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Article

Age and Density of Mated Females Affect Dispersal Strategies in a Spider Mite

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Simple Summary: The European native spider mite *Tetranychus ludeni* Zacher (Acari: Tetranychidae) is an invasive species which attacks many economically important crops. Investigation into how different factors affect its dispersal probability and distance helps better understand its population management and invasion evaluation. Here we tested the effect of age on dispersal probability and distance and subsequent reproduction and that of density on dispersal probability and distance. We show that older females with more eggs were more likely to disperse and moved longer distances than younger ones with fewer eggs. Older females spread most of their eggs out of the natal habitats and over longer distances, which reduced competition and increased offspring fitness. Our results also indicate that females significantly increased dispersal probability and distance with the increase of population density to avoid crowding. The synchronization of dispersal and reproduction, along with the positive density-dependent dispersal strategy, may facilitate habitat colonization and invasion speed of *T. ludeni*.

Abstract: Dispersal strategies of a species can affect its invasion success. Investigation into the dispersal strategies of invasive species in relation to different factors helps our understanding of the invasion mechanisms and provides knowledge for population management and invasion evaluation. *Tetranychus ludeni* Zacher (Acari: Tetranychidae) is an invasive species, which is native to Europe but now cosmopolitan. Here, we examined the effects of age and density on dispersal in mated females. Our results show that older females with more eggs were more likely to disperse and moved longer distances than younger ones with fewer eggs. Older females spread most of their eggs out of the natal habitats and over longer distances, which reduced competition and increased offspring fitness. Females significantly increased dispersal probability and distance with the increase of population density to avoid crowding. The synchronization of dispersal and reproduction, along with the positive density-dependent dispersal strategy, may facilitate habitat colonization and invasion speed of *T. ludeni*.

Keywords: age-specific reproduction; population density; resource competition; dispersal probability; dispersal distance

1. Introduction

Investigation into how different factors affect dispersal probability and distance helps better understand the dispersal process [1], knowledge of which is important for population management and invasion evaluation [2–6]. Population density and individual age are two important factors that can influence dispersal [1,3]. However, previous studies have mainly focused on how these factors modify dispersal probability [3,7–12]. To date, the effect of density and age on dispersal distance and subsequent reproduction is still poorly understood.

Many species exhibit age-specific dispersal due to the variance in reproductive costs or pressures with age [4]. Females at the age of high fecundity may have a higher dispersal probability and a longer dispersal distance to spread their eggs [13–16]. The availability of resources is the major factor affecting immature fitness, and thus mothers are expected to disperse and select habitats with sufficient resource supply or less resource competition, aiming to increase offspring fitness [1,11,17–21]. The intention of spreading offspring over habitats of high quality may be particularly strong for species whose immatures have limited dispersal ability and no parental care. Although many studies have investigated age-specific dispersal and reproduction, few examine the subsequent offspring competition intensity and fitness. Understanding age-specific dispersal strategies in relation to reproduction and offspring fitness is essential, because the reproductive output of dispersers at the age of dispersal can affect establishment success and population growth at new resident sites [22–26].

High population density often leads to intense resource competition, reducing individual fitness [3,27,28]. Increasing density would raise dispersal probability [2,3,9,29] over longer distance [30,31]. However, individuals living in high density may also have benefits such as reducing predation risk and increasing foraging efficiency [32], and thus individuals may tend to live in habitats with high density. It is predicted that density-dependent dispersal in animals depends on the balance of costs and benefits of group living, which warrants species-specific investigations.

The European native spider mite *Tetranychus ludeni* Zacher (Acari: Tetranychidae) has now invaded all continents except Antarctica [33,34]. It is a haplodiploid species where virgin females produce haploid sons and mated females produce haploid sons and diploid daughters [35,36]. *T. ludeni* attacks over 300 hosts, including many economically important crops such as beans, eggplant, hibiscus, apple, pumpkin, and other cucurbitaceous plants [33,37–39]. Spider mites have relatively short life cycle and high fecundity. They build up populations rapidly and exhaust resources very fast, and dispersal is common especially in overcrowding and food-depleted environments [2,30,40,41]. Previous research has demonstrated that high population density negatively affected individual reproduction and population growth of *T. ludeni* [42–44]. However, whether high density would promote the dispersal probability and distance of *T. ludeni* is unknown. Adult females of spider mites usually disperse after mating [45] through ballooning and walking (or aerial and ambulatory dispersal) [9,12]. Young females disperse more frequently than older females by ballooning [9]. However, the age-specific ambulatory dispersal is still not clear [12]. Moreover, mated females of *T. ludeni* gradually increase reproductive output until they are six days old [42]. So far, it is unknown whether age-specific ambulatory dispersal is related to reproduction.

In this study, we used *T. ludeni* to investigate the effect of age on dispersal probability and distance and subsequent reproduction and that of density on dispersal probability and distance. For the age experiment, we allowed females of different ages to disperse for 24 hours and then recorded their dispersal probability, dispersal distance, number of eggs laid, and the subsequent offspring survival. For the density experiment, we allowed females of different density to disperse for 24 hours and then recorded dispersal probability and distance. We hypothesized that (1) females at the age of high fecundity would have higher dispersal probability and longer dispersal distance and produce eggs over longer distances, and (2) high density would increase dispersal probability and distance.

2. Materials and Methods

2.1. Mite Colony and Experimental Conditions

We collected *T. ludeni* adults on *Passiflora mollissima* Kunth (Malpighiales: Passifloraceae) in Palmerston North, New Zealand and used 3- to 5-week-old common bean *Phaseolus vulgaris* L. (Fabales: Fabaceae) plants grown in pots to maintain the colony. We used leaf squares cut from the first expanded leaves of 1- to 2-week-old bean plants grown in pots for all experiments. Environmental conditions for the colony and experiments were set at 25 ± 1 °C, $40 \pm 10\%$ RH, and 14:10 h (L:D) photoperiod.

2.2. Preparation of Mated Females for Experiments

To obtain virgin males to inseminate females, we randomly selected 50 female deutonymphs from the colony and transferred them onto a clean fresh leaf square (4 × 4 cm) on wet cotton wool in a Petri dish (9 cm in diameter × 1 cm in height). We allowed these deutonymphs to develop into virgin adult females and lay unfertilised eggs for three days. We then removed adult females and allowed the eggs to develop to virgin male adults. We prepared a total of 46 Petri dishes for this purpose. We used 1-day-old virgin adult males to mate with females prepared below.

We randomly selected 50 adult females from the colony and transferred them onto a clean fresh leaf square (4 × 4 cm) on wet cotton wool in a Petri dish for egg laying for 24 hours. We then removed the female adults and allowed eggs to develop to the quiescent deutonymphal stage. We set up approximately 230 Petri dishes for this purpose. We transferred 15 ~ 20 virgin males prepared above to female quiescent deutonymphs in each Petri dish, allowing them to stay with the newly emerged females for 5 hours to ensure that all females mated at or soon after emergence (females mate immediately after emergence [35]). We transferred 50 newly emerged (< 24 hours old) and mated females from an above Petri dish onto a new leaf square (4 × 4 cm) on wet cotton wool in a Petri dish. We prepared 230 such dishes and replaced all leaf squares once a day to obtain mated females of 1, 3, 6, 9, and 12 days old for experiments. To keep female number on each leaf square consistent prior to experiments, we checked the leaf square every day and replaced dead females with live ones of the same age.

2.3. Dispersal and Reproduction of Females of Different Ages

We established a dispersal arena consisting of 21 leaf squares (2 cm × 2 cm for the first leaf square and 2 cm × 1 cm for others) placed on wet cotton in a tray (45 cm length × 36 cm width × 1.5 cm height) (Figure 1). We performed 17, 14, 14, 14, and 16 replicates for treatments of 1, 3, 6, 9, and 12 days old, respectively. For each replicate, we transferred 50 mated females of a desired age onto the first leaf square (primary leaf square) and allowed them to settle for 40 minutes. We then connected all leaf squares using Parafilm bridges (4.5 cm length × 1.5 cm width; Parafilm®, USA) and allowed mites to disperse freely for 24 hours, after which time, we removed all bridges, counted the number of adults on each leaf square, and removed all adults and counted the number of eggs on each leaf square. All leaf squares with eggs were individually transferred onto a wet cotton pad in a Petri dish, and the eggs were reared on their original leaf squares until adulthood.



Figure 1. Dispersal arena for experiments. The primary leaf square was used to introduce experimental mites. Leaf squares were connected by Parafilm bridges with approximately 1 mm overlapping to reduce the influence of water between leaf squares and bridges on mite dispersal.

We defined the dispersal probability as the percentage of mites that dispersed out of the primary leaf square. We used the number of leaf squares from the primary leaf square as dispersal distance and regarded the primary leaf square as zero distance. We recorded the mean dispersal distance of a

replicate by averaging the dispersal distance of all individuals in the replicate and the total number of eggs laid in a replicate by summing up the number of eggs on all leaf squares. We recorded the mean distance of eggs of a replicate by averaging the distance of all eggs laid on different leaf squares.

2.4. Dispersal of Females at Different Densities

Because spider mite density in nature ranges from 0.1 to 50 individuals/cm² [46,47], we set up four densities: 2.5, 12.5, 25.0 and 37.5 females/cm², with 10, 50, 100 and 150 mated females on the primary leaf square, respectively. There were 15, 15, 16, and 16 replicates, respectively, for above density treatments. For each replicate, we transferred 1-d-old mated females of a desired number according to treatments on the primary leaf square and allowed them to settle for 40 minutes. We then connected all leaf squares as above and allowed mites to disperse freely for 24 hours, after which time, we removed all bridges and counted the number of adults on each leaf square. We calculated dispersal probability and distance of a replicate as described above.

2.5. Statistical Analysis

We analyzed all data using SAS software (SAS 9.4, SAS Institute Inc., Cary, NC). Data on dispersal probability in age and density experiments, dispersal distance in age experiment, total number of eggs laid, and number of eggs laid outside the primary leaf squares were normally distributed (Shapiro-Wilk test, UNIVARIATE Procedure), and thus analyzed using ANOVA with Tukey's test for multiple comparison (GLM procedure). Data on distance of eggs, number of eggs laid on the primary leaf square, and dispersal distance in density experiment were square-root transformed to achieve normal distribution before ANOVA. The survival of offspring on and outside the primary leaf squares were not normally distributed even after transformation and analyzed using a non-parametric ANOVA (GLM procedure).

We used a logistic linear model, $y = \exp(a + bx)$, to determine the distribution of egg density (number of eggs/leaf square area) and changes of offspring survival rate over distance with a Gamma distribution and a log link function in all age treatments, where y is the response variable, x is distance, and a and b are constant parameters of the model (GLIMMIX Procedure).

3. Results

3.1. Dispersal and Reproduction of Females of Different Ages

Our results show that within 24-h experimental period, 1-d-old females were significantly less likely to disperse compared to the older females ($F_{4, 70} = 15.51$, $p < 0.0001$) (Figure 2a), and they dispersed significantly shorter distance than older females ($F_{4, 70} = 7.55$, $p < 0.0001$) (Figure 2b). Furthermore, 1-d-old females laid significantly fewer eggs than older females and 3-d-old females laid significantly more eggs than females of other ages ($F_{4, 70} = 25.73$, $p < 0.0001$) (Figure 2c).

Egg density significantly decreased over distance in all age treatments (Figure 3a; Table S1). Females of different ages laid similar number of eggs on the primary leaf squares ($F_{4, 70} = 1.62$, $p = 0.1797$) (Figure 4a), but older females laid significantly more eggs outside the primary leaf squares than 1-d-old females ($F_{4, 70} = 22.59$, $p < 0.0001$) (Figure 4b). In addition, older females produced eggs over significantly longer distance than 1-d-old females ($F_{4, 70} = 5.35$, $p = 0.0008$) (Figure 4c). Immature survival rate significantly increased over distance in all treatments (Figure 3b; Table S1). Offspring survival on or outside the primary leaf square were similar between females of different ages ($F_{4, 70} = 1.16$, $p = 0.3339$ for primary leaf square; $F_{4, 69} = 1.80$, $p = 0.1379$ for outside the primary leaf square) (Figure 4d,e).

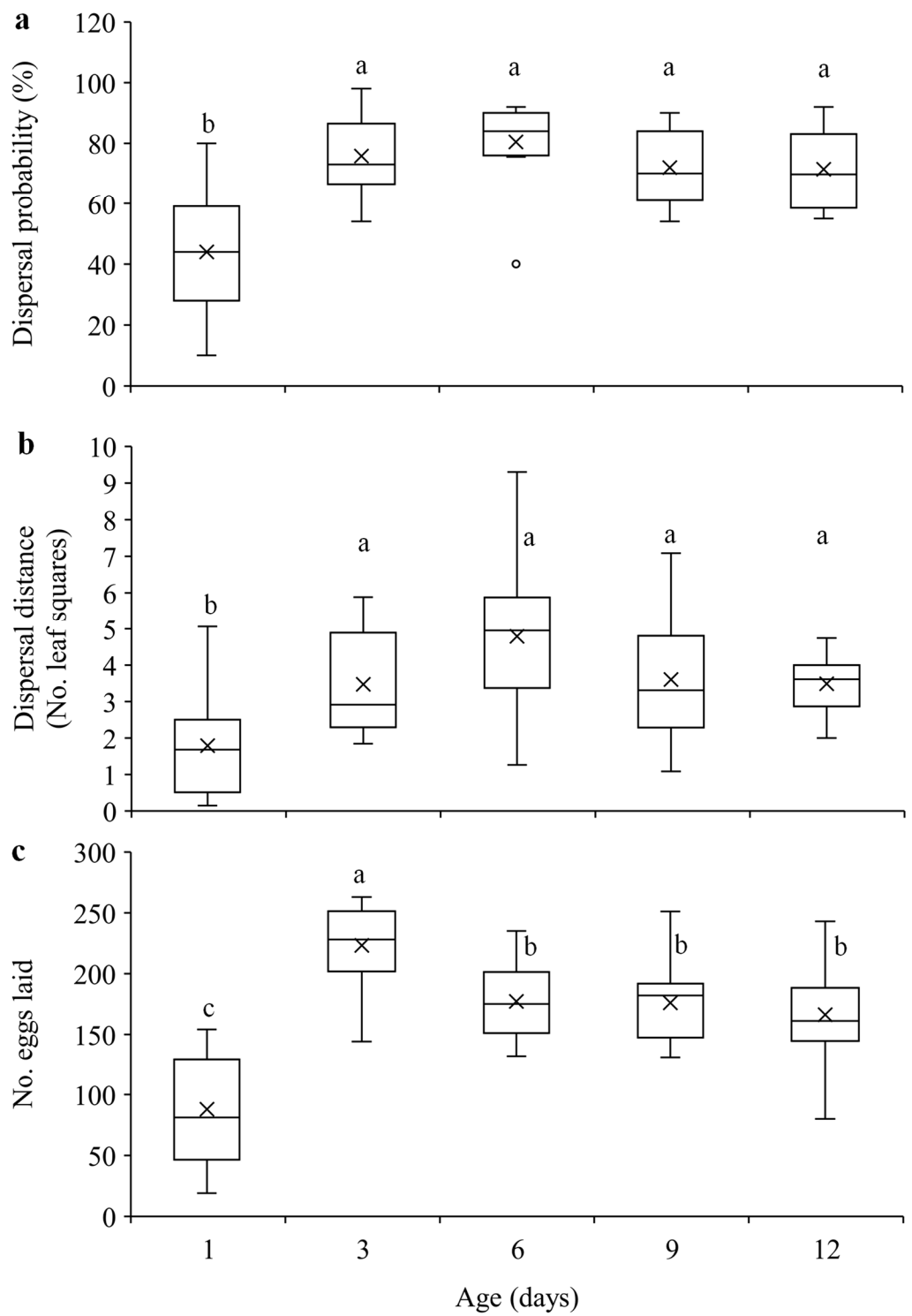


Figure 2. Mean dispersal probability (a), dispersal distance (b), and total number of eggs laid (c) within 24 hours in *T. ludeni* females of different ages. Each box plot shows the maximum (T) and minimum (L) scores, the range of upper and lower quartiles (the box), and the mean (x) and median (line in the box) scores; the circle is an outlier of minimum score. For each parameter, boxes with the same letters are not significantly different ($p > 0.05$).

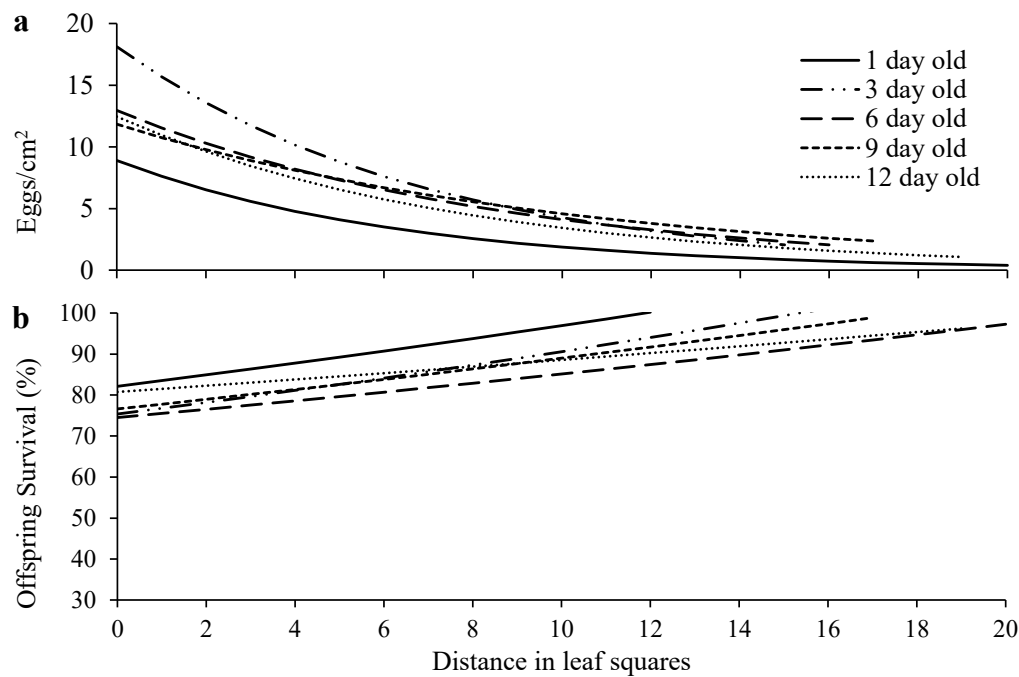


Figure 3. Egg density (a) and offspring survival (b) over distance in *T. ludeni* females of different ages. Raw data were subjected to analysis for all parameters.

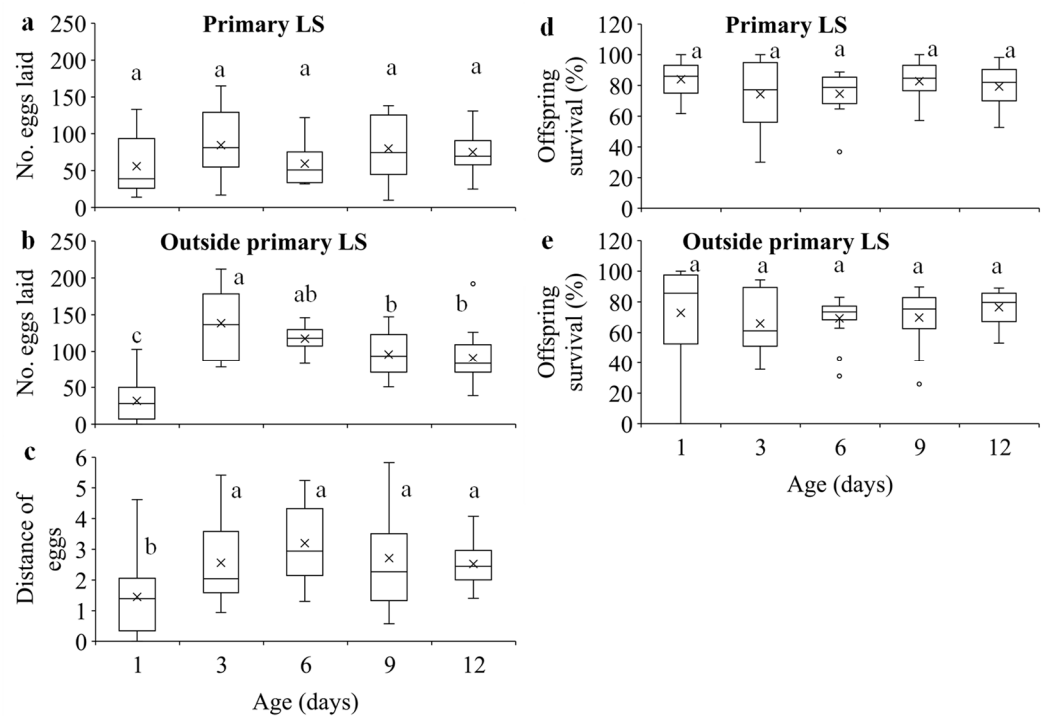


Figure 4. Number of eggs laid on (a) and outside (b) the primary leaf squares (LS), distance of eggs (c) within 24 hours, and offspring survival on (d) and outside (e) the primary leaf squares in *T. ludeni* females of different ages. Each box plot shows the maximum (T) and minimum (L) scores, the range of upper and lower quartiles (the box), and the mean (x) and median (line in the box) scores; the circles are outliers of minimum or maximum scores. For each parameter, boxes with the same letters are not significantly different ($p > 0.05$).

3.2. Dispersal of Females at Different Densities

Mites were significantly more likely to disperse with the increase of population density ($F_{3, 58} = 15.33$, $p < 0.0001$) (Figure 5a). Similarly, they dispersed significantly longer distance within 24 hours when population density was higher ($F_{3, 58} = 14.06$, $p < 0.0001$) (Figure 5b).

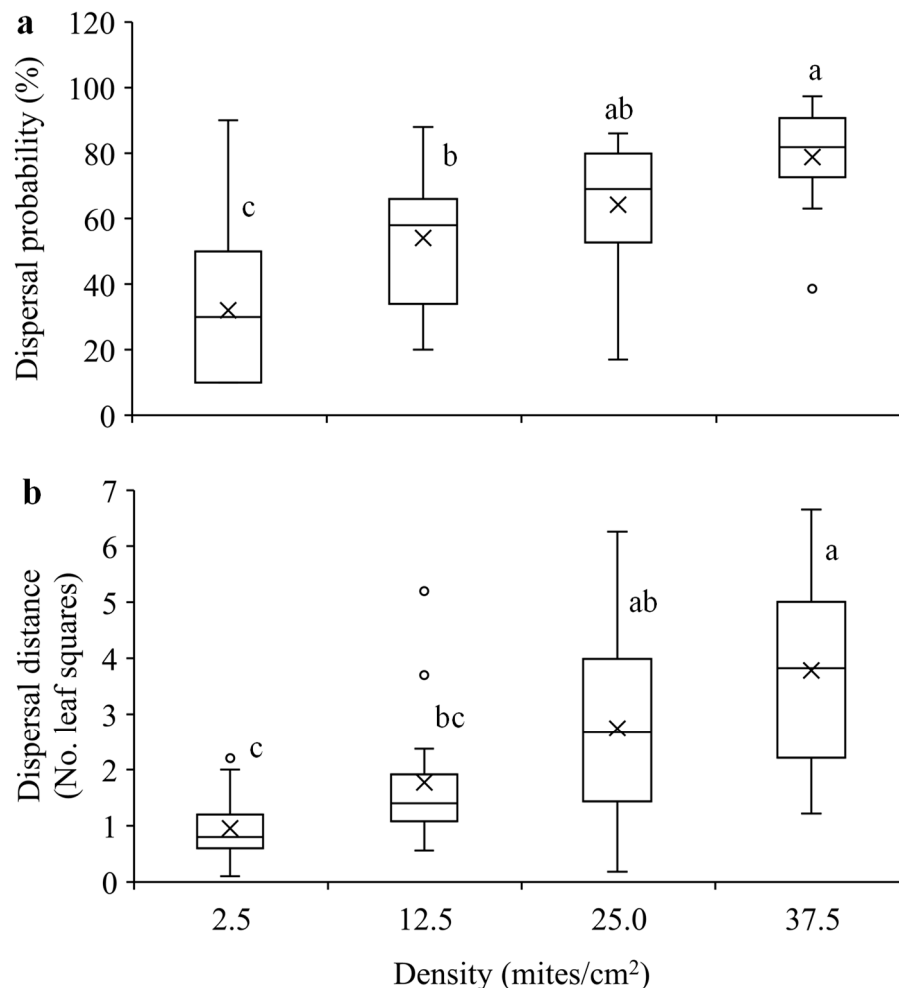


Figure 5. Mean dispersal probability (a) and dispersal distance (b) within 24 hours in *T. ludeni* females of different densities. Each box plot shows the maximum (T) and minimum (L) scores, the range of upper and lower quartiles (the box), and the mean (×) and median (line in the box) scores; the circles are outliers of minimum or maximum scores. For each parameter, boxes with the same letters are not significantly different ($p > 0.05$).

4. Discussion

Our results show that compared to 1-d-old females, older ones had significantly higher dispersal probability and longer dispersal distance and laid more eggs in *T. ludeni* (Figure 2). These findings strongly suggest that the age-dependent dispersal is caused by reproductive readiness when older females carry more mature eggs [42,48]. In general, females with more eggs tend to disperse in wider areas [1]. We also show that 3-d-old females laid the highest number of eggs under our experimental setting (Figure 2c) where the population density is relatively high [46,47]. This may be attributed to a resource allocation strategy of *T. ludeni* to maximizing reproduction during the early lifespan at high population densities [42]. The synchronization of dispersal and reproduction has also been reported in many insects such as the light brown apple moth, *Epiphyas postvittana* (Walker) (Lepidoptera: Tortricidae) [14], the tarnished plant bug, *Lygus lineolaris* (Palisot de Beauvois)

(Hemiptera: Miridae) [25,49], the rice bug, *Leptocorisa chinensis* Dallas (Hemiptera: Alydidae) [15], and the butterfly, *Maculinea (Phengaris) teleius* Bergsträsser (Lepidoptera: Lycaenidae) [11].

Higher dispersal probability and longer dispersal distance of females laden with more eggs may contribute to fast population growth at new sites, promoting pest outbreak and range expansion [22,24,49]. Furthermore, the similarly high dispersal probability and long dispersal distance among older females (3–12 days old) (Figure 2) suggest that *T. ludeni* females can quickly disperse and infest new host plants in most of their adult lives. In their study on dispersal behaviour of *T. urticae*, Li and Margolies reveal that the aerial dispersal is much more frequent in younger females because they are smaller in body size and may be more easily carried aloft by air currents [9]. We provide the first evidence that females of ≥ 3 days old carrying more eggs are more likely to disperse through walking.

Dispersal serves multiple functions such as avoiding harsh conditions and reducing competition for resources between their own [50,51] and their offspring [1,11,17–20,52]. We show that although older females had higher fecundity (Figure 2c), they laid similar number of eggs on primary leaf squares (Figure 3a) but produced significantly more eggs outside the primary leaf squares over longer distances compared to the 1-d-old ones (Figure 3b,c). The results suggest that older females of *T. ludeni* can reduce offspring competition for food resources by dispersing from their natal habitats to the new habitats (Figure 2a), and the spread of local population and increase of population size of spider mites within and between the adjacent habitats are through the females with high reproductive output.

We show that with the decrease of egg density over dispersal distance, offspring survival significantly increased (Figure 4), implying that crowding and competition negatively affect offspring fitness and rich resources available at their birthplace increases their fitness. This strategy is critical to offspring survival in spider mites because immatures have limited dispersal ability [9,45,53] and no parental care. Similar strategy has also been reported in the predatory mites, *Phytoseiulus persimilis* Athias-Henriot and *Neoseiulus californicus* McGregor (Acari: Phytoseiidae). Mothers of these two species produce eggs on a patch according to the number of prey eggs on the patch that are sufficient for their offspring to survive/develop before dispersing away [54,55].

The present study indicates that *T. ludeni* females significantly increased dispersal probability and distance with the increase of population density (Figure 5). This could be attributed to the fact that high population density reduces individual and population fitness [3,42–44,56,57]. In *T. ludeni*, high population density negatively affected individual reproduction and population growth [42–44]. Our results suggest that females would avoid overcrowding condition to reduce competition by dispersal with a longer distance. The density-dependent dispersal strategy has been reported in many animal species such as the butterflies *Maculinea nausithous* and *M. teleius* (Lepidoptera: Lycaenidae) [19], *T. urticae* [9,30], and the brown garden snail *Cornu aspersum* (Müller) (Gastropoda: Helicidae) [58]. This strategy allows individuals to leave high density habitats and emigrate into low density habitats and/or colonize new habitats [3,6,8,59], promoting the invasion speed of *T. ludeni*.

In conclusion, increasing age and density promote dispersal probability and distance in *T. ludeni*. Females with more mature eggs are more likely to disperse and move longer distances. These dispersal strategies spread eggs out of their natal habitats and over longer distances, which reduces competition and increases their own and offspring fitness. These strategies may facilitate pest outbreak and invasion success of *T. ludeni*.

Supplementary Materials: The following supporting information can be downloaded at the website of this paper posted on Preprints.org, Table S1: Statistical results of models of the egg density and offspring survival over distance in *T. ludeni* females of different ages in Figure 3.

Author Contributions: Conceptualization, P.Z., Q.W. and X.Z.H.; experimental design, P.Z., Q.W. and X.Z.H.; data collection, P.Z. and C.C.; data analysis, P.Z. and X.Z.H.; manuscript preparation and revision, P.Z., Q.W. and X.Z.H. All authors have read and agreed to the published version of the manuscript.

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Data Availability Statement: The raw data supporting the conclusions of this article will be made available by the authors on request.

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Conflicts of Interest: The authors declare no conflict of interest.

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