

Effects of blood meal source and seasonality on reproductive traits of *Culex quinquefasciatus* (Diptera: Culicidae)

Reviewed Preprint

Revised by authors after peer review.

[About eLife's process](#)

Reviewed preprint version 2
February 28, 2024 (this version)

Reviewed preprint version 1
September 15, 2023

Posted to preprint server
September 8, 2023

Sent for peer review
June 14, 2023

Kevin Alen Rucci , Mariana Pueta, Adrián Díaz 

Laboratorio de Arbovirus, Instituto de Virología “Dr. J. M. Vanella” (InViV), Facultad de Ciencias Médicas (FCM), Universidad Nacional de Córdoba (UNC) • Consejo Nacional de Investigaciones Científicas y Técnicas (CONICET), Argentina • Instituto de Investigaciones en Biodiversidad y Medioambiente (INIBIOMA), CONICET - Universidad Nacional de Comahue (UNCo)

 https://en.wikipedia.org/wiki/Open_access

 Copyright information

Abstract

Host selection by mosquitoes is a keystone to understanding viral circulation and predicting future infection outbreaks. *Culex* mosquitoes frequently feed on birds during spring and early summer, shifting into mammals towards late summer and autumn. This host switch may be due to changes in mosquito fitness. The aim of this study was to assess if the interaction effect of blood meal source and seasonality may influence reproductive traits of *Culex quinquefasciatus* mosquitoes. For this purpose, *Cx. quinquefasciatus* mosquitoes were reared in simulated summer and autumn conditions and fed on two different hosts, chickens and mice, in a factorial design. Fecundity, fertility, and hatchability during two consecutive gonotrophic cycles were estimated. We found greater fecundity and fertility for mosquitoes fed upon birds than mammals. Fecundity and fertility increased in autumn for chicken-fed mosquitoes, whereas they decreased for mouse-fed mosquitoes. These traits decreased in the second gonotrophic cycle for mouse-fed mosquitoes, whereas they did not vary between cycles for chicken-fed mosquitoes. The effect of hatchability on fertility was rather limited. These results indicate a statistically significant interaction effect of blood meal source and seasonality on fecundity and fertility. However, the pattern was opposite in relation to our hypothesis, suggesting that further studies are needed to confirm and expand our knowledge about mosquito biology and its relationship with seasonal host use shifting.

eLife assessment

This **useful** study provides the first assessment of the potentially interactive effects of seasonality and blood source on mosquito fitness, together in one study. However, the experimental approach is **incomplete** because it is limited without replication of the experiments and because of the small sample sizes for some groups. The work will be of interest to those studying mosquito biology.

Introduction

The southern house mosquito, *Culex quinquefasciatus* Say (Diptera: Culicidae), is a worldwide distributed vector of several pathogens, such as *Wuchereria bancrofti* Cobbold (Kasili et al., 2009 [\[1\]](#)), *Dirofilaria immitis* Leidy (Labarthe et al., 1998 [\[2\]](#)) and avian malaria (Farajollahi et al., 2011 [\[3\]](#)). Besides, this mosquito is also one of the main species responsible for the transmission of the arboviruses West Nile virus (WNV) and Saint Louis encephalitis virus (SLEV), among others (Farajollahi et al., 2011 [\[3\]](#)). Because of this, this species is of critical concern due to its impact on both public and veterinary health.

Since arboviruses are transmitted by the bite of infected mosquitoes between vertebrate hosts, activity of these pathogens is determined, at least in part, by the blood feeding habits of these vectors. Host preference of *Culex* mosquitoes is variable and it can be classified in four categories: (1) mammophilic (Muturi et al., 2008 [\[4\]](#)), (2) ornithophilic (Jansen et al., 2009 [\[5\]](#)), (3) herpetophilic (Janssen et al., 2015 [\[6\]](#)) and (4) generalist (Molaei et al., 2007 [\[7\]](#)). Additionally, some species may experience a host switch in its feeding habits, from avian to mammal hosts (Hancock & Camp, 2022 [\[8\]](#)). These species are of greater concern from an epidemiological standpoint because they can act as bridge vectors in the transmission of some arboviruses, such as SLEV and WNV (Kilpatrick et al., 2005 [\[9\]](#)).

Culex quinquefasciatus is generally considered an ornithophilic species in several areas around the globe (Takken & Verhulst, 2013 [\[10\]](#)). Despite its ornithophilic nature, this and other *Culex* species sometimes feed primarily on mammal hosts, including humans, in certain situations. Various studies have pointed out that in autumn there is an increased activity of SLEV and WNV in human populations (Kilpatrick et al., 2006 [\[9\]](#); Spinsanti et al., 2008 [\[11\]](#)), suggesting that viral activity spills over to mammals from birds as a result of the mosquito host switch (Edman & Taylor, 1968 [\[12\]](#); Kilpatrick et al., 2006 [\[9\]](#)). A few hypotheses have attempted to explain why primarily ornithophilic mosquitoes also feed upon mammals in some contexts. One of the most widely accepted hypothesis states that the shift on feeding behaviour of *Culex* mosquitoes is a consequence of the autumn migration of American robin (*Turdus migratorius*), the principal host in the northeastern US, generating an opportunistic change into mammal hosts (Kilpatrick et al., 2006 [\[9\]](#)). In South America in general and in Argentina in particular, studies regarding host switch are scarce. Nevertheless, a few limited studies have indicated a seasonal variation throughout the seasons (Stein et al., 2013 [\[13\]](#); Beranek, 2018 [\[14\]](#)). Despite that, Argentina has acknowledged seasonal variation in the activity of SLEV in humans, which can be attributed to variations in mosquito host-feeding pattern (Spinsanti et al., 2008 [\[11\]](#)). In Southern latitudes, species such as *T. migratorius* are lacking, hence the “migration hypothesis” is very unlikely for many austral areas, being restricted only for some regions of the US.

A second hypothesis proposed by Burkett-Cadena et al. (2011 [\[15\]](#)) suggests that host breeding cycles drive the host shift of mosquitoes. According to this hypothesis, in hosts with parental care, during reproductive seasons there is a greater investment of energy in assuring the offspring survivorship, consequently producing an increase in susceptibility of being bitten by mosquitoes as a result of a decrease in defensive behaviours. This event leads to the detection of peaks of host use during periods of reproductive investment, in summer for birds and autumn for mammals (Burkett-Cadena et al., 2011 [\[15\]](#)).

While some other minor hypotheses, unsupported by field data, have been proposed, none of them have been specifically focused on vector biology. Numerous biological factors, including stress, metabolic rate, and the source of a blood meal, as well as environmental variables such as temperature and photoperiod, exert an influence on mosquito physiology that might affect various reproductive traits such as fecundity, development rate or survivorship. This set of factors gives

rise to new nutritional requirements that, ultimately, may lead to seasonal variation in host choice by mosquitoes (Ciota et al., 2014 [↗](#); Costanzo et al., 2015 [↗](#); Gervasi et al., 2016 [↗](#); Yan et al., 2017 [↗](#), 2018 [↗](#)). In the literature, several studies have evaluated the impact of these biological and environmental variables on mosquito reproduction, as mentioned before. However, due to the complex nature of mosquito biology, there are likely multiple interactions among variables that give rise to new responses that may not be observed individually. Therefore, the aim of the present study was to assess whether exists an interaction effect between the source of the host blood meal and seasonality (in terms of temperature and photoperiod) on three reproductive traits of *Culex quinquefasciatus* mosquitoes, fecundity, fertility, and hatchability. Our hypothesis states that the interaction between these two variables influence the reproductive outcomes, potentially leading to a seasonal shift in host selection, driven by a reproductive advantage. Considering the reported host preference changes in *Cx. quinquefasciatus*, in autumn we expect a greater number of eggs (fecundity) and larvae (fertility) in mosquitoes after feeding on a mammal host compared to an avian host, and the opposite relationship in summer.

Materials and methods

Establishment and maintenance of mosquitoes

Egg rafts of *Culex quinquefasciatus* were collected from a drainage ditch at Universidad Nacional de Córdoba Campus, Córdoba city, in February 2021. Each raft was individually maintained in plastic containers with one liter of distilled water. The hatched larvae were fed with 100 mg of liver powder three times per week until pupation. Pupae were then transferred to plastic emerging cages (21 cm x 12 cm) covered with a tulle-like fabric, containing distilled water but no food. Adults emerging from each raft were identified morphologically (Darsie, 1985 [↗](#)) and molecularly (Smith & Fonseca, 2004 [↗](#)) to ensure they corresponded to *Cx. quinquefasciatus*, since *Cx. pipiens* and its hybrids are also present in Córdoba (Branda et al., 2021 [↗](#)). All adults were reared in a cardboard cage of 22.5 liters (28 cm x 36.5 cm) and provided *ad libitum* with a 10% sugar solution soaked in cotton pads placed on plastic cups. For long-term maintenance of the colony, 24-hour-starved mosquitoes were offered a blood meal from a restrained chicken twice a month. Four days after feeding, a plastic container with distilled water was placed inside the cage to allow engorged females to lay egg rafts. Batches of egg were collected and transferred to plastic containers (30 cm x 25 cm x 7 cm) in a proportion of 3 rafts per container, filled with three liters of distilled water. The hatched larvae were also fed with liver powder at the same proportion described above. Pupae were transferred to emerging cages, and adult mosquitoes were placed in the final cardboard cage. The colony has been maintained for over twenty generations in the Insectary of the Instituto de Virología “Dr. J.M. Vanella” (InViV). Room conditions were controlled at 28°C, with a 12L:12D photoperiod and 70% relative humidity.

Experimental design

Blood source

For experimental trials, avian hosts (live chicks of the species *Gallus gallus*) and mammalian hosts (live mice of the species *Mus musculus*, strain C57BL/6) were used to evaluate the effect of blood meal source on fecundity, fertility, and hatchability. Chicks were generously donated by the Bartolucci poultry farm (Córdoba, Argentina), while mice were commercially obtained from the Instituto de Investigación Médica Mercedes y Martín Ferreyra (CONICET - Universidad Nacional de Córdoba). In each trial, 24-hour-starved adult female mosquitoes were provided with a blood meal from restrained chicks or mice. Vertebrate hosts were offered to mosquitoes one hour before the lights were turned off and were kept for 3 hours during 2 consecutive gonotrophic cycles.

The experimental use of animals (*G. gallus* and *M. musculus*) was approved by the ethical committee at Facultad de Ciencias Médicas, Universidad Nacional de Córdoba (FCM-UNC) in compliance with the legislation regarding the use of animals for experimental and other scientific purposes (accession code: CE-2022-00518476-UNC-SCT#FCM).

Seasonality

To assess the effect of seasonality (photoperiod + temperature) on fecundity, fertility, and hatchability, a “typical” summer and autumn day from Córdoba city was simulated in an incubator, where mosquitoes were housed throughout the entire experiment. Summer conditions were as follows: $T^{\circ}_{\min} = 22^{\circ}\text{C}$, $T^{\circ}_{\max} = 28^{\circ}\text{C}$, photoperiod = 14L:10D. Autumn conditions were characterized by: $T^{\circ}_{\min} = 16^{\circ}\text{C}$, $T^{\circ}_{\max} = 22^{\circ}\text{C}$, and photoperiod = 10L:14D. The humidity levels were maintained at 60-70% for both conditions. Data for simulated conditions were obtained from Climate Data website (<https://es.climate-data.org/> .

Feeding trial design

The interaction effect of blood source (bird or mammal) and seasonality (autumn and summer) was evaluated during two consecutive gonotrophic cycles (I and II). Eight experimental colonies were established using egg rafts collected from the maintenance colony: mammal-autumn-I (MA-I), mammal-summer-I (MS-I), bird-autumn-I (BA-I), bird-summer-I (BS-I), mammal-autumn-II (MA-II), mammal-summer-II (MS-II), bird-autumn-II (BA-II), and bird-summer-II (BS-II).

The experimental colonies were maintained under controlled conditions to ensure that all adults were of the same size ([Kauffman et al., 2017](#) ). Adult mosquitoes were placed in cardboard cages of 8.3 liters (21 cm x 24 cm) and were provided with *ad libitum* access to a 10% sugar solution, which was soaked in cotton pads placed on plastic cups.

Feeding trials were conducted at two time points: 5 days post-emergence (first cycle) and 14 days post-emergence (second cycle). Following each blood meal, female mosquitoes were anesthetized using CO_2 and classified as fully engorged (VI Sella’s stage), partially fed (I-V Sella’s stage), or unfed (I Sella’s stage), following the classification by Silva Santos et al. (2019) . Fully engorged females were counted and separated into a separate cage to complete their gonotrophic cycle. Four days after feeding, a cup containing distilled water was placed inside the cage to facilitate oviposition. After each oviposition cycle, egg rafts were collected, counted, and transferred to a 12-well plate containing 3 mL of distilled water. They were then photographed to subsequently determine the number of eggs per raft. Rafts were maintained in each well until they hatched into L1 larvae, at which point they were preserved with 1 mL of 96% ethanol, and the number of L1 larvae per raft was counted.

Statistical analysis

The reproductive outputs measured for each experimental colony were fecundity, fertility, and hatchability. All analyses were performed using R Studio statistic software, v.4.2.1 ([R Core Team, 2022](#) .

Fecundity was defined as the number of eggs per raft and *fertility* as the number of L1 larvae hatched per raft. To evaluate the effect of blood meal source and seasonality on these two variables, a Generalized Linear Model (GLM; package MASS) with negative binomial error distribution and logarithmic link function was adjusted ([Venables & Ripley, 2002](#) ). Fecundity and fertility served as the response variables, while the explanatory variables included blood source, seasonality, and gonotrophic cycle, all in a three-way interaction.

Hatchability (= hatching rate) was defined as the ratio of the number of larvae to the number of eggs per raft. A GLM was applied using a quasipoisson error distribution and a logarithmic link function. The response variable was the number of larvae, while the explanatory variables included blood source, seasonality and gonotrophic cycle as interaction effects. Additionally, the logarithm of the number of eggs was incorporated as an offset in the model.

Multiple comparisons among treatments were conducted using the Tukey's honestly significant difference test (HSD Tukey), incorporating the [Kramer \(1956\)](#) [correction](#) for unbalanced data, also known as the Tukey-Kramer method. This approach allows to set the family-wise error rate to 0.05. These contrasts were performed using the *multcomp* package ([Hothorn et al., 2008](#) [correction](#)).

Model validation was assessed using a simulation-based approach to generate randomized quantile residuals to test goodness of fit (Q-Q plot), homoscedasticity, and outliers and influential points (Cook's distance and Leverage plots). These analyses were conducted with the DHARMA package ([Hartig, 2022](#) [correction](#)).

Model accuracy via simulations for the fecundity and fertility models

To assess the accuracy of the results, a simulation approach was conducted using a Monte Carlo algorithm to simulate data and model parameters according to the methodology proposed by [Carsey & Harden \(2013\)](#) [correction](#). For this purpose, parameters including coefficients, treatment means and dispersion parameter (referred to as theta, θ), were extracted from the fecundity and fertility models. Initially, a new dataset consisting of 1000 values (sample size of 125 per treatment) was generated. This was achieved using the *rnbinom* function from the MASS package, specifying the treatment means (computed from model coefficients) and the theta parameter. Subsequently, a negative binomial GLM was fitted to the simulated data, and the expected means for each of the eight treatments were obtained from the model. This simulation process was repeated 1000 times. The ratio between the number of iterations in which the simulated means fell within the observed 95% confidence interval divided by the total number of simulations was computed as an approximation of the robustness of results.

Results

Effect of blood source and seasonality on fecundity and fertility

Two interaction effects were found to be statistically significant in both the fecundity and fertility models (**Tables 1** [correction](#), **4** [correction](#)). The first interaction effect was found between blood source and seasonality (fecundity: LRT $X^2 = 25.0$, $p = 7.72 \times 10^{-7}$; fertility: LRT $X^2 = 13.82$, $p = 0.002$), meaning that the effect of blood meal source on the number of eggs and larvae per raft varied with season (**Tables 2** [correction](#), **3** [correction](#)). In general, mosquitoes that fed on avian blood sources exhibited greater fecundity and fertility than those that fed on mammalian blood source, regardless of the season. When treatments were examined by pairwise comparisons, fecundity and fertility were 13% and 18% greater in autumn compared to summer for bird-fed mosquitoes, while they were 17% and 19% lower for mammal-fed mosquitoes (**Fig. 1** [correction](#) A, C, **Table 1** [correction](#)). The second statistically significant interaction was between blood meal source and the gonotrophic cycle (fecundity: LRT $X^2 = 9.29$, $p = 0.0023$; fertility: LRT $X^2 = 14.4$, $p = 0.00073$). This indicated that the effect of blood type on the number of eggs and larvae per raft differed between the first and second feeding trial (**Tables 2** [correction](#), **3** [correction](#)). For mosquitoes that fed on mammalian blood, fecundity and fertility were 29% lower in the second cycle compared to the first. Although the difference was not statistically significant, fecundity and fertility showed a slight increase in the second cycle for bird-fed mosquitoes. During the second cycle, these outcomes were 41% and 40% higher, respectively, for bird-fed mosquitoes compared to mammal-fed mosquitoes (**Fig. 1** [correction](#) B, D, **Table 1** [correction](#)).

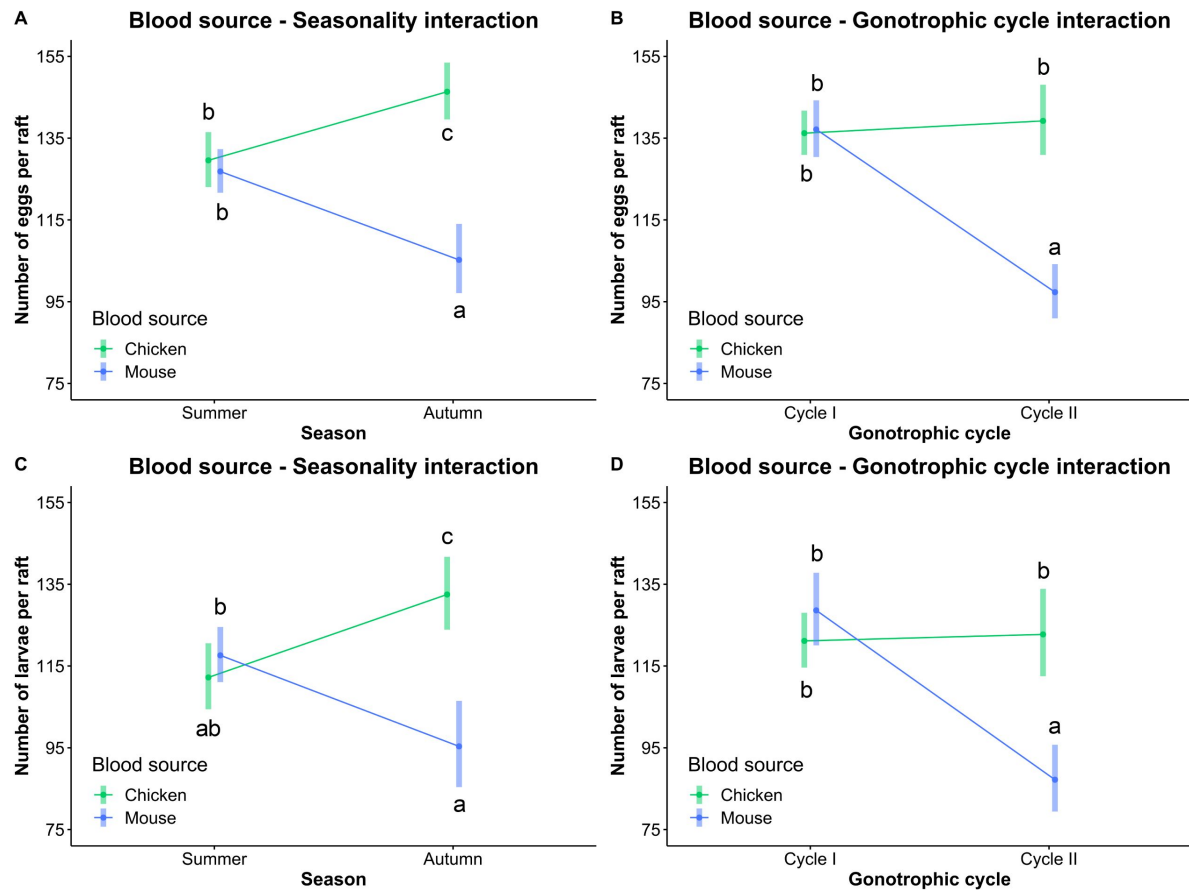


Figure 1.

Interaction plots showing the marginal effects of blood meal source-seasonality and blood meal source-gonotrophic cycle on (A, B) fecundity and (C, D) fertility. Predicted mean values and its corresponding confidence intervals are shown. Means sharing same letters are not statistically different.

Bloodmeal source	Season	Full engorged females	Numer of egg rafts	Fecundty (eggs/raft)	Fertility (larvae/raft)	Hatchability (larvae/eggs)
First gonotrophic cycle (first feeding)						
Mouse	Autumn	58	23	124.5 ± 20.9	120.7 ± 21.5	0.96 ± 0.046
Mouse	Summer	208	120	150.7 ± 29.4	141.8 ± 33.9	0.92 ± 0.10
Chicken	Autumn	163	76	147.5 ± 31.8	133.4 ± 38.4	0.89 ± 0.14
Chicken	Summer	116	74	125.6 ± 39.6	109.9 ± 44.0	0.85 ± 0.20
Second gonotrophic cycle (second feeding)						
Mouse	Autumn	23	15	89.1 ± 19.3	73.3 ± 28.0	0.82 ± 0.25
Mouse	Summer	92	45	106.7 ± 32.0	98.8 ± 31.4	0.91 ± 0.10
Chicken	Autumn	73	36	142.1 ± 36.6	129.2 ± 43.7	0.88 ± 0.18
Chicken	Summer	38	27	138.2 ± 30.3	117.3 ± 38.1	0.84 ± 0.18

Table 1.

Results of reproductive outputs of *Culex quinquefasciatus* for the first and second gonotrophic cycle. Mean values are accompanied with its respective standard deviation.

Factor	LRT χ^2	df	p-value
Seasonality	15.72	1	7.361×10^{-5}
Blood source	8.12	1	0.00438
Gonotrophic cycle	0.55	1	0.45904
Seasonality:Blood source	25.00	1	7.717×10^{-7}
Seasonality:Gonotrophic cycle	3.11	1	0.07764
Blood source:Gonotrophic cycle	9.29	1	0.00230
Seasonality:Blood source:Gonotrophic cycle	1.42	1	0.23340

Table 2.

Analysis of deviance table of the negative binomial model for the effect of blood meal source, seasonality and gonotrophic cycle on fecundity. Dispersion parameter (θ) = 18.75.

Factor	LRT χ^2	df	p-value
Seasonality	11.87	1	0.00057
Blood source	1.64	1	0.19964
Gonotrophic cycle	0.20	1	0.65066
Seasonality:Blood source	13.82	1	0.00020
Seasonality:Gonotrophic cycle	0.87	1	0.34972
Blood source:Gonotrophic cycle	11.40	1	0.00073
Seasonality:Blood source:Gonotrophic cycle	0.04	1	0.83479

Table 3.

Analysis of deviance table of the negative binomial model for the effect of blood meal source, seasonality and gonotrophic cycle on fertility. Dispersion parameter (θ) = 9.52.

Factor	LRT χ^2	df	p-value
Seasonality	5.52	1	0.37623
Blood source	73.06	1	0.00129
Gonotrophic cycle	29.37	1	0.04137
Seasonality:Blood source	7.56	1	0.30066
Seasonality:Gonotrophic cycle	41.02	1	0.01593
Blood source:Gonotrophic cycle	5.53	1	0.37590
Seasonality:Blood source:Gonotrophic cycle	1.20	1	0.67981

Table 4.

Analysis of deviance table of the quasipoisson model for the effect of blood meal source, seasonality and gonotrophic cycle on hatchability.

Effect of blood source and seasonality on hatchability

The overall hatchability reached 86%, with a range spanning from 76% for MA-II to a peak of 96% for MA-I, as outlined in [Table 1](#) and depicted in [Figure 2](#). The model found a statistically significant effect of blood meal source ($\text{LRT } X^2 = 73.06, p = 0.0012$) and an interaction effect between seasonality and gonotrophic cycle ($\text{LRT } X^2 = 41.02, p = 0.0015$) on hatching rates ([Fig. 3](#), [Table 4](#)). Pairwise comparisons revealed a statistically significant 15% decrease in hatchability for mosquitoes from autumn from the first to the second gonotrophic cycle. Additionally, mosquitoes fed on mammals had greater hatchability than those fed on birds. In summer, hatchability during the first cycle was also higher than during the second cycle. These two trends were not found to be statistically significant.

Model accuracy via simulations for the fecundity and fertility models

When comparing the simulated means for each of the eight treatments (combinations of blood meal source, seasonality, and gonotrophic cycle) with the means calculated from the observed data a high level of accuracy exceeding 90% was observed. Most of the simulated means fitted within the original 95% confidence interval, with an accuracy rate greater than 98%. The greatest mismatching among simulated and original data were found in the treatment MS-I, with an accuracy of 93% and 95% for the fecundity and fertility model, respectively. Accuracy remained high despite variations in sample sizes and data dispersion ([Figs. 4](#), [5](#)).

Discussion

Our results revealed that there exists an interaction effect of blood meal source and seasonality on reproductive outputs of *Cx. quinquefasciatus*. Although we expected a higher fecundity and fertility in summer for bird-fed mosquitoes and higher outputs in autumn for mammal-fed mosquitoes, we found the opposite pattern.

Most studies have reported a greater fecundity and fertility in *Cx. quinquefasciatus* fed on different bird species ([Akoh et al., 1993](#); [Richards et al., 2012](#); [Telang & Skinner, 2019](#)). This trend is also present in other species of *Culex* ([Shroyer & Silvery, 1972](#)) and *Aedes* mosquitoes ([Harrison et al., 2021](#)). The widely accepted hypothesis that has emerged to explain why avian blood produces a greater fecundity and fertility even in mosquitoes with a known mammal-preferred host choice focuses on the nutritional quality of host blood. Usually, it is addressed that because avian blood possesses nucleated and larger erythrocytes this offers greater quantities of proteins and energy than mammalian blood ([Alto et al., 2014](#)). However, this assumption lacks solid evidence due to studies regarding nutritional quality of avian blood and its impact in mosquito fitness are scarce. Actually, some studies have shown that mammalian blood are nutritionally richer than avian blood in terms of total protein content ([Pearce 1977](#), [Sant'Anna et al., 2010](#)). Pre-diuresis, a physiological process where red blood cells and proteins are concentrated in the midgut by excreting large amounts of water, has been postulated as a mechanism for explaining the higher fecundity and fertility observed in bird-fed mosquitoes ([Erram et al., 2022](#)). Furthermore, avian blood has higher concentrations of isoleucine than mammalian blood, a crucial limiting amino acid necessary for egg production ([Greenberg, 1951](#), [Harrison et al., 2021](#)).

The effect of temperature on fitness follows a unimodal relationship with a narrow optimal range. For mosquitoes of genus *Culex* and *Aedes*, this optimal band ranges from 20°C to 30°C, with an optimal at 28°C for species such as *Cx. quinquefasciatus* ([Mordecai et al., 2019](#)). Several studies have corroborated this trend, with higher fecundity and fertility towards the optimal of 28°C ([Abouzied, 2017](#); [Mogi, 1992](#)), regardless of the blood source (bird or mammal). Regarding

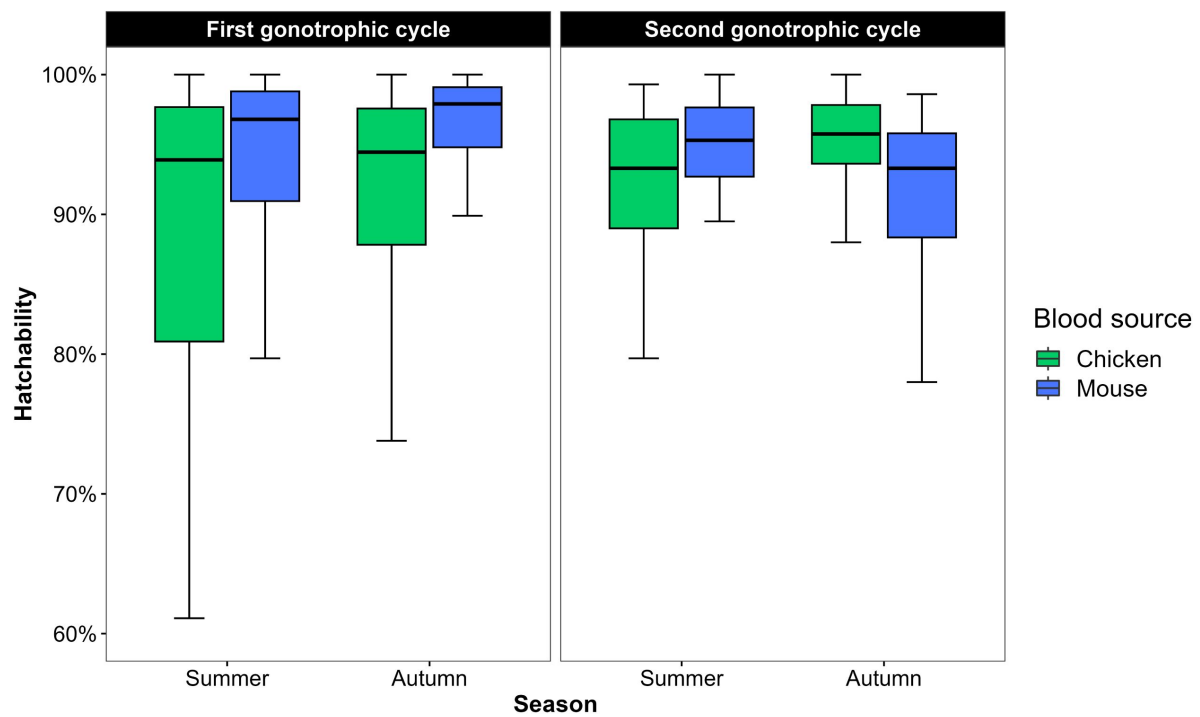


Figure 2.

Boxplots showing the effects of both blood meal source and seasonality on hatchability for the first and second gonotrophic cycle.

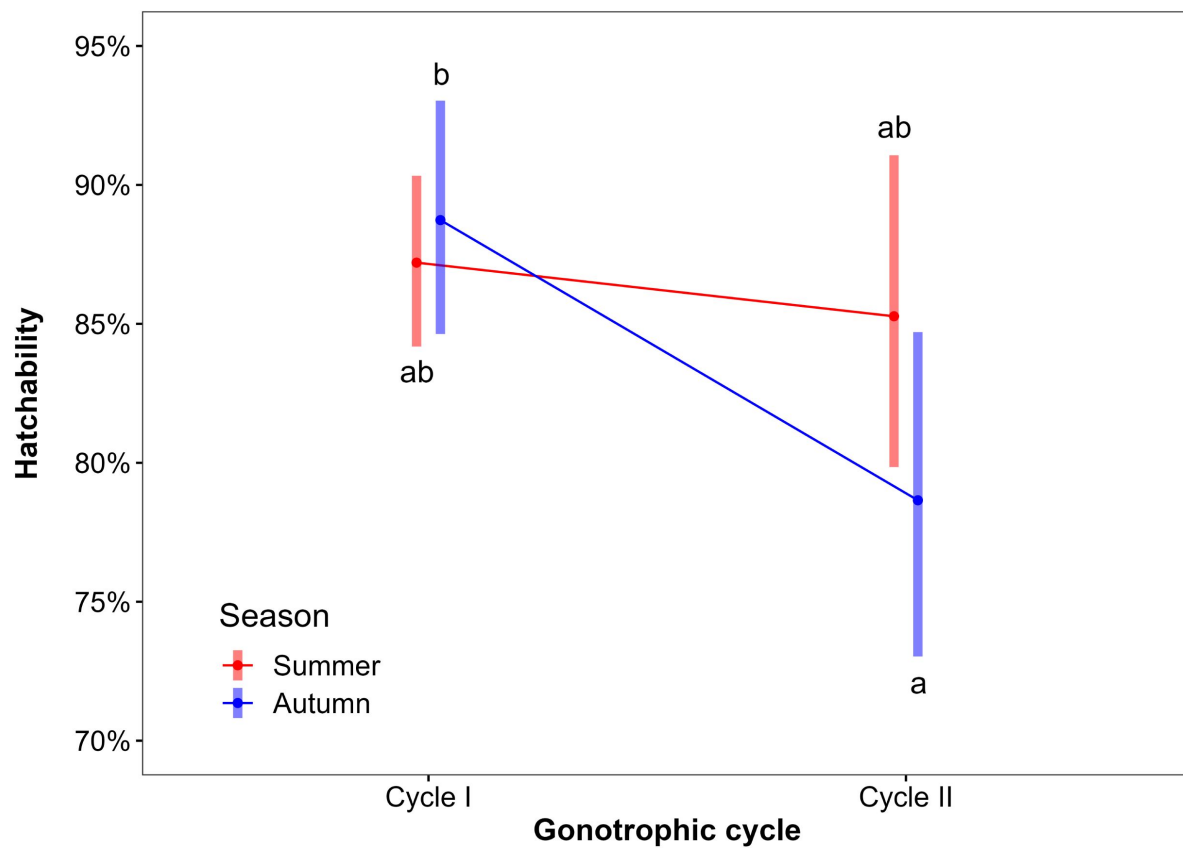


Figure 3.

Interaction plot showing the marginal effects of seasonality-gonotrophic cycle on hatchability.

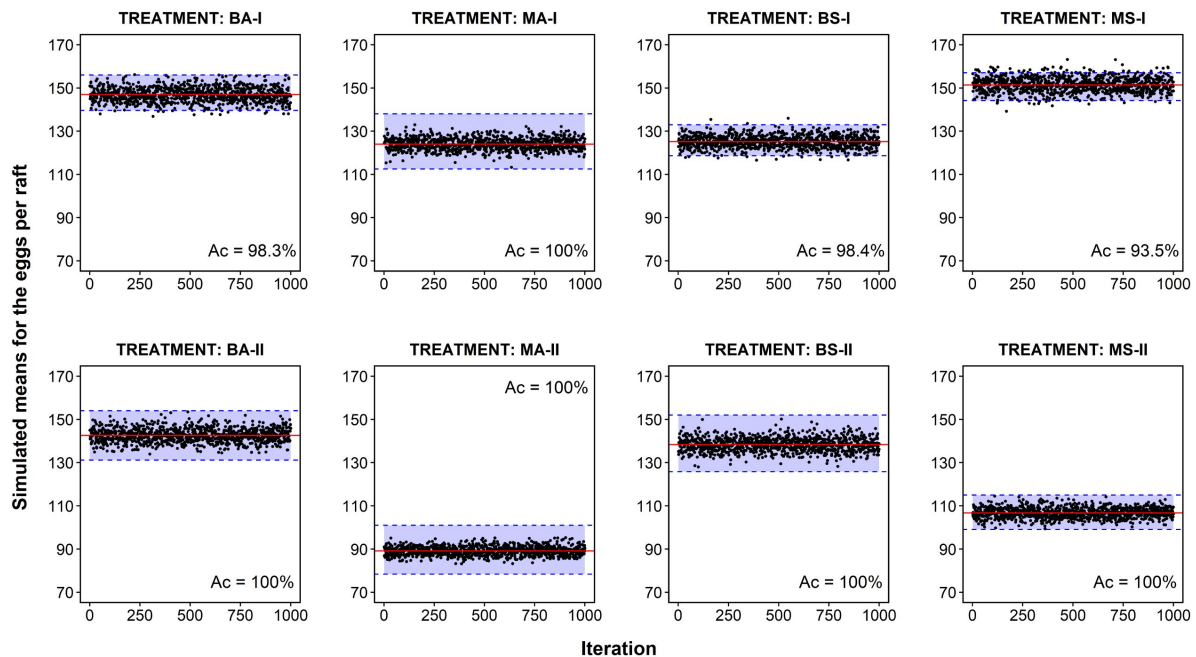


Figure 4.

Simulated means of fecundity (eggs per raft) of each treatment along 1,000 iterations. Abbreviations. B: avian blood; M: mammalian blood; A: autumn; S: summer; I: first gonotrophic cycle; II: second gonotrophic cycle; Ac: model accuracy.

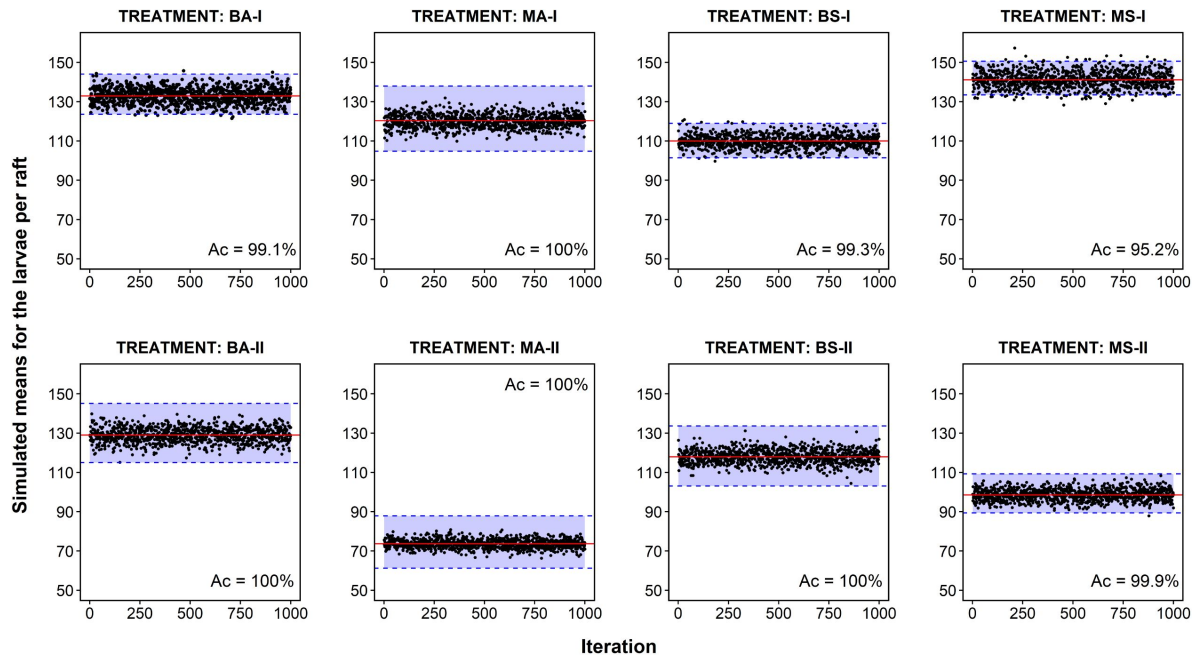


Figure 5.

Simulated means of fertility (eggs per raft) of each treatment along 1,000 iterations. Abbreviations same as [Figure 4](#).

photoperiod, the effects are not as evident as temperature since data are somewhat contradictory. While Mogi (1992) [found](#) that mid and long day lengths induced greater fecundity, Costanzo et al. (2015) [did not find](#) differences of day length on fecundity. Because temperature and photoperiod are always coupled, in general their effects follow the same unimodal trend. Although our experiment was conducted within this optimal range, the highest fecundity and fertility in optimal conditions (summer) was present only in mammal-fed mosquitoes, while higher fitness in bird-fed mosquitoes was detected in autumn. This difference may indicate an interaction of seasonality with blood source.

While our findings may have certain limitations due to moderate sample sizes in some treatments and their reliance solely on laboratory experiments, they verify an interaction effect between blood meal source and seasonality, which had not been previously investigated or documented in any other study. Nevertheless, the precise reasons behind the observed pattern remain inconclusive. The highest levels of fecundity and fertility observed in bird-fed mosquitoes during autumn, when conditions are more challenging, may confer a reproductive advantage, leading to increased offspring production as an adaptive response to these adverse conditions. In contrast, during the more favorable conditions of summer, this pattern may not be as readily discernible. This proposed hypothesis needs further studies to be confirmed.

A second interaction was observed between blood meal source and gonotrophic cycle. We found that fecundity and fertility during the first cycle were similar between mosquitoes fed on chickens and those fed on mammals, and this was also similar with bird-fed mosquitoes from the second gonotrophic cycle, regardless of the season. The only difference we observed was between the fitness of mammal-fed mosquitoes from the second gonotrophic cycle and the earlier mentioned treatments. This pattern is not different from those found in other studies, as the first and second cycles are generally the most productive (Awahmukalah & Brooks, 1985; Christiansen-Jucht et al., 2015). Although we observed a slight increase of fecundity and fertility in bird-fed mosquitoes between cycles, it was statistically non-significant. To our knowledge, only two studies have reported this increasing pattern: an older one by Bennett (1970) [and unpublished data](#) from Obholz (personal communication), both conducted with *Ae. aegypti*. On the contrary, a decrease in fitness between cycles has been reported in other studies for *Cx. quinquefasciatus* (Richards et al., 2012 [; Telang & Skinner, 2019](#) [\) fed upon birds or mammals. This is consistent with our findings regarding mouse-fed mosquitoes; however, because the fecundity and fertility of mouse-fed mosquitoes during the second cycle yielded the fewest egg rafts, this decreasing pattern might also be an effect due to the insufficient sample size.](#)

In addition to assessing fecundity, we also investigated whether hatchability could account for the variations in fertility observed among different treatments. The only statistically significant trend we identified was a decrease in the percentage of hatched eggs from the first to the second gonotrophic cycle, but this was observed only in autumn mosquitoes. This decline in hatching rate may be attributed to a constant exposure to cooling conditions, due to the challenging environmental conditions during this season. This temperature-related effect aligns with findings from other studies (van der Linde et al., 1990). However, although our data suggest that changes in fertility may be linked to variations in hatching rate, it becomes apparent that fecundity has a more pronounced impact on larval production due to the effects of blood meal source and seasonality than hatching rate. Hence, the influence of hatchability on the number of larvae per raft is rather limited.

In summary, our study investigates the critical issue of host shifting patterns in *Culex* mosquitoes, which holds significant importance due to their role as bridge vectors between bird and human hosts. Our model suggests that changes in photoperiod and environmental temperature induce a different physiological state that requires the mosquito to feed on a blood meal source that results in higher fitness (increased fecundity and fertility). We predicted that mosquitoes fed on mammals in autumn will perform higher fecundity and fertility than those fed on birds in the same season.

Unexpectedly, we observed the opposite pattern. Despite this, our study confirmed a significant interaction between blood meal source and seasonality, suggesting a complex interplay of factors affecting mosquito reproduction and its relationship with host use. Due to the observed opposite trend, we can consider three plausible explaining scenarios. Firstly, it is possible that additional factors, such as genetic mechanisms or variations in midgut microbiota composition, play a role in influencing host use patterns. Secondly, host switching might be explained by factors related to vector biology other than fitness. Lastly, it remains a possibility that Argentinian *Cx. quinquefasciatus* populations do not exhibit the same host-switching behaviour observed in US populations, and the observed interaction effects may be linked to different mosquito behaviours. Further investigations are warranted to corroborate our findings and gain a deeper understanding of the interaction between mosquito feeding pattern and seasonality. Ultimately, these investigations will provide valuable insights into mosquito-borne disease transmission and forecasting its emergence.

Acknowledgements

We would like to thank Dr. Agustín Quaglia for providing valuable help in statistical analysis. In addition, we greatly appreciate very helpful comments on the manuscript by anonymous reviewers. This study was supported by grant PICT 2018-1172 (FONCYT) and Consolidar Secyt-UNC (2018-2023).

References

- Akoh J.I., Aigbodion F.I., Onyeka J.O (1993) **Effect of source and size of bloodmeal on egg production by *Culex quinquefasciatus* (Diptera, Culicidae)** *Medical Entomology and Zoology* **44**:41–43 <https://doi.org/10.7601/mez.44.41>
- Alto B.W., Connolly C.R., O'Meara G.F., Hickman D., Karr N (2014) **Reproductive biology and susceptibility of Florida *Culex coronator* to infection with West Nile virus** *Vector-Borne and Zoonotic Diseases* **14**:606–614 <https://doi.org/10.1089/vbz.2013.1501>
- Bennett G.F (1970) **The influence of the blood meal type on the fecundity of *Aedes (Stegomyia) aegypti* L. (Diptera: Culicidae)** *Canadian Journal of Zoology* **48**:539–543 <https://doi.org/10.1139/z70-090>
- Beranek M. (2018) **Mosquitos del género *Culex* (Diptera: Culicidae) como vectores del virus Saint Louis encephalitis (Flavivirus) en ambientes urbanos de la ciudad de Córdoba**
- Branda M.F., Laurito M., Visintin A.M., Almirón W.R (2021) **Gonoactivity of *Culex* (Culex) (Diptera: Culicidae) Mosquitoes During Winter in Temperate Argentina** *Journal of Medical Entomology* **58**:1454–1458 <https://doi.org/10.1093/jme/tjaa295>
- Burkett-Cadena N.D., McClure C.J., Ligon R.A., Graham S.P., Guyer C., Hill G.E., Ditchkoff S.S., Eubanks M.D., Hassan H.K., Unnasch T.R (2011) **Host reproductive phenology drives seasonal patterns of host use in mosquitoes** *PLOS one* **6** <https://doi.org/10.1371/journal.pone.0017681>
- Carsey T.M., Harden J.J (2013) **Monte Carlo simulation and resampling methods for social science**
- Ciota A.T., Mataracchiero A.C., Kilpatrick A.M., Kramer L.D (2014) **The effect of temperature on life history traits of *Culex* mosquitoes** *Journal of Medical Entomology* **51**:55–62 <https://doi.org/10.1603/me13003>
- Costanzo K.S., Schelble S., Jerz K., Keenan M (2015) **The effect of photoperiod on life history and blood-feeding activity in *Aedes albopictus* and *Aedes aegypti* (Diptera: Culicidae)** *Journal of Vector Ecology* **40**:164–171 <https://doi.org/10.1111/jvec.12146>
- Darsie R.F., Argentina Mosquitoes of (1985) **Part I. Keys for identification of adult females and fourth stage larvae in English and Spanish (Diptera: Culicidae)** *Mosquito Systematics* **17**:153–253
- Edman J.D., Taylor D.J (1968) ***Culex nigripalpus*: Seasonal Shift in the Bird-Mammal Feeding Ratio in a Mosquito Vector of Human Encephalitis** *Science* **161**:67–68 <https://doi.org/10.1126/science.161.3836.67>
- Erram D., Vincent Black T., Burkett-Cadena N (2021) **Host Bloodmeal Source Has No Significant Effect on the Fecundity and Subsequent Larval Development Traits of the Progeny in *Culicoides furens* Poey (Diptera: Ceratopogonidae)** *Journal of Medical Entomology* **58**:2439–2445 <https://doi.org/10.1093/jme/tjab085>

- Farajollahi A., Fonseca D.M., Kramer L.D., Kilpatrick A.M (2011) **“Bird biting” mosquitoes and human disease: a review of the role of Culex pipiens complex mosquitoes in epidemiology** *Infection, genetics and Evolution* **11**:1577–1585 <https://doi.org/10.1016/j.meegid.2011.08.013>
- Gervasi S.S., Burkett-Cadena N., Burgan S.C., Schrey A.W., Hassan H.K., Unnasch T.R., Martin L.B (2016) **Host stress hormones alter vector feeding preferences, success, and productivity** *Proceedings of the Royal Society B* **283** <https://doi.org/10.1098/rspb.2016.1278>
- Greenberg J (1951) **Some nutritional requirements of adult mosquitoes (Aedes aegypti) for oviposition: Two figures** *The Journal of Nutrition* **43**:27–35 <https://doi.org/10.1093/jn/43.1.27>
- Hancock C., Camp J.V (2022) **Habitat-Specific Host Selection Patterns of Culex quinquefasciatus and Culex nigripalpus in Florida** *Journal of the American Mosquito Control Association* **38**:83–91 <https://doi.org/10.2987/21-7054>
- Harrison R.E., Brown M.R., Strand M.R (2021) **Whole blood and blood components from vertebrates differentially affect egg formation in three species of anautogenous mosquitoes** *Parasites & Vectors* **14**:1–19 <https://doi.org/10.1186/s13071-021-04594-9>
- Hartig F (2022) **DHARMa: Residual Diagnostics for Hierarchical (Multi-Level / Mixed) Regression Models** *R package version 0*
- Hothorn T., Bretz F., Westfall P (2008) **Simultaneous Inference in General Parametric Models** *Biometrical Journal* **50**:346–363
- Jansen C.C., Webb C.E., Graham G.C., Craig S.B., Zborowski P., Ritchie S.A., Russell R.C., van den Hurk A.F. (2009) **Blood sources of mosquitoes collected from urban and peri-urban environments in eastern Australia with species-specific molecular analysis of avian blood meals** *American Journal of Tropical Medicine and Hygiene* **81**:849–857 <https://doi.org/10.4269/ajtmh.2009.09-0008>
- Janssen N., Fernandez-Salas I., Díaz González E.E., Gaytan-Burns A., Medina-de la Garza C. E., Sanchez-Casas R.M., Börster J., Cadar D., Schmidt-Chanasit J., Jöst H. (2015) **Mammalophilic feeding behaviour of Culex quinquefasciatus mosquitoes collected in the cities of Chetumal and Cancun, Yucatán Peninsula, Mexico** *Tropical Medicine & International Health* **20**:1488–1491 <https://doi.org/10.1111/tmi.12587>
- Kasili S., Oyieke F., Wamae C., Mbogo C (2009) **Seasonal changes of infectivity rates of Bancroftian filariasis vectors in Coast Province, Kenya** *Journal of Vector Borne Diseases* **46**:219–224
- Kauffman E., Payne A., Franke M.A., Schmid M.A., Harris E., Kramer L.D (2017) **Rearing of Culex spp. and Aedes spp. mosquitoes** *Bio-protocol* **7**:e2542–e2542 <https://doi.org/10.21769/bioprotoc.2542>
- Kilpatrick A.M., Kramer L.D., Campbell S.R., Alleyne E.O., Dobson A.P., Daszak P (2005) **West Nile Virus Risk Assessment and the Bridge Vector Paradigm** *Emerging Infectious Diseases* **11**:425–429 <https://doi.org/10.3201/eid1103.040364>
- Kilpatrick A.M., Kramer L.D., Jones M.J., Marra P.P., Daszak P (2006) **West Nile Virus Epidemics in North America Are Driven by Shifts in Mosquito Feeding Behavior** *PLoS Biology* **4** <https://doi.org/10.1371/journal.pbio.0040082>

- Kramer C.Y (1956) **Extension of multiple range tests to group means with unequal numbers of replications** *Biometrics* **12**:307–310 <https://doi.org/10.2307/3001469>
- Labarthe N., Serrão M.L., Melo Y., José-Oliveira S., Lourenço-de-Oliveira R (1998) **Potential vectors of *Dirofilaria immitis* in Itacoatiara, Oceanic Region of Niterói Municipality, state of Rio de Janeiro, Brazil** *Memorias do Instituto Oswaldo Cruz* **93**:425–432 <https://doi.org/10.1590/s0074-02761998000400001>
- Mogi M (1992) **Temperature and photoperiod effects on larval and ovarian development of New Zealand strains of *Culex quinquefasciatus* (Diptera: Culicidae)** *Annals of the Entomological Society of America* **85**:58–66 <https://doi.org/10.1093/aesa/85.1.58>
- Molaei G. *et al.* (2007) **Host feeding pattern of *Culex quinquefasciatus* (Diptera: Culicidae) and its role in transmission of West Nile virus in Harris County, Texas** *American Journal of Tropical Medicine and Hygiene* **77**:73–81 <https://doi.org/10.4269/ajtmh.2007.77.73>
- Mordecai E.A. *et al.* (2019) **Thermal biology of mosquito-borne disease** *Ecology letters* **22**:1690–1708 <https://doi.org/10.1111/ele.13335>
- Muturi E.J., Muriu S., Shililu J., Mwangangi J.M., Jacob B.G., Mbogo C., Githure J., Novak R.J (2008) **Blood-feeding patterns of *Culex quinquefasciatus* and other culicines and implications for disease transmission in Mwea rice scheme, Kenya** *Parasitology Research* **102**:1329–1335 <https://doi.org/10.1007/s00436-008-0914-7>
- R Core Team (2022) **R: A language and environment for statistical computing**
- Richards S.L., Anderson S.L., Yost S.A (2012) **Effects of blood meal source on the reproduction of *Culex pipiens quinquefasciatus* (Diptera: Culicidae)** *Journal of Vector Ecology* **37**:1–7 <https://doi.org/10.1111/j.1948-7134.2012.00194.x>
- Shroyer D.A., Silvery R.E (1972) **A comparison of egg production of *Culex pipiens pipiens* L. fed on avian and mammalian hosts** *Mosquito News* **32**:636–637
- Santos Silva, Pie C., da Rocha M.R., Navarro-Silva T.C. (2019) **Molecular identification of blood meals in mosquitoes (Diptera, Culicidae) in urban and forested habitats in southern Brazil** *PloS one* **14** <https://doi.org/10.1371/journal.pone.0212517>
- Smith J.L., Fonseca D.M (2004) **Rapid assays for identification of members of the *Culex* (*Culex*) *pipiens* complex, their hybrids, and other sibling species (Diptera: Culicidae)** *The American Journal of Tropical Medicine and Hygiene* **40**:339–345 <https://doi.org/10.4269/ajtmh.2004.70.339>
- Spinsanti L.I, Díaz L.A, Glastein N., *et al.* (2008) **Human outbreak of St. Louis encephalitis detected in Argentina, 2005** *Journal of Clinical Virology* **42**:27–33 <https://doi.org/10.1016/j.jcv.2007.11.022>
- Stein M., Zalazar L., Willener J.A., Almeida F.L., Almirón W.R (2013) **Culicidae (Diptera) selection of humans, chickens and rabbits in three different environments in the province of Chaco, Argentina** *Memórias do Instituto Oswaldo Cruz* **108**:563–571 <https://doi.org/10.1016/j.actatropica.2015.12.002>
- Takken W., Verhulst N.O (2013) **Host preferences of blood-feeding mosquitoes** *Annual Review of Entomology* **58**:433–453 <https://doi.org/10.1146/annurev-ento-120811-153618>

- Telang A., Skinner J (2019) **Effects of host blood meal source on reproductive output, nutrient reserves and gut microbiome of West Nile virus vector *Culex quinquefasciatus*** *Journal of Insect Physiology* **114**:15–22 <https://doi.org/10.1016/j.jinsphys.2019.02.001>
- van der Linde, T.C. de K., Hewitt, P.H., Nel, A. & van der Westhuizen, M.C. (1990) **Development rates and percentage hatching of *Culex (Culex) theileri* Theobald (Diptera: culicidae) eggs at various constant temperatures** *Journal of the Entomological Society of Southern Africa* **53**:17–26 <https://doi.org/10.1093/jmedent/27.6.939>
- Venables W.N., Ripley B.D (2002) **Exploratory multivariate analysis** *Modern applied statistics with S* :301–330 https://doi.org/10.1007/978-0-387-21706-2_11
- Yan J., Gangoso L., la Puente Martínez-de, Soriguer J., Figuerola R. (2017) **Avian phenotypic traits related to feeding preferences in two *Culex* mosquitoes** *The Science of Nature* **104**:1–10 <https://doi.org/10.1007/s00114-017-1497-x>
- Yan J., Broggi J, la Puente Martínez-de, Gutiérrez-López J., Gangoso R., Soriguer L., Figuerola R. (2018) **Does Bird metabolic rate influence mosquito feeding preference?** *Parasites & Vectors* **11**:1–9 <https://doi.org/10.1186/s13071-018-2708-9>

Article and author information

Kevin Alen Rucci

Laboratorio de Arbovirus, Instituto de Virología “Dr. J. M. Vanella” (InViV), Facultad de Ciencias Médicas (FCM), Universidad Nacional de Córdoba (UNC), Consejo Nacional de Investigaciones Científicas y Técnicas (CONICET), Argentina

For correspondence: kevin.rucci@mi.unc.edu.ar

ORCID iD: [0000-0002-0940-2423](https://orcid.org/0000-0002-0940-2423)

Mariana Pueta

Instituto de Investigaciones en Biodiversidad y Medioambiente (INIBIOMA), CONICET - Universidad Nacional de Comahue (UNCo), Consejo Nacional de Investigaciones Científicas y Técnicas (CONICET), Argentina

ORCID iD: [0000-0002-9837-4718](https://orcid.org/0000-0002-9837-4718)

Adrián Díaz

Laboratorio de Arbovirus, Instituto de Virología “Dr. J. M. Vanella” (InViV), Facultad de Ciencias Médicas (FCM), Universidad Nacional de Córdoba (UNC), Consejo Nacional de Investigaciones Científicas y Técnicas (CONICET), Argentina

For correspondence: adrian.diaz@fcm.unc.edu.ar

ORCID iD: [0000-0001-5953-2907](https://orcid.org/0000-0001-5953-2907)

Copyright

© 2023, Rucci et al.

This article is distributed under the terms of the [Creative Commons Attribution License](https://creativecommons.org/licenses/by/4.0/), which permits unrestricted use and redistribution provided that the original author and source are credited.

Editors

Reviewing Editor

Bavesh Kana

University of the Witwatersrand, Johannesburg, South Africa

Senior Editor

Bavesh Kana

University of the Witwatersrand, Johannesburg, South Africa

Reviewer #1 (Public Review):

Summary: This study examines the role of host blood meal source, temperature, and photoperiod on the reproductive traits of *Cx. quinquefasciatus*, an important vector of numerous pathogens of medical importance. The host use pattern of *Cx. quinquefasciatus* is interesting in that it feeds on birds during spring and shifts to feeding on mammals towards fall. Various hypotheses have been proposed to explain the seasonal shift in host use in this species but have provided limited evidence. This study examines whether the shifting of host classes from birds to mammals towards autumn offers any reproductive advantages to *Cx. quinquefasciatus* in terms of enhanced fecundity, fertility, and hatchability of the offspring. The authors found no evidence of this, suggesting that alternate mechanisms may drive the seasonal shift in host use in *Cx. quinquefasciatus*.

Strengths: Host blood meal source, temperature, and photoperiod were all examined together.

Weaknesses: The study was conducted in laboratory conditions with a local population of *Cx. quinquefasciatus* from Argentina. I'm not sure if there is any evidence for a seasonal shift in the host use pattern in *Cx. quinquefasciatus* populations from the southern latitudes.

Comments on the revision:

Overall, I am not quite convinced about the possible shift in host use in the Argentinian populations of *Cx. quinquefasciatus*. The evidence from the papers that the authors cite is not strong enough to derive this conclusion. Therefore, I think that the introduction and discussion parts where they talk about host shift in *Cx. quinquefasciatus* should be removed completely as it misleads the readers. I suggest limiting the manuscript to talking only about the effects of blood meal source and seasonality on the reproductive outcomes of *Cx. quinquefasciatus*.

<https://doi.org/10.7554/eLife.89485.2.sa1>

Reviewer #2 (Public Review):

Summary:

Conceptually, this study is interesting and is the first attempt to account for the potentially interactive effects of seasonality and blood source on mosquito fitness, which the authors frame as a possible explanation for previously observed host-switching of *Culex quinquefasciatus* from birds to mammals in the fall. The authors hypothesize that if changes in fitness by blood source change between seasons, higher fitness on birds in the summer and on mammals in the autumn could drive observed host switching. To test this, the authors fed individuals from a colony of *Cx. quinquefasciatus* on chickens (bird model) and mice (mammal model) and subjected each of these two groups to two different environmental conditions reflecting the high and low temperatures and photoperiod experienced in summer and autumn in Córdoba, Argentina (aka seasonality). They measured fecundity, fertility, and

hatchability over two gonotrophic cycles. The authors then used a generalized linear model to evaluate the impact of host species, seasonality, and gonotrophic cycle on fecundity, fertility, and hatchability. The authors were trying to test their hypothesis by determining whether there was an interactive effect of season and host species on mosquito fitness. This is an interesting hypothesis; if it had been supported, it would provide support for a new mechanism driving host switching. While the authors did report an interactive impact of seasonality and host species, the directionality of the effect was the opposite from that hypothesized. The authors have done a very good job of addressing many of the reviewer concerns, with several exceptions that continue to cause concern about the conclusions of the study.

Strengths:

- (1) Using a combination of laboratory feedings and incubators to simulate seasonal environmental conditions is a good, controlled way to assess the potentially interactive impact of host species and seasonality on the fitness of *Culex quinquefasciatus* in the lab.
- (2) The driving hypothesis is an interesting and creative way to think about a potential driver of host switching observed in the field.
- (3) The manuscript has become a lot clearer and easier to read with the revisions - thank you to the authors for working hard to make many of the suggested changes.

Weaknesses:

- (1) The authors have decided not to follow the suggestion of conducting experimental replicates of the study. This is understandable given the significant investment of resources and time necessary, however, it leaves the study lacking support. Experimental replication is an important feature of a strong study and helps to provide confidence that the observed patterns are real and replicable. Without replication, I continue to lack confidence in the conclusions of the study.
- (2) The authors have included some additional discussion about the counterintuitive nature of their results, but the paragraph discussing this in the discussion was confusing. I believe that this should be revised. This is a key point of the paper and needs to be clear to the reader.
- (3) There should be more discussion of the host switching observed in the two studies conducted in Argentina referenced by the authors. Since host switching is the foundation for the hypothesis tested in this paper, it is important to fully explain what is currently known in Argentina.
- (4) In some cases, the explanations of referenced papers are not entirely accurate. For example, when referencing Erram et al 2022, I think the authors misrepresented the paper's discussion regarding pre-diuresis- Erram et al. are suggesting that pre-diuresis might be the mechanism by which *C. furens* compensates for the lower nutritional value of avian blood, leading to no significant difference between avian/mammal blood on fecundity/fertility (rather than leading to higher fecundity on birds, as stated in this manuscript). The study performed by Erram et al. also didn't prove this phenomenon, they just suggest it as a possible mechanism to explain their results, so that should be made clear when referencing the paper.
- (5) In some cases, the conclusions continue to be too strongly worded for the evidence available. For example, lines 322-324: I don't think the data is sufficient to conclude that a different physiological state is induced, nor that they are required to feed on a blood source that results in higher fitness.
- (6) There is limited mention of the caveat that this experiment performed with simulated seasonality that does not perfectly replicate seasonality in the field. I think this caveat should be discussed in the discussion (e.g. that humidity is held constant).

<https://doi.org/10.7554/eLife.89485.2.sa0>

Author Response

The following is the authors' response to the original reviews.

Reviewer #1 (Public Review):

Summary:

This study examines the role of host blood meal source, temperature, and photoperiod on the reproductive traits of Cx. quinquefasciatus, an important vector of numerous pathogens of medical importance. The host use pattern of Cx. quinquefasciatus is interesting in that it feeds on birds during spring and shifts to feeding on mammals towards fall. Various hypotheses have been proposed to explain the seasonal shift in host use in this species but have provided limited evidence. This study examines whether the shifting of host classes from birds to mammals towards autumn offers any reproductive advantages to Cx. quinquefasciatus in terms of enhanced fecundity, fertility, and hatchability of the offspring. The authors found no evidence of this, suggesting that alternate mechanisms may drive the seasonal shift in host use in Cx. quinquefasciatus.

Strengths:

Host blood meal source, temperature, and photoperiod were all examined together.

Weaknesses: The study was conducted in laboratory conditions with a local population of Cx. quinquefasciatus from Argentina. I'm not sure if there is any evidence for a seasonal shift in the host use pattern in Cx. quinquefasciatus populations from the southern latitudes.

We agree on the reviewers observation about the evidence on seasonal shift in the host use pattern in Cx. quinquefasciatus populations from southern latitudes. We include a paragraph in the Introduction section regarding this. Unfortunately, studies conducted in South America to understand host use by Culex mosquitoes are very limited, and there are virtually no studies on the seasonal feeding pattern. In Argentina, there is some evidence (Stein et al., 2013, Beranek, 2019) regarding the seasonal change in host use by Culex species, including Cx. quinquefasciatus, where the inclusion of mammals during the autumn has been observed. As part of a comprehensive study on characterising bridge vectors for SLE and WN viruses, our research group is currently working on the molecular identification of blood meals from engorged females to gain deeper insights into the seasonal feeding pattern of Culex mosquitoes. While the seasonal change in host use by Culex quinquefasciatus has not been reported in Argentina so far, there has been an observed increase in reported cases of SLE virus in humans between summer and fall (Spinsanti et al., 2008). It is based on this evidence that we hypothesise there is a seasonal change in host use by Cx. quinquefasciatus, similar to what occurs in the United States. This is also considering that both countries (Argentina and the United States) have regions with similar climatic conditions (temperate climates with thermal and hydrological seasonality). Since we work on the same species and in a similar temperate climate regimen, we assumed there is a seasonal shift in the host use by this mosquito species.

Reviewer #1 (Recommendations for the authors):

Abstract

Line 23: fed on two different hosts.

Accepted as suggested.

I think the concluding statement should be rewritten to say that immediate reproductive outcomes do not explain the shift in host use pattern of Cx. quinquefasciatus mosquitoes from birds to mammals towards autumn.

Accepted as suggested.

Introduction

No comments.

Materials and Methods

Please mention sample sizes in the text as well (n = ?) for each treatment.

Accepted as suggested.

Page 99:C. quinquefasciatus, since C. pipiens and its hybrids are present as well in Cordoba.

Accepted as suggested.

Results – Line 146: subsequently instead of posteriorly

Accepted all changes as suggested.

Line 148: were counted instead of was counted.

Accepted all changes as suggested.

Line 160: Subsequently instead of posteriorly

Accepted all changes as suggested.

Line 171: on fertility

Accepted all changes as suggested.

Line 174: there was an interaction effect on...

Accepted all changes as suggested.

Line 175: there were no differences in the number of eggs

Accepted all changes as suggested.

Discussion

I think the first paragraph in the discussion section is redundant and should be deleted.

The whole discussion was rewritten to be focused on our aims and results.

Line 282: this sentence needs to be rewritten.

Accepted as suggested.

Line 299: at 28{degree sign}C

Line 300: at 30{degree sign}C

Sorry, but we are not sure about your comment here. We checked. Temperatures are written as stated, 28°C and 30°C.

Line 363: *I think the authors need to discuss more about the bigger question they were addressing. I think that the discussion section can be strengthened greatly by elaborating on whether there is evidence for a seasonal shift in host use pattern in Cx. quinquefasciatus in the southern latitudes. If yes, what alternate mechanisms they believe could be driving the seasonal change in host use in this species in the southern latitudes now that they show the 'deriving reproductive advantages' hypothesis to be not true for those populations.*

Thanks for this observation. We agree and so the Discussion section was restructured to align it with our results, as suggested.

Reviewer #2 (Public Review):

Summary:

Conceptually, this study is interesting and is the first attempt to account for the potentially interactive effects of seasonality and blood source on mosquito fitness, which the authors frame as a possible explanation for previously observed host-switching of Culex quinquefasciatus from birds to mammals in the fall. The authors hypothesize that if changes in fitness by blood source change between seasons, higher fitness in birds in the summer and on mammals in the autumn could drive observed host switching. To test this, the authors fed individuals from a colony of Cx. quinquefasciatus on chickens (bird model) and mice (mammal model) and subjected each of these two groups to two different environmental conditions reflecting the high and low temperatures and photoperiod experienced in summer and autumn in Córdoba, Argentina (aka seasonality). They measured fecundity, fertility, and hatchability over two gonotrophic cycles. The authors then used a generalized linear mixed model to evaluate the impact of host species, seasonality, and gonotrophic cycle on fecundity and fertility and a null model analysis via data randomization for hatchability. The authors were trying to test their hypothesis by determining whether there was an interactive effect of season and host species on mosquito fitness. This is an interesting hypothesis; if it had been supported, it would provide support for a new mechanism driving host switching. While the authors did report an interactive impact of seasonality and host species, the directionality of the effect was the opposite of that hypothesized. While this finding is interesting and worth reporting, there are significant issues with the experimental design and the conclusions that are drawn from the results, which are described below. These issues should be addressed to make the findings trustworthy.

Strengths:

- (1) Using a combination of laboratory feedings and incubators to simulate seasonal environmental conditions is a good, controlled way to assess the potentially interactive impact of host species and seasonality on the fitness of Culex quinquefasciatus in the lab.*
- (2) The driving hypothesis is an interesting and creative way to think about a potential driver of host switching observed in the field.*

Weaknesses:

(1) There is no replication built into this study. Egg lay is a highly variable trait, even within treatments, so it is important to see replication of the effects of treatment across multiple discrete replicates. It is standard practice to replicate mosquito fitness experiments for this reason. Furthermore, the sample size was particularly small for some groups (e.g. 15 egg rafts for the second gonotrophic cycle of mice in the autumn, which was the only group for which a decrease in fecundity and fertility was detected between 1st and 2nd gonotrophic cycles). Replicates also allow investigators to change around other variables that might impact the results for unknown reasons; for example, the incubators used for fall/summer conditions can be swapped, ensuring that the observed effects are not artefacts of other differences between treatments. While most groups had robust sample sizes, I do not trust the replicability of the results without experimental replication within the study.

We agree egg lay is a variable trait and so we consider high numbers of mosquitoes and egg lay during experiments compared to our studies of the same topics. Evaluating variables such as fecundity, fertility, or other types of variables (collectively referred to as "life tables") is a challenging issue that depends on several intrinsic and extrinsic factors. Because all of this, in some experiments, sample sizes might not be very large, and in several articles, lower sample sizes could be found. For instance, in Richards et al. (2012), for *Culex quinquefasciatus*, during the second gonotrophic cycle, some experiments had 13 or even 6 egg rafts. For species like *Aedes aegypti*, the sample size for life table analysis is also usually small. As an example, Muttis et al. (2018) reported between 1 and 4 engorged females (without replicates). In addition, small sample size would be a problem if we would not have obtained any effect, which is not the case due to the fact that we were interested in finding an effect, regardless of the effect size. Because of this, we do find our sample sizes quite robust for our results.

Regarding the need to repeat the experiments in order to give more robustness to the study we also agree. However, after a review of the literature (articles cited in the original manuscript), it is apparent that similar experiments are not frequently repeated as such. Examples of this are the studies of Richards et al. (2012), Demirci et al. (2014) or Telang & Skinner (2019), which even they manipulate several cages at a time as "replicates", they are not true replicates because they summarise and manipulate all data together, and do not repeat the experiment several times. We see these "replicates" as a way of getting a greater N.

As was stated by the reviewer, repetition is a resource and time-consuming activity that we are not able to do. Replicating the experiment poses a significant time and resources challenge. The original experiment took over three months to complete, and it is anticipated that a similar timeframe would be necessary for each replication (6 months in total considering two more replicates). Given our existing commitments and obligations, dedicating such an extensive period solely to this would impede progress on other crucial projects and responsibilities.

Given the limitations of resources and time and the infrequent use of experimental replication in this type of studies, we performed a simulation-based analysis via a Monte Carlo approach. This approach involved generating synthetic data that mimics the expected characteristics of the original experiment and subsequently subjecting it to the same analysis routine. The main goal of this simulation was to evaluate the potential spuriousness and randomness of the results that might arise due to the experimental conditions. So, evaluating the robustness and confidence of our results and data.

*(2) Considering the hypothesis is driven by the host switching observed in the field, this phenomenon is discussed very little. I do not believe *Cx. quinquefasciatus* host switching has been observed in Argentina, only in the northern hemisphere, so it is possible that the species could have an entirely different ecology in Argentina. It would have been*

helpful to conduct a blood meal analysis prior to this experiment to determine whether using an Argentinian population was appropriate to assess this question. If the Argentinian populations don't experience host switching, then an Argentinian colony would not be the appropriate colony to use to assess this question. Given that this experiment has already been conducted with this population, this possibility should at least be acknowledged in the discussion. Or if a study showing host switching in Argentina has been conducted, it would be helpful to highlight this in the introduction and discussion.

Thanks for this observation. We agree. However, we conducted the experiment beside host use data from Argentina since we used the mosquito species, and the centre region of Argentina (Córdoba) has a similar temperate weather regimen that those observed in the east coast of US.

We are aware that few studies regarding host shifting in South America are available, some such that those conducted by Stein et al. (2013) and Beranek (2019) reported a moderate host switch for *Culex quinquefasciatus* in Argentina. We have already performed a study about seasonal host feeding patterns for this species. However, even though there are few studies regarding host shifting, our hypothesis is based mainly in the seasonality of human cases of WNV and SLEV, a pattern that has been demonstrated for our region, see for example the study of Spinsanti et al. (2008).

We include a new paragraph in the Introduction and Discussion sections. Please see answers Reviewer #1.

(3) The impacts of certain experimental design decisions are not acknowledged in the manuscript and warrant discussion. For example, the larvae were reared under the same conditions to ensure adults of similar sizes and development timing, but this also prevents mechanisms of action that could occur as a result of seasonality experienced by mothers, eggs, and larvae.

We understand the confusion that may have arisen due to a lack of further details in the methodology. If we are not mistaken, you are referring to our oversight regarding the consideration of carry-over effects of larvae rearing that could potentially impact reproductive traits. When investigating the effects of temperature or other environmental factors on reproductive traits, it is possible to acclimate either larvae or adults. This is due to the significant phenotypic plasticity that mosquitoes exhibit throughout their entire ontogenetic cycle. In our study, we followed an approach similar to that of other authors where the adults are exposed to experimental conditions (temperature and photoperiod). For a similar approach you can refer to the studies conducted by Ferguson et al. (2018) for *Cx. pipiens*, Garcia Garcia & Londoño Benavides (2007) for *Cx. quinquefasciatus* or Christiansen-Jucht et al. (2014, 2015) for *Anopheles gambiae*.

(4) There are aspects of the data analysis that are not fully explained and should be further clarified. For example, there is no explanation of how the levels of categorical variables were compared.

The methodology and statistical analysis were expanded for a better understanding.

(5) The results show the opposite trend as was predicted by the authors based on observed feeding switches from birds to mammals in the autumn. However, they only state this once at the end of the discussion and never address why they might have observed the opposite trend as was hypothesized.

The discussion was restructured to focus on our results and our model.

(6) Generally speaking, the discussion has information that isn't directly related to the results and/or is too detailed in certain parts. Meanwhile, it doesn't dig into the meaning of the results or the ways in which the experimental design could have influenced results.

As mentioned above, the discussion was restructured to reflect our findings. We also included the effect that our design might have influenced our results. However, as stated above we do not fully agree that the design is inadequate for our analysis, we performed standard protocols followed by other researchers and studies in this research field.

(7) Beyond the issue of lack of replication limiting trust in the conclusions in general, there is one conclusion reached at the end of the discussion that would not be supported, even if additional replicates are conducted. The results do not show that physiological changes in mosquitoes trigger the selection of new hosts. Host selection is never measured, so this claim cannot be made. The results don't even suggest that fitness might trigger selection because the results show that physiological changes are in the opposite direction as what would be hypothesized to produce observed host switches. Similarly, the last sentence of the abstract is not supported by the results.

We agree with this observation. However, we did not evaluate the impact of fitness on host selection in this study. Instead, we aimed to investigate the potential influence of seasonality on mosquito fitness as a potential trigger for a shift in host selection. We agree that we have incorrectly used the term “host selection” when we should actually be discussing “host use change”. Our results indicate a seasonal alteration in mosquito fitness in response to temperature and photoperiod changes. Building upon this observation, we re-discussed our hypothesis and theoretical model to explain this seasonal shift in host use.

(8) Throughout the manuscript, there are grammatical errors that make it difficult to understand certain sentences, especially for the results.

All English grammar and writing of the manuscript was revised and corrected to be easily understood.

This study is driven by an interesting question and has the potential to be a valuable contribution to the literature.

Reviewer #2 (Recommendations for The Authors):

I hope that the authors will consider the suggested revisions and experimental replication to improve the quality of the study and paper.

This study tests a very interesting hypothesis. I understand that additional replicates are difficult to conduct, but I do believe that fitness studies absolutely require experimental replicates. Unless you are able to replicate the observed effects, I personally would not trust the results of this study. I hope that you will consider conducting replicates so that this important question can be answered in a more robust manner. Below, I expand upon some additional points in the public review and also provide more specific suggestions. I provided some copy-editing feedback, but was not able to point out all grammatical mistakes. I suggest that you use ChatGPT to help you edit the English. For example, you can feed ChatGPT your MS and ask it to bold the grammatical errors or you can ask it to edit grammatical errors and bold the sections that were edited. I understand that writing in a second language is very difficult (from personal experience!), so I view ChatGPT as a great tool to help even the playing field for publishing. Below are line item suggestions. Apologies that wording is curt, I was trying to be efficient in writing.

20-21: *I suggest that you emphasize that you are investigating the interactive effect.*

Accepted as suggested.

22: *they weren't "reared" (from larvae) in different conditions, they were "maintained" as adults*

Accepted as suggested.

26-27: *increased/decreased is a bit misleading since you did not evaluate these groups sequentially in time. It might be more accurate to describe it as less than/greater than. Also, if you say increased/decreased or less than/greater than, you should always say what you are comparing to. The same applies throughout the MS.*

Accepted as suggested.

29-30: *"finding the" is not correct here; could be "with the lowest..."*

Accepted as suggested.

34-36: *I do not think that your results suggest this, even if you were to replicate the results of this experiment. You haven't shown metabolic changes.*

We understand the point. Accepted as suggested.

42-44: *"one of the main responsible" should be "one of the main species responsible..."*

Accepted as suggested.

48: *I think that "host preference" is better than selection here; -philic denotes preference*

Accepted as suggested.

50: *"Moreover" isn't the correct transition word here*

Accepted as suggested.

57: *"could" isn't correct here; consider saying "... species sometimes feed primarily on mammal hosts, including humans, in certain situations."*

Accepted as suggested.

58: *Different isn't correct word here*

Accepted as suggested.

60: *delete "feeding"*

Accepted as suggested.

66-68: *I am not familiar with any blood meal analysis studies in the southern hemisphere that show host switching for Culex species between summer and autumn. If this hasn't been shown, then this critique of the host migration hypothesis doesn't make sense.*

There are some studies pointing this out (Stein et al., 2013, Beranek 2019), and unpublished data from us). However, our hypothesis has supported by epidemiological data observed in human population which indicate a seasonal activity pattern. It was explained in depth in the Introduction section.

| 68: *ensures is not the right word; I suggest "suggests"*

Accepted as suggested.

| 68-70: *this explanation isn't clear to me; please revise*

It will be revised. Accepted as suggested.

| 70: *change cares to care*

Accepted as suggested.

| 76-77: *can you explain how they were not supported by the data for the benefit of those who are not familiar with these papers please?*

Accepted as suggested.

| 87-89: *I suggest the following wording: "In the autumn, we expect a greater number of eggs (fecundity) and larvae (fertility) in mosquitoes after feeding on a mammal host compared to an avian host, and the opposite relationship in the summer."*

Accepted as suggested.

| 99: *edit for grammar*

Accepted as suggested.

| 102: *suggest: "...offered a blood meal from a restrained chicken twice a month"*

Accepted as suggested.

| 107: *powder*

Accepted as suggested.

| 108: *inbred? Is this the term you meant to use?*

Changed as suggested.

| 109: *"several" cannot be used to describe 20 generations; suggest using "over twenty generations"; also, it would be good to acknowledge in your discussion that lab adaptation could force evolution, especially since mosquitoes are kept at constant temperatures and fed with certain hosts (with easy access) in the lab. Also, it would be good to know when the experiments were conducted to know the lapse of time between the creation of the colony and the experiments.*

Accepted as suggested.

110-111: Does humidity vary between summer and fall in Córdoba? If so, I suggest acknowledging in the discussion that if humidity differences are involved in a potential interaction between host species and seasonality, then this would not have been captured by your experimental design.

Several variables change during seasons. We were interested in capturing the effects of temperature and photoperiod, since humidity is a variable difficult to control.

113-116: I suggest combining into one sentence to make more concise.

Accepted as suggested.

135: You might be obscuring the true impact of seasonality by rearing the larvae under the same conditions. There may be signals that mothers/eggs/larvae receive that influence their behavior (e.g. I believe this is the case for diapause), so this limitation should also be acknowledged. I understand why you decided to do this to control for development time and size, but it is something that should be considered in the discussion.

As it was explained above, *Cx. quinquefasciatus* do not suffer diapause in our country. Maintaining mosquitoes from adults was an approach selected by us based on other studies.

138: edit: "with cotton pads soaked in... on plastic..."; what is plastic glass? Do you mean plastic dishes?

Accepted as suggested.

141: here and throughout paragraph, full should be "fully"

Accepted as suggested.

144: located should be "placed"

Accepted as suggested.

147: suggest editing to "at which point, they were fixed with 1 mL of 96% ethanol and the number of L1 larvae per raft was counted."

Accepted as suggested.

154-155: edit for grammar

Accepted as suggested.

157: Your GLM explanation doesn't say anything about how you made pairwise comparisons between your levels; did you use emmeans?

This revised version includes a more detailed methodology and statistical analysis. Accepted as suggested.

158-160: I don't understand why you took this approach - it seems strange to me to use this analysis, but I am not familiar with it, so it might be that I lack the knowledge to be able to adequately evaluate. Please provide more explanation so that readers can better

understand this analysis. A citation for this kind of application of the analysis would be helpful.

It was changed to be in accordance with the remaining analyses.

173: replace neither with either

Accepted as suggested.

174: this applies throughout; edit to : "An interaction effect was observed..."

Accepted as suggested.

175: "it was not found" is grammatically incorrect; instead : "We did not find ..." or "no differences in... were detected", etc

Accepted as suggested.

183: "it was detected" is grammatically incorrect

Accepted as suggested.

185-186: "being this treatment... in terms of fitness": I do not understand what this means. Please rephrase

Accepted as suggested.

170-199: you should provide the effect sizes and p values in text and/or in the figure for the pairwise comparisons

Accepted as suggested.

193-196. These two sentences are confusing and I am not sure what you mean, especially in the first sentence.

It was rewritten. Accepted as suggested.

*Figure 1: This figure is great and easy to read and interpret! Thank you for the comment!
218-219: it is important to state which mosquito species you are referring to here.*

Accepted as suggested.

226-227: you definitely should acknowledge the small sample size here.

Considered.

227: "it was observed" should be "We observed" or "A greater hatching rate.... was observed."

Accepted as suggested.

228-229: is the result really comparable even though you took very different approaches to the analysis for these outcomes?

Changed to be comparable.

230-278: *the discussion of these hypotheses is too long and detailed, especially since the comparison of mouse vs chicken wasn't your main question; you really wanted to understand this in the context of seasonality. I suggest cutting this down a lot and making room to dig into your results more, and also to discuss the potential impacts of your experimental design/limitations on the results.*

Discussion was changed to focus on our results and model. Accepted as suggested.

281: *Hoffman is an old citation; I suggest you cite a modern review.*

Accepted as suggested. We deleted it due to the re-writing of the manuscript.

282: *"It can be recognise".. I am not sure what you are trying to say here*

Accepted as suggested.

1. *After the first time you write a species name, you can abbreviate the genus in all future mentions unless it is at the beginning of a sentence.*

Accepted as suggested.

303-305: *Revise this sentence. E.g "Fewer studies are available regarding photoperiod and show mixed results; Mogi (1992) found that mid and long day lengths induced greater fecundity while Costanzo et al. (2015) did not find differences in fecundity by day length."*

Accepted as suggested.

315-316: *typically, unpublished data shouldn't be referenced; I'm not sure if eLife has a policy on this.*

We will check this with eLife guidelines. However, since the lack of evidence on this pattern we consider important to include this unpublished data.

316: *Aegypti should be lowercase*

Accepted as suggested.

328-330: *This sentence is redundant with the first sentence of the paragraph*

Accepted as suggested.

321-336: *You never reintroduced your hypothesis in your discussion. I suggest that you center your whole discussion more directly around the hypothesis that motivated the study. If you decide not to restructure your discussion, you should at least reintroduce your hypothesis here and discuss how your results do not support the hypothesis.*

Accepted as suggested.

337-348: *This paragraph is a bit confusing as you jump between fertility and hatchability*

Accepted as suggested.

353: *is viral transmission the right word to use here? I think you might mean bridge vector transmission to humans specifically?*

Accepted as suggested.

357: *you say "neither" but never define which traits you are referring to*

Accepted as suggested.

361: *I suggest "two variables previously analyzed separately..."*

Accepted as suggested.

General: There is no statement about the availability of data; it is eLife policy to require all data to be publicly available. Also, it would be helpful to share your code to help understand how you conducted pairwise comparisons, etc.

In the submission it was not mentioned anything about data availability. However, all data and scripts will be uploaded with the VOR if it is required.

Recommendations for the authors:

I found your study interesting and potentially promising. However, there are some fundamental problems with the study design and the hypothesis, including:

<(1) Seasonality simulation - Seasonality is strongly associated with time, so it is unusual to simulate seasonal factors without accounting for time. The actual factors associated with seasonal change in reproductive output may be neither a difference in host blood meal nor temperature and photoperiod. It is therefore, odd to reduce seasonality to a difference in photoperiod and temperature in summer and autumn without even mentioning the time of year when the experiment was carried (except for the mention of February as the time the stock samples were collected from the wild).

The temperature and photoperiod settings are established according to a representative day in both autumn and summer. To determine these settings, we utilized climate data spanning a 3-year period (2020-2022), encompassing the most frequently occurring temperatures and day lengths. The weather conditions remained notably consistent throughout this time frame, which is why the specific year was not mentioned. Moreover, including the year in laboratory experiment details is uncommon, as evident in various papers. This practice can be corroborated by referring to multiple sources (cited in the original manuscript). We mention this in the new version.

(2) Hypothesis - *While the hypothesis alludes to the 'reason' for seasonal host shift, the prediction is on the outcome of the interaction between blood meal type and season.*

*It might be nicer to frame your hypothesis to be consistent with the aim, which is, testing the partial contributions of blood meal type, versus photoperiod and temperature to seasonal change in the reproductive output of *Culex quinquefasciatus*. A hypothesis like that can be accompanied by alternative predictions according to the expected individual and interactive effects of both factors.*

It was rewritten in the revised version to be consistent with our predictions and findings.

*Blood meal type, temperature, and photoperiod are all components of seasonality, so the strength of the study is its potential to decouple the effect of blood meal type from that of temperature and photoperiod on the seasonal reproductive output of *Culex quinquefasciatus* by comparing the two blood meal types under simulated summer and winter conditions. Ideally, this should have been over a natural summer and winter because a natural time difference captures the effect of other seasonal factors other than temperature and photoperiod.*

*Furthermore, the hypothesis stemmed from field observations, while the study itself was conducted under laboratory conditions using a local population of *Culex quinquefasciatus* from Argentina. It remains uncertain whether there is supporting evidence for a seasonal shift in host usage in *Culex quinquefasciatus* from the stock population. Discussing the field observations within the stock population would provide valuable insights.*

It was considered in the new version.