Disorders of the Nervous System

Characterization of Seizure Induction Methods in Drosophila

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Abstract

Epilepsy is one of the most common neurologic disorders. Around one third of patients do not respond to current medications. This lack of treatment indicates a need for better understanding of the underlying mechanisms and, importantly, the identification of novel targets for drug manipulation. The fruit fly Drosophila melanogaster has a fast reproduction time, powerful genetics, and facilitates large sample sizes, making it a strong model of seizure mechanisms. To better understand behavioral and physiological phenotypes across major fly seizure genotypes we systematically measured seizure severity and secondary behavioral phenotypes at both the larval and adult stage. Comparison of several seizure-induction methods; specifically electrical, mechanical and heat induction, show that larval electroshock is the most effective at inducing seizures across a wide range of seizure-prone mutants tested. Locomotion in adults and larvae was found to be non-predictive of seizure susceptibility. Recording activity in identified larval motor neurons revealed variations in action potential (AP) patterns, across different genotypes, but these patterns did not correlate with seizure susceptibility. To conclude, while there is wide variation in mechanical induction, heat induction, and secondary phenotypes, electroshock is the most consistent method of seizure induction across known major seizure genotypes in Drosophila.

Key words: antiepileptic; Drosophila; epilepsy; insect; model; seizure induction

Significance Statement

Epilepsy is a neurologic disorder affecting one in 130 people globally, with a significant impact on patients, families, and society. Approximately one third of epileptics do not respond to currently available medication. Thus, better insights into underlying disease mechanisms and identification of new drugs are needed. Fruit flies (Drosophila melanogaster) are a powerful genetic model: a number of single gene mutant flies exhibit seizures, phenotypes that have been shown to respond to established antiepileptic drugs (AEDs). We compare methods of seizure induction and their utility, to establish which induction method is the most consistent across a range of different seizure-inducing genetic backgrounds. Adopting a common method for seizure analysis in this model will, we predict, speed identification of novel anti-convulsive treatments.

Introduction

Epilepsy is one of the most common neurologic disorders, affecting ~60 million people worldwide (Chen et al., 2018). While a variety of causes contribute to epilepsy, including traumatic brain injury and brain infections, the major contribution is from underlying genetic mutations...
It has been estimated that ~70% of epilepsies do not have a single known cause; of these, 60% have been associated with genetic mutations (Heron et al., 2007; Poduri and Lowenstein, 2011). There are around 700 identified gene mutations currently associated with epilepsy. These include genes contributing to planar cell polarity and the noncanonical WNT signaling pathway (e.g., PRICKLE1), autism spectrum associated genes (e.g., AUTS2), and mTOR signaling pathway genes (e.g., mTOR and TSC1; Bassuk et al., 2008; Citraro et al., 2016; Wang et al., 2017). However, a majority of epilepsy genes directly influence ion-channel function, specifically mutations in voltage-gated sodium, potassium, and calcium channels.

Clinicians have access to over 25 antiepileptic drugs (AEDs) to minimize epileptic seizures (Löschner and Schmidt, 2011). A majority of these drugs target ion channels or neurotransmitter signaling, in an attempt to re-establish an appropriate balance between excitatory and inhibitory signaling in the brain (Löschner and Schmidt, 2011; Vezzani et al., 2011). Although many new AEDs have been approved in recent years, many new drugs have proven no more effective in treating drug-resistant epilepsies than older compounds (Schmidt and Schachter, 2014; Moshé et al., 2015; Chen et al., 2018). It seems likely that treating drug-refractory epilepsy will require novel drug targets and therefore a deeper understanding of epilepsy at the level of basic mechanism(s).

In the genetic model Drosophila, both wild-type and mutant animals exhibit seizure-like behaviors; mutants undergo seizures with greatly reduced stimulus thresholds and/or seizure-like activity (SLA) lasts far longer (Ganetzky and Wu, 1982; Pavlidis et al., 1994; Kuebler and Tanouye, 2000; Parker et al., 2011a; Baines et al., 2017). In the adult fly, SLA includes repetitive proboscis extension, wing buzzing and loss of posture (Tan et al., 2004; Parker et al., 2011a). These SLAs have formed the basis of a variety of seizure-severity assays (Pavlidis and Tanouye, 1995). Several reviews have previously characterized fly electrophysiological and behavioral responses to specific seizure assays (Pavlidis and Tanouye, 1995; Kuebler and Tanouye, 2000; Parker et al., 2011a; Lee et al., 2019). In adult flies, seizure-susceptible mutants fall into two main categories based on seizure induction: mechanical (termed bang sensitive; BS) and temperature induced (Ganetzky and Wu, 1982; Kasbekar et al., 1987; Pavlidis and Tanouye, 1995; Burg and Wu, 2012). Mechanical induction has been standardized with the use of a laboratory vortexer to hyperstimulate sensory inputs and is termed the “vortex assay” (Kuebler and Tanouye, 2000; Fig. 1A). By contrast, the heat assay exploits temperature change to induce seizure (Burg and Wu, 2012; Saras and Tanouye, 2016; Fig. 1B). There is a third, and more involved seizure induction method in adult flies, known as high-frequency stimulation of the giant fiber (GF) pathway (Pavlidis and Tanouye, 1996). This method induces seizures in all BS and other seizure genotypes like shaker (Lee and Wu, 2006). While this assay allows investigation of synaptic transmission during seizures, stimulation of the GF pathway requires a more complex set up than either the vortex or temperature-shock assays and is not suitable for medium-throughput or high-throughput screening.

In larvae, seizures have been induced using a simplified electroshock assay, during which the whole body is subjected to electroshock (Marley and Baines, 2011; Fig. 1C). A particular advantage of larvae is that they are well-suited to drug screening and, moreover, provide unparalleled understanding of CNS structure and function (Choi et al., 2004; Worrell and Levine, 2008; Marley and Baines, 2011; Kadas et al., 2015; Lemon et al., 2015).

Genetic epilepsies include syndromes characterized by febrile (heat-“fever”-induced) seizures that often present in children. Genetic epilepsy with febrile seizures plus (GEFS+) is commonly caused by sodium channel mutations (Camfield and Camfield, 2015). Some extreme cases of GEFS+ are classified as Dravet syndrome (DS). This often affects children in their first year of life, and has additional comorbidities including motor and mental impairments (Ziobro et al., 2018). DS is typically resistant to AED treatment necessitating additional research. Use of simpler model organisms, including Drosophila, are well suited to investigate mechanisms of epilepsy, but also to identify novel treatments for drug-refractory epilepsies (Schutte et al., 2014; Griffin et al., 2017).

We aimed to establish which accessible seizure-induction methods are applicable across the known range of Drosophila seizure mutants. We found that electroshock applied to larvae is the most reliable method to induce seizures regardless of mutation. Since epilepsy is associated with many secondary phenotypes, we investigated whether locomotor activity is altered in a predictable fashion, which could be further explored to use for drug screens. We found that there was no significant trend in the daily adult activity levels across the range of mutants tested, but mechanically-sensitive mutant larvae showed reduced crawling. Since electroshock was the most widely applicable method of seizure induction, we investigated potential common underlying mechanisms predisposing larvae to seizures. We detected elongated bursts of motor neuron firing in all seizure mutants. However, the alteration was not exclusive to seizure mutants and did not have good predictive power for seizure severity measured as seizure duration.
Fly stocks
Flies were maintained on standard cornmeal medium at 25°C and a 12/12 h light/dark cycle. Fly stocks were obtained from the following sources: DS (S1231R), DS-C (S1231S), GEFS+ (K1270T), and GEFS-C (K1270K) from Diane O’Dowd (Sun et al., 2012; Schutte et al., 2014); w-, eae2F;1; pabs1;1, Canton-S and Oregon-R from Baines lab; w-, iusdefo7.3 from M. Tanouye; pk-sple (#422) from the Bloomington Drosophila Stock Centre.

Adult activity assay
Male flies, aged 4–10 d posteclosion, were placed in 65-mm glass tubes containing food in one end and a foam plug at the other. The experiments were conducted at 12/12 h light/dark cycle in an environment-controlled incubator at 25°C. Fly locomotor activity was monitored for 4 d, always excluding the first 24 h from data analysis. Animal activity was measured by distance moved per hour (mm/h) and obtained using custom-build image-acquisition software (CRITTA), written in LabView (National Instruments; Mohammad et al., 2016). All tracking experiments were repeated at least twice, fly N numbers are listed in the respective figures. Before tracking, flies were reared at 25°C, collected on the day of eclosion and kept in vials containing only males until the day of experiment.

Adult vortex assay
Adult flies were collected using CO2 3–4 d posteclosion into empty plastic vials. Ten flies per vial were left to recover from anesthesia for 1–2 h. To assess seizure, the vials were placed on a standard laboratory vortexer at maximum speed for 10 s. Seizure duration was measured as time required to regain posture and mobility for each fly.

Adult heat-shock assay
To measure seizure threshold and duration, adult flies were subjected to a heat assay. Male flies, aged 8–10 d posteclosion, were anesthetized using CO2 and placed into empty plastic fly vials, 5 flies per vial. Flies were given 1–2 h to recover. A water bath was heated to 40–41°C and kept constant at that temperature throughout the experiment. Vials containing flies were placed into the water bath and kept there for 120 s. Throughout the 2 min in the water bath, visual inspection occurred every 10 s and seizure-status noted. Seizures were identified by loss of posture and random wing buzzing. Vials were taken out of the bath after 120 s and the duration of seizure activity outside the water bath was also recorded (i.e., recovery period).

Larval locomotion tracking
Larval locomotion was tracked using the DanioVision Observation Chamber connected to a computer with EthoVision XT software (Noldus Information Technology); 2% agarose was poured into the lid of 96-well plate and four separate arenas were cut out. Grooves between the arenas were filled with 5 m NaCl to prevent larvae from crawling off their respective stages. Individual larvae were placed on the arenas and after a 30-s adaptation period their locomotion was tracked for 3 min. Total distance crawled in the tracking period was calculated by the EthoVision XT software using centroid tracking.

Larval electroshock assay
An electroshock assay induces seizure in third instar larvae. Wandering stage larvae were transferred from a vial containing food onto a plastic dish containing water to wash away food residue. Larvae were then transferred to an empty plastic dish and excess water gently removed with a paper towel. No more than four larvae were placed in a single dish. Once normal crawling behavior resumed a probe, composed of two conductive tungsten wires (0.1 mm in diameter, ~1–2 mm apart), was placed on the anterior dorsal surface of a larva, over the
approximate location of the CNS. A 7V DC current was applied for 2 s, generated by a DS2A Isolated Voltage Stimulator (Digimeter Ltd). In response to the shock, larvae exhibit sustained contractions of body wall muscles and occasional spasms, halting normal crawling behavior. Seizure duration was measured as a time period between the stimulus onset and resumption of normal crawling behavior. After seizure, the resumption of normal behavior involved a full peristaltic wave from either end of the animal which resulted in either forward or backward movement.

**Larval drug treatment**
Gravid females and males were kept on grape agar plates (Dutscher) at 25°C. The flies were fed live yeast paste supplemented by 25 mg/ml picrotoxin (PTX; Sigma-Aldrich) stock to a final concentration of 0.125 mg/ml for 3 d before embryo collection commenced. Exposed embryos were transferred onto standard food at the end of each collection day and allowed to develop normally.

**Larval electrophysiology dissection**
To record from identified aCC or RP2 motor neurons in third instar larvae, the CNS was dissected from a larva and placed in a droplet of external saline solution on a coverslip coated with a thin layer (1 μm) of cured SYLGARD Elastomer (Dow-Corning). The external saline solution was composed of 135 mM NaCl, 5 mM KCl, 4 mM MgCl2·6H2O, 2 mM CaCl2·2H2O, 5 mM N-Tris [hydroxy-methyl]methyl-2-aminoethanesulfonic acid (TES), and 36 mM sucrose at pH 7.15. The CNS was stabilized on the coverslip using tissue adhesive (Gluture, World Precision Instruments). The aCC or RP2 cells, located in the dorsal ventral nerve cord (VNC), were exposed for recordings by removing the overlying glial sheath using a glass pipette (GC100TF-10, Harvard Apparatus) filled with 1% (w/v) bacterial protease type XIV (Sigma-Aldrich).

**Loose patch recordings**
Loose patch recordings were performed using borosilicate glass electrodes (GC100TF-10; Harvard Apparatus) pulled to resistance between 1.0 and 1.5 MΩ. The electrodes were filled with external saline of the same composition as listed above. The aCC or RP2 neurons were identified based on location in the CNS, relative soma size and axon configuration (aCC has two axons originating from the soma, RP2 has just one). An electrode was placed on a cell soma without breaking into it. Slight negative pressure was applied to suck around 1/3 of the cell into the electrode. Recordings were conducted using Axopatch 200B amplifier controlled by pCLAMP (version 10.3) using a Digidata 1322A analog-to-digital converter (Molecular Devices, Molecular Devices) or MultiClamp 700B amplifier controlled by WinWCP (version 5.1) using a Digidata 1440A analog-to-digital converter (Molecular Devices, Molecular Devices). Data were sampled at 20 kHz and low pass filtered at 0.1 kHz. All recordings were made at room temperature (18–22°C).

**Electrophysiology trace analysis**
Analysis of loose patch traces was performed using a custom script written in MATLAB. Spike times were gathered using Clampfit (version 11.1). A burst is an event with a minimum of three spikes occurring within 100 ms from one another. A burst ends when no spike is detected within 100 ms following the last spike. The script provides summary data on mean burst duration and standard deviation, burst frequency, spike count per burst and firing frequency. Data were plotted using a custom Python script.

**Code accessibility**
MATLAB script described in the paper is freely available online at https://github.com/jm933/BurstAnalysis.

**Statistical analysis**
Data were analyzed with estimation methods using a Python custom script using the DABEST package (Ho et al., 2019). The effect size distributions shown are of the 5000 mean differences calculated in the bootstrap resampling procedure. To compare effect sizes across different assays a universal effect size measure Cohen’s d was used. Cohen’s d is an effect size where the difference has been standardized to the standard deviations of the two groups being compared (Cumming and Calin-Jageman, 2016). No significance testing was performed, p values were reported for legacy purposes only. Data are presented following best practices, showing observed values and effect sizes; means are provided with a standard deviation, effect-size errors are displayed as curves with 95% confidence intervals (CIs; generated with bootstrapping; Bernard, 2021).

Inferred linear relationships between various parameters were assessed using linear regression. The line was fitted and R² was calculated using scipy library in Python using linregress function, which finds the line with the least-squares regression. Each point represents the effect size of a genotype. Sample sizes (N) are the total number of animals for behavioral experiments or total number of cells for electrophysiology.

**Results**

**Mechanical seizure induction**
*Drosophila* mutants with increased susceptibility to mechanical shock are known as BS (Parker et al., 2011a). There are more than a dozen BS single gene mutants (Baines et al., 2017), including paraactive Naᵥ channel (Parker et al., 2011b), easily-shocked (eas) an ethanolamine kinase (Pavlidis et al., 1994), *julius seizure* (jus; also known as *slamdance*) with unknown protein function (Home et al., 2017; Zhang et al., 2002), and *pricklespin legs* (*pk-sple*), a protein involved in microtubule polarity (Tao et al., 2011; Ehaideb et al., 2014). Of these mutants, *bss* has the most severe seizure phenotype (Parker et al., 2011b). In our experiments with vortex induction (Fig. 1A), *bss* similarly exhibited the longest-duration seizures (168.4 ± 11.0 s; Fig. 2A). By
comparison, eas exhibited an average seizure duration of 107.7 ± 8.9 s and jus of 73.6 ± 3.9 s. The majority of pk-sple did not exhibit severe seizures.

Identical stimulation of wild-type lines (CS, OR) resulted in only brief disruption to posture, with a typical recovery of 0.2 ± 0.1 s (Fig. 2A). Both heat-sensitive mutants, GEFS+ or DS, did not exhibit any seizure-like behavior following mechanical stimulation (Fig. 2A).

**Temperature seizure induction**

Several seizure mutants are inducible by elevated temperature (Salkoff and Kelly, 1978; Kasbekar et al., 1987; Figure 2. Comparison of seizure induction methods in Drosophila adults and larvae. We tested BS pk-sple, eas, jus, and bss lines and heat-sensitive DS and GEFS+ lines. A, Vortex-induced seizures in pk-sple 15.22 s [95% CI: 11.37, 20.69], eas 107.54 s [95% CI: 88.71, 123.84], jus 73.35 s [95% CI: 64.60, 79.88], and bss 168.35 s [95% CI: 157.31, 179.39]. No SLA was observed in CS, DS-C, DS, GEFS-C, and GEFS+ flies. B, Seizure duration induced by the heat assay. DS showed seizures of average 71.82 s and GEFS+ of 431.03 s, with no seizures recorded for the respective matched controls (DS-C and GEFS-C). pk-sple showed seizures of 138.67 s [95% CI: 87.42, 191.84]. By comparison, CS, OR, jus, bss, or eas did not show heat-induced seizures. C, Cumulative fraction of flies seizing throughout the 120 s of heat assay. Only 100% of DS flies seized during the 2-min period. Other genotypes reached a maximum of: 90% for GEFS+, 83% for pk-sple, and 20% for eas. D, Electroshock induced seizures in all genotypes tested. Results of OR, jus, eas, bss, and pk-sple are reported as a ratio to CS seizure duration which was measured at 97.0 s [95% CI: 95.4, 101.6]. There were no differences between OR and CS. Pk-sple exhibited the weakest phenotype with seizure duration increased by 57.47%, whereas bss showed an increase of 244.11%. eas and jus had 112.62% and 104.95% longer seizures. DS and GEFS+ seizure durations are reported as a ratio to their respective controls (DS-C and GEFS-C). DS showed 119.78% and GEFS+ 70.67% increase in seizures as compared with controls. E, Seizure induction in larva and adult flies have similar effectiveness ($R^2 = 0.85$ [95% CI: 0.03, 1.0], shaded area indicates 95% CI for the linear regression model fit). Effect sizes for adults were derived from vortex assay for BS and heat assay for temperature-sensitive lines.
Burg and Wu, 2012; Hill et al., 2019). Significantly, there are two lines which model-specific human epilepsy mutations: GEFS+ (Sun et al., 2012) and DS (Schutte et al., 2014). These “humanized knock-in” mutants exhibit seizures after exposure to elevated temperatures of 38°C and higher (Schutte et al., 2014). We tested the same range of mutants in heat-shock (Fig. 2C) as described for mechanical-shock induction, finding that 90% of GEFS+ flies exhibit seizure after 120 s, while 100% of DS flies exhibit seizure after 50 s at 41°C (Fig. 2C). Seizure duration, measured after the heat application was terminated, revealed seizures of average 349 s for GEFS+ and 72s for DS (Fig. 2B). This shows that although DS has a lower threshold for seizure induction, the seizure event is less severe compared with GEFS+.

Some BS lines have been shown to have seizures on exposure to cold temperatures including bss and eas at 8°C, but not at 39°C (Burg and Wu, 2012). We find that eas, jus, and bss do not exhibit seizures in the heat assay (at 41°C; Fig. 2B). By comparison, 83% of pk-sple tested showed seizure-like behavior with an average time of recovery around 137 s (Fig. 2B,C).

Larval electroshock seizure induction

Larvae of Drosophila have a peristaltic motor action that has been exploited to study seizure. Application of a brief electric shock to the dorsal cuticle induces SLA (Marley and Baines, 2011; Fig. 1C). We tested whether this seizure-induction method could be used on both the mechanical-induced and heat-induced seizure mutants tested above. We found that, in response to electroshock, all the mutants exhibited long-lasting seizures (Fig. 2D). Of these, bss mutants exhibited the strongest phenotype having seizures 344% longer in duration than wild type (CS). Additional mutants, jus and eas, showed similar duration seizures at 105% and 113% longer than CS, whereas pk-sple seizures were 58% longer. Relative to their own genetic controls, GEFS+ and DS had seizure durations extended by 71% and 120%, respectively (Fig. 2D).

Seizure induction method comparison

Since electroshock is applied at a different stage of development (i.e., larval) as compared with mechanical and heat-shock (i.e., adult), we compared efficacy of seizure induction methods across the different stages. Since larval and adult seizure duration metrics with different scales were being compared, rather than raw differences, we used Cohen’s d, a standardized effect size (Cumming and Calin-Jageman, 2016). Cohen’s d for adults was derived from the vortex assay on BS mutants and heat assay on HS mutants. They were compared with the efficacy of e-shock in larvae (Fig. 2E). The higher the value of effect size the stronger seizure phenotype was observed in the mutant as compared with control. Although the seizure assay at the adult stage had slightly bigger effect size than at the larval stage, there is a good correlation between the efficacies of seizure induction across the two different stages. This means that seizure severity across the two stages, larvae and adult, is comparable. Thus, electroshock seems to be an efficient and easily applicable method to induce reliable seizures in both mechanical-induced and heat-induced mutants, making it a favorable method to compare seizure severity across many, if not all, seizure mutations.

Changes in adult locomotor activity

Alterations in activity patterns and/or sleep duration have been reported for Drosophila seizure mutants (Petruccelli et al., 2015). Some epilepsy patients, including DS sufferers present with associated ataxias (Oakley et al., 2011; Marcían et al., 2016). We hypothesized that fly seizure mutants could have similar secondary phenotypes in addition to seizure, and that these may provide “simpler” predictors of seizure susceptibility. To address this, we measured adult locomotor activity over a 3-d period (12/12 h cycle) using a custom video-tracking system (Fig. 3A). During the light phase (subjective day) BS mutants showed slightly lower activity (Fig. 3B). During darkness (i.e., subjective night) the same seizure-mutant flies tended to have an increased activity compared with wild-type controls (Fig. 3C). The largest difference from CS wild type during subjective day was exhibited by eas at 1012 mm/h reduction, the largest difference during subjective night was also exhibited by eas, where it was more active by 667 mm/h.

Of the heat-inducible lines, GEFS+ was slower by 654 mm/h during the day and by 313 mm/h during night relative to its own matched genetic control. However, DS was 361 mm/h more active in light but showed no difference to the controls in darkness. Out of the lines investigated in this study, DS is the only genotype to have increased activity during the day. GEFS+ was the only genotype to be consistently slower than control.

The stark differences between day-phase and night-phase activity changes for the differing mutants tested, and the absence of a consistent pattern between activity changes and seizure-induction type or severity, indicate the hypothesis is false: adult locomotor activity is not a reliable predictor of seizure severity.

BS mutants show reduced larval locomotion

There are previous reports showing alterations in either larval locomotion or peristaltic wave frequency in seizure susceptible Drosophila mutants (Stilwell et al., 2006; Graham et al., 2016; Streit et al., 2016). We decided to test for such in the range of mutants used for this study and compare that to the results of adult activity. We performed larval tracking and reported a total distance crawled by each genotype in a 3-min period (Fig. 3D,E). CS wild type had the most activity with a total distance traveled of 161.47 mm. The range of BS mutants showed a consistently lower locomotion level as compared with CS: pk-sple slower by 112.20 mm [95% CI: −129.94, −94.07], eas by 100 mm [95% CI: −117.24, −84.80], jus by 109.18 mm [95% CI: −124.02, −92.49] and bss being the slowest with a reduction of 129.05 mm [95% CI: −143.67, −114.46]. Unlike BS mutants, the range of HS flies did not show any meaningful differences in their...
locomotor activity among themselves, and in comparison to CS (Fig. 3E). These differences in phenotype reinforce the observation from adult fly activity levels, levels of larval locomotor activity are also not predictive of seizure severity.

Larval motor neuron activity patterns are not predictive of seizures

Observing that larval electroshock applies to the full panel of seizure mutants we have tested, we decided to investigate electrophysiological properties that may be associated with seizure susceptibility. Because the behavior we observed is because of altered muscular activity, we focused attention to motoneurons. We performed loose patch recordings from the aCC or RP2 motoneurons in third instar larvae. These motoneurons were selected because of their accessibility and prior extensive characterization (Baines and Bate, 1998; Choi et al., 2004). Recordings from aCC/ RP2, in wild-type larvae, show reduced locomotor activity (compared with appropriate controls), with eas and GEFS+ having the most prominent phenotypes; slower by 1012.09 mm/h [95% CI: −1211.17, −818.82] and 654.49 mm/h [95% CI: −1017.36, −267.84], respectively. DS showed increased activity by 361 mm/h [95% CI: 154.58, 592.13] over its control (DS-C) with an average speed of 727.60 mm/h. During lights off, several mutants showed a small increase in activity levels. Of these, eas had the most prominent phenotype at 667.02 mm/h [95% CI: 493.03, 870.27]. dus and bss showed similar activity levels with an increase of 261.51 mm/h [95% CI: 128.67, 404.60] and 285.95 mm/h [95% CI: 138.50, 458.49], respectively. GEFS+ is the only genotype remaining slower than its respective control (GEFS-C) by 313.14 mm/h [95% CI: −489.98, −154.34]. Other genotypes showed no meaningful differences. Fly activity was tracked for 3 d at 25°C at 12/12 h light/dark cycle. All genotypes were recorded at least twice. During the 12-h period of lights on, dus (1298.14 mm/h), eas (724.40 mm/h), pk-sple (1148.99 mm/h), and GEFS+ (1007.61 mm/h) show reduced locomotor activity (compared with appropriate controls), with eas and GEFS+ having the most prominent phenotypes; slower by 1012.09 mm/h [95% CI: −1211.17, −818.82] and 654.49 mm/h [95% CI: −1017.36, −267.84], respectively. DS showed increased activity by 361 mm/h [95% CI: 154.58, 592.13] over its control (DS-C) with an average speed of 727.60 mm/h. C, During lights off, several mutants showed a small increase in activity levels. Of these, eas had the most prominent phenotype at 667.02 mm/h [95% CI: 493.03, 870.27]. dus and bss showed similar activity levels with an increase of 261.51 mm/h [95% CI: 128.67, 404.60] and 285.95 mm/h [95% CI: 138.50, 458.49], respectively. GEFS+ is the only genotype remaining slower than its respective control (GEFS-C) by 313.14 mm/h [95% CI: −489.98, −154.34]. Other genotypes showed no meaningful differences. Fly activity was tracked for 3 d at 25°C at 12/12 h light/dark cycle. All genotypes were recorded at least twice. D, Schematic showing the set up for larval tracking. E, Total distances crawled by BS mutants pk-sple (49.27 mm), eas (60.51 mm), dus (52.29 mm), and bss (32.43 mm) were greatly lower in comparison to CS (161.47 mm). Heat-sensitive mutants did not show any meaningful reduction in the total distance crawled from their matched controls.

Figure 3. Adult seizure mutants do not exhibit obvious changes in locomotor activity. A, Schematic showing the locomotor assay. B, Average activity over 3 d at 25°C at a 12/12 h light/dark cycle. C, During the 12-h period of lights on, dus (1298.14 mm/h), eas (724.40 mm/h), pk-sple (1148.99 mm/h), and GEFS+ (1007.61 mm/h) show reduced locomotor activity (compared with appropriate controls), with eas and GEFS+ having the most prominent phenotypes; slower by 1012.09 mm/h [95% CI: −1211.17, −818.82] and 654.49 mm/h [95% CI: −1017.36, −267.84], respectively. DS showed increased activity by 361 mm/h [95% CI: 154.58, 592.13] over its control (DS-C) with an average speed of 727.60 mm/h. C, During lights off, several mutants showed a small increase in activity levels. Of these, eas had the most prominent phenotype at 667.02 mm/h [95% CI: 493.03, 870.27]. dus and bss showed similar activity levels with an increase of 261.51 mm/h [95% CI: 128.67, 404.60] and 285.95 mm/h [95% CI: 138.50, 458.49], respectively. GEFS+ is the only genotype remaining slower than its respective control (GEFS-C) by 313.14 mm/h [95% CI: −489.98, −154.34]. Other genotypes showed no meaningful differences. Fly activity was tracked for 3 d at 25°C at 12/12 h light/dark cycle. All genotypes were recorded at least twice. D, Schematic showing the set up for larval tracking. E, Total distances crawled by BS mutants pk-sple (49.27 mm), eas (60.51 mm), dus (52.29 mm), and bss (32.43 mm) were greatly lower in comparison to CS (161.47 mm). Heat-sensitive mutants did not show any meaningful reduction in the total distance crawled from their matched controls.
some variation in all of these parameters, burst duration is consistent with relatively small variation in wild-type but larger variation (in terms of observed SD) in seizure mutants (Table 1). Based on this insight, we classified bursting traces as normal if the average burst duration was the mean of CS burst duration ± 1.5 SD. Using this classification, we found all mutant lines tested had some normal bursting in addition to some elongated bursting traces (Fig. 4C–E). The fraction of elongated bursting was variable across genotypes, but consistently observed in both BS and HS seizure mutants. Of the mutants tested, jus had the highest fraction of elongated bursts among BS (42%; Fig. 4C) and DS among the HS mutants (58%; Fig. 4D). However, unlike the two wild types, DS and GEFS+ matched genetic controls also showed appreciable elongated burst firing. Thus, compared with their respective controls (and not true wild types), DS and GEFS+ showed increased elongated bursting of 17% and 150%, respectively.

Table 1: Analysis of activity recordings from larval motor neurons

| Genotype | N  | Burst duration (ms) | Burst duration SD (ms) | Burst frequency (Hz) | Firing rate (AP/s) | AP count per burst (AP/burst) | AP frequency per burst (Hz) |
|----------|----|---------------------|------------------------|----------------------|-------------------|------------------------------|-----------------------------|
| CS       | 13 | 174.57              | 91.54                  | 0.47                 | 6.68              | 14.25                        | 87.53                       |
| OR       | 10 | 155.40              | 84.80                  | 0.40                 | 6.38              | 16.24                        | 104.30                      |
| pk-sple  | 22 | 394.01              | 348.54                 | 0.46                 | 6.92              | 16.81                        | 51.48                       |
| jus      | 12 | 703.94              | 1211.46                | 0.31                 | 8.23              | 23.26                        | 58.88                       |
| eas      | 14 | 210.22              | 165.49                 | 0.49                 | 6.73              | 12.78                        | 69.78                       |
| bss      | 12 | 195.42              | 215.28                 | 0.43                 | 4.97              | 12.72                        | 77.89                       |
| GEFS-C   | 12 | 208.94              | 183.28                 | 0.37                 | 4.32              | 13.67                        | 76.16                       |
| GEFS+    | 12 | 1160.34             | 1827.30                | 0.46                 | 9.48              | 49.54                        | 362.66                      |
| DS-C     | 12 | 11,859.25           | 11,869.45              | 0.39                 | 6.21              | 166.21                       | 322.43                      |
| DS       | 12 | 3139.83             | 4338.58                | 0.23                 | 4.39              | 42.26                        | 34.13                       |
| CS –PTX  | 12 | 1964.82             | 5369.33                | 0.37                 | 13.99             | 65.30                        | 40.28                       |
| CS +PTX  | 12 | 170.16              | 89.99                  | 0.30                 | 4.07              | 12.59                        | 90.34                       |

Three-minute loose-patch recordings of spiking activity were analyzed using a custom-built MATLAB script. Bursts were defined as three or more spikes occurring within 100 ms from each other. The numbers are averages for all traces per genotype.
however, suggests that elongated bursting is not exclusive to seizure mutants.

PTX is known to induce SLA in wild-type flies. The advantage of this approach is that the genetic background is identical between control and PTX-fed animals. We tested for elongated bursting in a PTX-induced seizure model (CS fed PTX) and found that the fraction of elongated bursting was increased to 83%. By contrast unfed control animals showed no elongated bursts (Fig. 4E). PTX-induced seizures in larvae were verified using the e-shock assay, results of which show an increase in recovery time by 86% (Fig. 4F).

We further investigated how aberrations in bursting recordings might correlate with larval seizure (Fig. 5). We used mean burst duration, burst duration standard deviation, burst frequency, AP firing rate, AP per burst count and frequency. None of these parameters revealed any correlation between them and larval seizure severity (Fig. 5A–F). However, frequency with which abnormalities occur could have a more significant impact on seizure susceptibility than the absolute values of deviations in burst duration. We investigated whether the fraction of elongated bursting could be predictive of seizure duration. We found that there is no strong correlation ($R^2 = 0.08$) between

Figure 5. Larval motor neuron firing does not predict seizure susceptibility. Analysis of 3-min loose-patch recordings from larval RP2/aCC motor neurons revealed that there is no correlation between mean burst duration (A), SD of burst duration (B), burst frequency (C), AP count per burst (D), AP frequency per burst (E), AP firing rate (F), and fraction of elongated bursting (G) and larval seizure. (H) Fraction of elongated bursting from larval motor neurons showed no correlation with the effect size of adult seizure. Shaded areas indicate 95% CI for the linear regression model fit.
the two measures (Fig. 5G). The alterations in larvae development could potentially be predictors of adult phenotypes, however, when we compared adult seizure to bursting patterns, the measures were also not well-correlated ($R^2 = 0.14$; Fig. 5H). Thus, we conclude that while altered motor neuron activity is typical for seizure mutants, it is not diagnostic for seizure severity.

**Discussion**

Epilepsy is characterized by recurrent seizures, which can manifest as absence seizures, underlying periods of inattention, through to full generalized myoclonic seizures, resulting in muscle jerks to full collapse and unconsciousness. While *Drosophila* seizure mutants lack these full range of behaviors, seizures in both humans and flies nevertheless exhibit sufficient parallels to implicate that the underlying neuronal abnormalities are highly similar. Previous investigations have shown in *Drosophila* that seizures include (1) a defined seizure threshold, (2) genetic mutations that modify seizure-susceptibility, (3) electroshock therapy raises the threshold for subsequent seizures, (4) seizures spread throughout the CNS along defined neuronal tracts, (5) seizures can be localized to distinct regions of the CNS, and (6) seizures can be ameliorated by AEDs used to treat human epilepsy (Kuebler and Tanouye, 2000; Kuebler et al., 2001; Reynolds et al., 2004; Tan et al., 2004; Marley and Baines, 2011; Lin et al., 2017). Unlike human epilepsy, however, fly seizure mutants do not routinely exhibit spontaneous seizures which may reflect fewer numbers of neurons within the latter. However, there are a couple of notable exceptions: pk-sple (Tao et al., 2011; Ehaideb et al., 2016) and DS (Schutte et al., 2014). Nonetheless, seizures in flies can be successfully induced using either mechanical, temperature or electroshock induction in a variety of single gene mutant backgrounds, much as seizures can be induced in mammals by extreme external stimuli (light or sound), fever, and/or electroshock.

Our study reproduced earlier results validating that all of the lines tested exhibit induced-seizure behavior by a variety of methods (Pavlidis et al., 1994; Zhang et al., 2002; Parker et al., 2011a; Tao et al., 2011; Sun et al., 2012; Schutte et al., 2014; Horne et al., 2017). The length of seizure varies by mutation, with bss being the strongest BS phenotype in our experiments and, indeed, the strongest reported so far (Parker et al., 2011b). GEFS+ exhibited longer duration seizures than DS following heat-shock, although DS had a lower seizure induction threshold. This is similar to humans, where DS is normally a more severe syndrome than GEFS+ (Brunklus and Zuberi, 2014).

As in adults, seizures can be induced in *Drosophila* larvae. Electroshock induction has been successfully applied before (Marley and Baines, 2011). These authors reported the sda mutant larva (renamed to jus; Horne et al., 2017) exhibit seizure durations up to 6.6 times longer than CS wild-type controls. Likewise, our electroshock induction was effective in inducing seizures in all mutants studied. Seizure duration was variable by genotype, with the longest duration seizures seen in bss. Comparing the severity of seizures induced in either adults or larvae suggests that, although adults tend to have longer and more complex seizures, the chosen method for induction is critical.

Larval electroshock is comparable to adult electroshock. Adult electroshock successfully induces seizures in a range of BS mutants including eas, bss, and jus tested in our study (Pavlidis et al., 1994; Pavlidis and Tanouye, 1995). There is a stereotypical sequence of seizure in adults including two distinct phases of dorsal longitudinal muscle failure defined as a silent period lasting 35–40 s and abnormal responses during recovery which varied by genotype. However, adult electroshock does not always evoke seizures in temperature-sensitive mutants, e.g., seizure. By contrast, our results indicate that electroshock of larvae successfully induced seizures in all mutants and, moreover, generated a comparable hierarchy of seizure severity to adults. Similar to adult electroshock, the response is quite stereotyped in stages. It starts with larval paralysis of varied duration depending on genotype. This is followed by both, or either, partial peristaltic waves and mouth hook contractions of high frequency. Recovery is determined once several full peristaltic waves occur in the animal (Marley and Baines, 2011).

There are other methods available to reliably induce seizures in many mutants such as the proconvulsant PTX, which works in both larvae and adults (Stilwell et al., 2006; Lee et al., 2019). However, in this study we show that electroshock is a simple and reliable method, applicable to many seizure genotypes. We would suggest its use for studies of novel gene mutations considered to be seizure related.

We also explored whether adult seizure mutants exhibit pronounced alterations in locomotion, without seizure induction, as has been reported in some human epilepsies (particularly those associated with ataxias). Tracking adult fly activity over several days revealed that there are changes in daytime activity, although these were not consistent across the genotypes studied. During the night, activity was increased; this partly mimics human disorders where epilepsy impairs sleep (Jain and Kothare, 2015; Wang et al., 2018). However, there was again no obvious uniform trend. As such, our results only partially replicated previously observed changes in activity (Petrucelli et al., 2015), whereas the previous study shows an increase in activity during the night, we observed an overall decrease in GEFS+ activity in both day and night. This might be explained by accumulation of genetic modifiers over time. Alterations in food composition are also known to affect seizure phenotype in flies and potential differences in fly food could also impact secondary phenotypes observed (Stone et al., 2013; Fogle et al., 2019; Kasuya et al., 2019). On the other hand, the overall trend we observed matches previously reported data where seizure mutants are, overall, less active than controls (Radlicz et al., 2019).

Earlier research suggests that the reduced levels of sleep in flies could lead to increased seizure susceptibility (Lucey et al., 2015). The study showed that, in seizure prone BS flies ses$^{TS}$ and heat-sensitive sei$^{TS}$, there are no
alterations to sleep pattern, but sleep deprivation leads to increased seizure severity. However, another study by Petruccelli et al. (2015) reported reduced seizure likelihood and increased seizure threshold after sleep deprivation in GEFS+ flies. Clearly, fly models of epilepsy (like in human patients) are heterogenous and each mutation potentially has a distinct relationship to sleep and its effect on susceptibility to seizure. Both prior research and the present study suggest that sleep–seizure relationships could be investigated further for each genotype individually to better understand the underlying mechanisms.

Changes in larval locomotion exhibited a clear difference between mechano-sensitive and heat-shock-sensitive lines not evident in adult behavior. BS larvae showed a consistently lowered activity whereas HS larvae did not show any movement phenotype. A previous report shows that both GEFS+ and bss exhibit an increase in synchronicity in larval peristaltic waves (Streit et al., 2016); however, in our study, it did not translate to the same extent of alterations to locomotion. Moreover, there does not seem to be a clear relationship between locomotion and recovery time from electroshock in larvae. BS lines like pk<sup>opieh</sup>, /jus, and eae have reduced locomotion but their recovery time is comparable to DS and GEFS+ which display higher levels of locomotive activity.

Susceptibility to seizures may arise from mutations in a diverse range of genes including those that encode ion channels and/or microtubule polarity-associated genes. However, given a similar behavioral phenotype (i.e., seizure), we explored the possibility of a common underlying change in cellular activity. To investigate this idea, we performed loose-patch recordings from larval motor neurons aCC or RP2 (note, these two neurons are highly similar in their biophysical properties; Baines et al., 1999). In wild-type larvae (CS and OR), these neurons exhibit a robust and consistent bursting pattern. The same activity patterns were also present in the seizure mutants and their matched controls. However, in addition, seizure mutants also exhibited altered activity patterns that were not observed in the wild types. This altered activity showed increased burst duration accompanied by modest increases in AP firing frequency. Increased AP firing is a hallmark of epileptic seizures (Marley and Baines, 2011; Parker et al., 2011b; Ehaideb et al., 2016). However, our data clearly shows that the presence of elongated burst firing is not well-correlated to induced seizure duration. Thus, at present it is difficult to understand the causes of altered firing activity. The fact that such activity was also observed in both GEFS+ and DS control lines are suggestive that it is not exclusive to seizure mutants. To negate differences in genetic background, that may complicate our results, we fed PTX to CS and observed a greatly increased fraction of elongated bursting. Thus, while burst elongation might not be a sufficient explanation for seizure severity, these prolonged activity bursts are clearly associated with seizure per se. It is possible that the poor correlation with seizure severity could be because of differing efficiencies of neuronal homeostatic compensation present in the differing seizure mutations we investigated.

In conclusion, <i>Drosophila</i> seizure mutants form a diverse group of models of epilepsy syndromes. Mutants can be separated by seizure induction modes; of the methods tested here, larval electroshock is the induction method most broadly applicable. The more general adoption of larval electroshock could help reduce variability between reported studies that use differing methods of seizure induction. Thus, we recommend this method as a benchmark technique for testing known and novel <i>Drosophila</i> seizure mutations.

References

Baines RA, Bate M (1998) Electrophysiological development of central neurons in the <i>Drosophila</i> embryo. J Neurosci 18:4673–4683.
Baines RA, Giachello CNG, Lin WH (2017) <i>Drosophila</i>. In: Models of seizures and epilepsy, Chap 24, Ed 2 (Pitkänen A, Buckmaster PS, Galanopoulou AS, Moshé SL, eds), pp 345–358. San Diego: Academic Press.
Baines RA, Robinson SG, Fujikoa M, Jaynes JB, Bate M (1999) Postsynaptic expression of tetanus toxin light chain blocks synapto genesis in <i>Drosophila</i>. Curr Biol 9:1267–1270.
Bassuk AG, Wallace RH, Buhr A, Buller AR, Afawi Z, Shimojo M, Miyata S, Chen S, Gonzalez-Alpargue P, Griesbach HL, Wu S, Nashefsky M, Vladar EK, Antic D, Ferguson PJ, Cirak S, Voit T, Scott MP, Axelrod JD, Gurnett C, et al. (2008) A homoyzgous mutation in human PRICKLE1 causes an autosomal-recessive progressive myoclonic epilepsy-ataxia syndrome. Am J Hum Genet 83:572–581.
Bernard C (2021) Estimation statistics, one year later. eNeuro 8: ENEURO.0091-21.2021.
Brunklaus A, Zuberi SM (2014) Dravet syndrome—from epileptic en cephalopathy to channelopathy. Epilepsia 55:979–984.
Burg MG, Wu CF (2012) Mechanical and temperature stressor-induced seizure-and-paralysis behaviors in <i>Drosophila</i> bang-sensi tive mutants. J Neurogenet 26:189–197.
Camfield P, Camfield C (2015) Febrile seizures and genetic epilepsy with febrile seizures plus (GEFS+). Epileptic Disord 17:124–133.
Chen Z, Brodie MJ, Liew D, Kwan P (2018) Treatment outcomes in patients with newly diagnosed epilepsy treated with established and new antiepileptic drugs: a 30-year longitudinal cohort study. JAMA Neurol 75:279–286.
Choi JC, Park D, Griffith LC (2004) Electrophysiological and morphological characterization of identified motor neurons in the Drosophila third instar larva central nervous system. J Neurophysiol 91:2353–2365.
Citroaro R, Leo A, Constanti A, Russo E, De Sarro G (2016) mTOR pathway inhibition as a new therapeutic strategy in epilepsy and epileptogenesis. Pharmacol Res 107:333–343.
Cuming G, Calin-Jageman R (2016) Introduction to the new statistics: estimation, open science, and beyond. Ed 1. New York: Routledge.
Ehaideb SN, Iyengar A, Ueda A, Iacobucci GJ, Cranston C, Bassuk AG, Gubb D, Axelrod JD, Gunawardena S, Wu CF, Manak JR (2014) prickle modulates microtubule polarity and axonal transport to ameliorate seizures in flies. Proc Natl Acad Sci USA 111:11187–11192.
Ehaideb SN, Wignall EA, Kasuya J, Evans WH, Iyengar A, Koerselman HL, Lilienthal AJ, Bassuk AG, Kitamoto T, Manak JR (2016) Mutation of orthologous prickle genes causes a similar epilepsy syndrome in flies and humans. Ann Clin Transl Neurol 3:695–707.
Fogle KJ, Smith AR, Satterfield SL, Gutierrez AC, Hertzler JI, McCordell CS, Shon JH, Barile ZJ, Novak MO, Palladino MJ (2019) Ketogenic and anaplerotic dietary modifications ameliorate seizure activity in <i>Drosophila</i> models of mitochondrial encephalomyopathy and glycolytic enzvmopathy. Mol Genet Metab 126:439–447.
Ganetzky B, Wu CF (1982) Indirect suppression involving behavioral mutants with altered nerve excitability in *Drosophila melanogaster*. Genetics 100:597–614.

Graham S, Rogers RP, Alper RH (2016) An automated method to assay locomotor activity in third instar *Drosophila melanogaster* larvae. J Pharmacol Toxicol Methods 77:78–80.

Griffin A, Hamling KR, Knupp K, Hong S, Lee LP, Baraban SC (2017) Clemizole and modulators of serotonin signalling suppress seizures in Dravet syndrome. Brain 140:669–683.

Heron SE, Scheffer IE, Berkovic SF, Dibbens LM, Mulley JC (2007) Channelopathies in idiopathic epilepsy. Neurotherapeutics 4:295–304.

Hill AS, Jain P, Folan NE, Ben-Shahar Y (2019) The Drosophila ERG channel seizure plays a role in the neuronal homeostatic stress response. PLoS Genet 15:e1008288.

Ho J, Tumkaya T, Aryal S, Choi H, Claridge-Chang A (2019) Moving beyond P values: data analysis with estimation graphics. Nat Methods 16:556–566.

Home M, Krebshevsky K, Wells A, Tunio N, Jarvis C, Francisco G, Geiss J, Recknagel A, Deitcher DL (2017) julius seizure, a *Drosophila* mutant, defines a neuronal population underlying epileptogenesis. Genetics 205:1261–1269.

Jain SV, Kothare SV (2015) Sleep and epilepsy. Semin Pediatr Neurol 22:86–92.

Kadas D, Ryglewski S, Duch C (2015) Transient BK outward current enhances motoneurone firing rates during *Drosophila* larval locomotion. J Physiol 593:4871–4888.

Kasbekar DP, Nelson JC, Hall LM (1987) Enhancer of seizure: a new genetic locus in *Drosophila melanogaster* defined by interactions with temperature-sensitive paralytic mutations. Genetics 116:423–431.

Kasuya J, Iyengar A, Chen H-L, Lansdon P, Wu CF, Kitamoto T (2019) Milk-whey diet substantially suppresses seizure-like phenotypes of paraShu, a *Drosophila* voltage-gated sodium channel mutant. J Neurogenet 33:164–178.

Kuebler D, Tanouye MA (2000) Modifications of seizure susceptibility in *Drosophila*. J Neurophysiol 83:998–1009.

Kuebler D, Zhang H, Ren X, Tanouye MA (2001) Genetic suppression of seizure susceptibility in *Drosophila*. J Neurophysiol 86:1211–1225.

Lee J, Wu CF (2006) Genetic modifications of seizure susceptibility and expression by altered excitability in *Drosophila* Na(+) and K (+) channel mutants. J Neurophysiol 96:2456–2478.

Lee J, Iyengar A, Wu CF (2019) Distinctions among electroconvulsions and proconvulsant-induced seizure discharges and native motor patterns during flight and grooming: quantitative spike pattern analysis in *Drosophila* flight muscles. J Neurogenet 33:125–142.

Lemon WC, Pulver SR, Höckendorf B, McDole K, Branson K, Freeman J, Keller PJ (2015) Whole-central nervous system functional imaging in larval *Drosophila*. Nat Commun 6:7924.

Lin WH, Giachello CNG, Baines RA (2017) Seizure control through genetic and pharmacological manipulation of Pumilio in *Drosophila*: a key component of neuronal homeostasis. Dis Model Mech 10:141–150.

Löschner W, Schmidt D (2011) Modern antiepileptic drug development has failed to deliver: ways out of the current dilemma. Epilepsia 52:657–678.

Lucey BP, Leahy A, Rosas R, Shaw PJ (2015) A new model to study sleep deprivation-induced seizures. Sleep 38:777–785.

Marciani V, Filip P, Barès M, Brázdiil M (2016) Cerebellar dysfunction and ataxia in patients with epilepsy: coincidence, consequence, or cause? Tremor Other Hyperkinet Mov (Ny) 6:376.

Marley R, Baines RA (2011) Increased persistent Na+ current contributes to seizures in the siamdance bang-sensitive *Drosophila* mutant. J Neurophysiol 106:18–29.

Mohammad F, Aryal S, Ho J, Stewart JC, Norman NA, Tan TL, Eisaka A, Claridge-Chang A (2016) Ancient anxiety pathways influence *Drosophila* defense behaviors. Curr Biol 26:981–986.

Moshé SL, Perucca E, Ryvlin P, Tomson T (2015) Epilepsy: new advances. Lancet 385:884–898.

Oakley JC, Kalume F, Catterall WA (2011) Insights into pathophysiology and therapy from a mouse model of Dravet syndrome. Epilepsia 52 [Suppl 2]:59–61.

Parker L, Howlett IC, Rusan ZM, Tanouye MA (2011a) Seizure and epilepsy: studies of seizure disorders in *Drosophila*. Int Rev Neurobiol 99:1–21.

Parker L, Padilla M, Du Y, Dong K, Tanouye MA (2011b) *Drosophila* as a model for epilepsy: bss is a gain-of-function mutation in the para sodium channel gene that leads to seizures. Genetics 187:523–534.

Pavlids P, Tanouye MA (1995) Seizures and failures in the giant fiber pathway of *Drosophila* bang-sensitive paralytic mutants. J Neurosci 15:5810–5819.

Pavlids P, Ramaswami M, Tanouye MA (1994) The *Drosophila* easily shocked gene: a mutation in a phospholipid synthetic pathway causes seizure, neuronal failure, and paralysis. Cell 79:23–33.

Petruccelli E, Lansdon P, Kitamoto T (2015) Exaggerated nighttime sleep and defective sleep homeostasis in a *Drosophila* knock-in model of human epilepsy. PLoS One 10:e0137758.

Poduri A, Lowenstein D (2011) Epilepsy genetics—past, present, and future. Curr Opin Genet Dev 21:325–332.

Radlcz C, Chambers A, Oliis E, Kuebler D (2019) The addition of a lipid-rich dietary supplement eliminates seizure-like activity and paralysis in the *Drosophila* bang sensitive mutants. Epilepsy Res 155:106153.

Reynolds ER, Stauffer EA, Feeley L, Rojahn E, Jacobs B, McKeever C (2004) Treatment with the antiepileptic drugs phenytoin and gabapentin ameliorates seizure and paralysis of *Drosophila*bang-sensitive mutants. J Neurobiol 58:503–513.

Salkoff L, Kelly L (1978) Temperature-induced seizure and frequency-dependent neuromuscular block in a ts mutant of *Drosophila*. Nature 273:156–158.

Saras A, Tanouye MA (2016) Seizure suppression by high temperature via cAMP modulation in *Drosophila*. G3 (Bethesda) 6:3381–3387.

Schmidt D, Schachter SC (2014) Drug treatment of epilepsy in adults. BMJ 348:g254.

Schutte RJ, Schutte SS, Alagara J, Barragan EV, Gilligan J, Staber C, Savva YA, Smith MA, Reenan R, O’Dowd BK (2014) Knock-in model of Dravet syndrome reveals a constitutive and conditional reduction in sodium current. J Neurophysiol 112:903–912.

Stilwell GE, Sarasawti S, Littleton JT, Chouinard SW (2006) Development of a *Drosophila* seizure model for in vivo high-throughput drug screening. Eur J Neurosci 24:2211–2222.

Stone B, Evans L, Coleman J, Kuebler D (2013) Genetic and pharmacological manipulations that alter metabolism suppress seizure-like activity in *Drosophila*. Brain Res 1496:94–103.

Streit AK, Fan YN, Masullo L, Baines RA (2016) Calcium imaging of neuronal activity in *Drosophila* van identify anticonvulsive compounds. PLoS One 11:e0148461.

Suter L, Gillian J, Staber C, Schutte RJ, Nguyen V, O’Dowd DK, Reenan R (2012) A knock-in model of human epilepsy in *Drosophila* reveals a novel cellular mechanism associated with heat-induced seizure. J Neurosci 32:14145–14155.

Tan JS, Lin F, Tanouye MA (2004) Potassium bromide, an anticonvulsant, is effective at alleviating seizures in the *Drosophila*52. Nature 273:156–158.

Tan JS, Lin F, Tanouye MA (2004) Potassium bromide, an anticonvulsant, is effective at alleviating seizures in the *Drosophila*52. Nature 273:156–158.

Tao H, Manak JR, Sowers L, Mei X, Kiyonari H, Abe T, Dahdah NS, Yang T, Wu S, Chen S, Fox MH, Gurnett C, Montine T, Bird T, Shaffer LG, Rosenfeld JA, McConnell J, Madan-Khetarpal S, Cheung YK, Yarnette CM, Griesbach H, et al. (2011) Mutations in prickle orthologs cause seizures in flies, mice, and humans. Am J Hum Genet 88:138–149.

Vezzani A, French J, Bartfai T, Baram TZ (2011) The role of inflammation in epilepsy. Nat Rev Neurol 7:31–40.

Wang J, Lin ZJ, Liu X, Xu HQ, Shi YH, Yi YH, He N, Liao WP (2017) Epilepsy-associated genes. Seizure 44:11–20.
Wang YQ, Zhang MQ, Li R, Qu WM, Huang ZL (2018) The mutual interaction between sleep and epilepsy on the neurobiological basis and therapy. Curr Neuropharmacol 16:5–16.
Worrell JW, Levine RB (2008) Characterization of voltage-dependent Ca2+ currents in identified Drosophila motoneurons in situ. J Neurophysiol 100:868–878.
Zhang H, Tan J, Reynolds E, Kuebler D, Faulhaber S, Tanouye M (2002) The Drosophila slammedance gene: a mutation in an aminopeptidase can cause seizure, paralysis and neuronal failure. Genetics 162:1283–1299.
Ziobro J, Eschbach K, Sullivan JE, Knupp KG (2018) Current treatment strategies and future treatment options for dravet syndrome. Curr Treat Options Neurol 20:52.