ROYAL SOCIETY OPEN SCIENCE

rsos.royalsocietypublishing.org

Research





Cite this article: Hermann SL, Xue S, Rowe L, Davidson-Lowe E, Myers A, Eshchanov B, Bahlai CA. 2016 Thermally moderated firefly activity is delayed by precipitation extremes. *R. Soc. open sci.* **3**: 160712. http://dx.doi.org/10.1098/rsos.160712

Received: 19 September 2016 Accepted: 17 November 2016

Subject Category:

Biology (whole organism)

Subject Areas:

ecology/environmental science

Keywords:

lightning bug, Lampyridae, phenology, ecoinformatics, long-term ecological research

Author for correspondence:

Christie A. Bahlai e-mail: cbahlai@msu.edu

Thermally moderated firefly activity is delayed by precipitation extremes

Sara L. Hermann^{1,2,5}, Saisi Xue^{3,6}, Logan Rowe¹, Elizabeth Davidson-Lowe^{1,2}, Andrew Myers^{1,5}, Bahodir Eshchanov¹ and Christie A. Bahlai^{4,7}

CAB, 0000-0002-8937-8709

The timing of events in the life history of temperate insects is most typically primarily cued by one of two drivers: photoperiod or temperature accumulation over the growing season. However, an insect's phenology can also be moderated by other drivers like rainfall or the phenology of its host plants. When multiple drivers of phenology interact, there is greater potential for phenological asynchronies to arise between an organism and those with which it interacts. We examined the phenological patterns of a highly seasonal group of fireflies (Photinus spp., predominantly P. pyralis) over a 12-year period (2004–2015) across 10 plant communities to determine whether interacting drivers could explain the variability observed in the adult flight activity density (i.e. mating season) of this species. We found that temperature accumulation was the primary driver of phenology, with activity peaks usually occurring at a temperature accumulation of approximately 800 degree days (base 10°C); however, our model found this peak varied by nearly 180 degree-day units among years. This variation could be explained by a quadratic relationship with the accumulation of precipitation in the growing season; in years with either high or low precipitation extremes at our study site, flight activity was delayed. More fireflies were captured in general in herbaceous plant communities with minimal soil disturbance (alfalfa and no-till field crop rotations), but only weak interactions occurred between within-season responses to climatic variables and plant community. The interaction we observed between temperature and precipitation accumulation

¹Department of Entomology, ²Ecology, Evolutionary Biology and Behavior Program,

³Biomass Conversion Research Laboratory, Department of Chemical Engineering, and

⁴Department of Integrative Biology, Michigan State University, East Lansing, MI 48824, USA

⁵Department of Entomology, Pennsylvania State University, State College, PA 16803. USA

⁶DOE Great Lakes Bioenergy Research Center, East Lansing, MI 48824, USA

⁷Mozilla Science Laboratory, Mozilla, Mountain View, CA 94041, USA

suggests that, although climate warming has the potential to disrupt phenology of many organisms, changes to regional precipitation patterns can magnify these disruptions.

1. Introduction

Much can be learned about biological systems by observation alone [1], and observational data are often captured incidentally as a result of human activity [2]. Incidental data can range from the very informal and uncontrolled (e.g. comments on a topic in a Web forum) to highly controlled and meticulously collected (e.g. unused data from scientific experiments). Indeed, research activities can produce systemic observational data of very high quality; for instance, insect trapping systems seldom only capture target taxa. This 'by-catch' can provide data that support investigations into entirely uninvestigated phenomena. In this study, we examine one such 'by-catch' dataset: a 12-year time series of firefly observations in southwestern Michigan, for their responses to environmental and habitat conditions.

Over 2000 species of firefly (Coleoptera: Lampyridae) have been identified across various temperate and tropical environments around the world [3]. As larvae, species within the family Lampyridae spend much of their time living underground feeding on earthworms, molluscs and other subterranean invertebrates [4]. As adults, most species abstain from feeding [5], with the exception of the species *Photuris pennsylvanica*, of which the female is a voracious predator of both conspecifics as well as other insects [5–7]. Few studies have been conducted on firefly conservation and broader-scale ecology in relation to changing environments and land uses, and little is known about how environmental parameters drive firefly life history. It has been demonstrated that the life history of at least one species of firefly is temperature-dependent; researchers found that *P. pennsylvanica* adult emergence could be artificially accelerated by exposing larvae to increased soil temperature [8]. However, much of the primary research on fireflies has focused on the bioluminescent properties of the firefly [9–14], while research describing basic population and community ecology of this important family is lacking.

In addition to the scientific importance of the Lampyridae for their bioluminescent properties and as model organisms for evolutionary investigations, fireflies are also among the most widely recognized and culturally valued insect families among non-scientists. Two US states have designated the firefly as their 'State Insect' [15]. Notably, fireflies also feature prominently in Japanese culture, where they have been designated as national natural treasures in many districts and have been used to generate support for biodiversity conservation efforts in Japanese agricultural regions [16–18]. They have also been touted as useful classroom tools for sparking student interest in biology [19]. Because of their popular appeal, it is unsurprising that public concern has grown about apparent declines of firefly populations from regions around the world where they occur [20].

Considering the paucity of ecological information about fireflies, their widespread popularity, ease with which adults are observed and concerns about their population viability, fireflies represent an ideal species for citizen science investigations. Citizen science efforts are currently underway seeking to gain information about the status, geographical distribution and phenology of fireflies [21–23], and peer-reviewed publications on fireflies have already been produced based on these volunteer-generated data [24,25]. The popularity of fireflies gives them great potential as a flagship and umbrella conservation species and potentially an indicator species of ecological degradation in agricultural regions [26]. However, to our knowledge, no long-term systematic study of firefly phenology and responses to environmental drivers has been published.

Phenology plays a significant role in regulating species abundance, distribution and biodiversity [27,28]. The timing of phenological events in insect life histories is strongly linked to climatic conditions [29–31], such as temperature and precipitation [27,32,33]. Changes in phenology can have community-wide consequences, and differential responses among various species within a community can lead to trophic mismatches [28,30]. For example, the timing of larval winter moth (*Operophtera brumata*) emergence was formerly largely synchronized with oak (*Quercus robur*) bud burst. Caterpillars that emerge too early lack a sufficient food source and will starve, while caterpillars that emerge too late will be exposed to older, poor-quality leaves, leading to negative physiological implications [34]. Increased spring temperature has resulted in changes in the timing of oak bud burst. However, the winter moth has yet to adapt to changing temperatures, which has led to disrupted synchrony between these two species [34]. Thus, phenological shifts can have both top-down and bottom-up consequences extending throughout multiple trophic levels. Long-term observations are important for understanding ecological trends and the merit of phenology as a predictor of ecological consequences. A long-term study on the

Genji firefly (*Luciola cruciate*) in Japan found that population patterns changed in response to rainfall, potentially leading to early larval emergence and reduced foraging [35]. However, the ways in which climate change and other environmental events have impacted firefly species is less understood.

Developing a model for the emergence of adult fireflies is key to developing our understanding of firefly phenology, which can then be used to expand firefly conservation efforts, educational outreach and environmental research and to predict peak firefly display. In this study, we examine a 'by-catch' dataset documenting captures of fireflies at the Kellogg Biological Station over a 12-year period and place it in the context of other available data to gain insights into the long-term dynamics and phenology of this charismatic, but understudied taxon.

2. Material and methods

2.1. Data sources

Data were obtained through two publicly available datasets—a weather dataset that included daily maximum and minimum temperature and precipitation, as well as a dataset that focuses on ladybeetle observations, but also documents captures of the other insect species. Both datasets arise from Michigan State University's Kellogg Biological Stations (KBS), located in southwestern Michigan. The firefly abundance data were collected as a part of the KBS Long Term Ecological Research (LTER) Site within the Main Cropping System Experiment (MCSE) and forest sites starting in 2004. Fireflies were recorded to family alone; however, from spot-checks of the collected data, it appears that fireflies collected belonged to the genus *Photinus*, mainly *Photinus pyralis*, the big dipper firefly, although captures of other species cannot be excluded.

Within the MCSE, seven plant community treatments were established in 1989, ranging from a three-year rotation of annual field crops (maize, soyabean and wheat) under four levels of management intensity (conventional, no-till, reduced input or biologically based), to perennial crops including alfalfa, poplar and early successional vegetation (i.e. abandoned agricultural fields maintained in an early successional state by yearly burnings; table 1 and figure 1). Each of these treatments is replicated six times across the MCSE site with each replicate consisting of a 1 ha plot. We also included three forest sites in our analysis; these sites were established in 1993 within 3 km of the MCSE site on KBS and represent one of three plant community treatments: conifer forest plantations, late-successional deciduous forest and successional forest arising on abandoned agricultural land (table 1 and figure 1). Forested treatment plots are also 1 ha in size but are replicated three times for each treatment.

Observations were taken on a weekly basis throughout the sampling season at five sampling stations within each replicate (both MCSE and forest sites). These insect abundance data are available publicly, online at http://lter.kbs.msu.edu/datatables/67. Insect abundance monitoring was done using unbaited two-sided yellow cardboard sticky cards (Pherocon, Zoecon, Palo Alto, CA, USA) suspended from a metal post within each sampling station, 1.2 m above the ground. Cards were deployed each week for a one-week exposure for the duration of the growing season. Sampling start and end dates varied each year depending upon planting date of the various crops; however, the length of sampling was fairly consistent $(14 \pm 1 \text{ weeks}, \text{ on average}, \text{ per year})$.

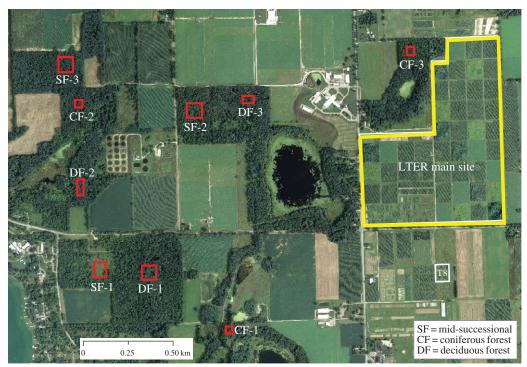
In addition to plant community treatment management information, we also included weather as an environmental factor to explain firefly abundance. These data were also obtained through a publicly available dataset, online at http://lter.kbs.msu.edu/datatables/7.

2.2. Data preprocessing

All analyses performed are available as an R script at https://github.com/cbahlai/lampyrid/. Analyses presented in this manuscript were run in R v. 3.3.1 'Bug in Your Hair' (R Development Core Team 2016). Firefly data were extracted from the database held at the KBS data archive and combined with relevant agronomic data (which are encoded in plot and treatment numbers in the main database) and are hosted at figshare at https://ndownloader.figshare.com/files/3686040.

Data were subjected to quality control manipulations to remove misspellings in variable names that had occurred with data entry. Observations with missing values for firefly counts were excluded from analysis. Because subsample data were zero-biased, we used reshape2 [36] to sum within date, within plot observations, and created an additional variable to account for sampling effort (which was usually consistent at five traps per plot per sampling period, but on occasion traps were lost or damaged).

Kellogg Biological Station LTER Main Cropping System Experiment successional and forested sites



airphoto date: 12 August 2011

Figure 1. Map of sites at the Kellogg Biological Station LTER (reproduced and modified from http://lter.kbs.msu.edu/maps/images/current-lter-forest-successional-sites.pdf). Red outlined areas indicate successional and forested sites, denoted by the key at the bottom right of the photo. Outlined in yellow is the MCSE (LTER main site) which houses seven treatments: conventional crop, no-till crop, reduced input crop, biologically based crop, poplar, alfalfa and early successional. Each replicate consists of a randomized block of 1 ha plots for each of the seven treatments, and the experiment is replicated six times in the MCSE.

Table 1. Description of the plant community treatments at the Kellogg Biological Station Long Term Echnological Research Site, in which insect sampling occurred.

crop type	plant community treatments	description
annual	conventional	rotated crop field (corn—soyabean—wheat), with conventional chemical input which is chisel ploughed
	no-till	rotated crop field (corn—soyabean—wheat), with conventional chemical input, with no tilling
	reduced input	biologically based rotated crop field (corn—soyabean—wheat), with low-input chemical control and a winter cover crop (leguminous). Plots are treated with banded herbicide and starter nitrogen at planting
	organic	biologically based rotated crop field (corn—soyabean—wheat), low input chemical control with winter cover crop (leguminous). Certified organic
perennial	poplar trees	10-year rotation cycle of a fast-growing <i>Populus</i> clone
	alfalfa	continuously grown alfalfa
	early successional	abandoned field from 1989, left to grow into native successional plants which are annually burned
forest	successional	40–60-year-old successional forest, left from former agricultural fields
	coniferous	40–60-year-old conifer plantations
	deciduous	late successional deciduous forest

Weather data (daily maximum and minimum temperatures were reported in degrees Celsius and daily precipitation in millimetres) were downloaded directly from the Kellogg Biological Station Data archive (http://lter.kbs.msu.edu/datatables/7.csv). To overcome errors in calculations requiring accumulated annual weather data caused by rare missing data points (most often occurring during winter, in periods of extreme cold leading to equipment malfunction), we created a function to replace missing values in the temperature data with the value that was observed for that variable from the day before the missing observation.

We created a dummy variable for 'start day' to enable the user to test the sensitivity of our conclusions to varying our within-year start of accumulation of environmental conditions. We empirically determined (by the Akaike information criterion (AIC)) that 1 March (start = 60) provided the best compromise between capturing early growing season weather variation and negating brief variation in winter conditions; however, the selection of the precise day did not dramatically influence the overall trends in the results unless changed by more than 15 days.

We then created a function to calculate daily degree-day accumulation and season-long degree-day accumulation based on Allen's [37] double sine function, using our daily maximum and minimum temperature data. We created a dummy variable for our minimum development threshold to facilitate sensitivity analysis, but set it to a default value of 10°C. We did not use a maximum development threshold in our calculation, assuming that temperatures exceeding its hypothetical value (often more than 30°C for temperate insects) were relatively rare. Accumulations were calculated from the start day variable, as described above. We also created functions that calculated the accumulation of precipitation over the sampling week, the accumulation of precipitation over the growing season, from the start date and the number of rainy days in a sampling period. Weather data were merged with firefly data to facilitate subsequent analyses.

2.3. Data analysis

We used ggplot2 [38] to visualize trends in captures of fireflies by plant community treatment over years. We then conducted a multivariate analysis to determine whether firefly plant community use patterns changed within or between years, and what environmental factors were associated with plant community use patterns. To accomplish this, data were cast as a date-by-treatment matrix at two resolutions (weekly observations and yearly observations), transformed using the Wisconsin standardization, and Bray–Curtis differences were subjected to non-metric multi-dimensional scaling (NMDS) in vegan [39]. Environmental parameters were fit to the NMDS plots using envfit to determine whether patterns were influenced by weather.

To examine patterns in firefly captures over time, and the interactions of these captures with environmental variables, we visualized trends in capture data by sampling week and degree-day accumulation. Noting that degree-day accumulation was associated with the clearest patterns in firefly captures (see Results), with some variation owing to plant community, we built a generalized linear model (GLM) with a negative binomial structure to explain these patterns. The model included the degree-day accumulation in linear and quadratic forms as continuous variables, year and plant community treatment as factors, and trapping effort as an offset variable (to account for lost or compromised traps). Model structure was determined empirically by AIC. After fitting the model, we used the resultant regression parameters to generate predicted values, so we could visually compare the performance of the model to the raw data.

Because the model found year-to-year variation in the activity peak that was not explained by degree-day accumulation, we extracted the activity peaks from each year as predicted by the model to a new data frame, and matched these data to other relevant environmental variables in the weather matrix (week the peak occurred in, precipitation variables corresponding to that week). We visualized the relationship between activity peak and other variables, and then constructed a generalized linear model for a quadratic relationship between the activity peak, by degree days, and the precipitation accumulation at the activity peak.

For all frequentist analyses, a significance level of $\alpha = 0.05$ was used.

3. Results

Over the 12-year study, 17 084 fireflies were captured in the trapping network. Visualizations of firefly capture data by treatment and time period revealed several patterns. Numbers of fireflies captured in each trap varied by plant community type and across samples (figure 2), but in general, more fireflies

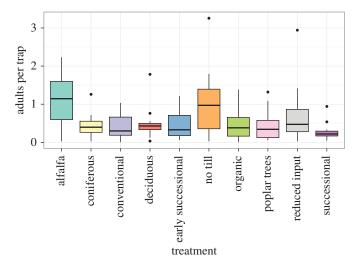


Figure 2. Box plot of average firefly captures, 2014–2015, by plant community treatment. Yearly average number of adult fireflies captured on weekly sampled yellow sticky cards across 10 plant community treatments at Kellogg Biological Station. Median firefly density in each treatment is represented by the bold line, and upper and lower margins of each box represent the upper and lower quartiles in that treatment, respectively.

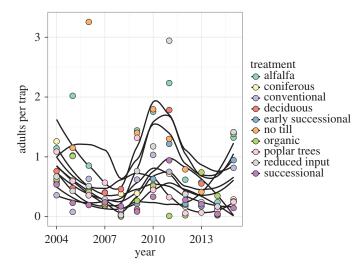


Figure 3. Average firefly captures, 2004–2015, by plant community treatment, by year. Yearly average number of adult fireflies captured on weekly sampled yellow sticky cards across 10 plant community treatments at Kellogg Biological Station. Loess smoother lines represent smoothed captures within a given treatment and are used to illustrate general trends in the population across treatments.

were captured in alfalfa and no-till row crop treatments. Average numbers of fireflies captured per trap also demonstrated variation by year independent of plant community treatment. Overlaid plots of average captures for all treatments against year (figure 3) suggest a 6–7-year firefly population cycle that appears uncorrelated with environmental variables.

NMDS revealed only weak trends in patterns of capture between plant community treatments at both the yearly and weekly resolutions. At the yearly resolution (figure 4a), plant community treatment use varied slightly with the number of rainy days in the growing season ($R^2 = 0.16$, p = 0.006, 2D NMDS stress = 0.14) with herbaceous habitat use generally associated with greater amounts of rainfall. At the weekly resolution (figure 4b), 2D NMDS stress was higher (0.19), but a general trend away from forest plots was observed with increasing degree-day accumulation ($R^2 = 0.15$, p = 0.001) and week ($R^2 = 0.15$, p = 0.001).

When plotting firefly abundance by week of capture, the timings of the peaks in firefly emergence show asynchrony among years (figure 5a), indicating that week of year (and, by proxy, day length)

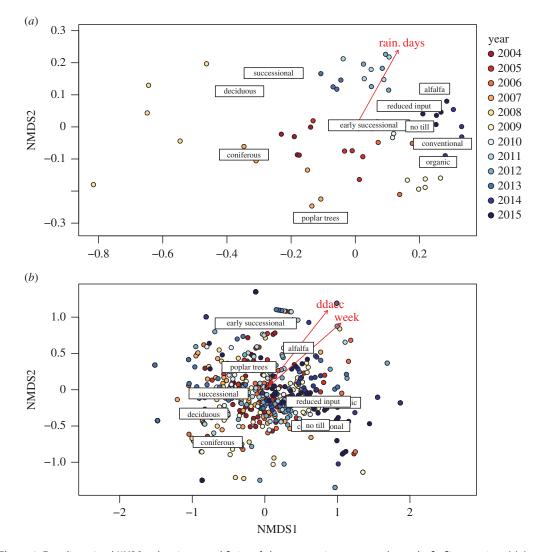


Figure 4. Two-dimensional NMDS and environmental fitting of plant community treatment plot use by fireflies over time. (*a*) At the yearly resolution, a 2D NMDS stress of 0.14 was observed. (*b*) At the weekly resolution, a 2D NMDS stress of 0.19 was observed.

is not a strong driver of firefly emergence. However, plotting firefly numbers instead against degreeday accumulation dramatically reduced the asynchrony of emergence peaks and indicated that a single activity peak occurred in each year (figure 5b). Thus, degree-day accumulation appears to be a better predictor of firefly populations than week of year or associated variables.

Our model for firefly activity incorporating degree-day accumulation, plant community treatment and year performed well at predicting the timing of the activity peaks (figure 6), accounting for more than 40% of the variation in the raw data. However, model selection favoured the inclusion of a year term as a factor, suggesting that another factor in addition to degree-day accumulation was varying from year to year and impacting firefly activity. Activity peaks varied from year to year by nearly 180 degree-day units, varying from 720 ± 38 DD in 2004 to 898 ± 55 DD in 2012 (figure 7). However, we found the year-to-year variation was well explained by precipitation accumulation: a quadratic relationship occurs between degree days at peak emergence and precipitation accumulation (pseudo- $R^2 = 0.456$, p = 0.026; figure 8).

4. Discussion

The greatest proportion of fireflies was captured in alfalfa and no-till plant communities (figure 2), indicating that areas with moderate soil disturbance and primarily herbaceous plant communities favoured firefly emergence. This result was unexpected; because fireflies spend much of their life cycle in the soil, it might be expected that plots with little soil disturbance (coniferous, deciduous and

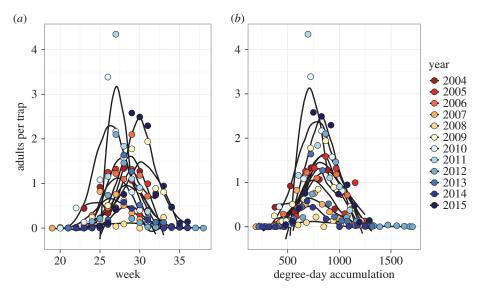


Figure 5. Average number of adult fireflies per trap across all sampled treatments at Kellogg Biological Station plotted by year. Samples were taken weekly over the growing season from 2004 to 2015, and plotted by (a) week of capture and (b) degree-day accumulation at capture. Loess lines represent smoothed capture trends for a given year and were used to assess consistency of response to a given variable between years.

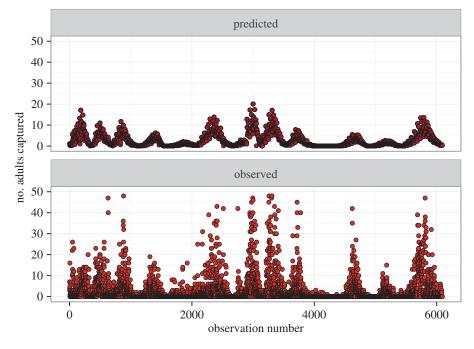


Figure 6. Number of firefly adults captured, as predicted by GLM and, as observed, by observation number. Predicted values were generated using GLM accounting for variability owing to plant community treatment degree-day accumulation, and year as a factor variable. Details of GLM can be found in the data analysis section in Material and methods.

successional forests) would foster the greatest populations of fireflies. However, these plots produced capture rates similar to those observed in the intensively managed and tilled conventional row crop plots. Our result contrasted with observations of another genus of fireflies in Malaysia (*Pteroptyx*), where researchers found that plant canopy structure was the most important determinant of abundance [40]. Also surprising was the relatively low capture rate in early successional plots, which are primarily herbaceous, with a no-till management regime. Thus, the yearly burnings may play a role in suppressing firefly populations in these plots. An alternative explanation for these variations in captures could be

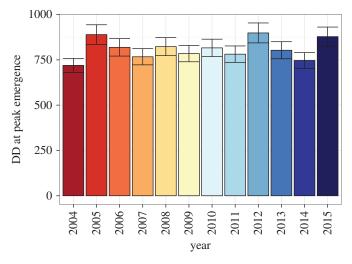


Figure 7. Degree-day accumulation at peak firefly activity by year. Degree-day accumulation (\pm s.e.m.) at peak emergence of firefly adults varied by sample year. Activity peaks were extracted from regression coefficients from GLM.

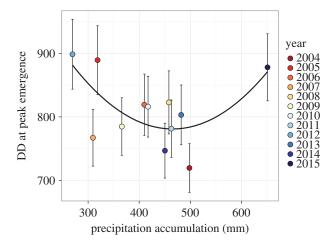


Figure 8. Firefly activity peaks by precipitation accumulation. Firefly activity per degree-day accumulation had a quadratic relationship with precipitation accumulation (pseudo- $R^2 = 0.456$, p = 0.026).

differences in trapping efficiencies between plant communities. However, if this were the case, we would expect trapping efficiencies in the three other row crop treatments (conventional, organic and reduced input management) not to differ appreciably from that of the no-till row crop plant community.

When plotted over sample years (figure 3), captures of fireflies by treatment seem to suggest an intriguing cyclical dynamic, with alternating peaks and troughs in captures on an approximately 6-year cycle. Our time series only spans 12 years, meaning more data will be required to elucidate this pattern and its drivers. Similarly, analysis of plant community use patterns was inconclusive (figure 4). At the weekly resolution there was a trend away from woody treatments over the growing season (figure 4b; i.e. with both increasing week and increasing degree-day accumulation). Although this pattern was not strong, it could result from fireflies overwintering in forest habitats and then moving to lower-canopy herbaceous habitats for mating displays. We observed very similar performance of both degree day and week, probably owing to autocorrelation between the two variables that cannot be resolved at the sampling resolution used over the course of the study.

The degree-day model GLM suggested that activity peaks occurred at a degree-day accumulation of approximately 800 DD, accumulated from March 1. A model for *Photinus carolinus* in the Great Smokey mountains found peak display occurred at approximately 1100 degree days (using a base of 10°C and the same start date as our model) [25]. The difference between heat units required for peak activity observed between this and our study may be a result of species or locality differences (i.e. more southern firefly

populations are probably adapted to warmer spring conditions). Similarly, differences in methodology for calculating degree days may account for some of these differences. However, both studies support the observation that degree-day accumulation is the dominant cue governing the activity patterns of temperate fireflies.

Although both photoperiod and degree-day accumulation can both play a role in the phenology of insects, our results suggest that degree-day accumulation is the dominant driver of firefly flight activity. The model was unable to account for between-trap variation within a single sampling day (figure 6), though it was able to capture the overall trends in activity quite well, using only degree-day accumulation, plant community treatment and year as predictors. Nevertheless, degreeday accumulation was not the sole driver in within-season variability. Our model found year-to-year variability in activity peaks that could not be explained by degree-day accumulation alone. We found that this variation in activity peak by degree-day accumulation had a quadratic relationship with precipitation, indicating that both drought and heavy rainfall in the time period leading up to their activity peak can delay the peak (figure 8). Assuming an approximate 20°C daily average temperature at this site in late June and early July, this could translate to a 10-or-more day change in activity peak owing to precipitation extremes in any given year. Yet, there are several alternate explanations for the pattern we observed, and some patterns detected may have been driven by statistical outliers. For example, very high rainfall in 2015 at the site strongly influences our conclusion that a quadratic relationship exists between degree-day accumulation and precipitation accumulation in explaining firefly activity peaks. Indeed, if observations from 2015 had not been included in our analysis, we would probably have concluded that degree-day accumulation had a negative, linear relationship with precipitation accumulation; that is, increasing rainfall would cause fireflies to emerge earlier, given constant temperatures. This result would align with previous work showing that firefly abundance in Japan is generally negatively correlated with rainfall [35]. However, considered within the context of firefly biology, it seems unlikely that the relationship between these parameters would be linear throughout the range of possible precipitation values, as soil-dwelling larvae of non-aquatic firefly species and/or their prey are probably adversely affected by abnormally waterlogged soils.

As the sampling at our study site continues, we will watch rainy years with particular interest to determine whether population data collected in these years support or refute this pattern, or if an alternate driver can explain more of the variation. Indeed, firefly activity may have been driven by factors not considered in this study. Although using a start date of 1 March was favoured in our analysis (i.e. the AIC of the model using this start date was minimized), when the start day was changed in a sensitivity analysis, the relationship between degree-day accumulation and precipitation in firefly activity changed or disappeared. This result could suggest that alternate drivers not accounted for in this study may be driving aspects of firefly activity. Factors such as winter snow cover and variations in winter temperature are known to affect the phenology of temperate insects [41], and thus these factors should be considered in subsequent work.

In this study, we have clearly demonstrated a taxon whose phenology varies in response to multiple drivers. Species with phenological responses to multiple drivers are not rare [42]. Yet ecological interactions among species with multiple drivers of phenology may be complex and unpredictable [43,44], potentially leading to dire consequences in a changing environment [45]. Our study examined the phenological responses to environmental conditions of adult fireflies; however, data on larvae or sex of the adults were unavailable. Adult *Photinus* fireflies are non-feeding [14], so shifts in their activity are unlikely to have direct consequences through phenological asynchronies. Shifts in adult activity probably correspond to shifts in development or activity among larvae, potentially leading to asynchronies between larvae-prey populations at this critical development time period. Resources acquired during the predaceous larval stage are important in determining mating success among adult fireflies: males provide an energetically costly nuptial gift to the female in the form of a spermatophore [46]. If sex differences in phenological responses to environmental conditions exist, asynchronies between males and females may additionally reduce mating success and fecundity [47]. Male fireflies were always observed earlier than females in Elkmont, TN, USA. In fact, in that system, females were often found during or after peak emergence of males and thus this should be an area of emphasis in future study [25]. Additionally, phenological shifts in fireflies may lead to consequences at other trophic levels. For example, generalist ground-dwelling predators like firefly larvae and other predaceous beetles are known to have dramatic effects on the establishment of agricultural pests early in the growing season [48]. Similarly, although distasteful and avoided by many predators, some birds, lizards and frogs are known to feed on adult fireflies [49], thus shifts in firefly activity may have dietary consequences for animals at higher trophic levels.

5. Conclusion

Fireflies are a charismatic and important taxon with ties to trophic function, economic importance and culture. Although empirical evidence of specific declines of *Photinus* fireflies has not been clearly demonstrated in longitudinal studies, naturalists and citizen scientists perceive a decline in their number [21], leading to interest in their conservation. Our study has offered new insight to support conservation efforts and to direct future research. *Photinus pyralis* appears to thrive in habitats with moderate soil disturbance. Thus, efforts to foster no-till and perennial agricultural systems [50,51] will probably benefit the species. Climate warming may advance the activity of fireflies to progressively earlier in the growing season, but other extremes of climate in the form of precipitation may introduce unpredictable elements to this, and add the possibility of inducing asynchrony with other systems.

The availability of long-term observational data, made freely accessible to the public, was an essential factor in the discoveries made in this study. Although the study that provided these data was not initiated with this purpose in mind, we were able to empirically demonstrate and disentangle the effect of multiple drivers on firefly phenology simply because we had the statistical power to do so. Although species that respond to multiple, interacting environmental drivers are relatively common, data supporting investigations of this kind are rare [52]. We therefore encourage all practising ecologists to curate their species observation data and make them publicly available, to foster long-term, broad-scale investigations in the future [53–55].

Data accessibility. All data and analysis code produced by this study are publicly available. Lampyrid abundance data are available at https://ndownloader.figshare.com/files/3686040. Weather station data are available at https://lter. kbs.msu.edu/datatables/7.csv. These data are automatically downloaded when the R script file at https://github.com/cbahlai/lampyrid/ is run.

Authors' contributions. S.L.H. contributed to conception, led study design and wrote major portions of the manuscript. S.X. conceived of analyses and wrote portions of the manuscript. L.R. led exploratory analysis and wrote portions of the manuscript. E.D.L., A.M. and B.E. provided critical commentary on the design and analysis of the study and wrote portions of the manuscript. C.A.B. conceived of the study, led the analysis, supervised the drafting of the manuscript and critically revised its content. All authors approved the final content of the manuscript. Competing interests. We declare no competing interests.

Funding. Data used in this study were produced with funding from the National Science Foundation Long Term Ecological Research program grant no. 1027253. C.A.B. was funded by a fellowship from the Mozilla Foundation and the Leona M. and Harry B. Helmsley Charitable Trust.

Acknowledgements. This paper was written as part of a course in Open Science and Reproducible Research offered through the Department of Entomology at Michigan State University. The authors would like to thank the Mozilla Science Laboratory and the rest of the open science community for their support in the drafting of this manuscript, and Stuart Gage, Manuel Colunga-Garcia and Douglas Landis for design and maintenance of the study that produced the data used herein. Monica Granados helpfully provided review of our analysis R code. Lynn Faust provided keen insight and invaluable commentary on an earlier version of this manuscript. We would like to thank Tyson Wepprich and one additional anonymous reviewer for helpful comments that helped us question our assumptions and improve the clarity of this manuscript.

References

- Rosenheim JA et al. 2011 Ecoinformatics for integrated pest management: expanding the applied insect ecologist's tool-kit. J. Econ. Entomol. 104, 331–342. (doi:10.1603/EC10380)
- Bahlai CA, Landis DA. 2016 Predicting plant attractiveness to pollinators with passive crowdsourcing. R. Soc. open sci. 3, 150677. (doi:10.1098/rsos.150677)
- Wang Y, Fu X, Lei C, Jeng M-L, Nobuyoshi O. 2007 Biological characteristics of the terrestrial firefly Pyrocoelia pectoralis (Coleoptera: Lampyridae). Coleopterists Bull. 61, 85–93. (doi:10.1649/907.1)
- Wing SR. 1989 Energetic costs of mating in a flightless female firefly, *Photinus collustrans* (Coleoptera: Lampyridae). *J. Insect Behav.* 2, 841–847. (doi:10.1007/BF01049405)
- Hess WN. 1920 Notes on the biology of some common Lampyridæ. *Biol. Bull.* 38, 39–76. (doi:10.2307/1536232)

- Williams FX. 1917 Notes on the life-history of some North American Lampyridæ. J. NY Entomol. Soc. 25, 11–33
- Eisner T, Goetz MA, Hill DE, Smedley SR, Meinwald J. 1997 Firefly 'femmes fatales' acquire defensive steroids (lucibufagins) from their firefly prey. Proc. Natl Acad. Sci. USA 94, 9723–9728. (doi:10.1073/ pnas.94.18.9723)
- Keiper RR, Solomon LM. 1972 Ecology and yearly cycle of the firefly *Photuris pennsylvanica* (Coleoptera: Lampyridae). J. NY Entomol. Soc. 80, 43–47.
- Branchini BR, Murtiashaw MH, Magyar RA, Portier NC, Ruggiero MC, Stroh JG. 2002 Yellow-green and red firefly bioluminescence from 5,5-dimethyloxyluciferin. J. Am. Chem. Soc. 124, 2112–2113. (doi:10.1021/ja017400m)
- Schaap AP, Gagnon SD. 1982 Chemiluminescence from a phenoxide-substituted 1,2-dioxetane: a

- model for firefly bioluminescence. *J. Am. Chem. Soc.* **104**, 3504–3506. (doi:10.1021/ja00376a044)
- Seliger HH, Buck JB, Fastie WG, McElroy WD. 1964
 The spectral distribution of firefly light. J. Gen. Physiol. 48, 95–104. (doi:10.1085/jgp.48.1.95)
- Branham MA, Wenzel JW. 2003 The origin of photic behavior and the evolution of sexual communication in fireflies (Coleoptera: Lampyridae). Cladistics 19, 1–22. (doi:10.1111/j.1096-0031.2003.tb00404.x)
- Lall AB, Seliger HH, Biggley WH, Lloyd JE. 1980 Ecology of colors of firefly bioluminescence. Science 210, 560–562. (doi:10.1126/science.210. 4469.560)
- Lloyd J. 1997 Firefly mating ecology, selection and evolution. In *The evolution of mating systems in insects and Arachnids* (eds JC Choe, BJ Crespi), pp. 184–192. Cambridge, UK: Cambridge University Press.

- National Museum of Natural History 2013 State Bugs. Encyclopedia Smithsonian. See https://www. si.edu/Encyclopedia_Sl/nmnh/buginfo/statebugs. htm (accessed 2 June 2016).
- Kobori H, Primack RB. 2003 Participatory conservation approaches for Satoyama, the traditional forest and agricultural landscape of Japan. Ambio 32, 307–311. (doi:10.1579/0044-7447-32.4.307)
- Takeda M, Amano T, Katoh K, Higuchi H. 2006 The habitat requirement of the Genji-firefly Luciola cruciata (Coleoptera: Lampyridae), a representative endemic species of Japanese rural landscapes. Biodivers. Conserv. 15, 191–203. (doi:10.1007/s10531-004-6903-y)
- Takada K. 2011 Popularity of different lampyrid species in Japanese culture as measured by Google search volume. *Insects* 2, 336–342. (doi:10.3390/ insects2030336)
- Faust LF. 2004 Fireflies as a catalyst for science education. *Integr. Comp. Biol.* 44, 264–265. (doi:10.1093/icb/44.3.264)
- Gardiner T. 2009 Glowing, glowing, gone? A history of glow-worm recording in the UK. Corby, UK: British Nationalists' Association. (http://www.bnanaturalists.org/gworm.pdf) (accessed 2 June 2016).
- Chow AT, Chong J-H, Cook M, White D. 2014
 Vanishing fireflies: a citizen-science project promoting scientific inquiry and environmental stewardship. Sci. Educ. Civic Engag. 6, 23–31.
- Firefly Watch. 2016 Museum of science. See https:// legacy.mos.org/fireflywatch/ (accessed 2 June 2016)
- Scagell, R. 2016 The UK glow worm survey. See http://www.glowworms.org.uk/ (accessed 2 June 2016)
- Picchi MS, Avolio L, Azzani L, Brombin O, Camerini G. 2013 Fireflies and land use in an urban landscape: the case of *Luciola italica* L. (Coleoptera: Lampyridae) in the city of Turin. *J. Insect Conserv.* 17, 797–805. (doi:10.1007/s10841-013-9562-z)
- Faust LF, Weston PA. 2009 Degree-day prediction of adult emergence of *Photinus carolinus* (Coleoptera: Lampyridae). *Environ. Entomol.* 38, 1505–1512. (doi:10.1603/022.038.0519)
- Guiney MS, Oberhauser KS. 2009 Insects as flagship conservation species. *Terr. Arthropod Rev.* 1, 111–123. (doi:10.1163/187498308X414733)
- Diez JM, Ibáñez I, Miller-Rushing AJ, Mazer SJ, Crimmins TM, Crimmins MA, Bertelsen CD, Inouye DW. 2012 Forecasting phenology: from species variability to community patterns. *Ecol. Lett.* 15, 545–553. (doi:10.1111/j.1461-0248.2012. 01765 x)
- 28. Miller-Rushing AJ, Høye TT, Inouye DW, Post E. 2010 The effects of phenological mismatches on

- demography. *Phil. Trans. R. Soc. B* **365**, 3177–3186. (doi:10.1098/rstb.2010.0148)
- Root TL, MacMynowski DP, Mastrandrea MD, Schneider SH. 2005 Human-modified temperatures induce species changes: joint attribution. *Proc. Natl Acad. Sci. USA* 102, 7465–7469. (doi:10.1073/pnas. 0502286102)
- Parmesan C. 2006 Ecological and evolutionary responses to recent climate change. *Annu. Rev. Ecol. Evol. Syst.* 37, 637–669. (doi:10.1146/annurev. ecolsvs.37.091305.110100)
- Rosenzweig C et al. 2008 Attributing physical and biological impacts to anthropogenic climate change. Nature 453, 353–357. (doi:10.1038/ nature06937)
- Crimmins TM, Crimmins MA, Bertelsen D, Balmat J. 2008 Relationships between alpha diversity of plant species in bloom and climatic variables across an elevation gradient. *Int. J. Biometeorol.* 52, 353–366. (doi:10.1007/s00484-007-0130-7)
- Crimmins TM, Crimmins MA, David Bertelsen C. 2010 Complex responses to climate drivers in onset of spring flowering across a semi-arid elevation gradient. J. Ecol. 98, 1042–1051. (doi:10.1111/j.1365-2745.2010.01696.x)
- Visser ME, Holleman LJM. 2001 Warmer springs disrupt the synchrony of oak and winter moth phenology. Proc. R. Soc. Lond. B 268, 289–294. (doi:10.1098/rspb.2000.1363)
- Yuma M. 2007 Effect of rainfall on the long-term population dynamics of the aquatic firefly Luciola cruciata. Entomol. Sci. 10, 237–244. (doi:10.1111/j.1479-8298.2007.00219.x)
- Wickham H. 2016 reshape2: Flexibly reshape data: a reboot of the reshape package, version 1.4.2. See https://cran.r-project.org/web/packages/ reshape2/index.html.
- Allen JC. 1976 A modified sine wave method for calculating degree days. *Environ. Entomol.* 5, 388–396. (doi:10.1093/ee/5.3.388)
- Wickham H, Chang W. 2016 ggplot2: an implementation of the grammar of graphics, version 2.1.0. See https://cran.r-project.org/web/ packages/ggplot2/index.html.
- Oksanen J et al. 2016 vegan: community ecology package, version 2.4-1. See https://cran.r-project. org/web/packages/vegan/index.html.
- Jusoh WFAW, Hashim NR, Ibrahim ZZ. 2010 Firefly distribution and abundance on mangrove vegetation assemblages in Sepetang Estuary, Peninsular Malaysia. Wetlands Ecol. Manag. 18, 367–373. (doi:10.1007/s11273-009-9172-4)
- Forrest JRK, Thomson JD. 2011 An examination of synchrony between insect emergence and flowering in Rocky Mountain meadows. *Ecol. Monogr.* 81, 469–491. (doi:10.1890/10-1885.1)

- Wolkovich EM et al. 2012 Warming experiments underpredict plant phenological responses to climate change. Nature 485, 494–497. (doi:10.1038/nature11014)
- Hua F et al. 2016 Community-wide changes in intertaxonomic temporal co-occurrence resulting from phenological shifts. Glob. Change Biol. 22, 1746–1754. (doi:10.1111/qcb.13199)
- Robinet C, Roques A. 2010 Direct impacts of recent climate warming on insect populations. *Integr. Zool.* 5, 132–142. (doi:10.1111/j.1749-4877.2010. 00196.x)
- McLaughlin JF, Hellmann JJ, Boggs CL, Ehrlich PR. 2002 Climate change hastens population extinctions. Proc. Natl Acad. Sci. USA 99, 6070–6074. (doi:10.1073/pnas.052131199)
- Lewis SM, Cratsley CK, Rooney JA. 2004 Nuptial gifts and sexual selection in *Photinus* fireflies. *Integr. Comp. Biol.* 44, 234–237. (doi:10.1093/icb/44.
- Calabrese JM, Ries L, Matter SF, Debinski DM, Auckland JN, Roland J, Fagan WF. 2008 Reproductive asynchrony in natural butterfly populations and its consequences for female matelessness. J. Anim. Ecol. 77, 746–756. (doi:10.1111/j.1365-2656.2008.01385.x)
- Safarzoda S, Bahlai CA, Fox AF, Landis DA. 2014 The role of natural enemy foraging guilds in controlling cereal aphids in Michigan wheat. *PLoS ONE* 9, e114230. (10.1371/journal.pone.0114230)
- 49. Lloyd JE. 1973 Firefly parasites and predators. *Coleopterists Bull.* **27**, 91–106.
- Werling BP, Meehan TD, Gratton C, Landis DA. 2011
 Influence of habitat and landscape perenniality on
 insect natural enemies in three candidate biofuel
 crops. Biol. Control 59, 304—312. (doi:10.1016/
 j.biocontrol.2011.06.014)
- Werling BP et al. 2014 Perennial grasslands enhance biodiversity and multiple ecosystem services in bioenergy landscapes. Proc. Natl Acad. Sci. USA 111, 1652–1657. (doi:10.1073/pnas.1309492111)
- Strayer DL, Eviner VT, Jeschke JM, Pace ML. 2006 Understanding the long-term effects of species invasions. *Trends Ecol. Evol.* 21, 645–651. (doi:10.1016/j.tree.2006.07.007)
- Michener WK, Jones MB. 2012 Ecoinformatics: supporting ecology as a data-intensive science. *Trends Ecol. Evol.* 27, 85–93. (doi:10.1016/j.tree. 2011.11.016)
- Hampton SE et al. 2015 The Tao of open science for ecology. Ecosphere 6, art120. (doi:10.1890/ES14-00402 1)
- Hampton SE, Strasser CA, Tewksbury JJ, Gram WK, Budden AE, Batcheller AL, Duke CS, Porter JH. 2013 Big data and the future of ecology. Front. Ecol. Environ. 11, 156–162. (doi:10.1890/120103)