Investigating the role of urban vegetation alongside other

environmental variables in shaping *Aedes albopictus*

presence and abundance in Montpellier, France

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Abstract

Urban greening helps address urbanization challenges and provides health benefits, but it may also favor mosquito species, vectors of pathogens causing human diseases. This study examines the relationship between urban vegetation and the presence and abundance of *Aedes albopictus* in Montpellier, the second greenest city of France, while accounting for meteorology, microclimate, air quality, human socio-demography, and landscape. From May to October 2023, adult mosquitoes were collected monthly in urban parks, residential areas, and the impervious city center using BG-Pro traps with odor and CO_2 attractants. Microclimate data were recorded on-site. Vegetation, land cover, meteorological, air quality, and human demographic data were gathered from open-access databases. *Ae. albopictus* presence and abundance were analysed according to environmental variables taken at different time lags and spatial distances using a two-stage modeling approach: bivariate analyses using generalized linear mixed models were conducted to select variables for inclusion in a multivariate random forest model, aiming to identify the factors that best explain *Ae. albopictus* presence and abundance.

While urban vegetation had a limited effect on *Ae. albopictus* presence, number of patches and the percentage of area covered by low vegetation were among the most important predictors of abundance. The main predictors for presence were minimum hourly temperature (24h-48h before sampling), minimum atmospheric pressure during sampling, and the weekly cumulated rainfall recorded six weeks before sampling. The most important predictors of abundance were maximum hourly temperatures during sampling, cumulated growing degree days (GDD) during two weeks preceding sampling and the length and patch

46 size of roads. To our knowledge, this is the first study examining urban vegetation's influence

47 on *Ae. albopictus* in France, offering insights for urban planning and suggesting further

48 research on vegetation's role in mosquito-borne disease transmission, particularly in the

49 context of increasing dengue incidence in Europe.

Author summary

Urban greening is often promoted as a solution to the challenges posed by increasing

urbanization, offering numerous environmental and societal benefits. However, it may also

create favorable conditions for mosquito vector species such as *Aedes albopictus*. Only two

studies in Europe—both conducted in Italy—have investigated the impact of urban vegetation

on the presence and abundance of *Ae. albopictus*. This study explores this relationship in

Montpellier, the second greenest city in France, while considering meteorology, microclimate,

air quality, human socio-demographic factors, and landscape characteristics. Using a two-

stage modeling approach, we first performed bivariate analyses with generalized linear mixed

models to select relevant variables, which were then integrated into a multivariate random

forest model to identify the key predictors of *Ae. albopictus* presence and abundance. Our

results indicate that urban vegetation does not significantly influence *Ae. albopictus* presence

but is among the most important predictors of its abundance. These findings provide valuable

insights for urban planning and highlight the need for further research on the role of

vegetation in mosquito-borne disease transmission.

Introduction

According to Hussain and Imitiaz [1], urbanisation is a complex, dynamic, and evolving social process that drives the emergence and growth of cities. It occurs unevenly across time and space, with disparities between continents and countries. Cities are mosaics of heterogeneous patches of diverse ecosystems with multiple interfaces [2]. Urbanisation can improve public health by enhancing living standards for citizens [3]. However, it also results in habitat loss and fragmentation, leading to a decline in biodiversity, as observed in China for birds species diversity [4]. Urbanisation also exacerbates air pollution [5], contributing to human health issues [6]. Moreover, social disparities and the lack of contact with nature favour mental health disorders [7]. Finally, the expansion of asphalt-covered areas that absorb energy during the day and release it as heat at night amplifies the urban heat island (UHI) effect [8], worsening its impact on human health [9].

Nature-Based Solutions (NBS), such as urban greening, are promoted to achieve the 11th Sustainable Development Goal: "to make cities and human settlements inclusive, safe, resilient, and sustainable" [10]. NBS also support the 12th target of the COP 15 Biodiversity Plan for Life on Earth, aiming to "enhance green spaces and urban planning for human well-being and biodiversity" [11], and align with the sustainability goals of the European Green Deal [12]. Urban greening involves deliberately integrating vegetation into urban areas (13), encompassing initiatives such as soil de-imperviousness, developing green and blue infrastructures, conserving urban green spaces (UGS), and promoting urban agriculture. UGS provide cultural and regular ecosystem services [13]. They help mitigate the UHI effect [14,15], and contribute to improving mental health [16]. Effective management of UGS can

help sequester carbon dioxide [17] through tree plantations and enhance animal biodiversity

by increasing the presence of indigenous vegetation [18].

Urban greening may also have negative side effects. It can increase pollen allergies [19] and

attract agricultural pests like tree moths [20]. UGS increase human-wildlife interactions,

raising the risk of zoonotic [21] and arthropod-borne diseases [22], as shown for the Lyme

disease pathogen (*Borrelia burgdorferi*) transmitted by ticks in urban parks across Europe [23].

In Europe, while numerous studies have explored the relationship between UGS and tick

populations, the relationship between urban greening and mosquitoes is less documented

[24], with only one study on *Culex pipiens* in the Netherlands [25] and two studies on the tiger

mosquito *Aedes albopictus* in Italy [26,27]. As shown in the scoping review by Mercat *et al*

(2025) [24], the effects of urban green infrastructures (UGIs) on mosquito populations are

highly dependent on the vector system studied and the environmental context. Urban

greening is a dynamic process, and its environmental changes may take time to manifest [28].

While it can create favorable conditions for vectors and increase human-vector interactions,

it may also promote the establishment of predators that could regulate vector populations.

Ae. albopictus is an invasive mosquito in Europe. It is native to Southeast Asia and a vector of

several infectious agents, including the dengue virus (DENV), chikungunya virus (CHIKV), and

Zika virus. The species adapted to antropogenic environments [29], including urban areas,

using natural and artificial breeding sites. It can survive extreme conditions through egg

diapause [30]. Its abundance is influenced by environmental factors such as weather [31],

microclimate [32], and landscape features [33]. Urban vegetation may favour *Ae. albopictus*

by creating cooler and more humid microclimates during warm season [34], facilitating access

to sugar meals, providing natural resting sites [35], and attracting blood sources (humans and

animals). In France, since its first detection in 2004 [36], *Ae. albopictus* spreads to 78 of 96

metropolitan France's departments by 2024, including all departments in the Occitanie region, where it has been linked to several autochthonous arboviral disease clusters since 2014. The Occitanie region is actively pursuing urban greening projects in its two largest cities, Toulouse and Montpellier, including plans to plant 150 000 trees. As Occitanie is a hotspot for both arboviral diseases and urban greening, it is crucial to assess the impact of these greening initiatives on *Ae. albopictus* population dynamics. With the goal to guide future research to improve the design and planning of urban green spaces, this study aims to evaluate the influence of urban vegetation in Montpellier in 2023 on the presence and abundance of *Ae. albopictus* females, using an explanatory approach taking into account land cover, microclimate, meteorology, air quality, and socio-demographics.

Material and methods

Mosquito sampling and study areas

Study areas

Montpellier is a city of 300 000 inhabitants located in the Occitanie region in southern France (Fig 1). The city has a Mediterranean climate with an average annual temperature of 15°C and an average annual precipitation of 739 mm [37]. The city features more than 10% of vegetated areas within densely urbanized areas. It ranks among the top three greenest cities in metropolitan France [38]. The study was conducted in nine areas belonging to three different environments: two urban parks (PRK), three residential neighborhoods (RES), and four densely built impervious areas (IMP) (Fig 1). Urban parks (Botanical Garden (PRK-BOT), and Park of

Aiguelongue (PRK-AGL)) and residential neighborhoods (Lemasson (RES-LEM), Aiguerelles

(RES-AGR), and Soulas (RES-SOUL)) were vegetated. The densely built impervious areas

included two areas with some vegetation (Saint Charles University (IMP-SCU) and Acapulco

Hotel (IMP-ACA)) and two with little to no vegetation (Buisson Bertrand Institute (IMP-BBI)

and Diderot School (IMP-DID)) (Fig 1). Detailed descriptions of study areas are provided in S1

Table, and images of study areas in S1 Fig. The minimum distance between study areas was

115 m. Although *Aedes* mosquitoes can disperse from 75 m [39] to 290 m [40] in urban areas

over 8 days, we treated the study areas as independent units due to impervious roads

between them, that are known to limit mosquito dispersal [41,42].

Fig 1: **Study areas**. Fig 1A and 1B show the location of Montpellier in France (map from

OpenStreetMap) and the land cover map of Montpellier with the study areas, respectively.

Sampling environments are shown in the legend of Fig 1B, delineating urban parks, residential

areas, and impervious built-up areas (boundaries from Montpellier Open Data). Land cover

categories are shown in the legend of Fig 1B. Fig 1C, 1D, and 1E show a residential area, the

southern part of the Botanical Garden (PRK-BOT), the Bouisson Bertrand Institute (IMP-BBI)

and the Diderot School (IMP-DID) respectively.

Mosquito sampling

Adult mosquitoes were captured using BG-Pro traps (BioGents, Regensburg, Germany)

modified according to L'Ambert *et al* [43], each equipped with a BG lure—synthetic human

odor mimic—and a daily replenished carbon dioxide source made from yeast, sugar, and water

(S1 Fig). Two BG-Pro traps were installed in each residential area (RES-LEM, RES-AGR, RES-

SOUL), and in the park of Aiguelongue (PRK-AGL). Due to their smaller size (S1 Table), only one

BG-Pro trap was deployed in each of the impervious built areas (IMP-ACA, IMP-BBI, IMP-DID

and IMP-SCU). Because of its larger size (S1 Table), three BG-Pro traps were deployed in the

Botanical Garden (PRK-BOT).

Traps were placed in shaded, wind-sheltered sites at a height of one meter. Therefore, 15 sites

were sampled: five in the two urban parks, six in the three residential areas, and four in

impervious built areas. Every trap was separated by a minimum distance of 50 m to prevent

interferences [44]. However, due to accessibility constraints, the two traps in RES-SOUL were

placed 22 m apart.

Every sampling session occurred for 48 hours each month from May to October 2023.

Mosquitoes were retrieved every morning, transported at 4°C to the laboratory (MIVEGEC

laboratory, IRD regional delegation, Montpellier), and stored at -20°C. Mosquito individuals

were then counted, sorted by sex, and morphologically identified using the multicriteria key

MosKeyTool [45]. These data are available on GBIF [46].

Environmental data collection and preparation

To assess the impact of urban vegetation on the presence and abundance of *Ae. albopictus*, it

was essential to consider all factors that may influence its population dynamics. For this task,

we adapted the conceptual model of human-*Anopheles* contact by Moiroux (2012) [47] to the

case of *Ae. albopictus*. Based on a literature review, we documented and hypothesised known

or potential predictors and their pathways to mosquito abundance [48]. This model guided

the selection of environmental data to be collected as described below.

Fine-scale vegetation and land cover data

The vegetation data were extracted from the fine-scale vegetation dataset of Montpellier

created in 2019 by Montpellier Mediterranée Métropole (S2 Table). The vegetation data were

categorised into low vegetation (< 3 metres in height) and high vegetation (> 3 metres in

height). In addition, a land cover map (Fig 2B) was generated by combining these vegetation

data with data from two open access databases: the Urban Atlas of the Copernicus European

Union's space program component and the French national database of Buildings (S2 Table).

The produced dataset included five categories: buildings, roads, low vegetation, high

vegetation, and "others" (that included railways, pathways, industrial areas, and sports

facilities). For each land cover category, the percentage of cover, total edge length, and

average patch size were calculated in buffer zones with radii of 20 m, 50 m, 100 m, and 250 m

around each trap location. For vegetation categories, the number of patches was also

extracted for each of the four buffer zones. The landscape diversity (Shannon index) was

calculated for all buffer zones (S2 Table). Buffer sizes were chosen according to previous works

investigating the effect of urban vegetation [27,49] on the dispersal of *Ae. albopictus* [40].

Table S3 provides the land cover variable values for each trap and buffer radii.

Microclimatic, meteorological and air quality data

We continuously collected hourly temperature and relative humidity near each trap using

Hygro Button data loggers from May to October 2023. The minimum (lowest recorded value),

maximum (highest recorded value), and mean temperatures, as well as relative humidity,

were calculated per 24-hour periods from 48h before the start of each sampling session to its

end (S2 Table).

Hourly temperature and rainfall data were obtained from the Observatoire Départemental de

l'Eau et de l'Environnement (ODEE) station located at Château d'Ô in the north-west of

Montpellier (S2 Table). Precipitations were recorded per 24-hour periods before and during

sampling sessions as for datalogger data. Relative humidity and wind speed data were

recorded every three hours from the Frejorgues airport station (belonging to the Meteo

France national meteorology agency network) (S2 Table). Growing Degree Days (GDD), a

known predictor of mosquito development, were calculated from daily mean temperatures

above 11°C, as this was shown to be the annual mean temperature limit for the establishment

of *Ae. albopictus* populations [50]. Temperatures above this level are conducive to mosquito

development. All weather data were aggregated for the sampling week and each of the six

weeks before sampling. Rainfall data and GDD were summed, while temperature, humidity,

and wind speed were averaged.

Atmospheric pressure can influence insects' flight [51] and locomotor activity [52], as shown

in *Drosophila* and beetles. Haufe, in 1954 [53], demonstrated that variations in atmospheric

pressure can affect the flight activity of *Aedes aegypti.* To evaluate the effect of daily

atmospheric pressure on *Ae. albopictus*, the minimum (lowest recorded value), maximum

(highest recorded value), and mean atmospheric pressure, measured every three hours at the

Frejorgues airport station, were extracted per 24-hours periods before and during sampling

sessions. The daily mean atmospheric pressure difference between two consecutive days was

calculated per 24-hours periods before and during sampling sessions. S2 Fig depicts the

temporal evolution of these meteorological variables (temperature, rainfall, wind speed,

relative humidity, and atmospheric pressure).

Air pollution can influence insect behaviour and abundance. Particulate matter (PM10, PM2.5)

affects pollinator orientation and reduces honeybee fitness [54]. Ozone (O_3) impairs

honeybee antenna perception [54] and lowers pollinator abundance [55]. Nitrogen monoxide

(NO) and nitrogen dioxide (NO2) reduce the pollinator performance [56]. To evaluate the

effect of air quality data on *Ae. albopictus*, for each trap location, we collected daily

concentrations of nitrogen monoxide (NO), nitrogen dioxide (NO2), particulate matter (of

diameter less than 10 (PM10) and 2.5 μm (PM2.5)), and ozone (O_3) from the closest station

of the Atmo Occitanie air quality observatory (S2 Table). Data were averaged weekly for up to

6 weeks, as for relative humidity, wind speed and temperatures, prior to each sampling

session.

Social and demographic data

Socioeconomic factors such as low living standards [57], aging or degrading infrastructure

[58], and population densities [59] can influence mosquito presence. Data on infrastructure

age and number of low-income households were obtained as a 200m spatial resolution grid

from the Filosofi database of the French Institut National de la Statistique et des Etudes

Economiques (INSEE). Values of the grid cells that intersect trap locations were retrieved.

Population density data were sourced from the Montpellier fine-scale population dataset,

created in 2016 by the Languedoc-Roussillon Geographic Information System (SIG-LR) (S2

Table) and aggregated within the same four buffer zones used for land cover data.

Statistical analyses of the presence and abundance of female *Ae. albopictus*

To study the relationships between the presence and the abundance of female *Ae. albopictus*

and environmental conditions, we analysed the data using a hurdle modeling approach, i.e.

considering the data responding to two distinguished processes: one causing zero vs. non-zero

(« presence » models) and one causing positive counts of females (« abundance » models).

This approach is justified because determinants of mosquito presence are expected to be

different than abundance ones [60].

For both presence and abundance analysis, we adopt a two-stage analysis method as

described by Taconet *et al* [61]. The first step involves bivariate analyses using Generalized

Linear Mixed Models (GLMMs), followed by the second step, a multivariate analysis employing

an interpretable machine learning method based on random forest models [62]. The bivariate

analyses aimed to improve our understanding of (i) the time-lagged effects of meteorological

and air quality variables, (ii) the impact of atmospheric pressure and microclimate on the tiger

mosquito, (iii) land cover-scale effects, (iv) effects of socio-economic and demographic

variables, and (iv) to identify and select key variables for multivariate analysis. The

multivariate analysis aimed to identify the most impactful determinants of the presence and

abundance of *Ae. albopictus* and the shape of their association while considering potentially

complex (e.g. non-linear) associations and interactions between variables, with the overall aim

of assessing more precisely the impact of urban vegetation. The modeling workflow is

summarised in S3 Fig.

Bivariate analysis using GLMMs

Every explanatory variable was tested using a GLMM with crossed random intercepts for trap

location, with sites nested within the sampling area, and for the sampling session. We fitted

binomial and zero-truncated negative binomial models to presence/absence and positive

counts data, respectively. Marginal R^2, representing the proportion of variance explained by

the fixed effects excluding the contribution of random intercepts [63], was calculated and

used for interpretation. Time-lagged effects of meteorological and air quality variables were

interpreted using cross-correlation maps (CCMs) [64] of marginal R^2 adjusted for the direction

(+/-) of the relationship.

Multivariate analysis using random forests

The results of the bivariate analyses were used to select the variables to be included in the

multivariate analysis. First, we excluded variables poorly correlated with the response

variables ($p > 0.2$). For each meteorological, microclimatic, air quality, land cover, and

demographic variable, we retained the time lag or radii with the higher marginal R^2.

Multicollinearity was addressed using Pearson's correlation coefficient with a threshold of 0.7.

Then, based on empirical knowledge, an initial selection of correlated variables was made.

Other collinear covariates were removed until the Variance Inflation Factor (VIF) of the

remaining variables was below three [65] (S3 Fig). Binary classification random forests (RFs)

and regression classification RFs were generated for the presence and abundance models,

respectively. To improve model efficiency and reduce the number of explanatory variables,

recursive feature elimination was performed for each model [66]. To account for spatial and

temporal autocorrelation, sampling variables (trap location, sampling day, and sampling

session) were included. For each set of hyperparameters and to assess the predictive power

of each model, both a spatial and temporal cross-validation framework [67] was used, taking

into account the potential spatial and temporal autocorrelation in the data. Spatial cross-

validation (leave-area-out) involved training the model recursively on data from eight out of

the nine sampled areas, testing it with data from the remaining area, and finally averaging the

performance metric for the nine areas [61]. The temporal cross-validation (leave-sampling-

session-out) involved training the model recursively on data from six out of the seven sampling

sessions and testing it with data from the remaining session. The performance metrics

selected were the Area Under the Curve (AUC) for the presence models [68] and the Mean

Absolute Error (MAE) for the abundance models. The model results were interpreted using

Variable Importance Plots (VIPs) [62] and Partial Dependence Plots (PDPs) [69]. Residual

spatial autocorrelation was assessed for each model.

Softwares used

The R version 4.2.3 programming language [70] was used. The land cover map creation and

landscape metrics calculation involved the use of the 'sf' [71], 'raster' [72], 'fasterize' [73], and

'landscapemetrics' [74] packages. GLMMs were fitted using the 'glmmTMB' package [75].

Marginal R^2 was calculated using the 'performance' package [76]. The 'correlation' package

was used to evaluate multicollinearity [77]. Multivariate analyses were performed using the

'caret' [78] and 'ranger' [79] packages. The 'CAST' package [80] was used for the cross-

validation and the interpretation plots were performed using the 'iml' [81], the 'pdp' [82], the

'MLmetrics' [83], the 'precrec' [84] and the 'patchwork' [85] packages. The package 'spatialRF'

[86] was used to assess spatial autocorrelation. All scripts are available on Software Heritage

[87].

Results

Entomological data

A total of 2344 mosquitoes were collected, predominantly *Ae. albopictus* (85.66%, N=2008),

with smaller proportions of *Culex pipiens* (13.91%, N=326), *Culiseta longiareolata* (0.39%,

N=9), and *Culiseta annulata* (0.04%, N=1). Among 2008 collected *Ae. albopictus*, there were

1106 females (55.08%) and 902 males (44.92%). The mean density of female *Ae. albopictus*

was 3.83 mosquitoes/trap/24h (SE = 1.97) in urban parks, 3.31 mosquitoes/trap/24h (SE =

1.65) in residential areas, and 1.99 mosquitoes/trap/24h (SE = 1.08) in impervious built-up

areas (Table 1). Mean mosquito densities (mosquitoes/trap/24 h), stratified by sampling

environment (or sampling areas), are presented in Table 1 (or S4 Table). The temporal

distribution of *Ae. albopictus* (S4 Fig) exhibited a bimodal pattern, with an initial emergence

in May, a primary peak in late June, and a secondary, smaller peak in late August.

Table 1: Mosquito density/trap/24h by species and sex in the three different sampling

environments.

Environment	Mosquito female /trap/24h (± SE)	Mosquito male /trap/24h (± SE)	*Ae. albopictus* female /trap/24h (± SE)	*Ae. albopictus* male /trap/24h (± SE)	*Cx. pipiens* female /trap/24h (± SE)	*Cx. pipiens* male /trap/24h (± SE)	*Cu. longiareolata* female /trap/24h (± SE)	*Cu. longiareolata* male /trap/24h (± SE)	*Cu. annulata* male /trap/24 h (± SE)

Urban parks	6.46 (± 2.80)	3.59 (± 2.89)	3.83 (± 1.97)	1.83 (± 1.67)	1.48 (± 0.79)	0.49 (± 0.38)	0.02 (± 0.02)	0.02 (± 0.02)	0.01 (± 0.01)
Residential areas	3.97 (± 1.64)	1.58 (± 1.14)	3.31 (± 1.65)	0.93 (± 0.83)	0.30 (± 0.16)	0.07 (± 0.05)	0.01 (± 0.01)	0.01 (± 0.01)	0
Impervious built areas	2.91 (± 1.37)	1.19 (± 0.87)	1.99 (± 1.08)	0.72 (± 0.69)	0.26 (± 0.18)	0.16 (± 0.12)	0.01 (± 0.01)	0.01 (± 0.01)	0

341   SE: Standard error

Bivariate analysis

Vegetation and other land cover variables

Fig 2 shows the vegetation and other land cover variables correlated with the probability of

presence or abundance of female *Ae. albopictus*. Two variables related to low vegetation were

significantly and positively associated with the probability of presence within a 20 m radius

buffer around the traps: the total length (in m) of low vegetation edges (OR =1.007, R^2 =0.042,

p =0.042, CI 95% [1.001; 1.014]) and the average patch size (in m²) of low vegetation (OR

=1.01, R^2 =0.07, p =0.006, CI 95% [1.003; 1.017]). The percentage of area occupied by buildings

within a 50 m radius buffer around the traps was negatively correlated with the probability of

presence (OR =0.959, R^2 =0.047, p =0.048, CI 95% [0.920; 0.999]). Within a 20 m radius buffer

around the trap, the total length edge of roads and the average size of a patch of roads were

negatively correlated with the probability of presence (Fig 2).

Regarding abundance, positive associations were observed with all vegetation-related

variables except for the average patch size of low vegetation within a 50 m radius buffer that

showed a negative and significant association (RR =0.996, R^2 =0.098, p =0.0001, CI 95% [0.994;

0.998]). The most remarkable positive association was for the total edges length of low

vegetation within a 100 m buffer (RR =1.0008, R² =0.190, p =0.007, CI 95% [1.0002; 1.0014]).

Variables representing the building land cover category consistently showed negative

relationships with female *Ae. albopictus* abundance, especially for the percentage area

variable (across all buffer radii) and for all variables within the 100m buffer radius. RR for the

percentage of buiding areas in the 100m buffer was 0.945 (R² =0.240, p =0.0007, CI 95%

[0.914; 0.976]) (Fig 2).

Fig 2: Bivariate relationships between *Ae. albopictus* and land cover variables for varied

buffer sizes. Relationships are analysed based on *Ae. albopictus* presence or abundance and

land cover variables for different buffer sizes around traps. The adjusted marginal R² reflects

the variance explained by the explanatory variable, adjusted for correlation direction. Boxes

are colored if the p-value was < 0.2 (No asterisk: p-value ∈ [0.05; 0.2], *: p-value ∈ [0.01,

0.05], **: p-value ∈ [0.001; 0.01]; ***: p-value ∈ [0; 0.001]). Box color depend on the direction

of the relationship (blue : negative, red : positive). Color intensity vary according to the

marginal R^2 value.

Microclimatic, meteorological, and air quality variables

Fig 3 shows the relationship between presence or abundance and microclimatic or

meteorological variables recorded from 48h before the start of sampling and during sampling.

Local relative humidity (RH) variable showed positive associations with presence when

recorded before sampling (with the strongest association for maximum hourly relative

humidity 24-48h before sampling; OR = 1.194, R^2 =0.201, p = 0.002, CI 95% [1.070; 1.332]) and

negative association when recorded during sampling (with the strongest association for

minimum RH; OR =0.924, R^2 =0.134, p = 0.001, CI 95% [0.880;0.969]). Significant associations

of RH data with abundance were negative whatever the period considered.

Fig 3: Bivariate relationships between *Ae. albopictus* and microclimatic and meteorological variables 48h before and during sampling. Relationships are analysed base on *Ae. albopictus* presence or abundance and variables recorded 48h before the start and during sampling sessions. Microclimatic variables include minimum hourly relative humidity (rhmin), maximum hourly relative humidity (rhmax), mean hourly relative humidity (rhmean), minimum hourly temperature (tmin), maximum hourly temperature (tmax) and mean hourly temperature (tmean). Meteorological variables include daily cumulated rainfall, minimum atmospheric pressure (Patmin), maximum atmospheric pressure (Patmax), mean atmospheric pressure (Patmean) and the difference of mean atmospheric pressure between two consecutive days (Patdiff). The adjusted marginal R^2 reflects the variance explained by the explanatory variable, adjusted for correlation direction. Boxes are colored if the p-value is < 0.2 (No asterisk: p-value $\in$ [0.05; 0.2], *: p-value $\in$ [0.01, 0.05], **: p-value $\in$ [0.001; 0.01]; ***: p-value $\in$ [0; 0.001]).

Box color depend on the direction of the relationship (blue : negative, red : positive). Color

intensity vary according to the marginal R^2 value. bef=before

Local temperatures during and before sampling consitently showed positive correlations with

presence, particularly for mean and maximum hourly temperatures recorded during sampling

(tmean: OR =1.592, R^2 =0.408, p =0.0002, CI 95% [1.251; 2.026]; tmax: OR =1.497, R^2 =0.402,

p =0.0001, CI 95% [1.226; 1.828]). Abundance was also positively associated with tmean (RR

=1.143, R^2 =0.197, p =0.0002, CI 95% [1.064; 1.228]) and tmax (RR =1.057, R^2 =0.061, p =0.07,

CI 95% [0.996; 1.122]) during sampling, but with lower R^2.

Daily total precipitation was negatively associated with presence during sampling (OR =0.863,

R^2 =0.137, p =0.003, CI 95% [0.783; 0.951]). *Ae. albopictus* presence was positively associated

with all variables describing atmospheric pressure recorded during the sampling hours but

negatively with lagged data. The strongest effect size was for the difference in mean daily

atmospheric pressure (Patdiff) between the sampling day and the previous day (OR = 1.008,

R^2 = 0.194, p = 0.008, CI 95% [1.002; 1.014]). Average (patmean) and maximum (patmax)

atmospheric pressure during the 24-48h period preceding sampling showed negative

associtaion with abundance (RR =0.997, R^2 =0.243, p =0.018, CI 95% [0.994; 0.999] and RR

=0.998, R^2 =0.177, p =0.0001, CI 95% [0.996; 0.999], respectively).

Local temperatures during and before sampling consitently showed positive correlations with

female presence, particularly for mean and maximum hourly temperatures recorded during

sampling (tmean: OR =1.662, R^2 =0.439, p =0.0001, CI 95% [1.298; 2.129]; tmax: OR =1.553, R^2

=0.424, p =0.0001, CI 95% [1.262; 1.991]). Abundance was also positively associated with

tmean (RR =1.134, R² =0.197, p =0.0001, CI 95% [1.055; 1.218]) and tmax (RR =1.052, R²

=0.050, p =0.098, CI 95% [0.991; 1.117]) during sampling, but with lower R².

Daily total precipitation was negatively associated with presence during sampling (OR =0.859,

R² =0.133, p =0.003, CI 95% [0.777; 0.949]). *Ae. albopictus* presence was positively associated

with all variables describing atmospheric pressure recorded during the sampling hours but

negatively with lagged data. The strongest effect size was for the difference in mean daily

atmospheric pressure (Patdiff) between the sampling day and the previous day (OR = 1.008,

R² = 0.217, p = 0.003, CI 95% [1.003; 1.015]). Average (patmean) and maximum (patmax)

atmospheric pressure during the 24-48h period preceding sampling showed negative

associtaion with abundance (RR =0.997, R² =0.217, p =0.049, CI 95% [0.994; 0.999] and RR

=0.998, R² =0.183, p =0.0003, CI 95% [0.996; 0.999], respectively).

Fig 4 shows the effect size and significance (p < 0.2) of weeks-lagged meteorological variables

on the presence and abundance of female *Ae. albopictus*. The relationship between weekly

cumulated rainfall (CUMRF) and both the probability of presence and abundance was positive,

with a statistically significant association for abundance and the highest effect size observed

the 6th week before sampling (for presence: OR = 1.430, R² = 0.334, p = 0.168, CI 95% [0.860;

2.375]; for abundance: RR = 1.075, R² = 0.127, p = 0.04, CI 95% [1.003; 1.152]). The probability

of presence was positively and significantly associated with minimum (TMIN), average

(TMEAN), and maximum (TMAX) temperatures. The highest effect size was observed for TMIN

one week before sampling (OR =1.620, R² =0.372, p =0.001, CI 95% [1.209; 2.172]. For

abundance, relationships with TMIN, TMEAN, and TMAX were also positive and significant for

time interval 0 to 2 weeks before sampling, with the highest effect size observed for TMAX

(RR = 1.161, R² =0.192, p = 0.0006, CI 95% [1.067; 1.267]). Both responses were positively and

significantly correlated with the weekly growing degree days (GDD). The highest effect on

presence and abundance was observed for GDD during the week preceding sampling (OR =

1.085, R^2 = 0.357, p = 0.002, CI 95% [1.031; 1.141]) and for time interval 0 to 2 weeks before

sampling (RR = 1.011, R^2 = 0.189, p = 0.0009, CI 95% [1.004; 1.017]), respectively. Both

responses were negatively and significantly correlated with relative humidity (RH), with the

highest effect observed for RH recorded 6 weeks before sampling (OR = 0.792, R^2 = 0.472, p

= 0.006, CI 95% [0.670; 0.937] for presence; RR = 0.948, R^2 = 0.183, p = 0.007, CI 95% [0.912;

0.986] for abundance). The relationship between wind speed (WS) and abundance was

negative, with the highest effect observed for time interval 0 to 5 weeks before collection (RR

=0.343, R^2 = 0.086, p =0.157, CI 95% [0.078; 1 .510]).

Fig 4: Cross correlation maps showing the lagged effect of meteorological variables on *Ae.*

***albopictus* presence or abundance**. Lagged meteorological variables include weekly

cumulated daily rainfall (CUMRF), mean daily temperature (TMEAN), daily minimal

temperature (TMIN), daily maximal temperature (TMAX), the weekly Growing Degree Days

(GDD), mean daily relative humidity (RH), and mean daily wind speed (WS). Time lags are

expressed in weeks prior to sampling. The adjusted marginal R^2 reflects the variance explained

by the explanatory variable, adjusted for correlation direction. Red-bordered squares highlight

the time lag with the highest marginal R^2, with the interval indicated in the top left corner.

Black-bordered squares denote marginal R^2 values close to the highest (within 10%). Gray

squares represent correlations with p-values > 0.2.

Daily nitrogen dioxide (NO2) concentrations were positively associated with the probability of

presence of female *Ae. albopictus*, with the highest effect size observed for the sampling week

(OR = 1.109, R^2 = 0.126, p = 0.163, CI 95% [0.959; 1.282]; S5 Fig). Particulate matter

concentrations (PM2.5 and PM10) were also positively correlated with the probability of

presence and with the abundance. However, PM2.5 and PM10 concentrations were strongly

correlated with all time-lagged temperature metrics (Pearson coefficient > 0.8). Tropospheric

ozone (O3) concentration (three weeks before sampling) was negatively correlated with the

probability of presence and positively (one week before sampling) with abundance. Daily O3

concentrations were also highly correlated with weekly cumulated rainfall, wind speed, and

daily relative humidity (Pearson coefficient > 0.8).

Socio-demographic variables

Human population density in a 250 m radius buffer around traps was positively correlated

with the *Ae. albopictus* probability of presence (OR=1.0013, R2=0.056, p=0.029, CI 95%

[1.0001; 1.0026]; S6 Fig).

Multivariate analysis

After bivariate analyses, multicollinearity assessments, and recursive feature elimination, ten

and seven variables, in addition to sampling design variables, were kept for the multivariate

random forest models of presence and abundance, respectively (S5 Table shows reason for

exclusion). Fig 5 and Fig 6 display the multivariate RF model interpretation plots for presence

and abundance, respectively. Fig 5A and Fig 6A illustrate the contribution of each variable in

explaining the response variable (VIPs). Fig 5B and Fig 6B depict the relationship between the

response variable and the explanatory variables (PDPs). Models of presence and abundance

did not show any residual spatial autocorrelation.

Fig 5: **Interpretation plots for the multivariate random forest model of** *Ae. albopictus*

presence. The left plot (Fig 5A) is the variable importance plot (VIP). Rights plots (Fig 5B) are

smoothed partial dependence plots (PDPs) with Confidence Intervals for each selected

variable in the models. The y-axis in PDPs represents the probability of catching at least one

mosquito. The range of values in the x-axis represents the range of values available in the data

for the considered variable. The rugs above the x-axis represent the actual values available in

the data for the variable. The black line represents the unsmoothed PDP, while the green line

shows the smoothed version. The shaded gray area corresponds to the 95% confidence

intervals, calculated using the standard deviation and adjusted for local data density. b/w:

between. bef.: before. tmin: minimum hourly temperature. rhmean: hourly average relative

humidity. Patmin: minimum daily atmospheric pressure. Patdiff: difference of mean daily

atmospheric pressure. CUMRF: Cumulated Rainfall.

Fig 6: Interpretation plots for the multivariate random forest model of *Ae. albopictus* abundance. The left plot (Fig 6A) is the variable importance plot (VIP). Rights plots (Fig 6B) are

smoothed partial dependence plots (PDPs) with Confidence Intervals for each selected

variable in the models. The y-axis in PDPs represents the number of *Ae. albopictus* females

caught/trap/24h. The range of values in the x-axis represents the range of values available in

the data for the considered variable. The rugs above the x-axis represent the actual values

available in the data for the variable. The black line represents the unsmoothed PDP, while

the green line shows the smoothed version. The shaded gray area corresponds to the 95%

confidence intervals, calculated using the standard deviation and adjusted for local data

density. b/w: between. bef.: before. tmax: maximum hourly temperature. GDD: Growing

Degree Day. WS: Wind Speed

Presence model

The most contributing variables to presence probability were a combination of real-time and

time-lagged microclimatic and meteorological variables (Fig 5A), with only a minimal

contribution of vegetation variables. The first contributing variable was the local minimum

hourly temperature recorded 24–48 hours before sampling. The probability of presence

showed a slight increase between 11.9°C and 16°C, rose sharply from 16°C to 25°C, and

plateaued beyond 25°C (Fig 5B). Minimum atmospheric pressure during sampling was the

second most influential variable, with an unimodal relationship: presence probability

increased up to 1012.91 HPa before declining. Cumulated rainfall 6 weeks before sampling

emerged as the third key variable, displaying a linear positive association with presence

probability, from 0 mm to 15 mm. The fourth major factor was the mean daily atmospheric

pressure difference between the sampling and previous day. Presence probabilities decreased

for descreasing pressures and was stable for stable or increasing pressures. Secondary

contributors included mean local hourly relative humidity 24–48 hours before sampling, daily

NO$_2$ concentrations, and vegetation metrics within a 20 m buffer around traps (average patch size of low vegetation, number of high vegetation patches, and total length of low vegetation edges). These variables slightly increased presence probability, whereas the total length of edges of roads in a 20 m radius buffer weakly reduced it. The model demonstrated strong predictive accuracy (AUC = 0.89; S7A Fig). Cross-validation with the leave-area-out method produced consistent predictions, except in the impervious Diderot School area (IMP-DID). In contrast, leave-sampling-session-out cross-validation exhibited greater variability between sampling areas (S7B Fig).

Abundance model

The most contributing variables to *Ae. albopictus* abundance comprised a combination of microclimatic, weeks-lagged meteorological, vegetation and other land cover variables. The first contributing variable was the local maximum hourly temperature during sampling (Fig 6A). A strong positive correlation with abundance was observed between 24°C and 38.7°C, followed by a negative correlation beyond 40.6°C (Fig 6B). The second most important variable was the weekly GDD between the sampling day and two weeks prior, which exhibited a linear positive correlation with abundance. The third key variable was the total length of road edges within a 250 m buffer around traps, with a strong negative correlation between 8230 m and 11 000 m. The fourth key variable was the average size of road patches within a 250 m buffer. It had no effect on abundance up to 1110 m² but exhibited a negatively correlation with abundance at sizes greater than 1630 m². The next variable was the average size of low-vegetation patches within a 50 m radius buffer around traps. Abundance increased markedly for areas between 0 and 120 m², plateaued up to 237 m², and subsequently decreased slightly. The percentage of low vegetation in a 250 m radius buffer around traps was positively

correlated with abundance between 10% and 23%, beyond which it had no observable effect.

Finally, average wind speed between day of sampling and five weeks before had no effect on

abundance between 3.7 m/s and 4.3 m/s, but showed a negative correlation beyond this

range. The abundance model demonstrated good overall accuracy, except for high abundance

cases (>11 mosquitoes, Mean Absolute Error (MAE)=7.86, S8A Fig). Spatial cross-validation

retained overall abundance trends, except in RES-LEM and PRK-AIG, and showed an

underestimation of high abundance and an overestimation of low abundance. Temporal cross-

validation, using leave-sampling-session-out, did not preserve abundance trends but captured

relative differences in *Ae. albopictus* abundance between areas (S8B Fig).

Discussion

This study examined, for the first time in France, the impact of urban vegetation on the

presence and abundance of *Ae. albopictus*, showing a minimal influence on the presence and

a moderate positive association with the abundance. The analytical methodology employed

in this study provides a nuanced evaluation of how different environmental factors contribute

to mosquito presence and abundance. It highlights the existence of non-linear relationships,

offering valuable insights for identifying effective vector control measures.

Influence of urban vegetation and other land cover variables

This study highlights that urban vegetation within a 20 m buffer radius around traps had a

minimal weakly positive influence on the probability of the presence of *Ae. albopictus*.

Regarding the abundance, urban vegetation demonstrated a moderate positive effect. This

positive effect of urban vegetation on the abundance of *Ae. albopictus* has also been observed

in other countries, including Italy [26], Brazil [88], Singapore [89], and China [33]. In our study,

the first vegetation metric influencing abundance was the average size of low-vegetation

patches within a 50 m buffer radius. These findings are consistent with those of Cianci *et al*

(2015) [26], who reported a positive relationship between vegetation cover within a 50 m

radius buffer around traps and *Ae. albopictus* eggs abundance in Italy. Manica *et al* (2016)

observed that, in Roma (Italy), vegetation cover within a 20 m radius around traps positively

influenced *Ae. albopictus* female abundance [27]. They hypothesised that the tiger mosquito

prefers "small green islands". Our findings refine this hypothesis by demonstrating that *Ae.*

albopictus abundance significantly increases with the average size of low-vegetation patches,

particularly within the 10 to 120 m² range. Beyond 120 m², the effect stabilises. This nuance

provides additional detail to the assumption proposed by Manica *et al*. This hypothesis, along

with the identified size range, requires further research to confirm its generalisability across

different urban contexts and mosquito populations.

The percentage of low vegetation cover within a 250 m radius around traps moderately

increased abundance. These results are consistent with the study of Sun *et al* (2021), which

demonstrated that, in Singapore, *Ae. albopictus* abundance increased with managed

vegetation within a 300 m radius buffer around traps [89]. However, Manica *et al* (2016) found

a negative relationship between vegetation cover within a 300 m radius buffer and mosquito

abundance [27]. These differences can be attributed to variations in city architecture and

study design. In our study, the percentage of low vegetation cover ranged from 10% to 23.5%,

whereas in Manica *et al*, it ranged from 30% to 55%. With different ranges of vegetation cover,

differing study designs, and contrasting urban environments, direct comparisons are challenging.

The other two land cover variables, the third and fourth contributors to abundance, are road metrics within a 250 m radius buffer around traps. The first was the total length of road edges, which was negatively correlated with abundance. These findings aligned with the study by Wang *et al* (2023) in Shanghai (China) which reported a negative correlation between road length and *Ae. albopictus* eggs density [33]. This supports our initial assumption that roads act as barriers to mosquito movement. This hypothesis was also consistent with Regilme *et al* (2021), who observed genetic differentiation in *Ae. aegypti* populations located north and south of a major road in Shanghai [42]. The second road-related variable was the average size of road patches. Roads may represent unfavorable habitats for host-seeking *Ae. albopictus* which could explain their limited abundance in areas dominated by large roads.

Influence of microclimatic and meteorological variables

The relationship between atmospheric pressure and the presence of *Ae. albopictus* in the field has, to our knowledge, been investigated for the first time in this study. The minimum atmospheric pressure during the 24-hour sampling period and the difference in daily mean atmospheric pressure between the sampling day and the previous day were the second and fourth main variables contributing to presence. Haufe (1954) demonstrated that atmospheric pressure affects the flight activity of *Ae. aegyp*ti, especially the change in atmospheric pressure after six hours of acclimatisation [53]. A study by Knop *et al* (2023) in Switzerland showed that the traffic rate of high-flying insects (number of insects per hour) had a positive correlation with air pressure [90]. Our results suggest an effect of the atmospheric pressure

metrics on the activity and behaviour of the tiger mosquito. Further studies under controlled

laboratory and field conditions are needed to investigate these findings in more detail.

Local hourly temperatures were the first variables to contribute to both presence and

abundance. Evans *et al* (2019) found an exponential relationship between the minimum

temperature during the 7 days preceding sampling and the abundance of females, as well as

an unimodal relationship between the maximum temperature and the abundance of females,

with an optimal daily maximum temperature around 35°C [32]. Our results suggest that

increasing daily maximum temperature increases the activity of *Ae. albopictus* up to 38°C.

The third variable influencing the presence of *Ae. albopictus* was the weekly cumulated rainfall

recorded six weeks before sampling. While other studies have identified a positive

relationship between *Ae. albopictus* dynamics and rainfall accumulated over several weeks

before sampling, none have specifically examined a six-week delay. Roiz *et al* (2015) observed

in Montpellier that strong rainfall, particularly that accumulated each week over the four

weeks before sampling, positively influenced abundance [31]. Similarly, Manica *et al* (2016)

reported a positive relationship between the cumulated rainfall one month before sampling

and mosquito abundance [27]. The positive effect of rainfall in our study may be explained by

the filling of larval habitats, benefiting previous mosquito generations that influenced the

sampled population. This hypothesis requires confirmation through additional temporal

replicates to validate the observed trend.

The second most influential variable of abundance was the weekly GDD during the two weeks

preceding sampling. Based on minimum, maximum, and average temperatures, the GDD is

widely used as a key predictor for many insect species [91]. For *Ae. albopictus*, the GDD is

calculated using a baseline temperature of 11°C, promoting mosquito development when

temperatures exceed this value [50]. This baseline corresponds to the average annual

temperature required for the spread of *Ae. albopictus* [92]. Jourdain *et al* (2020) identified the

GDD as the primary predictor of population dynamics for this species [93].

The last variable influencing abundance is the mean daily wind speed during the five weeks

preceding sampling. Le Goff *et al* (2024) observed a similar negative relationship between

monthly average daily wind speed and *Ae. albopictus* eggs abundance on the Réunion Island

[94]. Adeleke *et al* (2022) reported that in regions where the tiger mosquito is present, wind

speeds exceeding 4 m·s⁻¹ reduce the likelihood of occurrence [95]. The threshold identified in

our study (4.3 m·s⁻¹) can be considered comparable to the findings of Adeleke *et al*, although

direct comparisons between the studies are challenging due to differences in spatial and

temporal scales.

Limitations, perspectives and conclusion

In order to increase the resilience of Urban Green Infrastructures (UGIs) [96] and to adapt the

UGIs management to promote biodiversity [97], since 2017, studies have highlighted the need

to consider UGIs as socio-ecological ecosystems. This reflects the complexity of urban

greening, which must account for land cover, socio-demographic variables, and urban

architecture, as demonstrated in the systematic review by Menconi *et al* (2021) [98]. The

challenge of studying the relationship between urban greening and the risk of vector-borne

diseases is one of the key conclusions of the scoping review by Mercat *et al* (2025) [24]. The

difficulty stems from the need to integrate multiple variables essential to the analysis of this

socio-ecosystem, the time required to observe its effects [28], and the challenge of assessing

vector-borne disease risk using standardised protocols.

Due to limited financial and human resources, this study was limited to specific sampling areas

within vegetated environments representative of the city's UGS and non-vegetated

environments. A randomised sampling approach would have better captured the diversity of

land cover categories. In order to reduce interference between traps less than 50 m apart,

averages of meteorological, microclimatic, landscape and ethological variables were

calculated. The predicted abundance of *Ae. albopictus* in the RES-SOUL residential area was

therefore certainly underestimated. While the abundance model effectively captures

temporal variation, it does not account well for spatial variation. This suggests that key factors

such as environmental carrying capacity, defined as the 'equilibrium density of larvae in a

system' [99], and the availability of resting sites may have been overlooked (35). This is also

related to the complexity of describing urban vegetation using remote sensing [24]. In our

study, we employed a fine-scale characterisation of urban vegetation generated through deep

learning and photo interpretation, alongside a surface modelling metric derived from lidar

data. However, this approach does not capture the diversity of plant species or genera, which

can vary markedly between the Botanical Garden, the Park of Aiguelongue, and residential

areas.

The objectives of this study were to explore the relationships between urban vegetation and

other environmental factors with the presence and abundance of *Ae. albopictus*, to generate

hypotheses using an exploratory approach. The realisation of predictive model is another

objective, which requires a better understanding of the ecological and biological mechanisms

[100]. The results of the study are based on only one sampling year. To draw more robust

conclusions, additional years of sampling would be needed to assess inter-annual variation. In

addition, this study has focused solely on one aspect of the vector risk, the presence and

abundance of the vector, without considering its capacity to transmit pathogens, or human

exposure and vulnerability. The vector's capacity depends on the abundance of females, the

daily survival rate, and the extrinsic incubation period of pathogens [101]. It would be relevant

to investigate the effect of urban vegetation on these elements. Together with these

investigations, a socio-ecological approach is essential to gain a comprehensive understanding

of vector-borne disease risks in urban green spaces. Developing social surveys to assess

citizens' perceptions, alongside employing co-construction methods [102] with policymakers,

urban designers, and ecologists, could help formulate effective recommendations to mitigate

this risk. One example could be the development of green wind corridors, as implemented in

Stuttgart [103] and Lisbon [104], which can reduce urban heat islands and increase wind

speed, limiting mosquito abundance.

Urban greening appears to be a promising solution to address the environmental and health

challenges associated with increasing urbanisation. It offers numerous health and

environmental benefits, but can also contribute to an increased risk of vector-borne diseases.

This study, the first in France to investigate the effect of urban vegetation on *Ae. albopictus*,

provides a nuanced assessment of how urban vegetation influences both the presence and

abundance of mosquitoes. Our results indicate that urban vegetation moderately increases

abundance, but has no effect on presence. Urban greening is a dynamic and complex

phenomenon that requires a social-ecological approach in order to be studied. Further

research is needed to better elucidate the role of urban vegetation in the ecology of the tiger

mosquito and its influence on different aspects of risk.

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1012   vert-de-lisbonne/

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Supporting informations

1015

1016   **S1 Table: Table with sampling areas characterization**.

1017

1018   **S2 Table: Summary of the different variables used, with their processing, acquisition**

1019   **methods or source database and temporal/spatial scale.**

1020

1021   **S3 Table: Description of the landscape metrics for each landscape class for each trap, by**

1022   **buffer.**

1023

1024   **S4 Table: Mosquito density/trap/24h by species and sex in the nine different sampling areas.**

1025   The mosquito density represents the marginal mean of the number of mosquitoes caught per

1026   trap per 24 hours, calculated based on the sampling environment, with the standard error

1027   specified (SE).

1028

1029   **S5 Table: Variables Selected After Bivariate Analyses, with Justifications for Exclusion from**

1030   **Multivariate Analysis** (Correlation, VIF, Recursive Feature Elimination)

1031

1032 **S1 Fig: BG-PRO trap on the field and pictures of studied areas.** S1A Fig represents a BG-Pro

1033 on the field, S1B Fig the Park of Aiguelongue, S1C the Botanical Garden, S1D The Bouisson

1034 Bertrand Institute and 1E the Acapulco Hostel.

1035

1036 **S2 Fig: Analysis workflow: from the data acquisition to the multivariate analysis**

1037

1038 **S3 Fig: Time series of weekly cumulated precipitations (1A), mean (TMEAN), minimum**

1039 **(TMIN) and maximum (TMAX) temperature, (1B), growing degree-days (1B), mean relative**

1040 **humidity (1C), mean wind speed (1D) and mean (Patmean), minimum (Patmin) and**

1041 **maximum atmospheric pressure (Patmax), during the studing period.** Vertical dotted red

1042 lines indicate monthly mosquito sampling sessions.

1043

1044 **S4 Fig: Number of *Ae. albopictus* per trap per 24h during the study period, by sampling**

1045 **environment.** Boxes represent medians and interquartile ranges, whiskers indicate minimum

1046 and maximum values, and points show the abundance for each sampling session and day, for

1047 each trap, within the three environments.

1048

S5 Fig: Cross correlation maps showing the lagged effect of air-quality variables on *Ae.*

***albopictus* presence or abundance**. Lagged air-quality variables include daily concentration of

nitrogen monoxide (NO), of nitrogen dioxide (NO2), of particulate matter (PM10, PM2.5) and

of ozone (O3). Time lags are expressed in weeks prior to sampling. The adjusted marginal R^2

reflects the variance explained by the explanatory variable, adjusted for correlation direction.

Red-bordered squares highlight the time lag with the highest marginal R^2, with the interval

indicated in the top left corner. Black-bordered squares denote marginal R² values close to the

highest (within 10%). Gray squares represent correlations with p-values > 0.2.

1057

S6 Fig: Bivariate relationships between *Ae. albopictus* presence or abundance and the

sampling environment and the socio-demographic variables. The adjusted marginal R²

reflects the variance explained by the explanatory variable, adjusted for correlation direction.

Boxes are colored if the p-value was < 0.2 (No asterisk: p-value $\in$ [0.05; 0.2], *: p-value $\in$

[0.01, 0.05], **: p-value $\in$ [0.001; 0.01]; ***: p-value $\in$ [0; 0.001]). Box color depend on the

direction of the relationship (blue : negative, red : positive). Color intensity vary according to

the marginal R² value.

1065

S7 Fig: Evaluation of presence model. S7A Fig is the Receiver Operating Characteristics (ROC)

Curve for the presence model. It illustrates the model's sensitivity (its ability to predict the

presence of *Ae. albopictus*) against 1 - Specificity (its ability to detect the absence of *Ae.*

albopictus). The Area Under the Curve (AUC) represents the model's ability to discriminate

between the presence and absence of the mosquito, with values ranging from 0 to 1. The

closer the AUC is to 1, the better the model's predictive performance. S7A Fig shows that the

model demonstrated good predictive accuracy, with an AUC of 0.89. S7B Fig compares the

observed (in green) versus predicted presence probabilities for each out-of-sample leave-area

(yellow) and for each out-of-sample leave-sampling-session (in blue) . The y-axis represents

the probability of *Ae. albopictus* presence, while the x-axis corresponds to the sampling

session. Figure 10B indicates that the model accurately predicted the spatiotemporal trends

of presence/absence, although it tended to overestimate presence.

1078

S8 Fig: Evaluation of abundance model. S8A Fig is a violin plot showing the distribution of

residuals for the abundance model. The black dots indicate the median values. The Mean

Absolute Error (MAE) and the number of observations (n) are displayed in a square above the

plot. This figure demonstrates that the model tends to overestimate low counts of *Aedes*

albopictus (fewer than three caught in 24 hours per trap), accurately estimates counts

between 4 and 20, and underestimates higher counts (more than 20). S8B Fig compares the

observed versus predicted numbers of *Ae. albopictus* caught per site per sampling session (in

green), versus predicted numbers for each out-of-sample leave-area (yellow) and for each out-

of-sample leave-sampling-session (in blue). The y-axis represents the mean number of d *Ae.*

albopictus females caught per trap per area per sampling session, while the x-axis represents

the sampling session. This figure confirms the observations made in S8A Fig.

A. Location of Montpellier

C. Residential area

D. Botanical Garden

B. Studied areas

Sampling environments

- ☐ Urban parks (5 traps)
- ☐ Residential areas (6 traps)
- ☐ Impervious areas (4 traps)

Land cover categories

- ☐ Low vegetation (< 3 m)
- ☐ High vegetation (> 3 m)
- ☐ Buildings
- ☐ Roads
- ☐ Others

— 350 m

E. One impervous area

Fig1

Fig2

Fig3

Fig4

Fig5

Fig6