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#### **ORIGINAL PAPER**



# Recruitment, survival, and parasitism of monarch butterflies (*Danaus plexippus*) in milkweed gardens and conservation areas

Emily A. Geest<sup>1</sup> · L. LaReesa Wolfenbarger<sup>1</sup> · John P. McCarty<sup>1</sup>

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#### Abstract

Monarch butterflies (Danaus plexippus) are suffering from declining populations and conservationists have encouraged planting milkweed gardens in urban and suburban landscapes to help offset habitat loss across the breeding range. The effectiveness of gardens as a conservation strategy depends on their ability to attract ovipositing adults and the survival of monarch larvae in these gardens. Larvae are susceptible to a variety of predators as well as to parasitism by a tachinid fly (Lespesia archippivora) and a protozoan parasite (Ophryocystis elektroscirrha) which cause lethal or sublethal effects, yet the severity of these risks in gardens is not well understood. We compared egg abundance and larval survival in traditional conservation areas to gardens that incorporated milkweed to attract monarchs. Additionally, we collected late instar larvae and reared them in the lab to compare parasitism rates between monarch gardens and conservation areas. Both gardens and conservations sites varied widely in recruitment and survival of monarchs and there were no significant differences between the garden and conservation sites. Tachinid fly parasitism ranged from 30% of larvae from conservation sites in 2016 to 55% of larvae from gardens in 2017, but did not differ between the two categories of sites. Parasitism by O. elektroscirrha was detected in fewer than 2% of larvae. The density of milkweed had no effect on the number of monarch eggs in conservation areas or gardens in either year. Milkweed density had no effect on tachinid parasitism in conservation areas but had a significant effect in gardens with lower numbers of milkweed stems increasing tachinid parasitism in 2016. Gardeners planted a variety of species of milkweed and Asclepias syriaca was the most commonly used host plant for monarch larvae (85%). Overall, our results suggest that milkweed gardens have the potential to contribute to successful monarch reproduction. However, the variation among sites and the lack of recruitment from some gardens emphasizes that the realization of this potential contribution will depend on the quality of gardens.

**Keywords** Monarch butterfly  $\cdot$  *Danaus plexippus*  $\cdot$  Survival  $\cdot$  Parasitoids  $\cdot$  *Lespesia archippivora*  $\cdot$  Conservation  $\cdot$  Milkweed gardens

This work is derived from the master's thesis completed by EAGeest but has not otherwise been published

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# Introduction

Monarch butterflies (*Danaus plexippus*) have suffered severe population declines and are of increasing conservation concern (Marini and Zalucki 2017; Pleasants et al. 2017). Considerable attention has been focused on the largest and most well-known eastern population of monarchs that is known for migrating annually between Mexico and the Midwestern United States and southern Canada (Altizer et al. 2000; Shahani et al. 2015). During this migration, up to four generations of monarchs can occur. The first generation has overwintered in Mexico, breeds and moves north into Texas to oviposit. Each subsequent generation moves further north until the final generation emerges in late summer and early fall. This final generation is the migratory generation



that moves from southern Canada and the northern United States back to Mexico where they overwinter (Oberhauser et al. 2017b).

During their annual cycle monarchs encounter multiple threats contributing to their decline, including habitat loss and fragmentation in their breeding range (Gustafsson et al. 2015). For reproduction monarchs depend on the availability of milkweed (Asclepias spp.) as their only larval host plant, and access to nectar producing plants for adults (Zalucki and Suzuki 1987; Oberhauser et al. 2001). Historically, milkweed and nectar producing flowers existed in continuous grasslands throughout the Great Plains but now exist in a mosaic of patches in agricultural fields, roadsides, and remaining prairies fragments (Brower and Pyle 2004; Oberhauser et al. 2008). Recently, milkweed growing in agricultural fields has been reduced through the increased use of herbicide tolerant corn and soybeans, which has decreased the milkweed naturally occurring within these fields (Oberhauser et al. 2001, 2008; Pleasants and Oberhauser 2013; Jepsen et al. 2015; Stenoien et al. 2018). Roadside milkweed has also been reduced through frequent mowing and herbicide use.

The monarch population decline and habitat loss have spurred efforts to conserve monarchs at the local, state, national and international levels (Ward 2014; Shahani et al. 2015; Nguyen 2017; Panella 2017). As part of these initiatives numerous citizen scientist based programs have emerged, targeting topics varying from surveying and tracking migration (e.g. Journey North, the Western Monarch Thanksgiving Count), larval health (e.g. Monarch Larval Monitoring Project, Monarch Health), to increasing monarch habitat (e.g. MonarchWatch; Jepsen et al. 2015).

A number of programs encourage landowners residing in the migratory pathway to plant milkweed gardens to support conservation (i.e. MonarchWatch's "Monarch Waystations" Taylor 2017). The purpose of these gardens is to increase monarch habitat (Oberhauser et al. 2008) and to increase connectivity among milkweed patches (Zalucki and Lammers 2010; Zalucki et al. 2016). These citizen scientist programs have attracted thousands of participants (Nail et al. 2015; Taylor 2017) and influential groups such as the US Department of Agriculture list gardens as a way to help monarchs locally (USDA 2017).

While citizen scientist programs are widely advertised, promoted, and funded, little is known about what contribution to conservation these efforts actually make. Though they are intended to provide milkweed for larvae and, sometimes, nectar plants for adults, gardens differ from native habitats in key ways. Gardens are typically smaller, fragmented, and often planted with commercially sourced milkweed, compared to conservation areas that are larger, continuous, and have naturally occurring milkweed species present. The effects of such differences on monarch recruitment and

larval survival is not well understood but key to determining the value of milkweed gardens. Results of prior studies using experimental milkweed plantings intended to mimic conditions found in gardens and natural sites suggest that while gardens are attractive to ovipositing adults (Cutting and Tallamy 2015; Stenoien et al. 2015), a better understanding of the survival of eggs and larvae in gardens relative to natural areas is needed (Majewska et al. 2018). The efficacy of milkweed gardens as a conservation tactic is important both because of the investment being committed to promoting gardens, and the potential risk of attracting monarchs to oviposit in habitats where larval survival is unlikely.

We estimated survival of monarch caterpillars in milkweed gardens, with a special emphasis on evaluating the risk of parasitism by the tachinid fly Lespesia archippivora. Adult females lay eggs on monarch larvae. Upon hatching the maggots bore into the larva, ultimately causing mortality (Etchegaray and Nishida 1975). We also evaluated the potential for parasitism by the obligate protozoan (Ophryocystis elektroscirrha) that can leave monarchs with deformed wings and increased mortality (McLaughlin and Myers 1970; Altizer et al. 2000). Parasitism rates by L. archippivora and O. elektroscirrha can be influenced by habitat (Zalucki 1981; Prysby 2004; Tooker et al. 2006; Oberhauser et al. 2007). The more parasitized adults that visit the same milkweed the higher the spore load becomes (Bartel et al. 2011). Thus, the potential exists for parasitism rates in small gardens to become elevated.

Our approach was to compare monarch demography and parasitism rates in existing gardens to native prairie fragments and prairie restorations. For most species of conservation concern in the Great Plains, conservation efforts have focused on protecting, managing, and restoring grasslands—a strategy we will hereafter refer to as traditional conservation. Understanding the effect these gardens have on monarch recruitment and demography is vital to maximizing the conservation benefits of milkweed gardens. We hypothesized that: (1) egg numbers, (2) larval survival, and (3) parasitism rates differ between garden plantings and conservation areas. To evaluate these hypotheses, we quantified egg and larval abundance and demography at conservation areas and gardens in 2016 and 2017. We then collected and reared late instar monarch larvae to compare parasitism rates between gardens and conservation areas.

#### **Methods**

We compared monarch egg and larval density, larval survival, and larval parasitism on milkweed naturally occurring in conservation areas to milkweed planted in gardens. As the basis for comparison to gardens, we used five conservation areas within the Omaha, Nebraska metropolitan area



(Supplemental Material Fig. 4, Table A.1). The conservation sites were all tallgrass prairies set in urban or suburban landscapes. These sites varied from 5 to 178 ha and were actively managed through prescribed burning and/or mowing (Supplemental Material Table A.1).

We found gardens by describing the goals of the project to garden clubs and horticulture societies and requesting help finding gardens that could qualify as "Monarch Waystations" (Taylor 2017). The largest citizen science initiative focused on monarchs, Monarch Watch designates gardens of any size that contain at least two species of milkweed with a total of at least ten stems of milkweed as a "Monarch Waystation" (Taylor 2017). From gardeners who responded and who were willing to accept regular visits to their property, we chose fifteen sites based on accessibility of the gardens and proximity to our lab facilities to maximize the number of sites we could monitor. We eliminated two of the fifteen gardens a month after the start of the 2016 field season because the gardeners had removed all milkweed. The remaining 13 gardens within residential areas of the Omaha metropolitan area consisted of five private gardens and eight public gardens (Supplemental Material, Table A.2).

#### **Field observations**

Monarch larvae hatch from eggs approximately four days of being laid and pass through five developmental instars over a period of 9–14 days (Oberhauser 2004) though these rates vary depending on temperature and other environmental factors (Rawlins and Lederhouse 1981; Zalucki 1982). They then pupate and emerge from chrysalises as butterflies within 14 days. Instars are distinguishable from one another based on easily visible characteristics (Oberhauser and Kuda 1997; Urquhart 1998; Geest 2017).

We visited each field site every 4 days visually checking each milkweed stem for larvae and eggs and recording counts for each instar (Oberhauser et al. 2009; CEC 2017; Geest 2017). Using these data, we constructed survival curves to quantify differences in population demography between conservation sites and gardens. We collected all 4th and 5th instars and reared them to adulthood in the lab to estimate rates of parasitism (see below). We recorded the presence and instar of any depredated or deceased larvae that we found.

Because conservation areas are large relative to gardens, two patches of milkweed were located within each study site. These patches were chosen based on ease of access and the number of milkweed stems present. By choosing large patches of milkweed, we could efficiently maximize the number of stems checked at every visit. The area and number of milkweed stems varied among sites (Geest 2017) we recorded patch area (m²) and number of milkweed stems to include in analyses (see below). Within each of these patches

within the conservation areas we examined every milkweed stem at every visit. In gardens, due to the smaller number of plants, we examined every stem of milkweed in the entire garden during each visit. Due to the larger number of plants in prairies, we devoted more time and effort searching conservation areas relative to garden sites. We searched garden and prairie field sites during the period when adult monarchs were present in the area: May 21 through September 3 in 2016 and May 8 through August 25 in 2017.

#### Larvae rearing

All 4th and 5th instar larvae found were collected and reared in the lab to measure rates of parasitism. *L. archippivora* is unlikely to successfully parasitize larvae at instar five and collecting at instar four allows the maximum amount of time for the larvae to be parasitized (Etchegaray and Nishida 1975; Borkin 1982). Larvae were collected with the leaf they were found on to reduce overall larva stress and avoid the defensive act of immobilizing, falling into the leaf litter below, and escaping (Grace 1997; Mueller and Baum 2014).

We reared larvae until adulthood following protocols developed by Altizer and Oberhauser (1999). Each larva was housed in a ca 1 L (32 oz.) plastic container with a cheese-cloth lid and a wooden skewer to pupate on. The larvae were kept on a natural light and dark cycle, and checked daily for signs of parasitism. We provided fresh milkweed leaves to larvae daily. Every leaf was surface sanitized by rinsing in 10% bleach solution, rinsed in deionized water, and dried before being given to the larvae to prevent accidental spore and mold introduction (Altizer and Oberhauser 1999; Mueller and Baum 2014).

If parasitoids emerged during rearing, they were collected and identified to species (Wood 1987; O'Hara 2013). If a caterpillar died, we waited 24 h for maggots to emerge: if a larva died and no parasitoids emerged we dissected to determine if parasitoids were present (Oberhauser et al. 2007). If fly larvae were detected upon dissection, the maggots were collected and allowed to pupate normally. If a caterpillar larva stopped feeding and died with no noticeable cause it was classified as "Failure-to-thrive" syndrome (Oberhauser et al. 2007).

Butterflies that successfully eclosed were tested for *O. elektroscirrha* spores once their wings fully dried, following the methods outlined in Altizer et al. (2000). In brief, monarchs were grasped at the base of their wings exposing their abdomens and a clear, adhesive mailing seal (Avery®  $5248^{TM}$ ) was then placed along the abdomen, removed, and placed on a microscope slide. Slides were viewed under a microscope at  $\times 400$  magnification and a parasite load rating was given based on total spores counted. For example, a spore count of 0 = a score of 0, 1 = 1, 2-20 = 2, 21-100 = 3, 101-1000 = 4, 1001 + spores = a score of 5 (Altizer et al.



2000). All adults were subsequently released at the original collection site.

# Measuring milkweed density

We measured two components of milkweed density in conservation areas that occur at two different spatial scales: patch density and site density. First, we quantified milkweed density in the two patches surveyed at each site by measuring the area of each milkweed patch and counting the total stems of milkweed at each visit. Second, we quantified milkweed density across the entire conservation site for each prairie during the peak milkweed blooming period in late June. Site-level measurements were based on counts of milkweed stems in 1 m<sup>2</sup> quadrats placed every 10 m along 100 m transects. Ten transects were randomly placed in each prairie (Geest 2017).

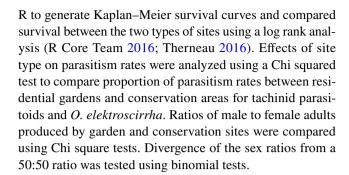
In gardens, we measured the total area of the gardens at the beginning of the study and then counted the stems of milkweed at every visit to quantify the number of stems available to monarchs at each site.

# **Analysis**

We did not detect every egg or larva present within each site at every visit. In order to estimate survival curves, we used a back counting approach to correct for missed observations (De Anda and Oberhauser 2015). The average time it takes for a larva to move from one instar to the next is less than 4 days under typical field conditions. Based on this timing we made the assumption that eggs should be first instars by the next visit, first instars should be second instar, etc. If all eggs and larvae were detected on visit one, then the numbers of the subsequent stages found on visit two should be less than or equal to those on the first visit. If we found the number of larvae in an instar had increased relative to the previous stage and visit, we assumed a missed detection had occurred in the prior visit and corrected for the missed detection by adding an individual to the previous count. With the corrections, the data represent a minimum level of mortality between stages. We tested whether there were differences in the adjusted number of eggs between years and between habitat types (residential gardens versus conservation sites) by using linear mixed models and least squared means comparisons using the lme4, lmtest, lmerTest packages from R (R Core Team 2016; Bates et al. 2015; Kuznetsova et al. 2017; Zeileis and Hothorn 2002).

# Survival and parasitism analysis

We compared survival from egg to adult in conservation areas and in residential gardens using the survival package in



# Effects of habitat variables on monarch demography and parasitism

We evaluated the influence of habitat variables on difference in number of eggs and incidence of parasitism among garden sites and among conservation sites using generalized linear mixed models (GLMM) using the lme4 package in R (Bates et al. 2015; R Core Team 2016). We used a Poisson distribution with site and visit number included as random effects. Prior to using the patch-level density in our models we confirmed that the measurements between patch-level and site level densities were not correlated (R Core Team 2016; Geest 2017). In prairies patch-level density was measured at each visit and the average measurement from the two closest visit dates to the transect survey used for the site-level density was used. For milkweed density in gardens, we averaged the total stems from each visit and used a mean milkweed density. We removed two gardens from the analysis in 2016 and one in 2017 due to low number of eggs recorded and because no larvae were collected from those sites. For each individual site, we also removed all visit data from each site until the first egg was counted, because it was unknown why monarchs were not at the site before eggs were detected. To interpret the direction and significance of relationships we reported the beta estimate of each GLMM.

#### Results

#### **Monarch recruitment**

The number of stems inspected varied among sites and within each site across the season. At conservation areas we surveyed an average of  $117.4 \pm 11.8$  milkweed stems per visit and at gardens we inspected an average of  $100.5 \pm 41.2$  stems per visit (Table 1). The number of monarch eggs varied among sites (Table 1) but did not differ significantly between residential and conservation sites. In 2016 conservation averaged  $14.0 \pm 7.4$  eggs (95% CI -1.8 to 29.8) and garden sites averaged  $26.5 \pm 4.6$  eggs (95% CI 16.8 - 36.3; df = 16, t = 1.43, p = 0.172). In 2017 conservation averaged  $38.8 \pm 30.2$  eggs (95% CI -25.3 to 1002.9) and garden sites



Table 1 Average stems of milkweed (Asclepias spp.) stems present and inspected at each visit across both 2016 and 2017 for all sites

Site name	Average stems ± SE	Sum eggs	Sum larvae collected	Eggs/stem	Larvae collected/ stem
Conservation areas					
Allwine Prairie	$154.9 \pm 14.6$	50	54	0.3	0.3
Audubon Society of Omaha Prairie Preserve	$87.1 \pm 6.3$	49	3	0.6	< 0.1
Bauermeister Prairie	$118.1 \pm 6.9$	45	5	0.4	< 0.1
Bluestem Prairie Preserve/Stolley Prairie	$128.1 \pm 13.2$	23	1	0.2	< 0.1
Schneekloth South Prairie	$98.5 \pm 7.2$	12	12	0.1	0.1
					$0.1 \pm 0.1$
Residential gardens					
105th street	$162.8 \pm 8.6$	127	25	0.8	0.2
46th street	$66.1 \pm 5.0$	25	11	0.4	0.2
80th street	$63.4 \pm 2.9$	43	27	0.7	0.4
Bellevue Public Library	$36.6 \pm 1.2$	57	15	1.6	0.4
Gemini Park	$91.6 \pm 6.1$	189	47	2.1	0.5
Heron Haven Sanctuary	$30.4 \pm 1.9$	5	2	0.2	0.1
Joslyn Art Museum's Children's Discovery Garden	$135.7 \pm 7.1$	90	27	0.7	0.2
Logan Middle School	$44.4 \pm 2.3$	33	14	0.7	0.3
New Life Baptist Church	$12.4 \pm 0.9$	0	0	0.0	0.0
Northern Hills Drive	$571.8 \pm 37.1$	89	122	0.2	0.2
St. Thomas More Catholic School	$17.3 \pm 1.4$	15	4	0.8	0.2
Sump Memorial Library	$31.3 \pm 2.6$	17	0	0.5	0.0
Waldo Circle	$42.3 \pm 1.9$	33	23	0.8	0.5

The total number of monarch eggs counted and total number of monarch larvae collected at each site are presented and then standardized by the average number of stems inspected. Number of stems represents the Mean ± SE across all visits

averaged  $68.1 \pm 18.8$  eggs (95% CI 28.3-107.8; df = 16, t = 0.823, p = 0.423). The number of eggs found across all sites was higher in 2017 than in 2016 [2016 =  $23.1 \pm 11.5$  eggs (95% CI -0.5 to 45.7)],  $2017 = 59.9 \pm 11.5$  eggs (95% CI 37.3-82.5; df = 17, t = -2.63, p = 0.017).

In 2016, we collected 27 fourth and fifth instar monarch larvae at conservation areas and 103 at gardens and reared them in the lab (Table 2). In 2017, we collected 48 monarch larvae at conservation areas and 214 larvae at gardens. The

number of fourth and fifth instar larvae found varied among sites with an average of  $15.0\pm9.9$  caterpillars per conservation site (95% CI -12.6 to 42.6) and  $24.4\pm9.0$  caterpillars per garden (95% CI 4.9–43.9; Table 1). Among the larvae reared in the lab, parasitism by tachinids accounted for 21 of 27 mortality events for larvae from conservation sites and 98 of 117 mortality events for larvae from gardens (Table 2). The ratios of males and females of monarchs that emerged as butterflies were similar for conservation areas

**Table 2** Fates of 4th and 5th instar monarch larvae collected and reared in the lab in 2016 and 2017 at conservation sites and gardens

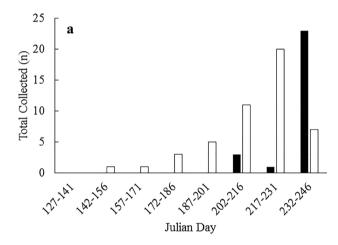
Year	Survived	Cause of death					
		Parasitism by fly	Failure-to- thrive	Failure to eclose	Unknown causes		
2016							
Conservation	20 (11 ♀, 9 ♂)	6			1		
Gardens	72 (40 ♀, 32 ♂)	27	3	1			
2017							
Conservation	28 (16 ♀, 12 ♂)	15	1		4		
Gardens	128 (61 ♀, 67 ♂)	71	7		8		

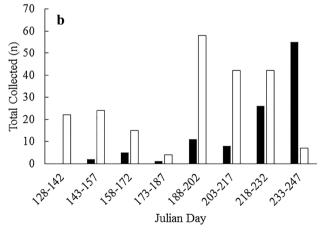
Mortality during the larval or chrysalis stage included parasitism by tachinid flies, larval death attributed to "failure to thrive" syndrome (*sensu* Oberhauser et al. 2007), and unexplained mortality. Individuals that survived to adulthood were sexed and released



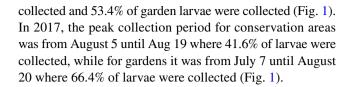
and gardens in 2016 [ $X^2$  (1, N=92)=0.002, p=0.965], in 2017 [ $X^2$  (1, N=156)=0.49, p=0.483], and with the years combined [ $X^2$  (1, N=248)=0.31, p=0.579; Table 2]. Sex ratios did not deviate from 50:50 for any group (binomial test, all p>0.20).

The timing of occurrence of monarch eggs did not differ between the types of sites in either year (Geest 2017). Late instar monarch larvae were found and collected in gardens before conservation areas in both years (Fig. 1). In 2016 in garden sites the first monarch larva collected was 49 days earlier than in conservation areas. Similar to 2016, in 2017 the first larva collected from residential gardens was 20 days earlier than the first larva collected from conservation areas. Peak collection periods were similar for conservation areas and gardens in 2016 but started earlier for gardens in 2017. In 2016, the peak collection period was from August 19 until September 3 where 85.2% of conservation larvae were





**Fig. 1** Timing of collection of monarch larvae (*Danaus plexippus*) from conservation areas (**a**) and residential gardens (**b**) in 2016 and 2017. Solid bars represent 2-week periods from May until September of 2016. Hollow bars are two-week periods from March until August of 2017. The *y*-axis is number of monarch larvae (4th and 5th instars) collected and the *x*-axis is the Julian day of collection. Note the difference in scale of *y*-axis



#### **Monarch survival**

In conservation areas survival from egg to adult was higher in 2016 (0.29  $\pm$  0.05 (SE)) than in 2017 (0.14  $\pm$  0.03; Log rank,  $X^2 = 6.2$ , df = 1, p=0.01; Fig. 2a). In gardens survival from egg to adult was also higher in 2016 (0.21  $\pm$  0.02) relative to 2017 (0.15  $\pm$  0.01; Log rank,  $X^2 = 4.0$ , df = 1, p=0.05, Fig. 2b). Survival did not differ between conservation and garden sites in 2016 (Log rank,  $X^2 = 1.5$ , df = 1, p=0.22; Fig. 2c) or 2017 (Log rank,  $X^2 = 1.1$ , df = 1, p=0.30; Fig. 2d). Across the 2 years, survival in conservation areas (0.18  $\pm$  0.02) and gardens (0.16  $\pm$  0.01) was similar (Log-rank test,  $X^2 = 0$ , df = 1, p=0.87).

# Lespesia archippivora parasitism

Parasitism rates by *L. archippivora* were similar between conservation areas and gardens in 2016 (conservation areas: 22.2% and gardens: 26.2%,  $X^2$  (1, N=130)=0.105, p>0.10; Table 2), in 2017 (conservation areas: 31.3% and gardens: 33.2%,  $X^2$  (1, N=262)=0.034, p>0.10; Table 2), and with the 2 years combined (conservation areas: 27.6% and gardens: 30.9%,  $X^2$  (1, N=392)=0.244, p>0.10; Table 2). There was no difference in parasitism rates between 2016 and 2017 in either conservation areas ( $X^2$  (1, N=75)=0.699, p>0.10) or gardens ( $X^2$  (1, N=317)=1.579, p>0.10).

Peak times of parasitism varied across both field seasons in both conservation areas and gardens but followed a general trend of high parasitism early in the season, followed by a period of no parasitism, with another wave of parasitism late in the field season (Fig. 3). In both years the period of no parasitism in conservation areas or in gardens occurred from June 14 to July 16.

# Ophryocystis elektroscirrha parasitism

In 2016, only 4 monarchs out of 92 (4.3%) adults tested were infected with *O. elektroscirrha*. Three of these butterflies came from a single garden and the remaining one was collected from a conservation area. The three butterflies from a single garden were heavily infected with a spore load of 4 while the conservation butterfly had a spore load of 3. In 2017, only 1 adult out of 156 (0.6%) was infected with spores with the single butterfly being heavily infected with a spore load of 4. This sole spore infected butterfly came from a garden. In both years, the butterflies with *O. elektroscirrha* occurred at the end of the field season in the migratory



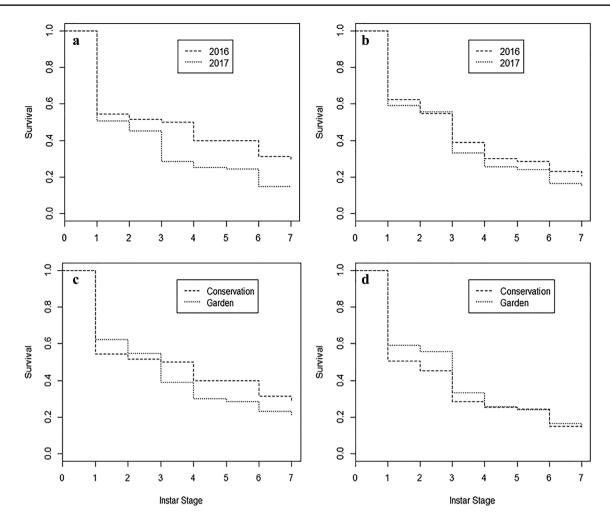


Fig. 2 Kaplan–Meier survival curves for monarch butterflies (*Danaus plexippus*) where the y-axis is the survival rate and the x-axis is the monarch life stage (0=egg, 1-5=instars, 6=chrysalis, 7=adult). In both conservation areas (a) and gardens (b) survival from egg to adult

is higher in 2016 than in 2017 (Log rank,  $X^2 = 6.2$ , df = 1, p = 0.01). Survival did not differ between conservation and garden sites in 2016 (c; Log rank,  $X^2 = 1.5$ , df = 1, p = 0.22) or 2017 (d; Log rank,  $X^2 = 1.1$ , df = 1, p = 0.30)

generation. There was no significant difference in parasitism by O. *elektroscirrha* between the two habitat types (Fishers Exact Test, p = 1.00).

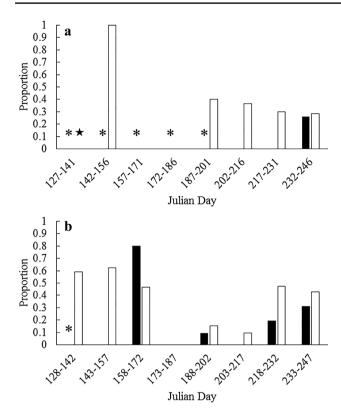
# Milkweed species and density

Site level and patch level milkweed density in conservation areas were not correlated in either year (2016: r=0.43, p=0.47, 2017: r=0.32, p=0.61; Supplemental Material Table A.3). Conservation areas had between 1 and 3 *Asclepias* species present with *A. syriaca* found at every site both years (Geest 2017). The mean number of *Asclepias* species per conservation area in 2016 and 2017 was  $1.2 \pm 0.03$  SE and  $1.7 \pm 0.08$  SE, respectively. Every garden site had at least one species of milkweed with one garden in 2016 having nine species and one garden in 2017 having eighteen species (Supplemental Material Table A.2). The mean number

of milkweed stems per garden in 2016 was  $68.9 \pm 25.2$  SE, however there was high variability from garden to garden with the minimum number of stems being 9.63 and the maximum 353.81. In 2017, the mean number of milkweed stems increased to  $134.74 \pm 11.07$  with the variability in gardens increasing as well with a minimum number of stems being 14.46 and the maximum 774.83 (Supplemental Material Table A.2).

Milkweed density at conservation areas varied among sites in 2016 and 2017 but this variation did not explain variation in monarch egg numbers or *L. archippivora* parasitism (all p-values > 0.05). Likewise, milkweed density did not influence monarch egg numbers at garden sites (all p-values > 0.05). However, in 2016 gardens with lower densities of milkweed had higher incidence of *L. archippivora* parasitism (GLMM, z = -2.93, p = 0.003, SE = 1.24,  $\beta = -3.63$ ). The size of sites did not explain variation among sites in the number of





**Fig. 3** Proportion of monarch larvae (*Danaus plexippus*) parasitized by *Lespesia archippivora* in conservation areas (**a**) and residential gardens (**b**) in 2016 and 2017. Solid bars are 2 week periods from May until September of 2016. Hollow bars are 2 week periods from March until September of 2017. The *y*-axis is proportion of monarch larvae parasitized and the *x*-axis is the Julian day of collection for the monarch larvae. An asterisk denotes period of time when no monarch larvae were collected in 2016. A star denotes period of time when no monarch larvae were collected in 2017

monarch eggs or incidence of parasitism in either conservation areas or milkweed gardens in either year (all p-values > 0.05; Supplementary material Table A.4).

Twelve different species of milkweed were recorded in 2016 with *A. syriaca* being the predominant milkweed species at every site except at two gardens where *A. incarnata* was the only species planted. In 2017, one garden planted an additional 6 species of milkweed bringing the total number of different milkweed species recorded to 18. *A. syriaca* remained the predominant species found in every garden except the two that solely had *A. incarnata*. In 2016 and 2017 the majority of collected monarchs came from *A. syriaca* (2016: 84.6% and 2017: 85.5%). Across both years monarchs were found to use 11% of available *A. syriaca* stems.



Significant effort and resources are being directed towards establishing milkweed gardens as part of the conservation strategy directed at monarch butterflies (Panella 2017; Pleasants et al. 2017). While these gardens attract adult butterflies, it is important to understand if reproduction in gardens is successful, relative to traditional conservation areas. If monarch eggs laid in gardens do not survive through to the adult stage investment in gardens may not be a good use of scarce conservation resources. Garden sites could be population sinks as has been suggested for some butterfly species (e.g. Levy and Connor 2004) or even act as ecological traps if adults are attracted away from sites where successful reproduction is more likely.

Our approach was to compare reproduction on milkweed found in existing gardens to that found in conservation areas. The previous studies that have most directly addressed the potential contribution of gardens to monarch conservation have used experimentally planted milkweed to mimic conditions in gardens and natural areas (Cutting and Tallamy 2015; Majewska et al. 2018). The advantage of this experimental approach is that it controls for other differences among sites such as milkweed variety, density and other management activities. In contrast, our approach of using existing gardens and conservation sites deliberately introduces variation in site characteristics and management activities known to be important determinants of butterfly reproductive success including plant choice, chemical use, and weed control (Levy and Connor 2004; Muratet and Fontaine 2014; Nail et al. 2015; Stenoien et al. 2015). While this observational approach is not a substitute for carefully controlled experimental studies, it provides an important complement to those results that encompasses the conditions butterflies experience in the field.

#### **Monarch recruitment**

To contribute to conservation, gardens must first attract adults to lay eggs. Female monarchs are selective about the milkweed stems they choose to oviposit on and do not use all of the host plants that are available to them (Borkin 1982; Stenoien et al. 2015). We used total number of eggs within a study site as one measure of monarch recruitment. Overall, we found more eggs in gardens than in conservation areas both years, in spite of the fact that conservation sites were larger and we spent more time searching those areas. However, the number of eggs found varied widely among both conservation and garden sites, producing broad and overlapping confidence intervals and the



differences were not significant. The lack of a consistent difference in eggs between conservation and garden sites suggests that gardens have the potential to recruit as many egg laying monarchs as traditional conservation sites. This conclusion is consistent with results from previous studies that found higher egg densities in gardens (Stenoien et al. 2015) and on milkweed planted in a residential setting compared to natural areas (Cutting and Tallamy 2015). Equally important, however, is the variability in egg density itself. The high variability we observed among both conservation areas and gardens emphasizes that other characteristics of sites produce large differences in the value of sites for monarchs that overwhelm any effects of conservation or garden status.

Monarch egg laying activity at all sites built up over the season, peaking with the final migratory generation in late summer. In 2017, monarchs arrived early relative to most years (Howard and Davis 2015), with adults being observed in late March and eggs first being detected in early May. In comparison, in 2016 the first sightings of adults were in mid to late May while eggs were first observed during the final week of May. By the end of May in 2016, 22 eggs had been detected across all study sites. In 2017, 257 eggs had been observed across all study sites by the end of May. This difference between years was not limited to our study area. In 2017, multiple observers noted monarchs arrived in Nebraska in mid to late March, representing the earliest recorded arrival of adult monarchs since at least 2000 (Graham 2017; Journey North 2017). The generation that arrived in Nebraska in early 2017 was the overwintering generation from Mexico which normally does not travel further north than southern Texas (Oberhauser et al. 2001; Graham 2017). It has been suggested that strong winds are the reason the monarchs traveled so far North so quickly in 2017 (Graham 2017). In contrast, in 2016 the initial generation that arrived was comprised of the offspring of the overwintering generation, also called the first adult generation. While this is the generation expected to reach Nebraska, a late February 2016 ice storm in the monarch's over-wintering region of Michoacán, Mexico killed over a million monarchs preparing to migrate north into Texas (Maeckle 2016) and may have delayed the arrival of monarchs to Nebraska and contributed to the small initial numbers of arriving monarchs in 2016.

#### Monarch survival

Our analysis of monarch survival found no difference between gardens with planted milkweed and conservation areas in either 2016 or 2017. When we combined data from the 2 years, we estimated survival for monarchs in garden areas was 16.2% and in conservation areas 18.2% though the differences were not significant. Survival was higher in 2016 than in 2017 in both conservation areas and residential

gardens. This is consistent with Cutting and Tallamy's (2015) experimental study that found no consistent difference in overall survival between monarchs using milkweed planted in residential and natural sites. Likewise, in their analysis of the long-term citizen-science based Monarch Larva Monitoring Project data, Nail et al. (2015) included site type in their analysis but a site's status as a garden did not emerge as one of the top models explaining survival. While this pattern starts to suggest that gardens can provide suitable habitat for monarch reproduction, Majewska et al. (2018) sounded a cautionary note based on their finding that larval survival was lower in their experimental garden sites relative to more natural sites outside the gardens.

The period of greatest mortality is from egg to 1st instar in both habitats in both years, as has been found in other studies of monarchs (e.g. Borkin 1982; Zalucki et al. 2001b; Prysby 2004; Oberhauser et al. 2007). Eggs and early instars may be especially vulnerable to predation due to a lack of protection from toxins that later instars have developed from consuming milkweed (Zalucki 1981) and can also suffer mortality from latex drowning (Zalucki and Brower 1992; Zalucki et al. 2001a). We produced survival curves that include transition from larval to adult stages. This portion of monarch survival is determined by rearing 4th and 5th instars indoors, rather than in the field because 5th instars leave milkweed to pupate and finding cryptically colored chrysalises in the wild are extremely rare and resource intensive (Borkin 1982; Oberhauser et al. 2007; Nail et al. 2015). Indoor rearing results in protecting larvae and pupa from predation, additional parasitoids, weather, exposure to insecticides, and other sources of mortality, producing estimates of survival that are almost certainly higher than occur naturally (Oberhauser et al. 2007; Nail et al. 2015). However, including this stage in analyses of survival is important because of the high mortality associated with parasitism that is not reflected in survival curves that end at the 3rd or 4th larval instar (Nail et al. 2015).

Our overall estimates of survival across life stages varied from 14.4 to 31.4%, placing our estimates at the high end compared to previous studies that reported survival rates between 2 and 24% (Borkin 1982; Oberhasuer et al. 2001; Prysby 2004; Cutting and Tallamy 2015; Nail et al. 2015). The substantial variation reported in survival is not surprising given that these studies were conducted across a species range, in varying habitats, and in different years (Zalucki and Kitching 1982b). As in our study, year to year variation has been found to be substantial within a site and estimates of survival benefit from long-term studies that quantify year-toyear variability (Nail et al. 2015). In addition, methodological differences may contribute to the variation in survival reported by different studies. For example, the time between checks of milkweed plants and larvae varied among studies from a day (Cohen and Brower 1982; Prysby 2004; De Anda

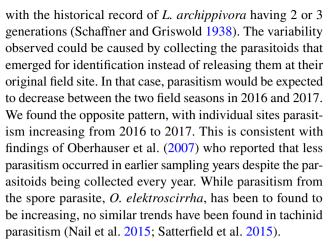


and Oberhauser 2015), to 3 or 4 days (Zalucki and Kitching 1982b, this study), to a week or more (Borkin 1982; Oberhauser et al. 2001; Nail et al. 2015; Majewska et al. 2018). More frequent checks should result in detecting a greater proportion of eggs laid and this produce lower estimates of survival relative to studies where plants are checked less frequently and some eggs are eaten before they are detected. Likewise, to account for imperfect detectability among the early instars we chose to correct data to account for missed detections (De Anda and Oberhauser 2015) and use instarbased survival curves, while other monarch larval survival studies bypass the issue of imperfect detectability of eggs and larvae by using the ratio of the number of eggs to instar five larvae to (Calvert 2004; Nail et al. 2015; Majewska et al. 2018), while still others limit their survival estimates to the first 2 or 3 instars due to concerns about detectability of older larvae (Cutting and Tallamy 2015; De Anda and Oberhauser 2015). These and other methodological differences highlight the importance of future studies that quantify the probability of detecting eggs and larvae and what factors drive detectability using double observer or other methods (Elphick 2008).

#### **Parasitism**

Estimates of survival from egg to late instar larvae will miss much of the mortality caused by tachinid fly parasitism that can be a major source of mortality in the pupal stage (Nail et al. 2015). Because so few larvae reach this stage, variation in parasitism between gardens and natural areas could greatly influence the number of adults produced. Parasitism rates at sites used in this study ranged from 0 to 100% parasitism of collected larvae. This level of parasitism by tachinids is not atypical (Zalucki 1981), and parasitism has been found to be widely variable year-to-year and patch-topatch (Oberhauser et al. 2007). While parasitism rates were variable, variation among sites in our study was not related to their status as gardens or natural areas. While a higher proportion of larvae from gardens were parasitized, the differences were not significant. L. archippivora is considered a generalist (Etchegaray and Nishida 1975; Prysby 2004) however some studies have suggested it shows a preference for monarch larvae and other recorded host species may be due to spill-over parasitism (Janzen and Hallwach 2009; Oberhauser 2012). All collected tachinid flies were identified as L. archippivora which is not unusual since L. archippivora is the most frequent tachinid parasitoid of monarchs (Schaffner and Griswold 1938; Oberhauser et al. 2017a).

We found one peak time period for *L. archippivora* in conservation areas and two peak time periods for parasitoids in residential gardens. This result might suggest that *L. archippivora* is bivoltine with a gap between generations occurring from mid-June to mid-July which is consistent



In contrast to the significant rates of parasitism by *L. archippivora*, *we* found *O. elektroscirrha* on only 2% of reared monarchs which is lower than the reported average of 8% infection rate for the eastern population (Altizer et al. 2000). Any monarch larvae parasitized by *L. archippivora* did not survive to be tested for *O. elektroscirrha*, so *O. elektroscirrha* spore infection rates may be higher. Not surprisingly given the small number of infections, we found no difference between garden and conservation sites. In 2016, three of the four infected *O. elektroscirrha* butterflies came from the same garden. Repeated visits from infected adults increase the number of spores that are distributed across the leaves and egg cases (Bartel et al. 2011). This suggests, that only a few infected adults are needed to cause heavy spore loads in offspring.

# Effects of milkweed density

One of the significant observations from our study is the large amount of variation among gardens and among prairie sites. This variability suggests that any differences that might exist between gardens and conservation areas is likely overshadowed by differences in quality among sites within each category. While the present study was not designed to determine why sites varied, we did ask how the density of milkweed might influence monarch survival or parasitism among conservation areas and residential gardens to begin to ask what makes a good prairie and a good garden for monarchs.

Milkweed density varied between sites and is known to influence monarch ovipositioning choice (Zalucki and Kitching 1982a; Zalucki and Suzuki 1987; Stenoien et al. 2015). We found a slight negative correlation between higher milkweed density and a lower number of eggs in 2017 in conservation areas and a slight positive correlation; however, neither was significant. Zalucki (1981) and Stenoien et al. (2015) found higher counts of eggs in sites with lower milkweed density. However, these studies occurred over a longer timespan than our study. Pitman (2017) similarly found



higher egg densities in sites with lower milkweed density. Pitman (2017) suggested that higher quality plants occurred if milkweed plant at low density experienced less competition and that quality of milkweed may explain monarch oviposit choice. Egg densities per plant vary considerable across regions and years with overall densities decreasing since 2006 (Lindsey et al. 2009; Stenoien et al. 2015). Our results reflect this variation at a smaller scale, with both egg and larval numbers per stem ranging from the low end of prior reports to average or slightly above for our most productive sites.

The characteristics of gardens and conservation sites have the potential to influence parasitism rates. We found a slightly positive correlation between L. archippivora parasitism and lower milkweed densities within conservation areas in 2017, but it was not significant. This may be due to milkweed patches being denser than the surrounding areas within conservation sites. In residential gardens we found a significant negative correlation in 2016 suggesting that sites with lower milkweed density had higher parasitism rates. A similar study in Australia with a different species of tachinid found the reverse and that parasitism was lower in smaller patches with lower densities of milkweed (Zalucki 1981). Most tachinid species find host larvae by sight (Prysby 2004). Milkweed within gardens was typically more exposed than milkweed within prairies. Similarly, in tallgrass prairies with high amounts of vegetation milkweed patches may be easier to locate for tachinids than isolated plants.

The most widespread and abundant milkweed species in both years was A. syriaca. Milkweed stems increased across all garden and conservation sites from 2016 to 2017, with the increase driven by an increase in stems of A. syriaca. This is unsurprising since A. syriaca can spread not only through wind dispersed seeds but also through rhizomes (Kaul et al. 2011). A. syriaca seedlings can produce large numbers of stems in short amounts of time with a single seedling creating an additional 56 stems and 94 seedlings in 4 years (Bhowmik 1978, 1994). Monarch larvae were found on approximately 11% of the A. syriaca stems available to them which was the most out of all native local milkweed species (Geest 2017). This is important as studies like Stenoien at al. (2015) list the loss of A. syriaca as one of the driving forces of population decline in the monarch. The two most commonly used host species after A. syriaca were A. incarnata and A. verticillate (Geest 2017). All three of these species have relatively low toxicity when compared to species across North America (Malcolm 1991). However, the cardenolide concentration in the leaves and shoots of the two most used host plants A. syriaca and A. incarnata are relatively high compared to the two least used host plants A. verticillata and A. tuberosa (Malcolm 1991; Rasmann and Agrawal 2011). Monarch oviposit choice has been shown to be influenced by species toxicity with female monarchs choosing species with intermediate levels of toxicity (Oyeyele and Zalucki 1990; Zalucki et al. 1990).

# **Conclusion**

We found similar recruitment, survival, and parasitism for monarch butterflies in conservation areas and gardens, suggesting that gardens have the potential to contribute to monarch butterfly conservation. However, two important factors emerge that emphasize the challenges of realizing the potential of milkweed gardens. First, our results document the variability in quality of among gardens. Additional information about what characteristics of gardens attract monarchs and contribute to their success is needed. Second, incorporating milkweed gardens into conservation strategies must recognize the critical role the human element will play in determining if gardens benefit. At the community level, gardens have been impacted by local laws that penalize gardeners for growing milkweed because it is classified as a weed by some states at the county level (Oberhauser et al. 2008; Shahani et al. 2015). Even more significant is gardener behavior. In order for gardens to be as effective a conservation strategy as possible, gardeners must practice best conservation methods on their land. Based on our interactions, some of the same gardeners who invested in planting milkweed were also most concerned about the appearance of their garden and were reluctant to forgo the use of pesticides and to leave milkweed into the late summer when monarch use peaked but the plants had passed their most attractive stages (Geest 2017). In addition, gardens are prone to alterations year-to-year and the impact of these changes on monarchs is not known but will need to be addressed. While the overall conclusion of our research is that there are potential benefits of using residential gardens as a conservation strategy for monarch butterflies, implementation of these programs needs to include a strong outreach and education component.

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#### Compliance with ethical standards

Conflict of interest None of the authors are aware of other potential conflicts of interest.

Research involving human participants and/or animals Insects and other invertebrates are not covered under U.S. animal care regulations and are not reviewed by the university's Institutional Animal Care and Use Committee (IACUC). Our research protocols followed standard methods for working with monarch butterflies. Human participants were not used in this study.

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