., 2003).

To disrupt the Ih gene in Drosophila, we used transposase-based excision of an intronic P element. In Figure 1B, the position of two intronic insertional mutants, the P element P[GSV2]GS50880 and the piggyBac element pBac(RB)[e01599], are shown relative to the Ih gene structure. These insertions were selected because they occur in an intron shared by most predicted Ih transcripts (Figure 1B). We mobilized the P[GSV2] element and screened a large set of independent excision lines for lethality, then used PCR to amplify regions of the Ih locus from the most viable and most lethal strains by PCR to identify the underlying molecular lesion. We isolated a viable precise excision strain, designated IhIE, for use as a revertant control. Several homozygous lethal excision alleles also failed to complement an Ih deficiency, IhIE. Of these, a lethal imprecise excision strain, designated IhIE, had a large deletion that ablated the entire Ih locus. All alleles were outcrossed five times to an isogenic control strain in order to remove second-site mutations.

Next, we confirmed that Ih mRNA is absent in the IhIE genotype by reverse-transcriptase polymerase chain reaction (RT-PCR) and in situ hybridization. Figure 1C shows the amplification of both genomic (left) and complementary DNA (right) purified from stage 17 embryos. The amplified region corresponds to the first three transmembrane domains of the Ih channel, and was chosen for analysis as it is immediately downstream of the insertion site of the P(GSV2) transposable element. The band corresponding to this region is absent from IhIE/IhIE embryonic gDNA and cDNA, and is present in IhIE/IhIE controls. Notably, the band is also present in cDNA from IhRB/IhRB and IhGS/IhGS embryos, suggesting that mRNA is spliced normally in spite of the large transposon insertions immediately upstream.

To confirm mRNA knockdown and observe the wild type distribution of Ih message, in situ hybridization was also performed on IhIE embryos. In IhIE/CyO, Twi-GFP heterozygotes, a DIG-labeled RNA probe to the same region as in Figure 1C generates a hybridization signal throughout the central nervous system (Figure 1D, left). By contrast, IhIE homozygotes exhibit a complete absence of neuronal DIG labeling (right), indicating that Ih mRNA is absent from these embryos, supporting our conclusion that IhIE is a null.

Ih Nulls are Late Embryonic Lethal

To determine the IhIE null lethal phase, we isolated dechorionated IhIE/IhIE embryos from their heterozygous siblings at embryonic stage 17 to observe hatching behavior. Homozygous embryos were observed for 24 h, and in all cases died without hatching (Figure 2A). When the vitelline membrane was removed manually, homozygotic larvae displayed severely reduced motility and did not develop beyond the first instar stage, dying within 24 h. This observation suggests that the Ih gene is essential and that channel may play a role in larval locomotion. Figure 2B shows that, within 6 h, the percentage of hatched eggs was severely reduced in the IhIE/IhIE and IhIE/IhIE genotype and only mildly reduced in the IhIE/IhIE and IhIE/IhIE genotypes.

Disruption of the Ih Gene Does Not Modify Larval Synaptic Morphology but Reduces the Number of Effective Vesicles Released by Presynaptic Terminals

We next examined the morphology of the larval NMJ to determine if it was altered in third instar larvae with a disrupted Ih gene. For these experiments, we focused on non-lethal hypomorphic allelic combinations (i.e., IhIE/IhIE, IhIE/IhIE), which survive to the third instar larval stage and can thus be analyzed at late larval stages. Figure 3A shows representative confocal images of muscle 4 NMJs in third instar larval fillets, stained with α-HRP to visualize neuronal membranes and α-Bruchpilot to reveal active zones. IhIE/IhIE hypomorphic NMJs (right) were similar in appearance to those of wild type larvae (left), a result that was also observed at muscle 6/7 (not shown). Neuronal branching and number of boutons were quantified for both muscle 4 and muscle 6/7 NMJs, revealing no significant differences between the IhIE/IhIE and IhIE/IhIE genotypes and only mildly reduced in the IhIE/IhIE and IhIE/IhIE genotypes.

To determine the role of the Ih gene in neurotransmission at the NMJ, we carried out electrophysiological recordings from muscle 6/7 of third instar larvae (representative electrophysiological traces, Figure 4A). For the heteroallelic loss-of-function mutant combinations, IhIE/IhIE and IhIE/IhIE, excitatory postsynaptic potentials (EPSPs) were significantly reduced compared to wild-type IhIE/IhIE controls (Figure 4B). By contrast, larval NMJs homozygous for the precise excision IhIE/IhIE allele had evoked potentials that were indistinguishable from wild type (Figure 4B). Together, these results suggest that loss-of-function lesions in the Ih locus impair neurotransmission. We noted that EPSP decay kinetics were unchanged between mutants and controls (Figure 4C). This result demonstrated unchanged timing for muscles to process a response to
neurotransmitter, which likely reflects unchanged postsynaptic development.

Nevertheless, we were uncertain if the diminished evoked amplitudes for Ih mutants reflected diminished presynaptic neurotransmitter release or a diminished capacity of the muscle to detect neurotransmitter. To address this question, we turned to additional electrophysiological measures. We found that the average amplitude of spontaneous miniature excitatory postsynaptic potentials (mEPSPs)—an indicator of muscle sensitivity to single vesicles of neurotransmitter—was not significantly altered when comparing Ih mutants either to wild type or to precise excision controls (Figure 5A). Interestingly, mEPSP frequency varied somewhat between genotypes (Figure 5B), but was not significantly changed for the Ih mutants. Additionally, we measured and plotted the cumulative probability of mEPSP amplitude for all genotypes (minimum 800 mEPSPs analyzed per genotype, see Methods). By this analysis, there was a slight—and statistically significant—decrease in the cumulative size distribution of mEPSPs for the Ih mutant combinations, showing a lower probability of release could be decreased in Ih mutants, we challenged NMJs of all recorded genotypes with short 10 Hz stimulus trains and analyzed the amplitude of the first two pulses to determine the paired pulse ratio (EPSP2/EPSP1, Figure 5F). Classically, one would expect synapses with a lower probability of release to have higher paired pulse ratios—due to a rapid accumulation of Ca$^{2+}$ in the presynaptic cleft and recruitment of presynaptic vesicles upon inducing a second pulse. Consistent with the EPSP data, both Ih mutant combinations showed a higher paired pulse ratio than wild-type and precise excision controls. For our data sets, only the Ih$^{IE}$/Ih$^{RB}$ combination achieved statistical significance ($p = 0.01$ by ANOVA vs. w$^{1118}$). By contrast, the Ih$^{IE}$/Ih$^{GS}$ combination achieved a similar elevated EPSP2/EPSP1 ratio vs. w$^{1118}$, but only trended toward statistical significance ($p = 0.18$).

**Ih Hypomorphs Exhibit Adult Motility Defects**

To examine how the Ih gene impacts motility in adult flies, we first examined flies containing transposon mutational insertions into the Ih gene. When we placed the two independent insertions, P[GSV2][GS580880 and pBac(RB)[e01599], over an Ih deficiency, Ih$^{P[RB]}$, we observed a similar motility phenotype in homozygous adult flies that included poor balance and spontaneous tremors as well as an abbreviated lifespan (Supplementary Figure 1). Due to the embryonic lethality of Ih$^{IE}$/Ih$^{IE}$ homozygotes, we focused on the Ih$^{IE}$/Ih$^{RB}$ genotype in order to characterize the effects of the Ih imprecise excision in adult flies. Figure 6A shows representative images of 15 day-old Ih$^{IE}$/Ih$^{RB}$ and Ih$^{RB}$/Ih$^{RB}$ adults, and highlights the hyperactive limb movement observed in these alleles compared to wild type adults. We found that the abnormal motility observed Ih$^{IE}$/Ih$^{RB}$ flies is comparable to that of Ih$^{RB}$/Ih$^{RB}$ flies (see Supplementary Videos 1–3).

In order to quantify the impaired motility of mutant Ih alleles, we assayed climbing behavior of adult flies (Figure 6B). A climbing index was established by measuring the percentage of adults able to climb four centimeters in 8 s. The climbing
indices of $Ih^{RB}/+$ and $Ih^{DF}/+$ heterozygotes, as well as that of precise excision homozygotes ($Ih^{PE}/Ih^{PE}$) were not significantly different from wild type flies throughout the course of the experiment ($p > 0.05$ on day 17). In contrast, all hypomorphic allelic combinations tested, included $Ih^{IE}/Ih^{RB}$, $Ih^{RB}/Ih^{RB}$, and $Ih^{DF}/Ih^{RB}$, had a dramatically lower climbing index than that of wild type flies by day 5 after eclosion (Figure 6B, $p < 0.001$ in all comparisons). As predicted, the $Ih^{IE}/Ih^{RB}$ allele had the weakest climbing index, dropping to zero by the second week. Our finding that climbing ability is strongly compromised in a series of adult $Ih$ mutant alleles is consistent with a role for the HCN channel in maintenance of basal synaptic function at the neuromuscular junction.

**DISCUSSION**

We examined the functional consequences of $Ih$ gene disruption in *D. melanogaster*. Notably, electrophysiological recordings show that $Ih$ gene hypomorphs exhibit reduced synaptic transmission at the NMJ in larvae because of a reduction in the number of effective vesicles released by presynaptic motor terminals. Previously, pharmacological inhibition of the presynaptic HCN channel by ZD7288 was shown to reduce the stimulation of neurotransmitter release by serotonin at the NMJ of *D. melanogaster* larvae, but this drug had no effect on basal neurotransmitter release in flies or in crayfish (Beaumont and Zucker, 2000; Cheung et al., 2006). A lack of an effect on basal
release by ZD7288 could have resulted from a compensating inhibition of other ion channels or from an incomplete inhibition of \( I_h \) that still allows enough HCN channel activity to maintain basal release. Alternatively, the reduced evoked release we observed may be due to alterations in other presynaptic ion channels resulting from the chronic loss of the \( I_h \) gene.

How might loss of presynaptic HCN channels reduce the number of effective vesicles released? Hyperpolarization-induced activation of the HCN channel normally depolarizes the membrane potential because of its permeability to both sodium and potassium ions. Thus, the activity of presynaptic HCN channels may depolarize the membrane voltage in the distal axon and motor terminal, and stimulate calcium influx by activating voltage-gated calcium channels, which then promotes release of neurotransmitter-containing vesicles. Alternatively, the channel may interact directly with synaptic release machinery upon activation and promote neurotransmitter release; this has been suggested previously as the mechanism by which serotonin augments neurotransmitter release in the NMJ of the crayfish (Beaumont and Zucker, 2000).

Using a combination of genetic approaches, presynaptic \( I_h \) has been shown to reduce glutamate release from amacrine cells in flies by limiting the activity of Cacophony (Cac)-containing (Cav2 alpha1 subunit) calcium channels (Hu et al., 2015). Cav2 channels are critical for appropriate evoked neurotransmission at the NMJ, as well as the execution of plastic modulations of neurotransmission, as seen with homeostatic synaptic plasticity (Rieckhof et al., 2003; Xing et al., 2005; Frank et al., 2006; Frank, 2014). Based on the data garnered in amacrine cells, \( I_h \) could promote neurotransmission through Cav2 regulation at the NMJ.

In the mouse, inhibition of low threshold voltage-gated T-type (Cav3.2) \( \text{Ca}^{2+} \) channels by presynaptic \( I_h \) has also been noted in synaptic terminals in entorhinal cortical layer III pyramidal neurons (Huang et al., 2011). Thus, in general, the presynaptic HCN channel may either stimulate or inhibit neurotransmitter release, depending upon the neuronal type and the context provided by other ion channels in the terminal axon and synapse.

Disruption of the \( I_h \) gene leads to a range of behavioral defects in adult flies, which include tremors, imbalance, decreased motility, and shortened lifespan. Homozygotic disruption of the \( I_h \) gene by imprecise excision produced a lethal phenotype, in which embryos did not hatch. Even when the vitelline membrane was removed manually, null \( I_h^{IE} \) homozygotic larvae moved slowly and did not survive beyond the first instar (within 24 h). Thus, the inability of the embryos to hatch was likely due to reduced motor activity. A previous study carried out a partial \( I_h \) deletion strategy which produced an allele that did not cause

**FIGURE 4 | \( I_h \) disruption reduces evoked potentials in larval motor terminals.** (A) Representative electrophysiological recordings from the neuromuscular junction. Scale bars for EPSPs (mEPSPs): 5 mV (1 mV); 50 ms (1000 ms). (B) Plots of average evoked EPSPs in muscle recorded in response to nerve stimulation vs. genotype. Each point represents the average of 30 EPSPs recorded from a single muscle (1 Hz), and each bar represents the average (± SEM) of all muscles recorded for the given genotype. EPSPs recorded from both experimental groups (\( I_h^{IE}/I_h^{RB} \) and \( I_h^{IE}/I_h^{GS} \)) are significantly smaller than those recorded from the wild type (**\( p < 0.001 \) for \( I_h^{IE}/I_h^{RB} \) vs. \( w^{1118} \) control; **\( p = 0.002 \) for \( I_h^{IE}/I_h^{GS} \) vs. \( w^{1118} \) control by one-way ANOVA with Tukey’s post-hoc to control for multiple group comparisons), whereas those recorded from the precise excision (\( I_h^{PE}/I_h^{PE} \)) are not significantly different from wild type EPSPs (\( p = 0.54 \) vs. \( w^{1118} \)). (C) Plots of average EPSP decay (90% peak to 10% peak) times for all recorded genotypes. Point data and averages displayed as in 4B. No genotype showed any statistically significant difference in evoked potential decay (\( p \) ranged from 0.44 to 0.98 for all possible pairwise comparisons by one-way ANOVA with Tukey’s post-hoc).
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FIGURE 5 | Ih disruption diminishes presynaptic vesicle release. (A) Plots of average quantal size (mEPSP amplitude) for each genotype. Each point represents 100–200 spontaneous mEPSP events measured and averaged for a single muscle; each bar represents the average (±SEM) mEPSP amplitude for each genotype. There are no significant differences in average quantal size among the genotypes (p ranges from 0.35 to 0.99 for all possible pairwise comparisons by one-way ANOVA with Tukey’s post-hoc). (B) Plots of average spontaneous mEPSP frequency for each genotype. Point data and averages displayed as described in 5A. Neither Ih hypomorphic combination significantly differs from controls (p > 0.1 vs. w1118 by one-way ANOVA with Tukey’s post-hoc). (C) Cumulative probability histograms of quantal size for each genotype examined. A minimum of 800 spontaneous mEPSP events over a minimum of 9 NMJs were analyzed for each genotype. Compared to control genotypes, IhIE/IhRB shows a slight diminishment in mEPSP size, assessed by cumulative probability (Kruskal-Wallis Test with Dunn’s multiple comparisons post-hoc). However, IhIE/IhGS shows no significant difference compared to controls. (D) Plots of quantal content (QC) determined from evoked potentials recorded in muscle in response to electrical stimulation vs. genotype. Each point represents the average of 30 EPSPs in one muscle divided by the average of 100–200 mEPSPs recorded from the same muscle. Bars are the average for each genotype (±SEM). Quantal content is reported as an uncorrected (average EPSP/average mEPSP). By this measure, both Ih hypomorphic combinations have diminished QC compared to w1118, but neither reaches statistical significance (p > 0.14 vs. w1118 by one-way ANOVA with Tukey’s post-hoc). (E) Plots of quantal content corrected for non-linear summation (NLS, See Methods). Points and averages plotted as in 5D. Both Ih hypomorphic combinations have diminished NLS QC compared to w1118 (***p = 0.006 for IhIE/IhRB; **p = 0.05 for IhIE/IhGS vs w1118 one-way ANOVA with Tukey’s post-hoc). (F) Paired-pulse plots (EPSP2/EPSP1) measured from short 10 Hz trains of evoked stimuli for each genotype. Point data are from the average of two separate paired pulse recordings per muscle and bars are the average (±SEM) for each genotype. The IhIE/IhRB combination achieved statistical significance (***p = 0.01 by ANOVA vs. w1118). The IhIE/IhGSV combination achieved a similar elevated EPSP2/EPSP1 ratio vs. w1118, but only trended toward statistical significance (p = 0.18).
embryonic lethality (Gonzalo-Gomez et al., 2012). The more robust phenotype observed in our null allele may be due to the fact that our approach using imprecise P-element excision ablated the majority of the \( \text{Ih} \) locus.

Adult hypomorphic mutant flies demonstrated an overall phenotype that is remarkably similar to that observed in mice either lacking the HCN2 isoform (Ludwig et al., 2003), or possessing a C-terminal truncated form of this isoform (Chung et al., 2009). Upon HCN2 knockout, mice are hypoactive, show a wide-based and abnormal gait, and exhibit a tremor (Ludwig et al., 2003). Video recordings and EEGs show that these mice suffer from absence epilepsy and frequent concomitant spike-wave discharges, which contribute to hypoactivity. Mice possessing a C-terminal truncated HCN2 channel possess a phenotype which is referred to as \textit{apathetic} (Chung et al., 2009); they exhibit ataxia, generalized spike-wave absence seizures, and tonic–clonic seizures. Homozygotic \textit{apathetic} mice cannot coordinate swimming when forced to do so, and have difficulty with balance. For example, they fall to one side when standing on their hind legs to feed. Furthermore, \textit{apathetic} homozygotes, but not heterozygotes, exhibit uncoordinated movement, unsteady gait and intention tremor. Finally, \textit{apathetic} homozygotes die prematurely from unknown causes. The mechanism by which the genetic disruption of the \( \text{Ih} \)/HCN channel leads to these phenotypes in mice remains unknown, but it could be related to its role in altering neurotransmission at the neuromuscular junction.

There are few published reports on the HCN channel in \( D. \text{melanogaster} \). Disruption of the \( \text{Ih} \) gene has been proposed previously to decrease life span and modify sleep and dopamine levels in adult flies (Chen and Wang, 2012; Gonzalo-Gomez et al., 2012). It seems reasonable to expect that the fly model will yield significant insights into structure, function and physiology of mammalian HCN channels, given the pertinent data contained in the few studies carried out to date (Marx et al., 1999; Gisselmann et al., 2005; Cheung et al., 2006; Gonzalo-Gomez et al., 2012; Hu et al., 2015) and the considerable impact that the fly has had on understanding closely-related voltage-gated ion channels (Littleton and Ganetzky, 2000; Cirelli et al., 2005). In particular, the fly offers a desirable model system to examine presynaptic \( \text{Ih} \) and its interactions with other presynaptic channels and proteins by exploiting genetic approaches and not relying exclusively on \( \text{Ih} \) inhibitor drugs, such as ZD7288.
ETHICS STATEMENT

This study was approved by the Office of Research Services at the University of British Columbia and the Office of The Institutional Animal Care and Use Committee at the University of Iowa.

AUTHOR CONTRIBUTIONS

AH, DA, and EA conceived the original idea and experiments. AH, CAF, MK, AB, DA, and EA designed the experiments, and contributed to the analysis and interpretation of data. AH, CAF, DA, and EA edited the manuscript. AH organized and prepared the first draft of the paper, prepared the figures and, with EA, guided the editing process and prepared the final manuscript. AH, CAF, AB, and DA carried out the experiments.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: http://journal.frontiersin.org/article/10.3389/fnmol.2017.00041/full#supplementary-material

Supplementary Figure 1 | Ih disruption reduces viability of Drosophila melanogaster adults. Graph showing the number of flies surviving over time for three genotypes. Flies with a P-element insertion in one copy (Ih/P+) have a normal lifespan of ~60 days, as do flies lacking the HCN locus on one chromosome (Ih/Df). In contrast, flies with disruptions in both copies of the HCN gene (Ih/IhDf) have a significantly shortened lifespan (~20-day median). Starting populations of 50 flies per genotype were followed daily until all had died.

Supplementary Videos 1–3 | Climbing ability in 15 day-old Ih/Eh/IhRB flies (video 2) and IhRB/IhRB flies (video 3) is comparable and both are compromised as compared to wild type control flies.

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Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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