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Reproductive diapause in *Hippodamia convergens* (Coleoptera: Coccinellidae)
and its life history consequences.

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1 **Running Head: REPRODUCTIVE DIAPAUSE IN HIPPODAMIA CONVERGENS**

2
3 **Abstract**

4 **Adult *Hippodamia convergens* (Guerin) in reproductive diapause**
5 **were collected from a spring cohort in western Kansas and held in pairs for**
6 **the duration of their lives to assess female reproductive schedules under**
7 **conditions of limited food availability. Environmental conditions were set**
8 **to mimic natural seasonal day lengths and diurnal temperature cycles for**
9 **the region. To approximate conditions of limited food availability typical of**
10 **summer conditions in western Kansas, beetles were provided continuous**
11 **access to sunflower petioles and periodic access to protein sources, both**
12 **animal (*Ephestia kuehniella* Zeller eggs) and vegetable (bee pollen). A total**
13 **of 113 out of 171 females (66.1%) became reproductive over the next five**
14 **months within a mean of 55.0 ± 3.0 d. These females lived an average of**
15 **134.5 ± 4.6 d and produced a mean of 106.9 ± 11.6 eggs in a mean of $6.6 \pm$**
16 **0.6 d of oviposition. Thus, reproductive diapause gradually decayed over**
17 **time even when females did not encounter a high quality food supply. Egg**
18 **production peaked every fourth day following provision of animal protein.**
19 **A subset of 20 females, randomly selected from among those still non-**
20 **reproductive on Aug. 14 and switched to an aphid diet (*ad libitum* provision**
21 **of *Schizaphis graminum* Rondani) produced a mean of 654.6 ± 109.7 eggs**
22 **each, almost 10 times as many as females on the maintenance diet with**
23 **similar reproductive schedules. However, the longevity of greenbug-fed**
24 **females was reduced by more than 30% compared to the latter group,**
25 **suggesting a tradeoff between reproductive effort and survival. The costs**
26 **of reproductive diapause were evident as an increased risk of mortality**
27 **prior to oviposition and declining fecundity and fertility with age. Our**
28 **results suggest a variable number of overlapping generations can occur**
29 **annually in western Kansas, potentially as many as five.**

30

1 **Key Words:** diapause, diet, fecundity-longevity trade-off, *Hippodamia*
2 ***convergens*, reproduction, *Schizaphis graminum***

3

4 **1. Introduction**

5

6 The convergent lady beetle, *Hippodamia convergens* (Guerin), is an
7 important predator of cereal aphids throughout much of the High Plains region of
8 the central United States (Michels et al. 2001, Nechols and Harvey 1998;
9 Michaud and Qureshi 2005). Principle prey species for *H. convergens* in western
10 Kansas include the Russian wheat aphid, *Diuraphis noxia* Mordvilko, and the
11 greenbug, *Schizaphis graminum* (Rondani). Other aphid species that
12 sporadically serve as alternative prey include the corn leaf aphid, *Rhopalosiphum*
13 *maidis* (Fitch), and the bird-cherry oat aphid, *Rhopalosiphum padi* L, along with a
14 number of other non-economic aphid species that feed on wild grasses and
15 herbaceous weeds. Other coccinellid species preying on these aphids in the
16 region include *Coleomegilla maculata* DeGeer, *Coccinella septempunctata* L.,
17 *Cycloneda munda* (Say), and *Scymnus* sp (Rice and Wilde 1988). Although
18 *Harmonia axyridis* Pallas has been present in the region for a number of years, it
19 has failed to rise to prominence in this coccinellid community as it has in many
20 other regions throughout North America over the past decade (Colunga-Garcia
21 and Gage 1998, Cottrell and Yeargen 1998, Michaud 2002).

22 For several years now, we have observed *H. convergens* dominate the
23 mixed species aggregations of coccinellids that form on sunflower plants
24 following emigration from maturing wheat fields in early summer where they have
25 completed their first generation feeding on cereal aphids. The relative
26 predominance of *H. convergens* among the assemblage of coccinellids in wheat
27 in this region was noted by Nechols and Harvey (1998) while evaluating natural
28 enemy responses to *Diuraphis noxia* (Mordvilko). As wheat fields dry down in
29 early summer, an abundance of wild *H. annuus* plants emerge in roadside
30 habitats, as do fields of cultivated varieties grown for oil and confection markets.
31 Coccinellid adults, along with many other insect families, can be directly

1 observed utilizing the succulent petioles of sunflower plants as a source of
2 hydration during summer months in this arid region. Michaud and Qureshi
3 (2005) demonstrated that access to sunflower sap sustained the survival of *H.*
4 *convergens* adults in the absence of food better than did access to distilled
5 water. Coccinellids feeding on aphid prey do not require supplementary water
6 (Hodek and Honek 1996), so the apparent importance of this 'sap drinking'
7 behavior to multiple coccinellid species likely reflects the fact that aphid prey can
8 be very difficult to find during hot summer months on the prairie. Sap-drinking
9 behavior is not limited to sunflower plants and other common weed species (eg.
10 *Kochia scoparia* L., *Amaranthus palmeri* S. Wats) are sometimes utilized,
11 although *H. annuus* plants seem to be the most attractive.

12 Coll and Guershon (2002) reviewed the various ways plants may be
13 utilized by omnivorous insects, but did not mention sap-feeding by predatory
14 species. Schmidt (1992) surveyed coccinellid abundance on 76 plant species in
15 Germany (mostly common weeds) and found strong evidence of patterns of plant
16 association and avoidance. Associations with plants did not seem to be driven
17 entirely by prey availability as fully 40% of the beetles surveyed occurred on
18 plants without any aphids. The use of plant sap by lady beetles is perhaps an
19 important aspect of coccinellid ecology in arid habitats that has been largely
20 overlooked. Sunflower sap, in particular, may be exceptionally suitable as a
21 source of hydration, and possibly some nutrition, for both predaceous and
22 parasitic insects. Royer and Walgenbach (1991) catalogued a large number of
23 predaceous arthropods in North Dakota associated with sunflower plants in
24 various growth stages. Suter (1988) found that sowing 2-3 sunflowers in every
25 100 m² of corn increased the abundance of beneficial insects (coccinellids,
26 syrphids, lacewings and predatory Hemiptera) compared to a corn monoculture.
27 Pilson (2005) documented the attractiveness of sunflower sap to a wide range of
28 insects once stems had been gouged open by *Euphora inda* (Coleoptera:
29 Scarabaeidae). We have now documented sap-drinking behavior on cultivated
30 sunflowers by adult lacewings (Neuroptera), flies (Diptera), beetles (Coleoptera)

1 of diverse families, and Hymenoptera, including a plethora of wasp and ant
2 species.

3 The predominance of *H. convergens* in the assemblage of coccinellids in
4 grassland ecosystems of the High Plains (Rice and Wilde 1980, Nechols and
5 Harvey 1998, Brewer and Elliott 2003) is suggestive of unique ecological
6 adaptations that provide it with substantial survival and/or reproductive
7 advantages over other species. Use of sunflowers is not unique to *H.*
8 *convergens* and virtually all coccinellid species in the region appear to engage in
9 this behavior. Aphids tend to be scarce during hot summer months but other,
10 less suitable, sources of animal protein are probably available, especially the
11 eggs of a wide range of lepidopteran species that are laid on the foliage of
12 various herbaceous plants throughout the season. Similarly, pollen of a variety
13 of wild and cultivated flowering plants is available periodically throughout the
14 summer and is known to serve as an important food supplement for adult
15 coccinellids during periods when animal prey are scarce (Hodek and Honek
16 1996). Although *H. convergens* females readily consume copious amounts of
17 pollen, this food alone is insufficient to support the maturation of their eggs (J.P.
18 Michaud, unpublished data).

19 The ability of first generation *H. convergens* adults to undergo
20 reproductive diapause in response to limitations in the quality and availability of
21 food may represent a significant adaptation not shared by other species
22 (Michaud and Qureshi 2005). Induction of reproductive diapause in response to
23 food limitation has also been demonstrated for *Harmonia sedecimnotata* F.
24 (Zaslavski et al. 1998). Adult *H. convergens* can use limited food resources to
25 develop fat bodies in their haemolymph and delay onset of reproduction until a
26 high quality food source, such as a large colony of a suitable aphid species, is
27 encountered. Previously, Michaud and Qureshi (2005) demonstrated that
28 immediate provisioning of greenbug, *S. graminum*, to first generation adult
29 females prevented most from entering diapause, whereas provisioning of
30 greenbugs after several weeks on a maintenance diet caused a significant
31 proportion to break diapause within a few days. Here we examine complete

1 reproductive and life history schedules of a first generation (spring) cohort of *H.*
2 *convergens* adults under environmental conditions replicating seasonal norms for
3 western Kansas. The insects were subjected to a diet treatment designed to
4 simulate limited availability of plant and animal protein sources using types of
5 food likely available to adult beetles throughout summer and fall (pollen and eggs
6 of Lepidoptera). Partway through the summer, a subsample of beetles was
7 switched to an ad libitum diet of greenbug to simulate encounter with a rich prey
8 patch so that its effects on reproductive diapause, oviposition, and overall
9 reproductive success could be observed.

10

11 **2. Materials and methods**

12

13 On 5 June, 2005, we estimated relative coccinellid species abundance
14 locally by counting adult beetles in naturally occurring aggregations on cultivated
15 sunflowers, *H. annuus*, at the Agricultural Research Station at Hays, Kansas.
16 Subsequently, we collected more than 500 adult *H. convergens* by tapping them
17 off plant petioles into ventilated Plexiglass containers, transported them to the
18 laboratory, and then transferred them to 9 cm diameter plastic Petri dishes in
19 groups of 10-15 per dish. We knew from previous work (Michaud and Qureshi
20 2005) that adult females collected from sunflowers at this time of year are not yet
21 gravid, having recently emigrated from mature wheat fields as newly emerged
22 adults. Mating pairs were isolated as quickly as they formed and transferred to
23 5.5 cm diameter plastic Petri dishes until a total of 247 pairs were isolated. Each
24 dish was immediately provisioned with an excised segment of sunflower petiole
25 (ca. 4 cm long) and a measure of frozen eggs of *Ephestia kuehniella* Zeller (ca.
26 10 mg.). The sunflower petiole segments were replaced every two days with
27 material harvested from cultivated fields. On every third day, each dish received
28 a measure of bee pollen (ca. 10 mg.), and every sixth day, another measure of *E.*
29 *kuehniella* eggs. All insects were transferred to clean dishes every sixth day.
30 This feeding cycle was designed to simulate sporadic access to limited sources
31 of vegetable and animal protein that we anticipated would represent the field

1 situation for a majority of beetles during summer months in the High Plains. This
2 feeding cycle is henceforth referred to as the 'maintenance diet'. Sunflower
3 stalks were discontinued on 11 September and replaced with water encapsulated
4 in polymer beads (Hydrocapsules[®], ARS Laboratories, Gainesville, Florida) since
5 little succulent plant material remains available in fall, although beetles may still
6 have periodic access to moisture in the form of dew. Provision of pollen was
7 discontinued after 7 October on the rationale that natural pollen sources would
8 no longer be available in the field beyond this date.

9 All dishes were labeled with a number corresponding to the female. When
10 any female died, she was immediately weighed, dissected to determine the
11 presence or absence of mature eggs in her ovaries, and the male held separately
12 until it could be re-paired with a widowed female. In this manner, all females had
13 virtually continuous access to a male for the entire course of the experiment and
14 no female was ever held without a male for more than a few days.

15 Dishes containing beetle pairs were laid out on a series of trays and held
16 in a climate-controlled growth chamber. Day length was adjusted weekly to
17 correspond to that of particular dates at 40° latitude according to Beck (1968)
18 and an oscillating temperature cycle was maintained with an amplitude of ca. 10
19 °C corresponding to long-term average values for the region obtained from
20 weather records at Agricultural Research Center - Hays, albeit without the
21 occasional extreme temperatures often experienced in High Plains summer
22 conditions. These climatic conditions were adjusted every two weeks to
23 correspond to average seasonal values and are summarized in Table 1 for the
24 entire experiment. All dishes were examined daily for oviposition. Eggs were
25 usually attached directly to the surface of the plastic dishes and were harvested
26 by simply changing the beetle pair to a new dish inscribed with the female's
27 reference number. Eggs were labeled with date and female number and then
28 held until eclosion under the same environmental conditions as the adults,
29 whereupon the total number of eggs and hatching larvae were counted. Female
30 fertility was calculated as the proportion of her eggs that hatched.

1 Large colonies of greenbug, *S. graminum* 'biotype I', were reared on trays
2 of sorghum, *Sorghum bicolor* (L.) Moensh, in a greenhouse. On 14 August, a
3 series of 20 pairs of beetles were selected at random from among those in which
4 females had yet to oviposit. These beetle pairs were henceforth provided with an
5 *ad libitum* diet of greenbug (introduced on excised sorghum leaves) daily for the
6 next two months, or as long as the female remained alive.

7 Voucher specimens of beetles were placed in the Kansas State University
8 Museum under Lot. 177. Life history and reproductive data were compared
9 between various treatment groups by one-way ANOVA, followed by a Tukey test
10 for means separation in cases where three groups were compared. Longevity
11 and reproduction data for greenbug-fed females were compared to two groups of
12 maintenance diet females, those reproducing earlier than the mean of greenbug-
13 fed females (18 August), and those beginning oviposition on or after 18 August).

14 15 **3. Results**

16 17 *3.1 Relative Abundance*

18
19 A total of 454 adult coccinellids were tallied on cultivated sunflower plants
20 on 6 June, 2005. The relative composition of species was 74.4% *H. convergens*,
21 20.3% *C. septempunctata*, 2.6% *C. munda*, 1.8% *C. maculata*, and 0.9% *H.*
22 *axyridis*.

23 24 *3.2 Longevity*

25
26 Larvae of the parasitoid *Dinocampus coccinellae* (Shrank) (Hymenoptera:
27 Braconidae) emerged from a total of 90 beetles (18.2 %) over a period of 31 days
28 after collection (43 males, 47 females). The sex of parasitized beetles was
29 determined by dissection and their surviving mates were re-paired with others of
30 the opposite sex. Data for parasitized beetles were excluded from all analyses.
31 The cumulative survival of unparasitized beetles on the maintenance diet varied

1 between males and females, the former tending to die somewhat sooner (Fig. 1).
2 Females receiving the greenbug diet demonstrated a steeper mortality curve
3 than did their counterparts that had been non-reproductive for the same period
4 (up to Aug. 18) but remained on the maintenance diet. There were significant
5 differences in longevity among groups of females that were transferred to the
6 greenbug diet ($n = 20$), reproductive on the maintenance diet beginning
7 oviposition earlier than greenbug-fed females ($n = 113$), and 3) reproductive on
8 the maintenance diet but not beginning oviposition earlier than greenbug-fed
9 females ($n = 28$) ($F = 14.742$; $df = 2,130$; $P < 0.001$). Greenbug-fed females
10 lived an average of 118.1 ± 8.4 days, significantly less than the 171.1 ± 6.0 day
11 lifespan of females on the maintenance diet that were non-reproductive up to the
12 same date (Tukey test, $\alpha < 0.05$). Females on the maintenance diet that
13 reproduced on an earlier schedule than greenbug-fed females were not
14 significantly different in longevity from them, but were significantly shorter-lived
15 than the group of maintenance diet females with late reproductive schedules
16 (Tukey test, $\alpha < 0.05$). Twelve of the 20 females fed greenbug (60%) died with
17 fully developed eggs in their ovaries, compared with only 17 (15%) of
18 reproductive females on the maintenance diet.

19

20 3.3 *Reproduction*

21

22 A total of 113 out of 171 females (66.1%) on the maintenance diet became
23 reproductive during the course of the experiment and their onset of oviposition
24 over the course of the experiment is depicted in Fig. 2. Total egg production by
25 all females on the maintenance diet is depicted throughout the course of the
26 experiment in Fig. 3. Egg production followed a distinct periodicity with respect
27 to food supply, an ovipositional peak occurring every four days after provisioning
28 with *Ephestia* eggs (Fig. 4). These females lived an average of 134.5 ± 4.6 d
29 and produced a mean of 106.9 ± 11.6 eggs in a mean of 6.6 ± 0.6 d of
30 oviposition that yielded a mean of 72.3 ± 7.9 larvae. Female oviposition on the

1 greenbug diet, summed over all females, is depicted as a function of female
2 reproductive life in Fig. 5.

3 Females receiving the greenbug diet beginning Aug 14 began oviposition
4 a mean of 3.8 ± 0.2 d days later. There was significant variation among groups of
5 females in the mean numbers of eggs ($F = 54.007$; $df = 2,130$; $P < 0.001$) and
6 larvae ($F = 40.292$; $df = 2,130$; $P < 0.001$) they produced. Females receiving the
7 greenbug diet produced a mean (\pm SEM) of 654.6 ± 109.8 eggs and a mean (\pm
8 SEM) of 451.0 ± 92.2 larvae, significantly more (Tukey test, $\alpha < 0.05$) than
9 females on the maintenance diet that began reproduction prior to Aug. 18 (mean
10 \pm SEM = 121.0 ± 14.8 eggs and 82.7 ± 10.1 larvae) or after Aug. 18 (mean \pm
11 SEM = 64.2 ± 9.7 eggs and 40.7 ± 6.7 larvae). Differences between the latter
12 two groups were not significant for either eggs or larvae (Tukey test, $\alpha > 0.05$).
13 There was no difference in fertility among the three groups ($F = 1.534$; $df = 2,130$;
14 $P = 0.220$).

15 Date of first reproduction among females on the maintenance diet was
16 negatively correlated with both lifetime fecundity ($F = 20.23$; $df = 111$; $P < 0.001$;
17 $r^2 = 0.154$) and mean fertility ($F = 3.98$; $df = 111$; $P = 0.048$; $r^2 = 0.035$). There
18 was no significant relationship between fecundity and fertility ($F = 1.70$; $df = 111$;
19 $P = 0.195$), between fecundity and longevity ($F = 3.51$; $df = 111$; $P = 0.64$), or
20 between fecundity and death weight ($F = 0.73$; $df = 111$; $P = 0.395$). However,
21 there was a strong positive correlation between fecundity and number of
22 reproductive days for both females on the maintenance diet ($F = 1201.82$; $df =$
23 111 ; $P < 0.001$; $r^2 = 0.915$) and those receiving greenbug ($F = 60.88$; $df = 18$; P
24 < 0.001 ; $r^2 = 0.772$). The last date on which any viable eggs were laid was 3
25 November, suggesting that a mean temperature of 12°C might represent a
26 threshold below which oviposition does not occur.

27

28 **4. Discussion**

29

30 Aestivation has been inferred to represent an adaptation to hot dry
31 summers (Masaki 1980). The reproductive diapause of *H. convergens* falls short

1 of true aestivation in that the beetles remain active, but a similar function can be
2 inferred, that of enhancing adult survival during periods of low prey availability
3 that are typically associated with very hot weather in the region. Food limitation
4 has been found to be a regulating factor for aestival diapause in relatively few
5 insects (Nechols et al. 1999). Zaslavski et al. (1998) demonstrated a similar
6 effect for *H. sedecimnotata* and Tauber and Tauber (1973) found that food was
7 an important determinant of diapause induction and termination in a California
8 strain of *Chrysoperla carnea* (Neuroptera: Chrysopidae), another relatively
9 specialized aphid predator. Thus, food-regulated diapause could be a particular
10 adaptation of predators specializing on prey such as aphids that exhibit
11 seasonally unpredictable cycles of abundance.

12 Because of the facultative nature of reproductive activity in *H. convergens*,
13 only the spring generation of overwintered adults is likely to reproduce as a
14 comparatively synchronous cohort. Our data suggest that subsequent
15 generations overlap as a function of two factors: 1) variation in the foraging
16 success of first generation females in finding patches of suitable prey sufficient to
17 terminate reproductive diapause directly and, 2) variation among females in
18 spontaneous diapause termination when such patches are not encountered.
19 Thus, *H. convergens* could potentially complete as many as five generations per
20 year if suitable prey remained continuously available throughout the season, a
21 highly unlikely circumstance in most years. These inferences are supported by
22 the previous observations of Hagen (1962) that *H. convergens* can have a
23 variable number of generations per year in California depending on the
24 availability of suitable aphid prey.

25 Although some non-reproductive females on the maintenance diet were
26 able to survive to an age of at least eight months, all were dead by mid-January.
27 Our experimental beetles were subjected to a relatively moderate range of winter
28 temperatures and at least two additional months survival would be needed for
29 beetles to reach spring conditions suitable for reproduction. Therefore, it seems
30 unlikely that first generation adults, even those able to maintain reproductive
31 diapause throughout summer and fall, could survive to overwinter successfully

1 and contribute offspring to the next spring generation. We conclude that a
2 minimum of two generations per year appear necessary for the persistence of *H.*
3 *convergens* populations in the High Plains region of the United States.

4 Largely as a result of the great variation in onset of reproduction among
5 individual females experiencing food limitation (Fig. 2), we would expect *H.*
6 *convergens* populations to normally exhibit an extended distribution of
7 reproductive activity through summer and fall, somewhat similar to that depicted
8 in Fig. 3. However, food sufficient to induce egg production is not necessarily
9 sufficient to ensure female reproductive success – the local food supply at
10 oviposition sites must be adequate to support the completed development of
11 larvae. Even in years when suitable prey are never sufficiently abundant to
12 directly induce diapause termination, variation in onset of reproduction under
13 conditions of food-limitation will ensure that the reproductive effort of some
14 subset of females coincides with the period of best prey availability for larvae,
15 whenever this happens to occur in the course of the growing season. Thus, in
16 certain years, a small proportion of spring generation females may contribute a
17 large proportion of progeny to subsequent generations.

18 A wide range of coccinellid species can be observed exploiting sunflowers
19 for their sap during summer months and this behavior has been previously
20 documented (Michaud and Qureshi 2005). Three of every four coccinellid adults
21 tallied in our sunflower field were *H. convergens*, reflecting the relative
22 dominance of this species among the assemblage of aphidophagous coccinellids
23 inhabiting western Kansas (Rice and Wilde 1988, Nechols and Harvey 1999). To
24 the best of our knowledge, summer diapause in response to food limitation has
25 not been reported in any of the other five species commonly found in our region
26 of study. Reproductive diapause in response to food limitation may therefore
27 comprise an important life history trait that contributes to the success of *H.*
28 *convergens* in the arid High Plains region of the central United States and its
29 significance as an important predator of cereal aphids (Brewer and Elliott 2003).

30 Pairs of *H. convergens* were observed mating frequently from their date of
31 collection until late in life, and in many cases long after the female ceased

1 oviposition, suggesting that sexual activity is entirely unaffected by reproductive
2 diapause. The fitness benefits of female promiscuity have been previously
3 demonstrated in other aphidophagous coccinellids (Majerus 1994) and such
4 promiscuity could give rise to sperm competition. By re-pairing widowed females
5 in the experiment, we reduced the likelihood that female fecundity or fertility
6 would ever be limited by sperm availability, although we were unable to control
7 for possible variation in male fitness that may have affected female reproductive
8 performance via effects on sperm quality.

9 *Hippodamia convergens* females that were switched to an *ad libitum*
10 greenbug diet after more than two months of reproductive diapause initiated a
11 burst of reproductive effort, beginning an average of four days later, that
12 subsequently tapered off through the remainder of their reproductive life (Fig. 5).
13 This distribution of reproductive effort is consistent with the triangular fecundity
14 function inferred by Dixon and Agarwala (2002) for coccinellid females, reflecting
15 a maximal reproductive effort shortly after onset of reproduction that declined
16 gradually with age. Similarly, ovipositing females on the maintenance diet
17 required four days to effectively convert animal protein into eggs (Fig. 4),
18 suggesting that this period may reflect the time required by the female
19 reproductive system to respond to a favorable change in diet. Regression
20 analyses revealed that fecundity varied as a function of the number of days a
21 female produced a clutch, but was not correlated with either female weight or
22 longevity, suggesting that body size bore no relation to female fitness.

23 Fully one third of females on the maintenance diet never became
24 reproductive even though they survived for an average of 147 days. These could
25 represent a sub-group of strongly diapausing females that will only become
26 reproductive if and when an adequate supply of the preferred prey is
27 encountered. Alternatively, they may represent a sub group of individuals
28 intrinsically unable to mature eggs on the sub-optimal maintenance diet.

29 The strategy of delaying onset of reproduction through reproductive
30 diapause (even though resources are available for some egg production) can
31 only comprise an adaptive strategy if two conditions are met: 1) females are able

1 to extend their life by doing so, and thus increase their chances of encountering
2 an optimal prey patch, and 2) if the reproductive success obtainable in a rich
3 patch of optimal prey is sufficiently large to balance the risks associated with
4 delayed onset of reproduction. Individuals of both sexes died at a relatively
5 constant rate throughout the course of the experiment, although males began
6 dying earlier than females (Fig. 1), suggesting that the risk of mortality was
7 relatively constant throughout adult life even in the absence of exposure to
8 predation. The steeper mortality curve of females on the greenbug diet in
9 comparison with those on the maintenance diet is likely indicative of the
10 physiological costs associated with their high reproductive effort. Comparison of
11 longevity between greenbug-fed females and those that also diapaused for at
12 least two months but never received greenbug prey revealed that the latter lived
13 an average of 45 % longer than the former, confirming that females on the
14 maintenance diet increased their lifespan by forgoing or reducing reproduction
15 under sub-optimal conditions. Furthermore, greenbug-fed females were four
16 times more likely to die with mature eggs left in their ovaries, suggesting that
17 their cause of death was often due to failure of physiological processes unrelated
18 to reproduction.

19 The negative correlations observed between date of onset of reproduction
20 and both fecundity and fertility are likely indicative of the costs of reproductive
21 diapause that appear to increase with female age. The longer females delay
22 onset of reproduction, the lower their intrinsic reproductive potential and the
23 greater the risk of dying without progeny. The fact that two thirds of females on
24 the maintenance diet eventually terminated diapause even though optimal
25 conditions were not encountered suggests a tendency for reproductive diapause
26 to decay over time as a function of female age, independently of other factors.
27 However, females that received the greenbug diet after more than two months in
28 diapause produced more than five times the eggs of those on the maintenance
29 diet that initiated reproduction earlier, and almost 10 times as many as those
30 initiating reproduction later. The most fecund female on the greenbug diet
31 produced more than four times the eggs of the most fecund female on the

1 maintenance diet even though she began reproduction more than two months
2 later in life.

3 Clearly, the reproductive diapause of *H. convergens* meets the criteria
4 necessary for it to be considered an adaptive female strategy. Even though the
5 strategy is time-limited because of the inevitable decline in female reproductive
6 potential with age, female lifespan can be effectively extended by delaying the
7 onset of reproduction and female fitness can be increased by 5-10 fold provided
8 a rich patch of suitable prey are eventually encountered. Variation among
9 females in the strength of reproductive diapause likely results in a broad temporal
10 distribution of reproductive activity in the population throughout the summer and
11 fall in most years when prey availability is sporadic. Flexibility in the reproductive
12 schedule of female *H. convergens*, at both the level of the individual and the
13 population, may well be a key life history feature determining its relative
14 importance as a biological control agent of cereal aphids in the arid High Plains
15 region of the United States.

16

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18

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24

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1 **Table 1.** Monthly means of diurnal temperature oscillations and daylengths experienced
 2 by pairs of *Hippodamia convergens* adults monitored for reproductive activity over a six
 3 month period. Daylengths were obtained from Beck (1968) and temperatures
 4 approximated long-term seasonal values for the region.

Month	Mean minimum temperature (°C)	Mean maximum temperature (°C)	Mean median temperature (°C)
June	13.9	24.3	19.1
July	14.0	24.4	19.2
August	12.6	24.1	18.3
September	11.4	23.7	17.6
October	7.2	20.3	13.7
November	6.5	16.3	11.8

5

Date	Daylength
6/6 – 6/14	14:50
6/15 – 6/30	15:01
7/1 – 7/14	14:58
7/15 – 7/31	14:43
8/1 – 8/14	14:18
8/15 – 8/31	13:46
9/1 – 9/14	13:03
9/15 – 9/30	12:31
10/1 – 10/14	11:44
10/15 – 10/31	11:14
11/1 – 11/14	10:29
11/15 – 11/30	9:57

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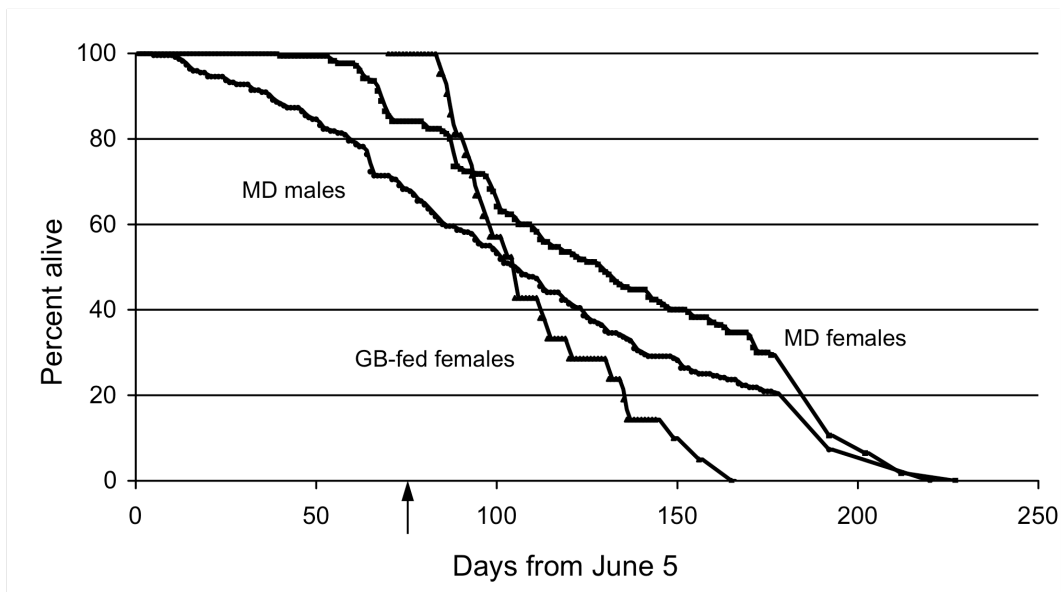
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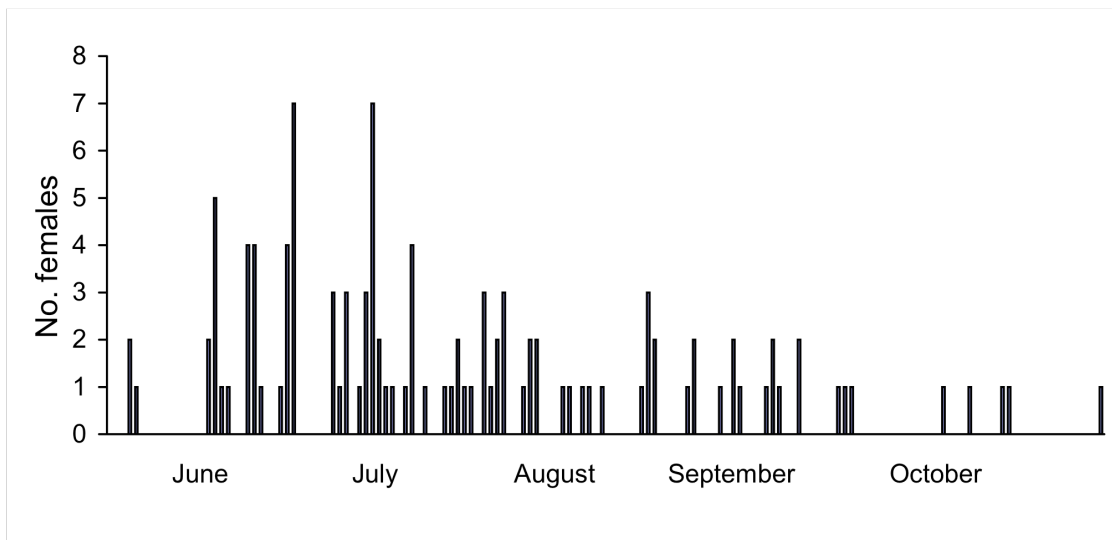
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1 Fig.1. Cumulative mortality of male and female *Hippodamia convergens*
 2 receiving a maintenance diet (MD-females and MD-males) comprising limited
 3 access to sub-optimal food (eggs of *Ephestia kuehniella* once every six days
 4 followed by pollen three days later, with continuous access to sunflower petioles).
 5 GB-fed females were switched to an *ad libitum* diet of greenbug, *Schizaphis*
 6 *graminum*, refreshed daily, on Aug 14 (arrow), prior to any oviposition.
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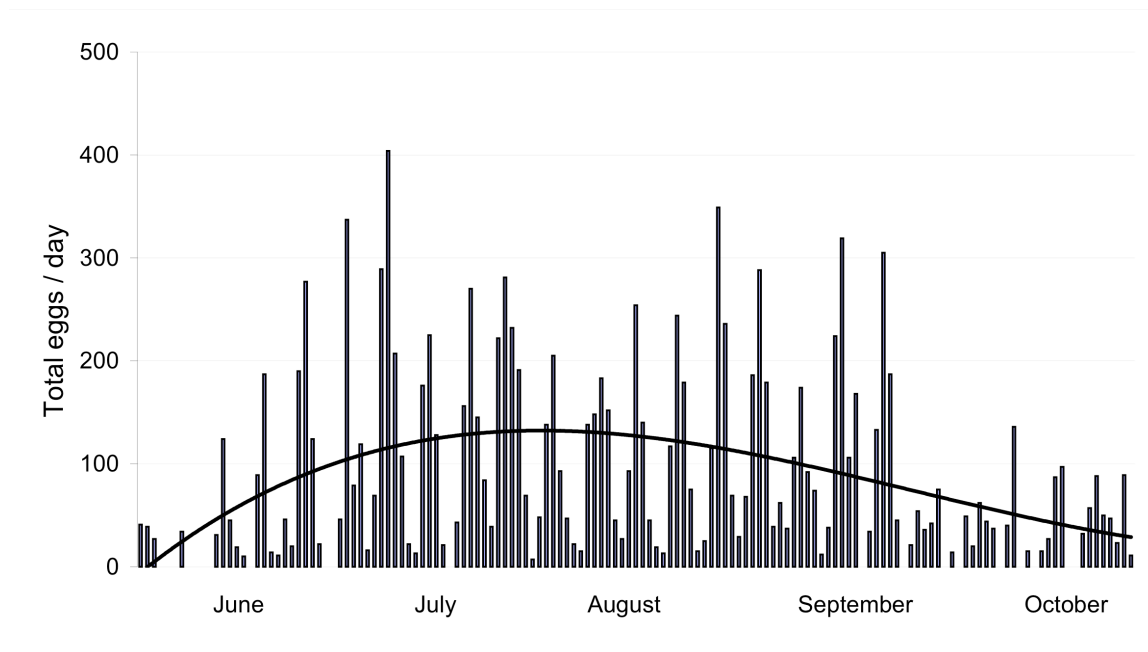
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1 Fig. 2. Numbers of females receiving a maintenance diet (eggs of *Ephestia*
2 *kuehniella* once every six days followed by pollen three days later) that initiated
3 oviposition on particular dates over the course the experiment (n = 113 females
4 total). The first date of oviposition was 9 June, three days after collection and the
5 last day was 3 November, 152 days after collection.



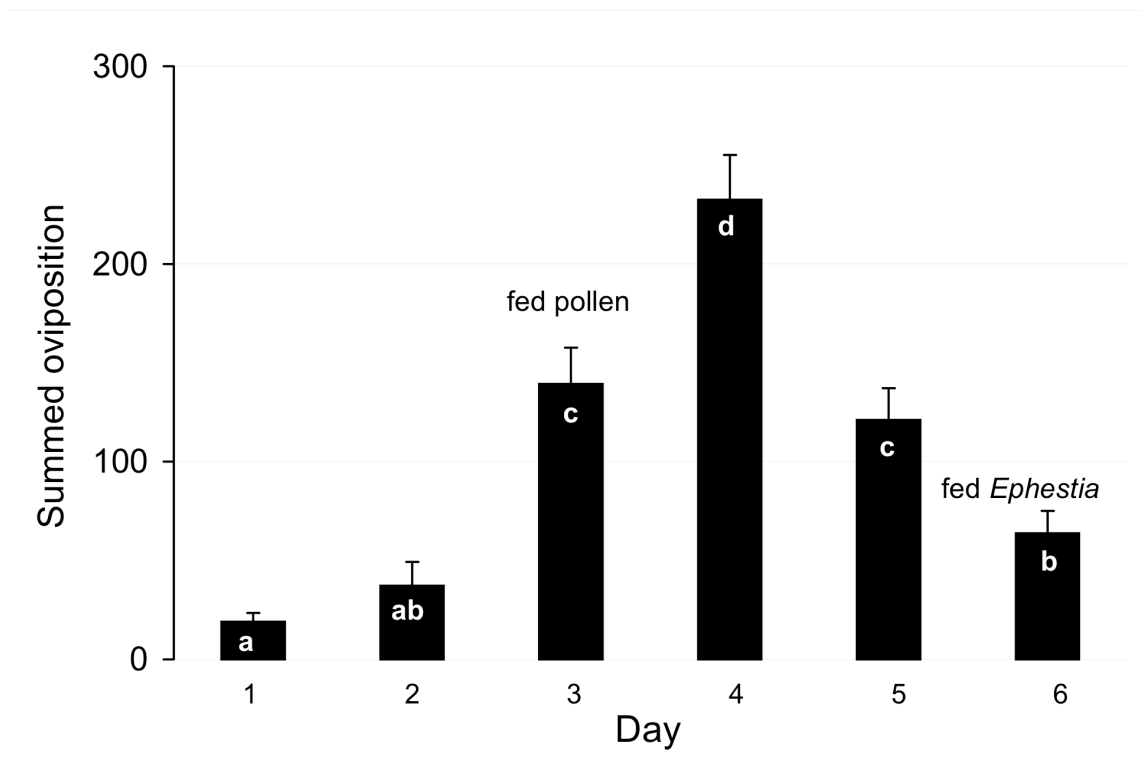
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1 Fig. 3. Temporal distribution of reproductive effort (total number eggs/day) by a
2 cohort of *Hippodamia convergens* females on a maintenance diet (eggs of
3 *Ephestia kuehniella* once every six days followed by pollen three days later) over
4 a period of five months. The data were generated by a total of 113 females with
5 variable longevity and onset of reproduction.



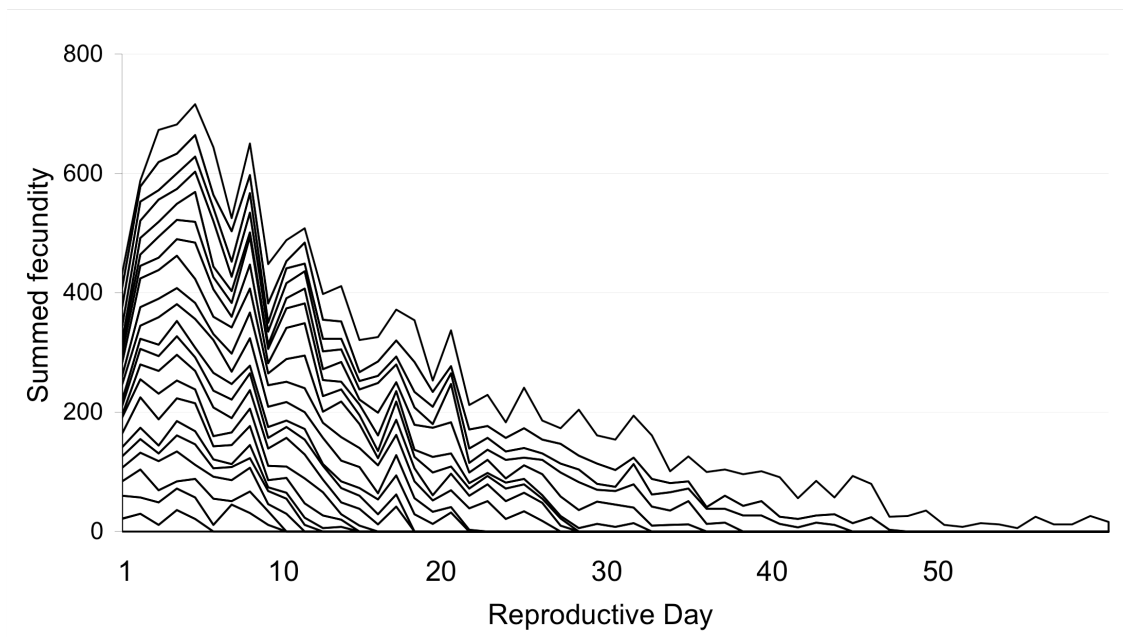
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1 **Fig. 4.** Summed egg production (Mean \pm SEM) by *Hippodamia convergens*
2 females (n = 113) over 135 days as a function of food provisioning showing a
3 peak in oviposition every fourth day following provision with animal protein.
4 Females paired with males were held on a six day feeding cycle (eggs of
5 *Ephestia kuehniella* once every six days followed by pollen three days later) with
6 continuous access to sunflower petioles. Means bearing the same letter were
7 not significantly different (Tukey test, $\alpha > 0.05$).



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- 1 Fig. 5. Stacked line graph of egg production by *Hippodamia convergens* females
- 2 ($n = 20$) that received an *ad libitum* diet of greenbug, *Schizaphis graminum*. Data
- 3 for various females are aligned according to their first date of reproduction and
- 4 summed on the y-axis in order of fecundity to demonstrate the triangular shape of
- 5 the female fecundity function under conditions of unlimited food supply.



6