

Research Paper

Cite this article: Bengoa M, Rotger A, Luzón R, Barceló C (2021). Larvae ecology and adult activity of *Aedes mariae* (Diptera: Culicidae) in a touristic rock-pool area of the Balearic Islands (Western Mediterranean). *Bulletin of Entomological Research* 1–8. <https://doi.org/10.1017/S0007485321001024>

Received: 7 February 2021
Revised: 12 September 2021
Accepted: 5 November 2021

Keywords:

Abundance; algae; human landing; Majorca
pH; salinity; Spain; tourism

Author for correspondence:

Carlos Barceló, Email: carlos.barcelo@uib.es

Larvae ecology and adult activity of *Aedes mariae* (Diptera: Culicidae) in a touristic rock-pool area of the Balearic Islands (Western Mediterranean)

Mikel Bengoa¹, Andreu Rotger², Raúl Luzón³ and Carlos Barceló⁴ 

¹Anticimex, Department of Research, Innovation and Development, Barcelona, Spain; ²Animal Demography and Ecology Unit, GEDA – IMEDEA (CSIC/UIB), Esporles, Spain; ³Vectobal, Department of Development and Science, Palma, Spain and ⁴Applied Zoology and Animal Conservation Group, UIB, Palma, Spain

Abstract

Mosquitoes are vectors of several diseases of medical concern such as malaria or dengue and can also negatively affect tourism and the life-quality of the neighbourhood. The species *Aedes mariae* (Sergeant and Sergeant, 1903) is a poorly studied mosquito that breeds in rock-pools of the Mediterranean coast. General Linear Mixed Models (GLMM) were used to determine drivers affecting the presence and abundance of this species. Abiotic and biotic factors were recorded in rock-pools with the presence of *Ae. mariae* sub-adults across a supralittoral area of Majorca Island (Balearic Islands, Spain) from July 2018 to June 2019. We tested how abiotic factors affected the presence of larvae, while the biotic factors were used to check their effect on larvae abundance. human landing collection was also conducted to assess the adult activity of this species. Valuable data were recorded to improve our knowledge about the bioecology of *Ae. mariae* in a touristic area of the island of Majorca. Salinity and pH were the most explanatory variables for the presence of *Ae. mariae* larvae. The presence of *Posidonia oceanica* (L.) Delile 1813 leaves negatively affected the abundance of *Ae. mariae* larvae while the presence of other fauna enhanced it. Adult females of *Ae. mariae* were active for 26 min after sunset in June and its host-seeking activity decreased during autumn months. Control methods against this species should be focussed on rock-pools and planning treatments according to tides, waves and precipitation.

Introduction

The relationship between humans and mosquitoes (Diptera: Culicidae) has been turbulent since the origin of humanity (Becker *et al.*, 2010). The World Health Organization (WHO) considers this group of insects as the deadliest animals in the world for their capacity to transmit several pathogens such as malaria, dengue and yellow fever. In addition, mosquitoes can also reduce neighbourhood life-quality and the host-seeking behaviour of anthropophilic species can also have a negative impact on the tourism industry (UNDP, 2017).

Mosquito sub-adult stages can breed in a wide range of aquatic habitats (Clements, 1999). These habitats are usually changing environments where the presence and abundance of mosquito larvae are modulated by environmental factors of their breeding sites (Juliano, 2009; Yee *et al.*, 2010).

Most of the tourist residences in the Spanish Mediterranean coast are located near natural wetlands, where mosquitoes are commonly present. Knowledge of these sites is important to understand the drivers that could increase the risk of nuisance biting by adult mosquitoes or potential outbreaks of vector-borne diseases (Burroni *et al.*, 2013; Cardo *et al.*, 2013; Estallo *et al.*, 2018).

From a total of 65 species of mosquitoes recorded in Spain, 17 species are present in the Balearic Islands (Robert *et al.*, 2019). One of them is the Asian tiger mosquito, *Aedes albopictus* (Skuse, 1894), an invasive mosquito with strong anthropophilic behaviour (Muñoz *et al.*, 2011). However, other opportunistic species can also feed on humans in areas nearby to their habitats. These species (e.g. *Aedes caspius* (Pallas, 1771) and *Aedes mariae* (Sergeant and Sergeant, 1903)) usually breed in natural areas (e.g. wetlands, salt marshes and rock-pools) affecting human settlements within their flight range (Rioux *et al.*, 1968; Porretta *et al.*, 2007).

Currently, *Ae. mariae* is not a vector of human diseases (Ribeiro *et al.*, 1988) but can play a role in the transmission of avian malaria, *Plasmodium relictum* (Grassi and Feletti, 1891) (summarized in Schaffner *et al.* (2001)). This species was first recorded in the Balearic Islands in 2011 (Bueno-Marí and Jiménez-Peydró, 2011) and belongs to the *Ae. mariae* complex (Coluzzi and Sabatini, 1968). This complex includes other two sibling species: *Aedes zammitii* (Theobald, 1907) and *Aedes phoeniciae* (Coluzzi and Sabatini, 1968). *Aedes mariae* is mainly restricted to

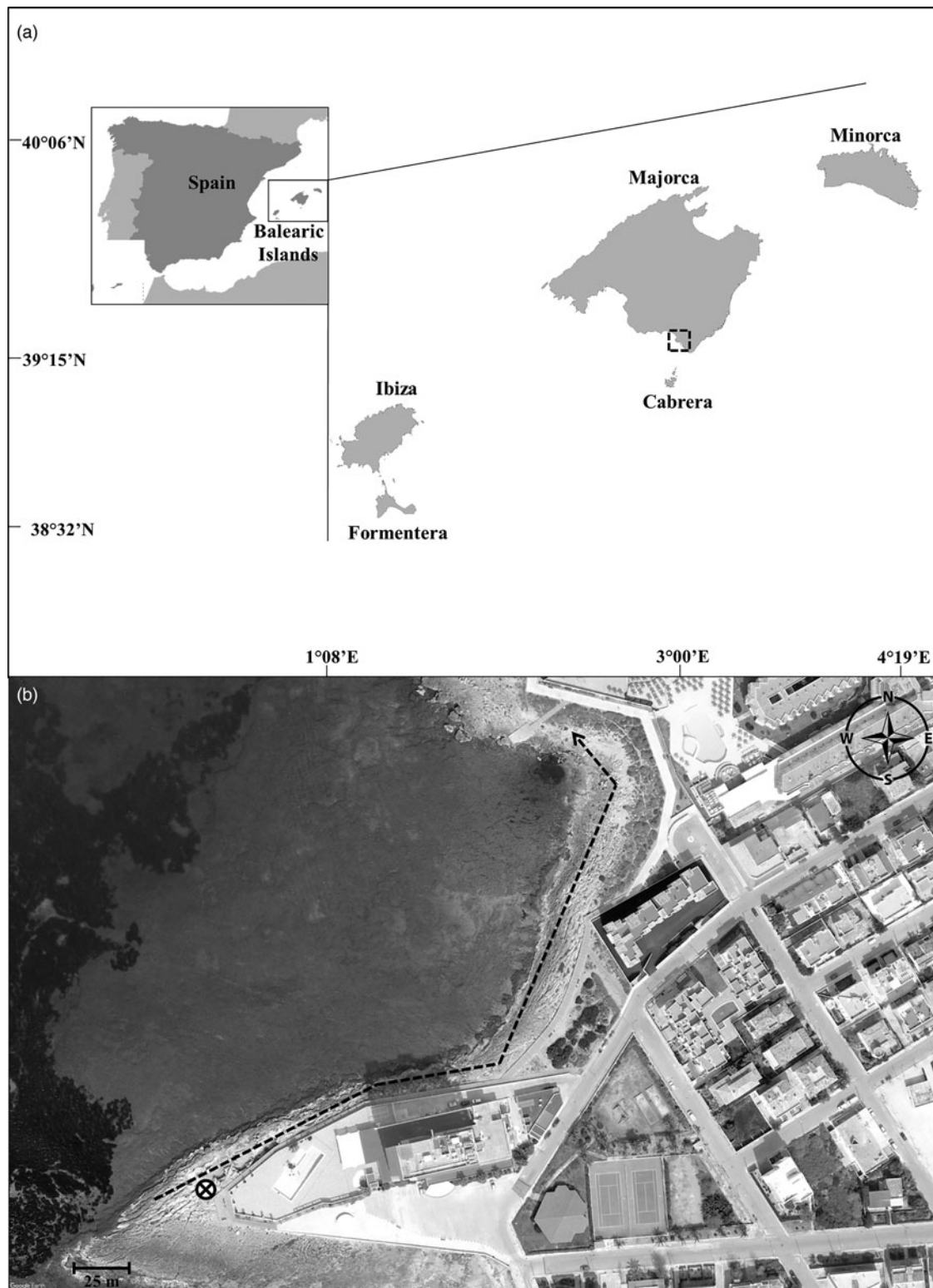


Figure 1. (a) Map of the Balearic Islands. Dotted square indicates the municipality of the study (*Sa Colònia de Sant Jordi*). (b) Location of the study. The dotted arrow indicates the transect along the supralittoral rock-pools. The cross icon indicates the HLC site.

the Western Mediterranean basin but in Spain can be also recorded in Huelva and Cantabria (Bueno-Marí and Serna-Mompeán, 2015). *Aedes zammitii* and *Ae. phoeniciae* are distributed within central and Eastern Mediterranean coasts (Coluzzi and Sabatini, 1968; Coluzzi *et al.*, 1974; Mastrantonio *et al.*, 2015). Immature stages

of these species complex breed in supralittoral rock-pools of the Mediterranean basin (Schaffner *et al.*, 2001; Mastrantonio *et al.*, 2015) tolerating wide ranges of salinity concentration (Rioux, 1958).

Previous research, such as that of Margalef (1949), studied the biology of *Ae. mariaae*; however, these studies mainly focused on

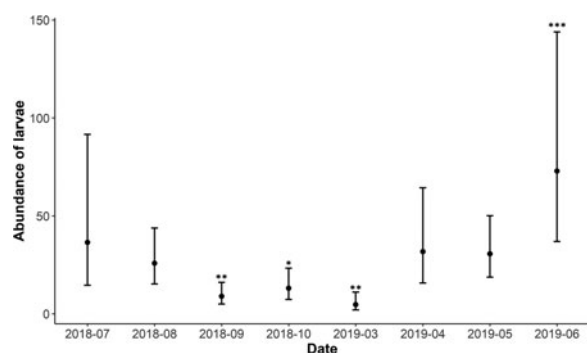


Figure 2. Phenology of the estimated abundance of *Ae. mariae* during 2018 and 2019. Significance codes with respect to July 2018: *** $P < 0.001$; ** $P = 0.01$; * $P = 0.05$.

molecular systematics, hybridization, natural selection and evolution of the *Ae. mariae* complex (Urbanelli *et al.*, 2014; Mastrantonio *et al.*, 2015; Rosenfeld *et al.*, 2018; Delgado-Serra *et al.*, 2021), or related to the larvae bioecology of the sibling species *Ae. phoeniciae* (Rosenfeld *et al.*, 2019). Hence, there is a lack of information about adult activity and abiotic/biotic requirements for *Ae. mariae* sub-adult stages.

The aim of this study was to determine the factors affecting the abundance and presence of *Ae. mariae* focusing on its breeding sites near a hotel area of Majorca island. In addition, its host-seeking activity was also evaluated. This information could be useful for predicting mosquito responses and improve future control strategies.

Materials and methods

Area of study

Majorca is the largest island of the Balearic archipelago (3640 km²) with 896,000 inhabitants in 2019. The island is an important tourist destination, recording 16.5 million visitors in 2019 (IBESTAT, 2020). Its climate is characterized by hot dry summers and wet winters. Mean annual temperatures ranged from 27.7°C during the hottest month (August) to 5.0°C during the coldest month (February). The mean annual precipitation on the island is 411 mm (AEMET, 2020).

The study was conducted in *Sa Colònia de Sant Jordi* municipality located in the South of Majorca island (39°19'00"N 2°59'34"E) (fig. 1a). This coastal municipality has 2692 inhabitants and an important tourist destination with more than 3500 hotel rooms (FEHM, 2019). In fact, one of the most visited beaches in Majorca, *Es trenc*, is located in this municipality.

Mosquito nuisance biting is a common problem during the high tourism season, reducing the quality of outdoor activities and neighbourhood lifestyle. This nuisance biting is commonly related to adults of four different species: the wetland mosquito *Ae. caspius* that produce high population densities when the soil phreatic surface increases; the common mosquito *Culex pipiens* s. l. (Linnaeus, 1758) from unattended swimming pools in single-family houses; the Asian tiger mosquito *Ae. albopictus* with a population increasing year by year (Sanz-Aguilar *et al.*, 2018), and the target species of the current study, *Ae. mariae*, in residential areas nearby supralittoral rock-pools.

Analysis of *Ae. mariae* breeding sites

A maximum of 15 rock-pools with the presence of *Ae. mariae* immature stages were randomly selected throughout a transect

of 350 m across the supralittoral area (fig. 1b). These rock-pools were weekly checked from 27/07/2018 to 16/11/2018 (12 consecutive weeks) and from 22/03/2019 to 07/06/2019 (8 consecutive weeks).

Three main factors were determined in each rock-pool: the number of individuals of *Ae. mariae* immature stages (larvae and pupae) hereafter called 'larvae', abiotic parameters and biotic factors. The physical and chemical (abiotic) parameters analysed were: pH, temperature (°C), total dissolved solids (TDS) (ppt), conductivity (µS) and salinity (g l⁻¹). These parameters were determined with a multiparametric PC-5 tester (XS Instruments, Italy). The biotic factors were determined by observation according to the authors' criteria. These factors were: algae cover (0: no algae, 1: 25% covered by algae, 2: 50% covered by algae, 3: 75% covered by algae and 4: all the rock-pool covered by algae); presence of other fauna (1: presence, 2: absence) and presence of the seagrass *Posidonia oceanica* (L.) leaves (1: presence, 2: absence).

The abundance of larvae per rock-pool was estimated to describe the variations of immature stages of this species during the year and how this abundance may vary by month. This analysis was carried out using a Generalized Linear Mixed Model (GLMM) with a zero-inflated Poisson model to handle excess zero counts, considering rock-pools sampled on the same date as a random effect in order to avoid pseudo replication. Models were run using the R Development Core Team (2017) package 'glmmTMB' (Brooks *et al.*, 2017).

In order to define a viability range of this species during their larval stage, we investigated how water physicochemical characteristics (abiotic parameters) affect the presence or absence of larvae. This analysis was carried out using GLMM with the R package 'lme4' (Bates *et al.*, 2015) with binomial distribution (logit link function), being the sample date of the rock-pool the random effect to consider the time variance. To avoid multicollinearity problems with the highly related abiotic variables such as conductivity, salinity and TDS, the variance inflation factor (VIF) was previously checked to detect multicollinearity in regression analysis (Dodge, 2008). VIF ranges from 1 upwards showing what percentage the variance (i.e. the standard error squared) is inflated for each coefficient. Following Ringle *et al.* (2015), all variables with a VIF value >5 were immediately discarded. The best model indicated that the three abiotic factors tested (salinity, temperature and pH) had an important effect on the presence of larvae (table S1).

The number of larvae in the rock-pool may be associated with competition or predation and resource availability, hence the association between larval abundance of *Ae. mariae* and biotic factors were assessed. The number of larvae presented by each rock-pool was considered as an abundance measure. This analysis was carried out using a GLMM with a Poisson distribution (log link function) considering as well the sample date of the rock-pool as a random effect to consider the time variability. Abundance was influenced by the three biotic factors as indicated the best model (table S2).

Model selection in all analyses was performed using the Akaike's information criterion (AIC), considering models to be equivalent when the difference in AIC with the best model (Δ AIC) was <2 (Burnham and Anderson, 2004; tables S1 and S2).

Adult mosquito sampling

Human landing captures (HLC) were performed weekly by two authors of the current study: MB and CB (same two authors

Table 1. Variation in the abiotic factors of the sampled rook-pools and variation in the mean number of larvae depending on the biotic factors tested

Abiotic factors	Mean $\pm$ SD	Min	Max
pH	8.41 $\pm$ 0.64	5.54	10
Conductivity	70.19 $\pm$ 40.10	1.1	166
Temperature	26.44 $\pm$ 5.76	14.5	41.8
TDS	49.25 $\pm$ 28.29	1	104.5
Biotic factors	Mean number of larvae $\pm$ SD	Min	Max
Algae cover			
No algae	2.88 $\pm$ 11.11	0	60
25% covered	12.17 $\pm$ 34.63	0	300
50% covered	33.37 $\pm$ 68.29	0	300
75% covered	63.83 $\pm$ 100.13	0	400
100% covered	25.83 $\pm$ 47.58	0	120
Other fauna			
Presence	16.74 $\pm$ 54.63	0	400
Absence	28.03 $\pm$ 51.00	0	200
<i>Posidonia oceanica</i> leaves			
Presence	17.65 $\pm$ 53.73	0	400
Absence	29.10 $\pm$ 55.52	0	250

during all collection events) from 28/06/2019 to 29/10/2019. People sampled mosquitoes from each other with a mouth aspirator (Silver, 2008; WHO, 2013) from 30 min before sunset till 1 h after sunset, remaining 1:30 min exposed. The time of capture (minutes after the sunset) for each mosquito was recorded.

We examined the relationship between time (in minutes) when females of *Ae. mariae* landed after the sunset and the month of the year (June to October 2019) using a General Linear Model (GLM). Finally, we examined the variation in landing times after sunset and between months.

Ethical considerations

The human collectors consent to participate in the current study. During the study period, no transmission of mosquito-borne diseases was reported by local authorities for the location in which the study was conducted. Human landing collections were conducted in an area without registered cases of vector-borne diseases. The species sampled are not a potential vector of human pathogens.

Results

Breeding sites

Between July 2018 and June 2019, a total of 282 rock-pools were analysed recording a total of 4861 larvae of *Ae. mariae*. Some rock-pools were without water or without larvae, therefore the number of rock-pools checked varied between 12 and 15 per sampling period. June 2019 was peak larval abundance (67.84 $\pm$ 0.47 SD) showing a significant difference in comparison to the first sampling month ($P < 0.0001$). The lowest larval abundance was recorded in March, September and October (fig. 2). These three months also showed significant differences ($P_{\text{September}} = 0.004$,

$P_{\text{October}} = 0.03$ and $P_{\text{March}} < 0.0001$) to July 2018. November was discarded from the model due to the low number of observations (only three larvae in the study area).

The abiotic values varied between months (table 1). The physical and chemical characteristics of conductivity, salinity and TDS in the rock-pools water were highly correlated since they measure similar properties. Using VIF value, conductivity and TDS were discarded, which presented VIF values of 11.35 and 11.45, respectively. Algae cover of 75% in rock-pools showed the maximum number of larvae as the presence of other fauna (see below) and the presence of *P. oceanica* leaves (table 1).

Salinity, temperature and pH had a great effect on the presence of larvae, with salinity and pH the most important variables (fig. 3 and figs S1 and S3). Both variables had a positive effect on the presence of larvae in the rock-pools, however, salinity was a limiting factor since high concentration of salt in the rock-pools (>75 g l⁻¹) drastically reduce the probability of larval presence.

The three biotic factors considered had a strong effect on the abundance of larvae. A relatively high cover of algae (75%) was important to maintain a high abundance of larvae; however, this abundance decreased when algae covered the entire bottom of the rock-pool or when there was no algae coverage (fig. 4). Other fauna recorded in the rock-pools consisted of insects from orders Coleoptera and Diptera but no other mosquito species were recorded in rock-pools. This fauna observed had a significantly positive effect on the abundance of the larvae of *Ae. mariae* ($P < 0.0001$), while the presence of *Posidonia* leaves had the contrary effect, significantly decreasing the number of larvae ($P < 0.0001$; fig. 4).

Adult activity

A total of 32 female *Ae. mariae* were collected by HLC. Only three individuals of *Ae. caspius* were recorded on October 29 but were

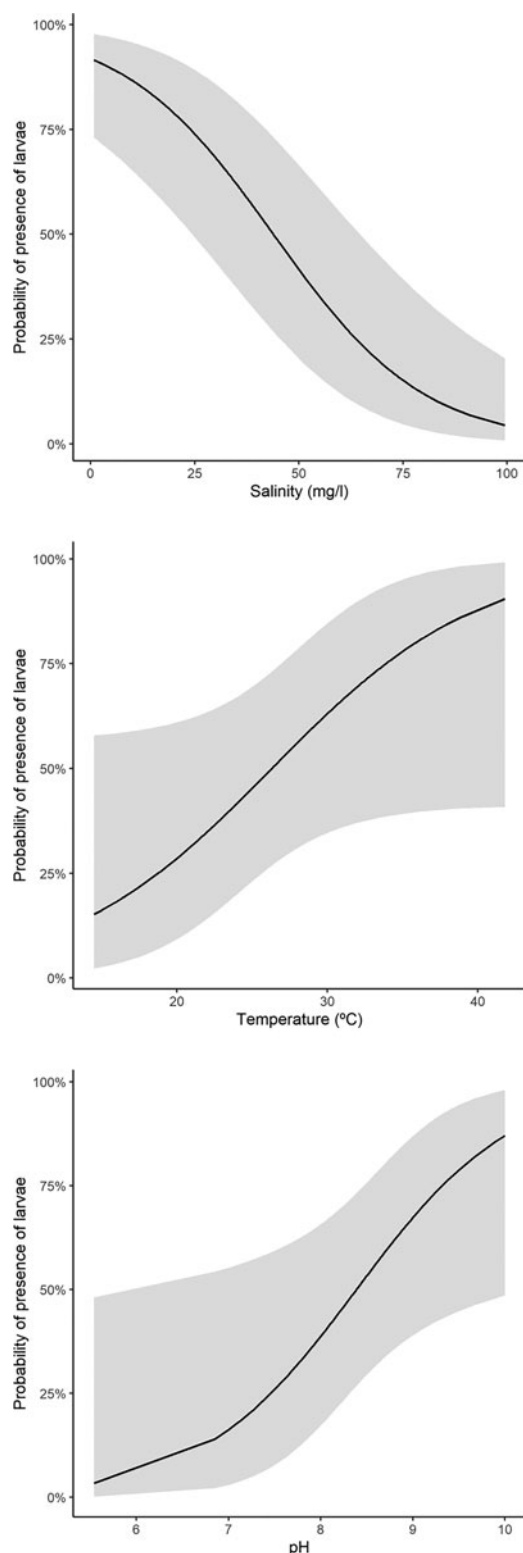


Figure 3. Relationship between larvae presence in the rock-pools and abiotic factors. Salinity (on the top), temperature (middle), pH (on the bottom). In shaded grey the confidence intervals.

not included in the analysis. Results showed that *Ae. mariaae* activity was recorded up to 43 min after sunset (fig. 5). During the first HLC on June 28, females remained active for a period of 26 min after sunset (from minute 17 to minute 43), while on

October 29, the activity was concentrated for 10 min after sunset (from minute 12 to minute 22). In fact, there were significant differences between the timing of *Ae. mariaae* activity (GLM (Poisson); $P < 0.0001$), showing a reduction from 17 min after the sunset in June to 9 min in October. No HLC were recorded in September since the weather was not favourable for adult mosquito flight.

Discussion

Breeding sites of *Ae. mariaae* immature stages were highly heterogeneous since variables vary from one month to another. These kinds of habitats are continuously altered due to rainfall, storms, waves, tides, insolation, etc., changing its environmental conditions (e.g. food source, microorganism communities, drying and physicochemical characteristics). The influence of these abiotic drivers is important since climate is becoming more suitable for mosquitos due to climate change (Coumou and Rahmstorf, 2012; Campbell-Lendrum *et al.*, 2015), including the increase of salinity of the Mediterranean Sea (Ozer *et al.*, 2016).

The seasonal activity of this species occurred from March to October (Becker *et al.*, 2010). An increase in the larval population in spring and the highest abundance were recorded in June during the current study. The best model for larvae presence included variables of temperature, salinity and pH.

Seawater has 37.5 g l^{-1} of salt and could increase in rook-pools due to evaporation and sea sprays, especially during months with high insolation. In addition, rock-pools with larvae present can be filled with rainwater, seawater or mixed, showing values ranging from 1 to 93.4 g l^{-1} , highlighting the adaptive capability of this species. Rioux (1958) found *Ae. mariaae* larvae in rock-pools with salinity up to 200 g l^{-1} , but we have not recorded these values in pools with larvae; however, we found a strong relationship between the number of observed larvae and the water temperature and salinity measured in the rock-pools (figs S1 and S2). High values of salinity reduced the probability of larval presence, similar to the results obtained with the sibling species *Ae. phoeniciae* (Rosenfeld *et al.*, 2019). Interestingly, we observed a complete absence of larvae when salt was solidified on the surface of rock-pools. We hypothesize that this salt layer prevents larvae from breathing through the surface layer of the pool.

A mild positive effect of pH was recorded in our study, with *Ae. mariaae* capable of tolerating a wide range of pH (fig. S3). Mosquito larvae can tolerate pH values ranging from 3 to 11 (Clements, 1992). In fact, pH values could determine the species composition. Previous studies observed that *Ae. phoeniciae* and species from genus *Culex* have a preference for basic pH values (Grech *et al.*, 2019; Rosenfeld *et al.*, 2019) while others recorded negative or neutral effects (Burroni *et al.*, 2007; Gardner *et al.*, 2013).

Regarding the biotic factors, the presence of algae, *Posidonia* leaves and other fauna play an important role to maintain the presence of larvae in rock-pools. Margalef (1949) studied the ecological limitations of this species and found that the absence of algae in the rock-pools was also a limiting factor, as did Rosenfeld *et al.* (2019) for *Ae. phoeniciae* breeding sites. Algae in these sites could be a food source for mosquito larvae or even a shelter for other organisms that favour the presence of *Ae. mariaae*. Conversely, a high coverage of algae decreased the number of larvae. This phenomenon could be related to an increase of other dipteran larvae that share a similar diet with *Ae. mariaae* (e.g. Chironomids) (Naeem, 1988). This increase of

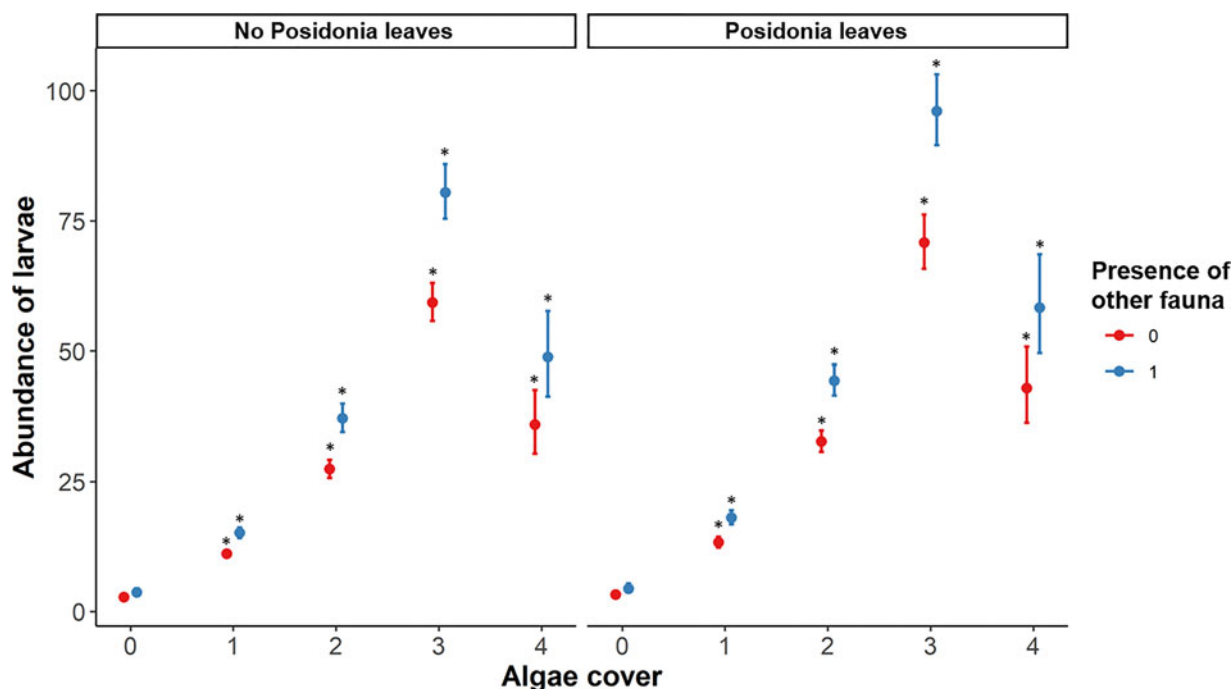


Figure 4. Relationship between the number of larvae found in the rock-pools with the algae cover, and presence of fauna and *Posidonia* leaves. (*): when P values ≤ 0.05 . Bars indicate standard deviation.

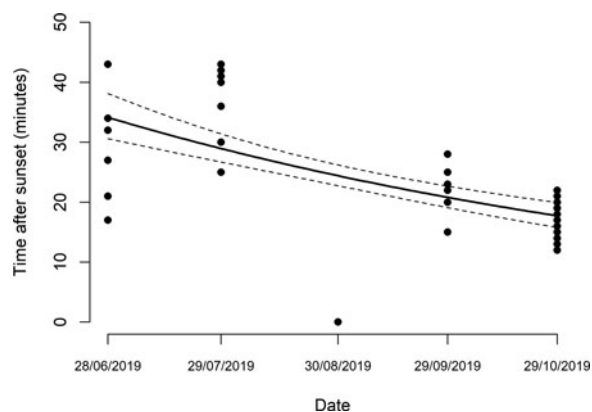


Figure 5. Number of adult *Ae. mariaae* collected with human landing captures (HLC). Each dot is a collected mosquito. n , number of mosquitoes landed in each sampling session.

dipteran larvae could also attract other fauna that can colonize the rock-pool and be potential predators or competitors of *Ae. mariaae* larvae (Kolasa and Romanuk, 2005). The presence of *Posidonia* leaves in the rock-pools was a factor that slightly decreased the abundance of larvae, suggesting that floating *Posidonia* leaves could reduce the water surface for larvae breathing.

Margalef (1949) also found a possible predation competence between larvae and other fauna. Fauna found in breeding sites of the current study were mainly specimens of Coleoptera of genus *Ochthebius* (Leach, 1815), Chironomidae and shore flies (family Ephydriidae) which can compete or predate on *Ae. mariaae* larvae (Linthicum *et al.*, 1985; Rosenfeld *et al.*, 2019). However, it was determined that the presence of other fauna had a positive effect on the abundance of *Ae. mariaae* larvae suggesting a

successful co-inhabit of these species in the rock-pool ecosystem (Rosenfeld *et al.*, 2019). Further studies addressing this issue must be recommended to better understand the degree of these interactions.

The effort to perform HLC was justified due to the number of mosquitoes captured and the information obtained. It is important to highlight the reduction of time when this species is actively seeking hosts in accordance with the photoperiod, being aggressive during 26 min after sunset in June and a narrower interval of only 10 min in October. Previous studies demonstrated that winter diapause of *Ae. mariaae* was determined by the photoperiod (Coluzzi *et al.*, 1975), however there is a lack of information about the relation of the photoperiod with the host-seeking behaviour of this species. Studies with *Ae. albopictus* and *Aedes aegypti* (L., 1762) recorded that this reduction could be determined by the short sunset and lower temperature during autumn months (Camara, 2010; Costanzo *et al.*, 2015).

This is the first study of the bioecology of *Ae. mariaae* in the Mediterranean basin including a wide number of environmental drivers, thus, valuable data were obtained to improve our knowledge about the activity of this species. Control methods against *Ae. mariaae* should focus on their biology and the heterogeneity of the larvae habitat. Integrated mosquito control is the best approach to reduce the nuisance to tourists and citizens, and a reduction of their larval habitat should be prioritized. The biotic and abiotic variations observed in this study should be considered in the planning of control methods, taking into account tides, waves and precipitation. Hence, the application of *Bacillus*-based formulations is the control method recommended by the authors. Control methods like physical barriers (polydimethylsiloxane, perlite, etc.) or filling these pools with expanding foam like in tree holes (CDC, 2016) were not recommended due to the aforementioned heterogeneity of this environment. Finally, the role of *Ae. mariaae* as a vector of pathogens is poorly studied,

therefore further studies addressing this subject should be considered in order to evaluate the importance of its host-seeking activity.

Supplementary material. The supplementary material for this article can be found at <https://doi.org/10.1017/S0007485321001024>

Acknowledgements. The authors thank the council of the municipality of Sa Colònia de Sant Jordi for letting us perform the present study.

Conflict of interest. None.

References

- Agencia Estatal de Meteorología (AEMET) (2020) *Valores climatológicos normales en Palma de Mallorca*. Retrieved from AEMET website http://www.aemet.es/es/serviciosclimaticos/datosclimatologicos/efemerides_extremos?w=0&k=bal&l=B278&datos=det&x=B278&m=13&v=todos2 (Accessed 29 December 2020).
- Bates D, Mächler M, Bolker B and Walker S (2015) Fitting linear mixed-effects models using lme4. *Journal of Statistical Software* 67, 1–48.
- Becker N, Petric D, Zgomba M, Boase C, Madon MB, Dahl C and Kaiser AA (2010) *Mosquitoes and Their Control*, 2nd Edn, Berlin: Springer-Verlag.
- Brooks ME, Kristensen K, van Benthem KJ, Magnusson A, Berg CW, Nielsen A, Skaug HJ, Maechler M and Bolker BM (2017) glmmTMB: Balances speed and flexibility among packages for Zero-inflated Generalized Linear Mixed Modeling. *The R Journal* 9(2), 378–400.
- Bueno-Marí R and Jiménez-Peydró R (2011) First confirmed record of *Ochlerotatus mariae* (Sergeant & Sergeant, 1903) in the Balearic Islands (Spain) and its significance in local mosquito control programmes. *European Mosquito Bulletin* 29, 82–87.
- Bueno-Marí R and Serna-Mompeán JP (2015) Primera cita de *Ochlerotatus mariae* (Sergeant & Sergeant, 1903) (Diptera, Culicidae) en la Comunidad Valenciana: implicaciones en el control de esta especie singular. *Revista gaditana de Entomología* 6, 63–66.
- Burnham KP and Anderson DR (2004) Multimodel inference understanding AIC and BIC in model selection. *Sociological Methods & Research* 33, 261–304.
- Burroni N, Loetti V, Freire G, Jensen O and Schweigmann N (2007) New record and larval habitats of *Culex eduardoi* (Diptera: Culicidae) in an irrigated area of Patagonia. *Argentina. Memórias do Instituto Oswaldo Cruz* 102(2), 237–240.
- Burroni NE, Loetti MV, Marinone MC, Freire MG and Schweigmann NJ (2013) Larval habitat of *Ochlerotatus albifasciatus* (Diptera: Culicidae) in the southern edge of the Americas, Tierra del Fuego Island. *Open Journal of Animal Sciences* 3, 5–10.
- Camara TN DL (2010) Activity patterns of *Aedes aegypti* and *Aedes albopictus* (Diptera: Culicidae) under natural and artificial conditions. *Oecologia Australis* 14, 737–744.
- Campbell-Lendrum D, Manga L, Bagayoko M and Sommerfeld J (2015) Climate change and vector-borne diseases: what are the implications for public health research and policy? *Philosophical Transactions of the Royal Society B: Biological Sciences* 370, 1–8.
- Cardo MV, Vezzani D and Carabajo AE (2013) The role of the landscape in structuring immature mosquito assemblages in wetlands. *Wetlands Ecology and Management* 21, 55–70.
- Centre for Disease Control and Prevention (CDC) (2016) *What you need to know about filling tree holes*. Retrieved from CDC website <https://www.cdc.gov/zika/pdfs/TreeHoles-FactSheet.pdf> (Accessed 21 December 2020).
- Clements AN (1992) *The Biology of Mosquitoes Volume 1: Development, Nutrition and Reproduction*. London: Chapman and Hall.
- Clements AN (1999) *The Biology of Mosquitoes. Volume 2: Sensory Reception and Behaviour*. Wallingford: CABI publishing.
- Coluzzi M and Sabatini A (1968) Morphological divergences and sterility barriers in the complex of *A. mariae*. *Rivista di Parassitologia* 29, 49–70.
- Coluzzi M, Sabatini A, Bullini L and Ramsdale C (1974) Further data on the distribution of the species of the *mariae* complex of the genus *Aedes*. *Rivista di Parassitologia* 35, 321–330.
- Coluzzi M, Di Deco M and Gironi A (1975) Effect of the photoperiod of the site of oviposition of *Aedes mariae* (Diptera, Culicidae). *Parassitologia* 17, 121.
- Costanzo KS, Schelble S, Jerz K and 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.
- Coumou D and Rahmstorf S (2012) A decade of weather extremes. *Nature Climate Change* 2, 491–496.
- Delgado-Serra S, Viader M, Ruiz-Arroondo I, Miranda MA, Barceló C, Bueno-Marí R, Hernández-Triana LM, Miquel M, Lester K, Jurado JA and Paredes-Esquivel C (2021) Molecular characterization of mosquito diversity in the Balearic Islands. *Journal of Medical Entomology* 58, 608–615.
- Dodge Y (2008) *The Concise Encyclopedia of Statistics*. University of Neuchâtel, Switzerland: Springer & Bussines media.
- Estallo EL, Sangermano F, Grech M, Ludueña-Almeida F, Frías-Céspedes M, Ainete M, Almirón W and Livdahl T (2018) Modelling the distribution of the vector *Aedes aegypti* in a central Argentine city. *Medical and Veterinary Entomology* 32, 451–461.
- Federación Empresarial Hotelera de Mallorca (FEHM) (2019) *Colònia de Sant Jordi*. Retrieved from FEHM website <http://www.fehm.info/asociaciones/colonia-sant-jordi> (Accessed 29 December 2020).
- Gardner AM, Anderson TK, Hamer GL, Johnson DE, Varela KE, Walker ED and Ruiz MO (2013) Terrestrial vegetation and aquatic chemistry influence larval mosquito abundance in catch basins, Chicago, USA. *Parasites & Vectors* 6, 1–11.
- Grech MG, Manzo LM, Epele LB, Laurito M, Claverie AN, Ludueña-Almeida FF, Miserendino ML and Almirón WR (2019) Mosquito (Diptera: Culicidae) larval ecology in natural habitats in the cold temperate Patagonia region of Argentina. *Parasites and vectors* 12(1), 1–14.
- Institut d'Estadística de les Illes Balears (IBESTAT) (2020) Flujo de turistas (FRONTUR). Retrieved from IBESTAT website <https://www.caib.es/ibestat/estadistiques/economia/turisme/fluxe-turistes-frontur/043d7774-cd6c-4363-929a-703aaa0cb9e0> (Accessed 29 December 2020).
- Juliano SA (2009) Species interactions among larval mosquitoes: context dependence across habitat gradients. *Annual Review of Entomology* 54, 37–56.
- Kolasa J and Romanuk TN (2005) Assembly of unequals in the unequal world of a rock pool metacommunity. In Holyoak M, Leibold MA and Holt RD (eds), *Metacommunities: Spatial Dynamics and Ecological Communities*. Chicago: University of Chicago Press, pp. 212–232.
- Linthicum KJ, Davies FG and Kamau J (1985) Predation on emerging adult mosquitoes by *Brachydeutera munroi* (Diptera: Ephydriidae). *Proceedings of the New Jersey Mosquito Control Association annual meeting*, USA.
- Margalef R (1949) Sobre la ecología de las larvas del mosquito *Aedes mariae*. *Publicaciones del Instituto de Biología Aplicada* 6, 83–102.
- Mastrantonio V, Porretta D, Bellini R, Nascetti G and Urbanelli S (2015) Molecular systematics and origin of the Mediterranean Sea rock-pool mosquitoes of the *Aedes mariae* (Diptera: Culicidae) complex. *Annals of the Entomological Society of America* 108, 593–599.
- Muñoz J, Eritja R, Alcaide M, Montalvo T, Sorriquer SC and Figuerola J (2011) Host-feeding patterns of native *Culex pipiens* and invasive *Aedes albopictus* mosquitoes (Diptera: Culicidae) in urban zones from Barcelona, Spain. *Journal of Medical Entomology* 48, 956–960.
- Naeem S (1988) Predator-prey interactions and community structure: chironomids, mosquitoes and copepods in *Heliconia imbricata* (Musaceae). *Oecologia* 77, 202–209.
- Ozer T, Gertman I, Kress N, Silverman J and Herut B (2016) Inter-annual thermohaline and nutrient dynamics (2002–2014) in the Levantine surface and intermediate water masses. *Rapport Communication Internationale Mer Méditerranée* 41, 131.
- Porretta D, Canestrelli D, Bellini R, Celli G and Urbanelli S (2007) Improving insect pest management through population genetic data: a case study of the mosquito *Ochlerotatus caspius* (Pallas). *Journal of Applied Ecology* 44, 682–691.
- R Development Core Team (2017) *R: A Language and Environment for Statistical Computing*. Vienna, AT: R Foundation for Statistical Computing. Retrieved from R website <http://www.R-project.org/>.
- Ribeiro H, Ramos HC, Pires CA and Capela RA (1988) An annotated checklist of the mosquitoes of continental Portugal (Diptera: Culicidae). In III

- Congreso Ibérico de Entomología: actas. Granada, Universidad de Granada, pp. 233–253.
- Ringle CM, Wende S and Becker JM (2015) *SmartPLS 3*. Bönningstedt: SmartPLS. Retrieved from SmartPLS website <http://www.smartpls.com>.
- Rioux JA (1958) Les culicides du midi Méditerranéen. Etude systématique et écologique. *Encyclopédie Entomologique* 35, 303.
- Rioux JA, Croset H, Corre JJ, Simonneau P and Gras G (1968) Phyto-ecological basis of mosquito control: cartography of larval biotopes. *Mosquito News* 28, 572–582.
- Robert V, Günay F, Le Goff G, Boussès P, Sulesco T, Khalin A, Medlock JM, Kampen H, Petric D and Schaffner F (2019) Distribution chart for Euro-Mediterranean mosquitoes (western Palaearctic region). *Journal of European Mosquito Control Association* 37, 1–28.
- Rosenfeld S, Porretta D, Rahav E, Mastrantonio V, Duchet C and Blaustein L (2018) Molecular identification of *Aedes phoeniciae* (Diptera: Culicidae) in rock-pools along the northern Israeli coast. *Journal of Vector Ecology* 43, 344–346.
- Rosenfeld S, Blaustein L, Kneitel J, Duchet C, Horwitz R, Rybak O, Polevikov A and Rahav E (2019) The abundance and larval performance of *Aedes phoeniciae* in supralittoral rock-pools. *Hydrobiologia* 846, 181–192.
- Sanz-Aguilar A, Rosselló R, Bengoa M, Ruiz-Pérez M, González-Calleja M, Barceló C, Borràs D, Paredes-Esquível C, Miranda MA and Tavecchia G (2018) Water associated with residential areas and tourist resorts is the key predictor of Asian tiger mosquito presence on a Mediterranean island. *Medical and Veterinary Entomology* 32, 443–450.
- Schaffner F, Angel G, Geoffroy B, Hervy JP, Rhaïem A and Brunhes J (2001) *The Mosquitoes of Europe: An Identification and Training Programme*. Montpellier: IRD editors.
- Silver JB (2008) *Mosquito Ecology: Field Sampling Methods*, 3rd Edn. Dordrecht: Springer.
- United Nations Development Programme (UNDP) (2017) *A Socio-economic Impact Assessment of the Zika Virus in Latin America and the Caribbean: With a Focus on Brazil, Colombia and Suriname* (UNDP Synthesis report), Apr 3. New York, USA: United Nations Development Programme.
- Urbanelli S, Porretta D, Mastrantonio V, Bellini R, Pieraccini G, Romoli R, Crasta G and Nascetti G (2014) Hybridization, natural selection, and evolution of reproductive isolation: a 25-years survey of an artificial sympatric area between two mosquito sibling species of the *Aedes mariaae* complex. *Evolution* 68, 3030–3038.
- World Health Organization (WHO) (2013) *Malaria Entomology and Vector Control. WHO Guide for Participants*. Geneva, Switzerland: WHO.
- Yee DA, Kneitel JM and Juliano SA (2010) Environmental correlates of abundances of mosquito species and stages in discarded vehicle tires. *Journal of Medical Entomology* 47, 53–62.