



A review on interspecific competition and population dynamics of mosquitoes

Diksha Raghav¹, Gauravi Yadav¹, Anil Kumar^{*1}, Hridayesh Arya¹,

¹Department of Zoology, N.R.E.C. College, Khurja, Bulandshahr. Affiliated to Chaudhary Charan Singh University, Meerut, Uttar Pradesh, India.

*Corresponding Author E-mail: anilkashyapnrec@gmail.com

DOI: <https://doi.org/10.59436/jsiane.381.2583-2093>

Abstract

As carriers of several deadly illnesses, including dengue, chikungunya, malaria, the Zika virus, and lymphatic filariasis, mosquitoes are hematophagous insects of major medical significance. They are members of the Culicidae family and have intricate life cycles and ecological relationships that affect their capacity for survival and spread. The taxonomy, disease correlations, and vectorial potential of the main mosquito genera—*Aedes*, *Culex*, and *Anopheles*—are examined in this paper. It also looks at population control techniques, emphasizing the drawbacks of chemical pesticides, including their damage to the environment, development of resistance, and non-target impacts, and advocating for environmentally benign substitutes including microbiological agents and herbal larvicides. A significant focus is placed on interspecific competition, particularly at the larval stage, where mosquito species compete for food, space, and oxygen. These interactions influence morphological traits such as wing length and adult body size, directly affecting survival, fecundity, and vector competence. Environmental variables, including both biotic (predators, microbial flora) and abiotic (temperature, pH, resource availability) factors, further modulate these outcomes. The review synthesizes data on how interspecific and intraspecific competition impact mosquito population dynamics, larval development, and disease ecology. Understanding these multidimensional interactions is crucial for implementing sustainable vector control strategies. By integrating ecological principles with innovative, environmentally responsible technologies, future mosquito management can be both effective and ecologically sound.

Keywords: Mosquito-borne diseases, Interspecific competition, Vector ecology, larval development, insecticide resistance.

Received 23.03.2025

Revised 18.04.2025

Accepted 13.06.2025

Online Available 20.06.2025

Introduction

There are more than 3,500 species of mosquitoes worldwide, particularly in tropical and subtropical areas, making them one of the most significant insect carriers. Mosquitoes are members of the class Insecta, phylum Arthropoda, kingdom Animalia, and suborder Nematocera. The main human disease vectors, *Anopheles*, *Aedes*, and *Culex*, are members of the subfamilies Anophelinae and Culicinae, which are subfamilies of the family Culicidae. Their importance stems from their ability to spread deadly illnesses like dengue, chikungunya, malaria, Zika virus, and lymphatic filariasis, especially in areas with little resources (WHO, 2023).

Vector competence depends on species biology, environmental conditions, and human interactions (Becker *et al.*, 2010). Urbanization and climate change are expanding mosquito habitats and increasing disease risk (Kraemer *et al.*, 2019). Larval competition—often overlooked—is crucial in mosquito ecology. When species share breeding sites, they compete for limited resources, affecting development and vector potential. For example, *Aedes albopictus* often outcompetes *Aedes aegypti* in shared environments (Juliano & Lounibos, 2005).

While chemical control methods are still widely used, issues like resistance and ecological damage have led to the exploration of alternatives such as microbial agents and plant-based larvicides (Ghosh *et al.*, 2012). A detailed understanding of mosquito taxonomy, ecology, and interspecific interactions is essential for designing sustainable vector control strategies.

Taxonomic Diversity

There are three subfamilies within the Culicidae family:

1) *Anophelinae*: This family includes the only human malaria vectors, *Anopheles* spp.

2) *Culex*, *Aedes*, *Mansonia*, and other species are members of the Culicinae family. Zika and dengue viruses are mostly spread by *Aedes aegypti* and *Aedes albopictus*.

3) *Toxorhynchitinae*: A distinct subfamily of mosquitoes with non-hematophagous members and predatory larvae that frequently feed on other mosquito larvae.

Life Cycle and Habitat:

Mosquitoes have four different life stages: egg, larva, pupa, and adult. They develop holometabolically. Every immature stage is aquatic, usually growing in bodies of water that are still or move slowly. Depending on the species, eggs might be placed single or in rafts. Often referred to as "wrigglers," mosquitoes have four unique life stages: egg, larva, pupa, and adult. They develop holometabolically. Every immature stage is aquatic, usually growing in bodies of water that are still or move slowly. They lay eggs. Larvae commonly called wrigglers, and pupae ("tumbler") remain in water and are vulnerable to environmental changes and predators.

Even though only a small percentage of the approximately 3,500 species of mosquitoes are vectors, they are crucial in the transmission of infectious

diseases, especially in tropical areas. *Aedes* (dengue, Zika, chikungunya, yellow fever), *Culex* (filariasis, West Nile virus, Japanese encephalitis), and *Anopheles* (malaria) are important genera. Malaria alone causes around 200 million cases annually, while dengue affects about 390 million people each year. Diseases like Zika and chikungunya are associated with neurological and joint complications, and lymphatic filariasis leads to severe disability. These mosquito-borne illnesses not only threaten public health but also place heavy economic and social burdens, necessitating comprehensive and sustained control efforts.

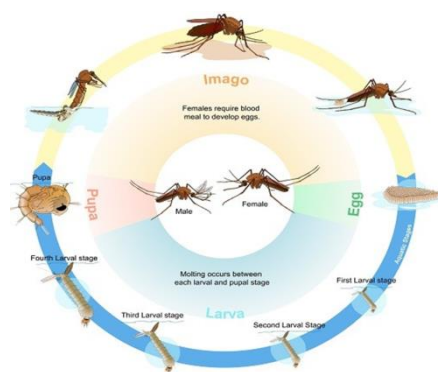


Figure 1. Insect life cycle.

(Source: <https://eu.biogents.com/life-cycle-mosquitoes/>)

Effect of Interspecific Competition on Population Dynamics of Mosquito.

Mosquito population dynamics are significantly shaped by interspecific competition, particularly in areas where several species coexist in the same habitat. Their populations are impacted by this ecological interaction in the following ways over time:

Alters Species Abundance and Distribution: -

Each mosquito species frequently gains dominance when several species vie for the same scarce resources (such as food, habitat, and breeding grounds). For example:

- *Aedes albopictus* is more competitive than *Aedes aegypti* in many environments, often leading to a decline or displacement of *Ae. aegypti* populations (Juliano & Lounibos, 2005).

- This displacement alters species composition in urban and suburban areas, potentially changing disease transmission patterns.

Influences Reproductive Success and Growth Rates: -

Competition at the larval stage reduces access to food and space, which in turn affects:

- Larval survival
- Developmental time
- Adult size and fecundity

Smaller adult mosquitoes resulting from intense competition tend to have lower reproductive success, reducing the population's growth rate (Noden *et al.*, 2016).

Drives Local Extinction or Coexistence: -

If one species consistently outcompetes another in a shared habitat, the weaker competitor may experience:

- Population decline
- Local extinction
- Or adapt by niche partitioning (using different resources or breeding at different times)

This dynamic can lead to unstable or shifting population equilibria, where dominant species vary based on environmental conditions and resource availability (Alto & Juliano, 2001).

Modifies Vectorial Capacity and Disease Risk: -

Changes in species dominance can affect which diseases are prevalent in a region. For example:

- Though often less effective than *Ae. aegypti*, *Ae. albopictus* is a capable vector for dengue and chikungunya.
- A shift in dominance from *Ae. aegypti* to *Ae. albopictus* may lead to changes in outbreak frequency or intensity (Juliano & Lounibos, 2005).

Affects Population Resilience and Recovery: -

Populations exposed to strong interspecific competition may:

- Recover more slowly after environmental disturbances (e.g., droughts, insecticide spraying)
- Become more vulnerable to additional stressors (e.g., predation or habitat loss)

This makes competitive interactions a key factor in mosquito population resilience and long-term stability.

Competition for Resources and Environmental Interactions: -

In watery environments, mosquito larvae frequently struggle for few resources like food, space, and opportunities for movement.

These interactions significantly influence their growth, development, and survival.

- Food:** Mosquito larvae feed on microorganisms and detritus. In resource-limited habitats, competition intensifies, leading to slower growth and smaller adult size (Gimnig *et al.*, 2002).

- Space and Locomotion:** Dense populations reduce physical space and oxygen availability, restricting movement and access to the water surface for respiration (Juliano & Lounibos, 2005).

- Biotic Interactions:** Presence of predators, parasites, or competing mosquito species (e.g., *Ae. albopictus* outcompeting *Ae. aegypti*) alters behavior and survival outcomes (Alto & Juliano, 2001).

- Abiotic Factors:** Environmental variables like temperature, pH, and water quality influence larval metabolism and modify the strength of competition. For instance, high temperatures can exacerbate food scarcity by increasing metabolic demands (Tun-Lin *et al.*, 2000).

These ecological pressures shape mosquito community structure, affect population dynamics, and influence disease transmission potential.

Interspecific Competition In Mosquitoes

Interspecific competition refers to the interaction between individuals of different species competing for the same limited resources, such as food, breeding sites, or habitat. In mosquitoes, this competition can significantly affect survival, development, and population dynamics, examples in mosquitoes:

Larval Stage Competition: -

Larvae of various mosquito species, including *Aedes aegypti*, *Aedes albopictus*, and *Culex quinquefasciatus*, may compete for the following in aquatic environments like ponds or containers:

- Microbial food sources
- Space
- Oxygen

For instance, it has been demonstrated that in certain container habitats, *Aedes albopictus* outcompetes *Aedes aegypti*, which causes the latter to be displaced in some areas (Juliano & Lounibos, 2005).

Adult Competition: -

Though less direct, adults of different species may compete for:

- Nectar or sugar sources
- Hosts for blood meals (especially in regions with limited host availability).

Ecological Impact:

- Alters species distribution in overlapping habitats
- Affects vector competence (i.e., ability to transmit disease)
- May reduce the overall population size of weaker competitors

Impact Of Interspecific Competition On Mosquito Morphology And Larval Morphology And Their Survival

Interspecific competition among mosquito larvae—particularly between species like *Aedes aegypti*, *Aedes albopictus*, and *Culex pipiens*—can significantly affect larval morphology, survival, and adult traits like :-

Impact on Mosquito Fitness

Interspecific competition among mosquito species significantly reduces larval survival, delays development, and leads to smaller adult sizes. For instance, *Anopheles* species showed decreased survival and delayed pupation when cohabiting with *Aedes aegypti*, consistent with previous studies on larval competition effects (Courret *et al.*, 2014; Alomar *et al.*, 2023; Romeo Aznar *et al.*, 2018).

Developmental Delay and Resource Limitation

Extended development time under competitive conditions is often due to limited food availability and chemical cues released by larvae. Growth-inhibiting factors released by *Ae. aegypti* have been reported to prolong *Anopheles* larval development (Moore & Whitacre, 1972; Bédhomme *et al.*, 2005). Additionally, increased physical contact within dense habitats may disrupt feeding behavior (Renshaw *et al.*, 1993).

Niche Partitioning through Foraging Behavior

Differences in larval positioning reduce direct competition. *Ae. aegypti* larvae forage near the container bottom, while *Anopheles* larvae feed at the surface, allowing spatial separation (Yee *et al.*, 2004; Van de Wolfshaar *et al.*, 2006). Such partitioning supports coexistence despite competition.

Morphological Effects of Competition

Crowding and limited resources during larval development result in smaller wing length and body size in adult mosquitoes. These morphological traits can impair flight, mating, and host-seeking efficiency, reducing vector competence (Gimnig *et al.*, 2002; Takken *et al.*, 1998; Suwanchaichinda & Paskewitz, 1998).

Cannibalism and Predation Dynamics

The observed disappearance of larvae suggests cannibalism or predation, particularly by *Ae. aegypti*, which develops faster and may prey on cohabitants. These behaviors have been reported even in the presence of adequate food, indicating facultative predation (Koenraadt & Takken, 2003; Muturi *et al.*, 2010; Shoukry, 1980).

Competitive Dominance of *Aedes aegypti*- Because of its higher foraging efficiency and quicker development, *Ae. Aegypti* outcompeted *An. arabiensis*, *An. gambiae*, and *An. funestus*. This is consistent with other research showing that *Ae. aegypti* outperforms other species when food is scarce (Armistead *et al.*, 2008; Yee & Juliano, 2006).

Coexistence and Ecological Distribution

Despite competitive disadvantages, *Anopheles* species can coexist with *Ae. aegypti* in urban and suburban habitats due to behavioral and spatial differentiation (Lawal *et al.*, 2011; Mahgoub *et al.*, 2017; Mbanzulu *et al.*, 2022). However, interspecific competition may still shift species dominance and affect local disease transmission patterns (Juliano *et al.*, 2004; Juliano & Lounibos, 2005).

Study Limitations and Future Directions

This semi-field study lacked natural predators and used second-instar laboratory-reared larvae, possibly limiting ecological validity. Cannibalism and predation were inferred from larval disappearance without molecular confirmation (e.g., PCR or gut analysis). Future studies should examine fitness outcomes such as fecundity, vector competence, and host-seeking behavior under natural environmental conditions (Alto & Bettinardi, 2015; Romeo Aznar *et al.*, 2018).

Larval Survival and Development Time: -

When different mosquito species share breeding sites, competition for food and oxygen often reduces survival and delays development. For instance, *Ae. albopictus* shows reduced emergence rates and longer development periods under competitive stress (Noden *et al.*, 2016). Similarly, *Ae. aegypti* experiences extended larval duration and lower survival when reared with *Anopheles stephensi* (Juliano & Lounibos, 2005).

Effects on Larval and Adult Morphology: -

Reduced Body Size and Wing Length

Increased larval competition often leads to smaller adults. *Aedes aegypti* and *Ae. albopictus* reared under crowded conditions develop smaller wing lengths and body sizes, impairing flight, mating, and host-seeking ability (Edgerly *et al.*, 1999; Noden *et al.*, 2016). *Anopheles gambiae* also shows reduced adult size when larval density increases (Gimnig *et al.*, 2002).

Morphological Changes in Larvae

Although less frequently documented, some studies suggest that intense larval competition can result in modified mouthparts or siphons, potentially altering feeding and breathing efficiencies (Relyea, 2002).

Carry-Over Effects on Adult Survival and Vector Competence: -

Adult Survival- Adult mosquitoes emerging from highly competitive larval environments tend to have shorter lifespans. For example, *Ae. aegypti* adults from dense larval groups showed reduced survival under dry condition (Alto & Bettinardi 2015)

Disease Transmission Potential

Smaller, nutritionally stressed mosquitoes may transmit disease differently. For instance, *Ochlerotatus triseriatus* showed altered susceptibility to

arboviruses under larval stress (Grimstad & Walker, 1991). Such effects vary by vector and pathogen.

Species-Specific Competitive Interactions: -

- *Aedes albopictus* typically outcompetes *Ae. aegypti*, especially in well-resourced habitats (Juliano & Lounibos, 2005).
- Interspecific competition between *Cx. pipiens* and *Ae. albopictus* reduces adult emergence and alters morphology (Noden et al., 2016).
- *Ae. aegypti* generally survives better than *An. stephensi* under food-limited co-rearing conditions (Juliano & Lounibos, 2005).

Ecological and Public Health Implications; -

Interspecific competition has various ecological and public health implications, like Larval competition may restructure mosquito communities and influence vector competence. Smaller, shorter-lived mosquitoes may bite less frequently or transmit disease differently. These ecological effects must be considered when planning vector control interventions (Juliano & Lounibos, 2005).

Table 1. Generalized linear mixed model for the effects of competition, habitat, and their interactions on the developmental time for *Aedes aegypti* and *Anopheles arabiensis*, *Aedes aegypti* and *Anopheles gambiae*, and *Aedes aegypti* and *Anopheles funestus*.

Population	Species	Effects	RR (95% CI)	p-Value
<i>Ae. aegypti</i> and <i>An. arabiensis</i>	<i>Ae. aegypti</i>	Competition	Alone	1
			Mixed	0.54 (0.25, 1.56)
		Habitat	Small	1
			Large	1.12 (0.54, 2.30)
	<i>An. arabiensis</i>	Competition	Alone	1
			Mixed	6.11 (2.59, 14.45)
		Habitat	Small	1
			Large	0.93 (0.44, 1.97)
<i>Ae. aegypti</i> and <i>An. gambiae</i>	<i>Ae. aegypti</i>	Competition	Alone	1
			Mixed	0.30 (0.12, 0.73)
		Habitat	Small	1
			Large	1.12 (0.49, 2.57)
	<i>An. gambiae</i>	Competition	Alone	1
			Mixed	1.74 (0.74, 4.09)
		Habitat	Small	1
			Large	1.57 (0.65, 3.80)
<i>Ae. aegypti</i> and <i>An. funestus</i>	<i>Ae. aegypti</i>	Competition	Alone	1
			Mixed	0.07 (0.02, 0.28)
		Habitat	Small	1
			Large	1.75 (0.81, 3.79)
	<i>An. funestus</i>	Competition	Alone	1
			Mixed	1.40 (0.95, 2.07)
		Habitat	Small	1
			Large	1.00 (0.50, 1.99)

RR stands for risk ratio. The interaction row was left out of the table because it didn't show any significant impacts, but it is still included in the text for clarity.

The experimental configuration of larval densities under various competitive conditions is displayed in Table 1 (Lushasi et al., 2025). The effects of

competition, habitats, and their interactions on the survival of larvae to adulthood for *Aedes aegypti*, *Anopheles arabiensis*, *Anopheles gambiae*, and *Anopheles funestus* are examined in Table 2 using a generalized linear mixed model.

Table 2 displays the effects of habitat size, interspecific and intraspecific competition, and their interaction on larval survival to adulthood

Population	Species	Effects	RR (95% CI)	p-Value
------------	---------	---------	-------------	---------

Population	Species	Effects	RR (95% CI)	p-Value
<i>Ae. aegypti</i> and <i>An. arabiensis</i>	<i>Ae. aegypti</i>	Competition	Intraspecific	1
			Interspecific	0.40 (0.30, 0.55) <0.001
		Habitat	Small	1
			Large	0.88 (0.66, 1.16) 0.359
	<i>An. arabiensis</i>	Competition	Intraspecific	1
			Interspecific	0.23 (0.15, 0.35) <0.001
		Habitat	Small	1
			Large	0.84 (0.56, 1.29) 0.441
<i>Ae. aegypti</i> and <i>An. gambiae</i>	<i>Ae. aegypti</i>	Competition	Intraspecific	1
			Interspecific	0.50 (0.34, 0.74) 0.001
		Habitat	Small	1
			Large	0.89 (0.63, 1.27) 0.55
	<i>An. gambiae</i>	Competition	Intraspecific	1
			Interspecific	0.43 (0.26, 0.71) 0.001
		Habitat	Small	1
			Large	0.82 (0.52, 1.28) 0.393
<i>Ae. aegypti</i> and <i>An. funestus</i>	<i>Ae. aegypti</i>	Competition	Intraspecific	1
			Interspecific	0.26 (0.17, 0.39) <0.001
		Habitat	Small	1
			Large	0.65 (0.45, 0.94) 0.901
	<i>An. funestus</i>	Competition	Intraspecific	1
			Interspecific	0.19 (0.13, 0.28) <0.001
		Habitat	Small	1
			Large	1.02 (0.69, 1.52) 0.024

Note: RR stands for risk ratio. The interaction row was left out of the table because it had no discernible impacts, but it is still there in the text to give a clear explanation.

For *Aedes aegypti*, *Anopheles arabiensis*, *Anopheles gambiae*, and *Anopheles funestus*, Table 2 displays the effects of habitat size, interspecific and intraspecific competition, and their interaction on larval survival to adulthood (Lushasi *et al.*, 2025).

Table 3. Generalized linear model of the effects of competition, habitat, and their interactions on the adults' wing length (mm) for *Aedes aegypti* mixed with either *Anopheles arabiensis*, *Anopheles gambiae*, or *Anopheles funestus*.

Population	Species	Effects	RR (95% CI)	p-Value
<i>Ae. aegypti</i> and <i>An. arabiensis</i>	<i>Ae. aegypti</i>	Competition	Alone	1
			Mixed	0.76 (0.72, 0.80) <0.001
		Habitat	Small	1
			Large	1.16 (1.09, 1.22) <0.001
		Competition × Habitat	Mixed × Large	
				1.21 (1.11, 1.31) <0.001

Population	Species	Effects	RR (95% CI)	p-Value
<i>Ae. aegypti</i> and <i>An. arabiensis</i>		Competition	Alone	1
			Mixed	0.47 (0.44, 0.50) <0.001
	<i>An. arabiensis</i>	Habitat	Small	1
			Large	1.13 (1.08, 1.19) <0.001
		Competition × Habitat	Mixed × Large	1.61 (1.48, 1.74) <0.001
<i>Ae. aegypti</i> and <i>An. gambiae</i>		Competition	Alone	1
			Mixed	0.80 (0.76, 0.84) <0.001
	<i>Ae. aegypti</i>	Habitat	Small	1
			Large	0.20 (1.15, 1.26) <0.001
		Competition × Habitat	Mixed × Large	1.04 (0.98, 1.12) 0.196
		Competition	Alone	1
			Mixed	0.56 (0.54, 0.58) <0.001
	<i>An. gambiae</i>	Habitat	Small	1
			Large	1.09 (1.04, 1.14) <0.001
		Competition × Habitat	Mixed × Large	0.29 (0.21, 1.37) <0.001
<i>Ae. aegypti</i> and <i>An. funestus</i>		Competition	Alone	1
			Mixed	0.77 (0.73, 0.82) <0.001
	<i>Ae. aegypti</i>	Habitat	Small	1
			Large	1.22 (1.17, 1.29) <0.001
		Competition × Habitat	Mixed × Large	1.13 (1.05, 1.22) 0.001
		Competition	Alone	1
			Mixed	0.58 (0.56, 0.60) <0.001
	<i>An. funestus</i>	Habitat	Small	1
			Large	1.18 (1.14, 1.22) <0.001
		Competition × Habitat	Mixed × Large	0.28 (1.21, 1.35) <0.001

Note: RR = risk ratio.

For *Aedes aegypti* co-raised with *Anopheles arabiensis*, *Anopheles gambiae*, or *Anopheles funestus*, Table 3 shows the effects of competition, habitat size, and their interaction on mature wing length (mm) (Lushasi et al., 2025).

Table 4: *Aedes aegypti*, *Anopheles arabiensis*, *Anopheles gambiae*, and *Anopheles funestus* cannibalistic and predacious behavior in relation to competition, food, and habitats using a generalized linear mixed model

Population	Species	Effects	RR (95% CI)	p-Value
------------	---------	---------	-------------	---------

Population	Species	Effects	RR (95% CI)	p-Value
<i>Ae. aegypti</i> and <i>An. arabiensis</i>	<i>Ae. aegypti</i>	Competition	Alone	1
			Mixed	0.54 (0.38, 0.79)
		Habitat	Small	1
			Large	1.15 (0.82, 1.62)
		Food	No	1
			Yes	0.001 (0.0001, 0.005)
	<i>An. arabiensis</i>	Competition	Alone	1
			Mixed	8.24 (4.91, 13.83)
		Habitat	Small	1
			Large	1.28 (0.79, 2.06)
		Food	No	1
			Yes	0.13 (0.07, 0.21)
<i>Ae. aegypti</i> and <i>An. gambiae</i>	<i>Ae. aegypti</i>	Competition	Alone	1
			Mixed	0.49 (0.36, 0.66)
		Habitat	Small	1
			Large	0.86 (0.64, 1.16)
		Food	No	1
			Yes	0.02 (0.01, 0.03)
	<i>An. gambiae</i>	Competition	Alone	1
			Mixed	6.35 (4.34, 9.29)
		Habitat	Small	1
			Large	1.16 (0.83, 1.63)
		Food	No	1
			Yes	0.16 (0.11, 0.24)
<i>Ae. aegypti</i> and <i>An. funestus</i>	<i>Ae. aegypti</i>	Competition	Alone	1
			Mixed	0.71 (0.49, 1.03)
		Habitat	Small	1
			Large	0.98 (0.67, 1.45)
		Food	No	1
			Yes	0.01 (0.003, 0.013)
	<i>An. funestus</i>	Competition	Alone	1
			Mixed	14.09 (8.55, 23.22)
		Habitat	Small	1
			Large	1.99 (1.27, 3.11)

Population	Species	Effects	RR (95% CI)	p-Value
		No	1	<0.001
		Food	0.26 (0.16, 0.42)	

Table 4 summarizes the percentage of missing larvae (indicative of predation or cannibalism) across treatments with and without food (Lushasi *et al.*, 2025).

Conclusion

- Interspecific competition significantly affects mosquito population dynamics, influencing survival, development, reproduction, and species distribution through resource limitation and ecological overlap (Juliano & Lounibos, 2005).
- *Aedes albopictus* often outcompetes *Aedes aegypti* in shared habitats due to its adaptability and faster development, altering vector dominance and disease risk in various regions (Juliano *et al.*, 2004).
- Environmental elements that affect competition intensity include temperature, food availability, and habitat size to niche partitioning, coexistence, or local extinction (Alto & Juliano, 2001; Tun-Lin *et al.*, 2000).
- Larval competition results in smaller adult mosquitoes, which affects fecundity, flight ability, host-seeking behavior, and ultimately vector competence (Noden *et al.*, 2016; Gimnig *et al.*, 2002).
- Cannibalism and predation are frequent under competitive stress, especially in resource-limited environments, and may further influence species survival and community structure (Muturi *et al.*, 2010; Koenraadt & Takken, 2003).
- Understanding these ecological interactions is essential for vector control, as competition-driven shifts in mosquito populations can directly impact disease transmission and the effectiveness of public health interventions (Harrington *et al.*, 2008; WHO, 2023).

Reference

- Alomar, A., Chelbi, I., Hsairi, M., Zhioua, E., & Bouattour, A. (2023). Interspecific larval competition between *Culex pipiens* and *Aedes albopictus* (Diptera: Culicidae) under laboratory and semi-field conditions in Tunisia. *Parasites & Vectors*, 16(1), 1–13. <https://doi.org/10.1186/s13071-023-05606-y>
- Alto, B. W., & Bettinardi, D. (2015). Temperature and density effects on survival and mosquito susceptibility to arboviral infection. *Journal of Medical Entomology*, 52(6), 1033–1038.
- Alto, B. W., & Juliano, S. A. (2001). Precipitation and temperature effects on populations of *Aedes albopictus* (Diptera: Culicidae): Implications for range expansion. *Journal of Medical Entomology*, 38(5), 646–656.
- Armistead, J. S., Nishimura, N., Arias, J. R., & Lounibos, L. P. (2008). Interspecific larval competition between *Culex quinquefasciatus* and *Culex nigripalpus* (Diptera: Culicidae) in Florida. *Journal of Medical Entomology*, 45(4), 629–637. <https://doi.org/10.1093/jmedent/45.4.629>
- Barrera, R. (1996). Competition and resistance to starvation in larvae of container-inhabiting *Aedes* mosquitoes. *Ecological Entomology*, 21(2), 117–127. <https://doi.org/10.1111/j.1365-2311.1996.tb01176.x>
- Bédhomme, S., Agnew, P., Sidobre, C., & Michalakos, Y. (2005). Kin selection, sib competition and the evolution of virulence. *Heredity*, 94(1), 73–79. <https://doi.org/10.1038/sj.hdy.6800579>
- Becker, N., Petric, D., Zgomba, M., Boase, C., Madon, M., Dahl, C., & Kaiser, A. (2010). *Mosquitoes and their control* (2nd ed.). Springer.
- Bhatt, S., Gething, P. W., Brady, O. J., Messina, J. P., Farlow, A. W., Moyes, C. L., ... & Hay, S. I. (2013). The global distribution and burden of dengue. *Nature*, 496(7446), 504–507. <https://doi.org/10.1038/nature12060>
- Clements, A. N. (1999). *The biology of mosquitoes: Volume 2. Sensory reception and behaviour*. CABI Publishing.
- Couret, J., Dotson, E., & Benedict, M. Q. (2014). Temperature, larval diet, and density effects on development rate and survival of *Aedes aegypti* (Diptera: Culicidae). *PLoS ONE*, 9(2), e87468. <https://doi.org/10.1371/journal.pone.0087468>
- Edgerly, J. S., Willey, M. S., & Livdahl, T. (1999). Intraguild predation among larval treehole mosquitoes, *Aedes albopictus*, *Aedes aegypti*, and *Aedes triseriatus* (Diptera: Culicidae), in laboratory microcosms. *Journal of Medical Entomology*, 36(3), 394–399.
- Fernandes, L., & Briegel, H. (2005). Reproductive physiology of *Anopheles gambiae* and *An. atroparvus*. *Journal of Vector Ecology*, 30(1), 11–18. <https://www.sove.org/SOVE%20folder/journal/journal.htm>
- Ghosh, A., Chowdhury, N., & Chandra, G. (2012). Plant extracts as potential mosquito larvicides. *Indian Journal of Medical Research*, 135(5), 581–598.
- Gimnig, J. E., Ombok, M., Otieno, S., Kaufman, M. G., Vulule, J. M., & Walker, E. D. (2002). Density-dependent development of *Anopheles gambiae* (Diptera: Culicidae) larvae in artificial habitats. *Journal of Medical Entomology*, 39(1), 162–172.
- Grimstad, P. R., & Walker, E. D. (1991). *Aedes triseriatus* (Diptera: Culicidae) and La Crosse virus. IV. Nutritional deprivation of larvae affects the adult barriers to infection and dissemination. *Journal of Medical Entomology*, 28(3), 378–386.
- Harrington, L. C., Edman, J. D., & Scott, T. W. (2008). Why do female *Aedes aegypti* (Diptera: Culicidae) feed preferentially and frequently on human blood? *Journal of Medical Entomology*, 38(3), 411–422. <https://doi.org/10.1603/0022-2585-38.3.411>
- Hemingway, J., Hawkes, N. J., McCarrroll, L., & Ranson, H. (2002). The molecular basis of insecticide resistance in mosquitoes. *Insect Biochemistry and Molecular Biology*, 34(7), 653–665.
- Hoffmann, A. A., Montgomery, B. L., Popovici, J., Iurbe-Ormaetxe, I., Johnson, P. H., Muzzi, F., ... & O'Neill, S. L. (2011). Successful establishment of *Wolbachia* in *Aedes* populations to suppress dengue transmission. *Nature*, 476(7361), 454–457.
- Ho, B. C., Ewert, A., & Chew, L. M. (1989). Interspecific competition among tropical *Aedes* mosquitoes. *Population Ecology*, 31(1), 135–144. <https://doi.org/10.1007/BF02515419>
- Isman, M. B. (2006). Botanical insecticides, deterrents, and repellents in modern agriculture and an increasingly regulated world. *Annual Review of Entomology*, 51, 45–66.
- Jaga, K., & Dharmani, C. (2003). Sources of exposure to and public health implications of organophosphate pesticides. *Reviews on Environmental Health*, 18(2), 115–128.
- Juliano, S. A., & Lounibos, L. P. (2005). Ecology of invasive mosquitoes: Effects on resident species and on human health. *Ecology Letters*, 8(5), 558–574. <https://doi.org/10.1111/j.1461-0248.2005.00755.x>
- Juliano, S. A., Lounibos, L. P., & O'Meara, G. F. (2004). A field test for competitive effects of *Aedes albopictus* on *Aedes aegypti* in South Florida: Differences between sites of coexistence and exclusion? *Oecologia*, 139(4), 583–593. <https://doi.org/10.1007/s00442-004-1540-7>
- Kraemer, M. U. G., Reiner, R. C., Brady, O. J., Messina, J. P., Gilbert, M., Pigott, D. M., ... & Hay, S. I. (2019). Past and future spread of the arbovirus vectors *Aedes aegypti* and *Aedes albopictus*. *Nature Microbiology*, 4(5), 854–863.
- Lawal, O. A., Adeleke, M. A., Olatunji, A. O., & Sam-Wobo, S. O. (2011). Observation on *Anopheles* and *Aedes* mosquitoes cohabiting breeding containers in some urban areas of Abeokuta, Southwestern Nigeria. *Journal of Vector Borne Diseases*, 48(3), 157–160. <https://pubmed.ncbi.nlm.nih.gov/21985802/>
- Leishnam, P. T., & Juliano, S. A. (2012). Impacts of climate, land use, and biological invasion on the ecology of immature *Aedes* mosquitoes: Implications for La Crosse emergence. *EcoHealth*, 9(2), 217–228.
- Lizuin, A. A., Stein, M., & Almirón, W. R. (2022). Interspecific competition between *Aedes aegypti* and *Aedes albopictus* (Diptera: Culicidae): A review. *Journal of Medical Entomology*, 59(4), 1359–1370. <https://doi.org/10.1093/jme/tjac056>
- Lyimo, E. O., & Takken, W. (1993). Effects of adult body size on fecundity and the pre-gravid rate of *Anopheles gambiae* females in Tanzania. *Medical and Veterinary Entomology*, 7(4), 328–332. <https://doi.org/10.1111/j.1365-2515.1993.tb00701.x>
- Mahgoub, M. M., Sulaiman, S. M., & Elaigip, A. H. (2017). Seasonal abundance and breeding habitats of *Aedes aegypti* in urban areas of Khartoum State, Sudan. *Tropical Biomedicine*, 34(4), 792–802. <https://pubmed.ncbi.nlm.nih.gov/29641925/>
- Mbanzulu, K. M., Mapatano, M. A., & Mavoko, H. M. (2022). Co-breeding and spatio-temporal dynamics of *Aedes aegypti* and *Anopheles* mosquitoes in Kinshasa, Democratic Republic of the Congo. *Parasites & Vectors*, 15(1), 1–12. <https://doi.org/10.1186/s13071-022-05485-4>
- Moore, C. G., & Whitacre, D. M. (1972). Competition in mosquitoes. 2. Production of inferior *Culex tarsalis* (Diptera: Culicidae) adults from larvae raised with *Culex pipiens quinquefasciatus*. *Annals of the Entomological Society of America*, 65(4), 914–918. <https://doi.org/10.1093/aesa/65.4.914>
- Murrell, E. G., & Juliano, S. A. (2008). Detritus type alters the outcome of interspecific competition between *Aedes aegypti* and *Aedes albopictus* (Diptera: Culicidae). *Journal of Medical Entomology*, 45(3), 375–383. <https://doi.org/10.1093/jmedent/45.3.375>
- Muturi, E. J., Costanzo, K., Kesavaraju, B., & Alto, B. W. (2010). Can pesticides stress alter mosquito predator-prey interactions? *Ecotoxicology*, 19(7), 1475–1481. <https://doi.org/10.1007/s10646-010-0536-5>
- Ng'habi, K. R., Knols, B. G., Killeen, G. F., & Ferguson, H. M. (2005). Predicting the productivity of mosquito larval habitats using landscape and remotely sensed indicators. *Malaria Journal*, 4, 7. <https://doi.org/10.1186/1475-2875-4-7>
- Noden, B. H., O'Neal, P. A., Fader, J. E., & Juliano, S. A. (2016). Impact of inter- and intra-specific competition among mosquito larvae on adult longevity, size, and emerging population patterns. *Ecological Entomology*, 41(1), 76–86.
- Paaijmans, K. P., Blanford, S., Bell, A. S., Blanford, J. I., Read, A. F., & Thomas, M. B. (2009). Influence of climate on malaria transmission depends on daily temperature variation. *Proceedings of the National Academy of Sciences*, 107(34), 15135–15139. <https://doi.org/10.1073/pnas.1006422107>
- Petersen, L. R., Jamieson, D. J., Powers, A. M., & Honein, M. A. (2016). Zika virus. *New England Journal of Medicine*, 374(16), 1552–1563.
- Pocock, M. J. (2007). Can traits predict species' vulnerability to extinction? A test with New Zealand birds. *Animal Conservation*, 10(3), 199–207. <https://doi.org/10.1111/j.1469-1795.2007.01118.x>
- Ranson, H., N'Guegan, R., Lines, J., Moiroux, N., Nkuni, Z., & Corbel, V. (2011). Pyrethroid resistance in African anopheline mosquitoes: What are the implications for malaria control? *Trends in Parasitology*, 27(2), 91–98.
- Relyea, R. A. (2002). The many faces of predation: How induction, selection, and thinning combine to alter prey phenotypes. *Ecology*, 83(7), 1953–1964.
- Renshaw, M., Service, M. W., & Birley, M. H. (1993). Density-dependent regulation of *Aedes cantans* (Diptera: Culicidae) in natural and artificial populations. *Ecological Entomology*, 18(3), 223–229. <https://doi.org/10.1111/j.1365-2311.1993.tb01198.x>
- Romeo Aznar, V., Alto, B. W., & Lounibos, L. P. (2018). The effects of larval competition on the fitness of *Aedes aegypti* and *Aedes albopictus* mosquitoes. *Medical and Veterinary Entomology*, 32(1), 70–77. <https://doi.org/10.1111/mve.12266>
- Rueda, L. M., Patel, K. J., Axtell, R. C., & Stinner, R. E. (1990). Temperature-dependent development and survival rates of *Culex quinquefasciatus* and *Aedes aegypti* (Diptera: Culicidae). *Journal of Medical Entomology*, 27(5), 892–898. <https://doi.org/10.1093/jmedent/27.5.892>
- Suleman, M. (1982). Competition between *Aedes aegypti* Linnaeus and *Aedes albopictus* Skuse (Diptera: Culicidae) in the laboratory. *Mosquito News*, 42(3), 366–371.
- Tun-Lin, W., Burkot, T. R., & Kay, B. H. (2000). Effects of temperature and larval diet on development rates and survival of the dengue vector *Aedes aegypti* in north Queensland, Australia. *Medical and Veterinary Entomology*, 14(1), 31–37. <https://doi.org/10.1046/j.1365-2915.2000.00207.x>
- Vinogradova, E. B. (2000). *Culex pipiens pipiens* mosquitoes: Taxonomy, distribution, ecology, physiology, genetics, applied importance and control. Pensoft Publishers.
- World Health Organization. (2020). Vector-borne diseases. <https://www.who.int/news-room/fact-sheets/detail/vector-borne-diseases>
- Yee, D. A., & Juliano, S. A. (2006). Consequences of detritus type in an aquatic microsystem: Effects on water quality, micro-organisms and performance of the dominant consumer. *Freshwater Biology*, 51(3), 448–459. <https://doi.org/10.1111/j.1365-2427.2005.01509.x>
- Yee, D. A., & Juliano, S. A. (2007). Abundance matters: A field experiment testing the more-individuals hypothesis for richness-productivity relationships. *Oecologia*, 153(1), 153–162. <https://doi.org/10.1007/s00442-007-0713-1>