

Distribution of *Bactrocera oleae* (Rossi, 1790) throughout the Iberian Peninsula based on a maximum entropy modeling approach

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Abstract

The Iberian Peninsula (Portugal and Spain) is a great production area of olives. The fruit production can be severely affected by the olive fruit fly, *Bactrocera oleae* (Rossi, 1790) (Diptera). Detailed geographical distribution maps of key pests, such as *B. oleae*, are essential for their integrated management. Although different sources reporting the occurrence of *B. oleae* are available for sub-regions of Portugal and Spain, the data

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available are dispersed and centralization of this information considering the Iberian Peninsula as a faunistic geographical unit is currently lacking. In this work, we built two distribution maps of *B. oleae* throughout the Iberian Peninsula, one based on occurrence sites and another based on its bioclimatic habitat suitability. After modeling the bioclimatic suitability of *B. oleae* using a maximum entropy model, three potential distribution areas beyond the previously known occurrence range of the olive fruit fly were identified corresponding to the autonomous community of Galicia (Spain), the Spanish and Portuguese sides of the International Douro Natural Park, and the autonomous community of Castilla y León (Spain). Interestingly, each region houses nowadays autochthonous olive cultivars. The drivers that most contributed to the model were the precipitation of the coldest quarter and the precipitation of driest month which agrees with the *B. oleae* bioecology. Although our approach is not fully-comprehensive in terms of occurrence sites, we show how a maxent modeling approach can be useful to identify potential risk areas of *B. oleae* occurrence throughout a target geographical extent such as the Iberian Peninsula.

Keywords

Olea europaea, olive fruit fly, pest distribution, maxent, WorldClim.

The olive tree (*Olea europaea* L.) cultivation strongly influences the human life across the Mediterranean basin from economic to landscape aspects (Loumou & Giourga, 2003). The Mediterranean basin encompasses 91.83% of the world olive harvested area (FAOSTAT, 2019). This region produces and consumes 99% and 87% of the world's olive oil respectively. Southwestern Europe (i.e. the Iberian Peninsula housing Portugal and Spain) represents 29.36% of the olive harvested area within the entire Mediterranean basin (FAOSTAT, 2019).

The olive tree is susceptible to the attack of different insect pests such as *Bactrocera oleae* (Rossi, 1790) (Diptera), *Prays oleae* Bernard, 1788 (Lepidoptera), *Saissetia oleae* (Olivier, 1791) (Hemiptera), *Euphyllura olivina* Costa, 1839 (Hemiptera), and *Phloeotribus scarabaeoides* (Bernard, 1788) (Coleoptera) (Haniotakis, 2003; Haber & Mifsud, 2007). Among them, the olive fruit fly is the most damaging pest across the Mediterranean basin because of direct pulp destruction caused by the larvae development, premature drop of the infested fruits and general reduction in olive oil quality, which may cause a loss up to 40% of the production (Pereira et al., 2004; Boccaccio & Petacchi, 2009).

Accordingly, monitoring schemes have been implemented by the countries as an informative and predictive-like tool such as the Spanish monitoring web of olive pests "Red DACUS". However, efficient networks of monitoring stations are usually expensive (e.g. in terms of human resources) and the information provided by them does not extend beyond their geographical position.

In contrast, species distribution modeling is a widely used method to investigate the potential occurrence and spread of species. Species distribution models (SDMs) are based

on the relationship of a pool of georeferenced observations of a species occurrence (or abundance) with different environmental predictors across a determined geographical extent (Franklin, 2013). SDMs have been used in several fields of ecology such as research on invasive species (e.g. Dark, 2004), climate change impacts (e.g. Thuiller et al., 2006), conservation (e.g. Dubuis et al., 2011), and modeling of pest species distribution and spread (see Kehlenbeck et al., 2012, and Sutherst, 2013). Among these modeling techniques, the Maxent models (i.e. maximum entropy models) raised as a general-purpose machine learning method that allows modeling a species distribution using presence-only data (Phillips et al., 2006). However, although open-source and user-friendly tools are already available to follow this approach (e.g. Phillips et al., 2017) and detailed presence/absence occurrence data are freely accessible from different resources (e.g. global species repositories), there is still a lack of detailed stand-alone distribution maps and species distribution models for *B. oleae* in the Iberian Peninsula.

The objectives of this work were (1) to provide an occurrence-based distribution map of B. *oleae* for the Iberian Peninsula (i.e. Spain and Portugal), and (2) to assess the climatic suitability across the same geographical extent based on bioclimatic drivers.

2 | Material and methods

The compilation of occurrence sites of *B. oleae* and *O. europaea*, the selection of bioclimatic drivers, and modeling processes were carried out using the environment for statistical computing R (R Core Team, 2018).

2.1 | Occurrence-based distribution map

Occurrence sites of *B. oleae* were collected through a bibliographic review that included scientific papers, Spanish government technical reports of *B. oleae* monitoring stations, the Global Biodiversity Information Facility online database (GBIF, 2018), and personal observations. Analogously, occurrence sites for *O. europaea* were obtained from the former database only. The GBIF occurrence datasets (*Bactrocera oleae* (Rossi, 1790) in GBIF Secretariat (2017) & *Olea europaea* L. in GBIF Secretariat (2017)) were obtained using the *gbif* function of the {rgbif} package. The list of occurrence coordinates and corresponding references are provided in Table S1. A digital elevation model from Reuter et al. (2007) was used to project the occurrences sites for the Iberian Peninsula.

2.2 | Bioclimatic variables

The bioclimatic variables used were obtained from the WorldClim database (Fick & Hijmans, 2017). WorldClim is a gridded climate database that provides a set of bioclimatic variables derived from the monthly temperature and rainfall values representing annual trends, seasonality, and extreme or limiting environmental factors. The pool of 19 variables was downloaded at 2.5 minutes spatial resolution (~ 4.5 km at the equator) and is described in Table 1. A driver selection was then carried out to avoid multicollinearity among the bioclimatic variables using the *cor* and *findCorrelation* functions of the {stats} and {caret} packages respectively. For this purpose, a pool of 1000 random points across the Mediterranean basin extent was generated and those variables exceeding a threshold of 0.8 for the Pearson pairwise correlation were excluded for modeling.

The model was developed using an R implementation of Maxent (Phillips et al., 2006), a machine-learning method that uses the principle of maximum entropy to approximate the unknown probability distribution of a species based on presence-only data (Phillips et al., 2006) using the *maxent* function of {dismo} package (Hijmans et al., 2017). Since *B. oleae* is associated to the olive tree cultivation, the geographical extent of the selected bioclimatic drivers was focused on the Mediterranean basin (15°W, 40°E, 27°S, 50°N), the highest olive producing region worldwide (FAOSTAT, 2019) using the *crop* function of {raster} package (Fig. S1).

The optimal model was selected following Muscarella et al. (2014) towards a balance of goodness-of-fit with model complexity and evaluation of models with spatially independent data. The {ENMeval} package allows creating data sets for k-fold cross-validation using different methods for partitioning occurrence data, building a series of candidate models using the function *ENMevaluate* (calling the algorithm of {maxnet} package that uses {glmnet} for model fitting based on "Maxent" Java package, version 3.4.0), and provides evaluation metrics to select the optimal model settings. We used the "checkerboard1" method for partitioning occurrence data to build a series of 40 models corresponding to five combinations of feature classes (linear, quadratic, product, threshold, and hinge) and eight regularization multipliers (β) (0.5, 1, 1.5, 2, 2.5, 3, 3.5, and 4). Duplicate presence points (i.e. occurrence points that fell in the same grid cell) were removed and 10000 random background localities were used across the study extent (Fig. S2). The occurrences records of B. oleae used in this work correspond to presence-only data thus the background data represent pseudo-absences towards AUC (area under the receiver operating characteristic curve) calculations (see below) (Phillips & Dudík, 2008; Barbet-Massin et al., 2012).

The optimal model was assessed by selecting the one with the lowest AIC (Akaike Information Criterion). The AIC corrected for small samples sizes (AICc) reflects both model goodness-of-fit and complexity thus the model with the lowest AICc is considered the best model across the combinations of feature classes and regularization multipliers (see Muscarella et al. (2014) for details). Once selected, the model was refitted using the optimal model tuning and the AUC was calculated. The AUC is a threshold independent measure of predictive accuracy based only on the ranking of locations (Merow et al., 2013) and ranges from 0.5 (i.e. random) for models with no predictive ability to 1.0 (i.e. perfect discrimination between suitable and unsuitable cells) for models with a perfect prediction. According to Swets (1988), a model with an AUC > 0.7 provides "useful" discrimination ability, > 0.8 means "good" model performance, and > 0.9 results in a "very good" model performance.

The bootstrapped standard error of AUC, its 95% confidence interval, and the ROC (Receiver operating characteristic) curve were calculated following Steven & Phillips (2009) and using the {ROCR} package (Sing et al., 2005). Finally, the number of predicted values falling at each step of 0.1 within the range of predicted climatic suitability (i.e. 0.0– 0.1, 0.1–0.2, 0.2–0.3, 0.3–0.4, 0.4–0.5, 0.5–0.6, 0.6–0.7, 0.7–0.8, 0.8–0.9 and 0.9–1.0) was calculated and 20 replicates of the predictions were run for plotting purposes. A response curve was fitted for each bioclimatic variable representing the average prediction for all data points that were used to fit the model. The response curves were calculated using the *response* function of {dismo} package. Finally, the chorological map of *O. europaea* of Caudullo et al. (2017) (file *Olea_europaea_europaea_plg_clip.shp*) was used to graphically assess the amount of extended climatic suitability of *B. oleae* beyond the known distribution of the olive tree occurrence area in the Iberian Peninsula.

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3 | Results

3.1 | Occurrence-based distribution map

The compilation of occurrence records of *B. oleae* resulted in a total of 762 records (Table S1) of which 219 presence records were used for training and 73 for testing after reducing the geographical extent to the target region (the Mediterranean basin). In the case of *O. europaea*, 37,556 records were downloaded from the GBIF database and used only for plotting purposes. The occurrence-based distribution map of *B. oleae* for the Iberian Peninsula is presented in Fig. 1.

3.2 | Bioclimatic variables

The multicollinearity analysis resulted in the exclusion of the bioclimatic variables bio1, bio5, bio10, bio11, bio12, bio13, bio16, bio17, and bio18 (Fig. 2) so that the mean diurnal range, isothermality, temperature seasonality, minimum temperature of coldest month, temperature annual range, mean temperature of wettest quarter, mean temperature of driest quarter, precipitation of driest month, precipitation seasonality, and precipitation of coldest quarter were used as predictors for subsequent modeling purposes.

3.3 | Species distribution modeling

Among the five feature combinations used to select the optimal model (L, LQ, LQH, LQHP, and, LQHPT) and the eight regularization multipliers (ranging from 0.5 to 4.0), the one that achieved the lowest AICc and a Δ AICc = 0 was the linear and quadratic model (LQ) with a regularization multiplier (β) = 0.5, resulting in 17 parameters (Table

S2). The LQ model gave an AUC = 0.923 (Fig. S3), with a bootstrapped standard error = 0.0146, and a 95% confidence interval of [0.894, 0.951]. In terms of prediction within the Iberian Peninsula (10°W, 5°E, 35°S, 15°N) (Fig. 3), 19.44% of cells fell within the [0.0, 0.1] interval of p (predicted climatic suitability), 9.21% within [0.1, 0.2], 8.41% within [0.2, 0.3], 8.59% within [0.3, 0.4], 9.42% within [0.4, 0.5], 9.81% within [0.5, 0.6], 9.68% within [0.6, 0.7], 8.97% within [0.7, 0.8], 8.15% within [0.8, 0.9], and 8.31% within [0.9, 0.10] (Fig. S4). Among the selected bioclimatic variables, the drivers that contributed most to the Maxent model were the precipitation of coldest quarter, precipitation of driest month, and mean diurnal range whereas the mean temperature of the wettest quarter, the mean temperature of driest quarter, and isothermality were the less contributing variables (Table 2). The response curve for each bioclimatic variable is presented at Fig. 4.

The average omission and predicted area for species, and the sensitivity versus specificity results (ROC) for the 20 replication runs of the optimal model are provided at the supplementary material as the Fig. S5 and Fig. S6 files respectively.

Three potential distribution areas beyond the previously known occurrence range of the olive fruit fly were identified (Fig. 5), each one representing a region where the olive tree is currently being cultivated: (1) the autonomous community of Galicia (Spain), (2) the Spanish and Portuguese sides of the International Douro Natural Park, and (3) the autonomous community of Castilla y León (Spain).

4 | Discussion

In this work an occurrence-based distribution map of *B. oleae* for the Iberian Peninsula was built using different occurrence sources. The occurrence records were used

subsequently used to model the climatic suitability of *B. oleae* using a maximum entropy modeling approach and the AUC obtained (0.923) resulted in a very good model performance according to Swets (1988). Maxent models are commonly used to assess the probability of occurrence of a given species according to its potential distribution and based on the discrimination of suitable versus unsuitable areas for the species. In addition, nichebased models are usually projected on a target geographical extent, thus giving a theoretical area of predicted presence for the species (Phillips et al., 2006). However, several factors such as biotic interactions (Anderson et al., 2002) and historical factors (Peterson et al., 1999) can lead a species to be unable to occupy a certain area even if favorable. In fact, in our case study, the "probability of occurrence" of *B. oleae* is determined *a priori* by the presence or absence of *O. europaea* thus we prefer to interpret our results strictly in terms of climatic suitability or potential distribution area instead of occurrence probability (see Yackulic et al., 2013).

The Maxent model prediction agrees in general with Müller et al. (2019) and with the pattern found by Ponti et al. (2014) through mechanistic physiologically-based demographic models (PBDMs) for the olive fruit fly abundance in the Mediterranean basin and extends its potential distribution area (i.e. areas with $P \ge 0.5$) to the autonomous communities of Galicia, Asturias, and Castilla y León, as well as to the whole Ebro river valley within Spain (Figs 3 & 4).

These results are of capital importance since Galicia (northwestern Spain) had recently emerged as a new olive-growing region housing two autochthonous cultivars (*Brava* and *Mansa de Figueredo*) with potential relevant health benefits due to high concentration levels of phenolic compounds (Reboredo-Rodríguez et al., 2018; Figueiredo-González et al., 2019).

Analogously, the olive tree is one of the most extensively cultivated perennial species in the International Douro Natural Park. The Park encompass the border section of the Douro River, a deep, embedded valley with cliff margins that separate Portugal from Spain across more than 100 km. The Spanish side of the Valley runs along the provinces of Zamora and Salamanca (Castilla y León) (Fig. 4). In this region, the olive tree cultivation, with its own autochthonous cultivar (*Zorzal*) is of economic and social importance attending to its traditional use as excipient, medicinal properties, and ecotourism (including "olive-oil tourism") incoming (González et al., 2010; Marcos, 2010).

The autonomous community of Castilla y León houses 2% of the olive crop area of Spain (Muriel & Marcos, 2012). Within this region, the olive tree cultivation in the province of Valladolid has been increasing during the last years. Cropping areas that formerly did not support the growing of *O. europaea* are nowadays being exploited taking advantage of increased temperatures due to climate change and irrigation (Moreno et al., 2005; Tanasijevic et al., 2014).

In terms of the predictive bioclimatic variables selected, our results are in line with the current knowledge on the life history of *B. oleae* and *O. europaea*. The olive tree is the most abundant perennial tree historically adapted to Mediterranean conditions (Loumou & Giourga, 2003), i.e. a Csa-type climate according to the Köppen-Geiger classification characterized by hot and dry summers and mild and moist winters (Peel et al., 2007). The drivers that most contributed to the model were the precipitation of the coldest quarter and the precipitation of driest month which suggest that the habitat requirements of *B. oleae* are directly or indirectly driven by boundary climatic conditions (i.e. limiting environmental factors for both the pest and the host species, *O. europaea*).

Although Podgornic et al. (2013) did not found a significant relationship between precipitation and trap captures in Slovenia, Yokohama & Miller (2007) found in a series of laboratory and field studies at California that exposure to 5°C and 85% RH caused 100% mortality of eggs and three larval instars. Moreover, 3^{rd} instar larvae and 0–4 d-old pupae reached almost 100% of mortality after immersion in water saturated sand for 3 days. These agree with the predictions of our model since the climatic suitability for *B. oleae* decreased at high values of the precipitation during the coldest quarter. In addition, the decreasing pattern in climatic suitability found for precipitation during the driest month agrees with the adaptation of *B. oleae* and it host to the hot and dry summers across the Mediterranean region. However, the survival of overwintering pupae of the olive fruit fly could be affected by soil moisture as Wang et al. (2013) found in California suggesting that mortality could result from pupae dehydration before the winter rains.

On the other hand, the following most contributing variable to the model prediction was the mean diurnal range of temperature. Almost all ecophysiological parameters of *B. oleae* are strongly affected by temperature (Wang et al., 2009; Pappas et al., 2013; Ordano, 2015; Baratella et al., 2017). For example, our model predicted a strong decrease in the climatic suitability for *B. oleae* at a mean diurnal range higher than 10°C and an optimal for the mean temperature of coldest month at *c*. 4.5°C. Marchi et al. (2016) found a consistent relationship between the degree of infestation and the temperature-based indices calculated for the previous period using decade-long agrometeorological time series whereas Gonçalves et al. (2012) estimated that daily minimum temperature could explain the mortality of overwintering *B. oleae* in olive groves in the "Terra Quente" region of Trás-os-Montes (northeastern Portugal).

Our Maxent model could reliably capture the bioclimatic information involved in the lifehistory of B. oleae using presence-only data. Of course, our modeling approach can be improved by considering different common model limitations such as sampling bias, representativeness of the selected geographical extent, optimal model selection (e.g. testing different methods of data partitioning, feature class combinations, and regularization multipliers) as well as the consideration of further explanatory variables (e.g. pest dispersion ability, presence/absence of olive trees, and the application of pesticides and management practices). Dealing with these issues implies a huge effort in terms of comprehensive sampling programs, bibliographic review and model development; however, despite some criticism to Maxent models (based on the narrow set of circumstances that allow for an unbiased equivalence between the logistic output and probability of occurrence (Yackulic et al., 2013)), we showed how this approach can help identify potential risk areas of pest occurrence or colonization with high economic implications. In any case, despite the hypothesis testing aim, model performance, and interpretation of a species distribution model, the process implicitly generates an occurrence-based distribution map for the target species which is itself a useful contribution.

In conclusion, the spread of the occurrence (actual or potential) of *B. oleae*, the amount of areas with predicted climatic suitability higher than 0.5, and the extension of olive crop areas in the Iberian Peninsula make this region an excellent geographical area to investigate the processes involved in the potential dispersion/colonization of *B. oleae*. For example, the increasing temperatures in response to climate change predicted by climate models (IPCC, 2014) makes mandatory to study the effect of climate change on the relationships between the olive tree and its obligate pest *B. oleae*.

Although using a Maxent approach to model shifts in a species distribution (e.g. anticipation of pest invasion purposes) due to climate change (e.g. under different representative concentration pathways (RCPs), i.e. different greenhouse gas concentration trajectories adopted by the IPCC) could be tempting, different aspects such as biotic interactions, evolutionary change, dispersal ability, and emergence of new fundamental niches should be taken into account beyond regarding only climate variables (Pearson & Dawson, 2003; Veloz et a., 2012; Guisan et al., 2014). For this, powerful individual-based approaches such as climate envelope and physiologically based demographic models (PBDM) (Gutiérrez et al., 2008, 2009) and recent eco-evolutionary models (Moya-Laraño et al., 2014; Grimm et al., 2017) have been used and rising respectively and could help to deal with this task in the near future. We hope this work would represent a stimulus for further research towards a deeper knowledge of a biological system as important as the olive grove agroecosystem.

Contribution of authors

J.B.-M, P.B., and J.A.P. conceived the idea, J.B.-M. developed the model, and J.B.-M, P.B., J.A.P., and S.A.P.S. contributed to writing the paper.

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References

Anderson, R.P., Peterson, A.T. & Gómez-Laverde, M. (2002). Using niche-based GIS modeling to test geographic predictions of competitive exclusion and competitive release in South American pocket mice. *Oikos*, **98**, 3–16.

Bactrocera oleae (Rossi, 1790) in GBIF Secretariat. (2017). *GBIF Backbone Taxonomy*. Checklist dataset https://doi.org/10.15468/39omei Accessed via GBIF.org: 09 Feb 2019.

Baratella, V., Pucci, C., Paparatti, B. & Speranza, S. (2017). Response of *Bactrocera oleae* to different photoperiods and temperatures using a novel method for continuous laboratory rearing. *Biological Control*, **110**, 79–88.

Barbet-Massin, M., Jiguet F., Albert, C.H., & Thuiller, W. (2012). Selecting pseudoabsences for species distribution models: how, where and how many? *Methods in Ecology and Evolution*, **3**, 327–338.

Boccaccio, L. & Petacchi, R. (2009). Landscape effects on the complex of Bactrocera oleae parasitoids and implications for conservation biological control. *BioControl*, **54**, 607–616. Caudullo, G., Welk, E., & San-Miguel-Ayanz, J. (2017). Chorological maps for the main European woody species. *Data in Brief*, **12**, 662–666.

Dark, S.J. (2004). The biogeography of invasive alien plants in California: an application of GIS and spatial regression analysis. *Diversity and Distributions*, **10**, 1–9.

Dubuis, A., Pottier, J., Rion, V., Pellissier, L., Theurillat, J.P. & Guisan, A. (2011). Predicting spatial patterns of plant species richness: a comparison of direct macroecological and species stacking modelling approaches. *Diversity and Distributions*, **17**, 1122–1131.

FAOSTAT. (2019). Food and Agriculture Organization of the United Nations. *FAOSTAT Statistics Database*. Available at: http://www.fao.org/faostat/en/#home. Accessed: 09 Feb 2019.

Fick, S.E. & Hijmans, R.J. (2017). Worldclim 2: New 1-km spatial resolution climate surfaces for global land areas. *International Journal of Climatology*. Available at http://worldclim.org/version2. Accessed: 18 Feb 2018.

Figueiredo-González, M., Reboredo-Rodríguez, P., González-Barreiro, C., Carrasco-Pancorbo, A., Cancho-Grande, B. & Simal-Gándara, J. (2019). The involvement of phenolic-rich extracts from Galician autochthonous extra-virgin olive oils against the α glucosidase and α -amylase inhibition. *Food Research International*, **116**, 447–454.

Franklin, J. (2013). Species distribution models in conservation biogeography: developments and challenges. *Diversity and Distributions*, **19**, 1217–1223.

GBIF. (2018). The Global Biodiversity Information Facility. *What is GBIF?* Available at: https://www.gbif.org/what-is-gbif Accessed: 01 Jan 2019.

Gonçalves, F.M., Rodrigues, M.C., Pereira J.A., Thistlewood, H. & Torres, L.M. (2012). Natural mortality of immature stages of *Bactrocera oleae* (Diptera: Tephritidae) in traditional olive groves from north-eastern Portugal. *Biocontrol Science and Technology*, **22**, 837–854.

González, J.A., García-Barriuso, M. & Amich, F. (2010). Ethnobotanical study of medicinal plants traditionally used in the Arribes del Duero, western Spain. *Journal of Ethnopharmacology*, **131**(2), 343–355.

Grimm, V., Ayllón, D., & Railsback, S.F. (2017). Next-Generation Individual-Based Models Integrate Biodiversity and Ecosystems: Yes We Can, and Yes We Must. *Ecosystems*, **20**, 229–236.

Guisan, A., Petitpierre, B., Broennimann, O., Daehler, C., & Kueffer, C. (2014). Unifying niche shift studies: insights from biological invasions. *Trends in Ecology & Evolution*, **29**(5), 260–269.

Gutierrez, A.P., Ponti, L. ,& Cossu, Q.A. (2009). Effects of climate warming on Olive and olive fly (*Bactrocera oleae* (Gmelin)) in California and Italy. *Climatic Change*, **95**, 95–217. Gutierrez, A.P., Ponti, L., d'Oultremont, T., & Ellis, C.K. (2008). Climate change effects on poikilotherm tritrophic interactions. *Climatic Change*, **87** (Suppl 1), S167–S192.

Haber, G., & Mifsud, D. (2007). Pests and diseases associated with olive trees in the Maltese islands (Central Mediterranean). *The Central Mediterranean Naturalist*, **4**(3), 143–161.

Haniotakis, G.E. (2003). Olive pest control: present status and prospects. In *Proceedings of the IOBC/WPRS Conference on Integrated Protections of Olive Crops, Chania, Crete*.

Hijmans, R.J., Phillips, S., Leathwick, J., & Elith, J. (2017). dismo: Species distribution Modeling. *R package*, version 1.1-4. Available at: https://CRAN.Rproject.org/package=dismo Accessed: 01 Jan 2019.

IPCC. (2014). Climate Change 2014: Synthesis Report. *Contribution of Working Groups I, II and III to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change*. Core Writing Team, Eds R.K. Pachauri and L.A. Meyer. Geneva, Switzerland: IPCC. 151 pp. Kehlenbeck, H., Robinet, C., van der Werf, W., Kriticos, D., Reynaud, P., & Baker, R. (2012). Modelling and mapping spread in pest risk analysis: a generic approach. *OEPP/EPPO Bulletin*, **42**(1), 74–80.

Loumou, A., & Giourga, C. (2003). Olive groves: "the life and identity of the Mediterranean". *Agriculture and Human Values*, **20**, 87–95.

Marchi, S., Guidotti, D., Ricciolini, M., & Petacchi, R. (2016). Towards understanding temporal and spatial dynamics of *Bactrocera oleae* (Rossi) infestations using decade-long agrometeorological time series. *International Journal of Biometeorology*, **60**, 1681–1694.

Marcos, D. (2010). Arribes del Duero: Olivicultura al límite. www.variedadesdeolivo.com. Boletín nº 7. Available at: http://www.variedadesdeolivo.com/reportajes /7%20Arribes%20del%20Duero,%20olivicultura%20al%20limite.pdf. 16 pp. Accessed: 11 Feb 2019.

Merow, C., Smith, M.J., & Silander, J.A. (2013). A practical guide to Maxent for modeling species' distributions: what it does, and why inputs and settings matter. *Ecography*, **36**, 1058–1069.

Moreno, J.M. *et al.* (2005). A Preliminary Assessment of the Impacts in Spain due to the Effects of Climate Change. *ECCE PROJECT – Final Report*. Ministerio del Medio Ambiente - Universidad de Castilla la Mancha. 786 pp.

Moya-Laraño, J., Bilbao-Castro, J.R., Barrionuevo, G., Ruiz-Lupión, D., Casado, L.G., Montserrat, M., Melián C.J., & Magalhães, S. (2014). Eco-evolutionary spatial dynamics: rapid evolution and isolation explain food web persistence. In *Advances in ecological research*, Vol. 50, pp. 75–143. Eds J. Moya-Laraño, J. Rowntree and G. Woodward. Oxford: Oxford Academic Press. Accepted Article

Müller, F.A., Dias, N.P., Gottschalk, M.S., Mello, F.R., & Nava, D.E. (2019). Potential distribution of *Bactrocera oleae* and the parasitoids *Fopius arisanus* and *Psyttalia concolor*, aiming at classical biological control. *Biological Control*, **132**, 144–151.

Muriel, C. & Marcos, D. (2012). *El cultivo del olivo en Valladolid.* www.variedadesdeolivo.com. Available at: <u>http://www.variedadesdeolivo.com/reportajes</u> /<u>El%20olivo%20en%20Valladolid.pdf</u>. Reportaje nº 30. 8 pp. Accessed: 11 Feb 2019.

Muscarella, R., Galante, P.J., Soley-Guardia, M., Boria, R.A., Kass, J.M., Uriarte, M., & Anderson, R.P. (2014). ENMeval: An R package for conducting spatially independent evaluations and estimating optimal model complexity for Maxent ecological niche models. *Methods in Ecology and Evolution*, **5**, 1198–1205.

Olea europaea L. in GBIF Secretariat. (2017). GBIF Backbone Taxonomy. Checklist dataset https://doi.org/10.15468/39omei Accessed via GBIF.org: 09 Feb 2019.

Ordano, M., Engelhard, I., Rempoulakis, P., Nemny-Lavy, E., Blum, M., Yasin, S., Lensky, I.M., Papadopoulos, N.T., & Nestel, D. (2015). Olive fruit fly (*Bactrocera oleae*) population dynamics in the Eastern Mediterranean: Influence of exogenous uncertainty on a monophagous frugivorous insect. *PLoS ONE*, **10**(5), e0127798.

Pappas, M.L., Broufas, G.D., Koufali, N., Pieri, P., & Koveos, D.S. (2011). Effect of heat stress on survival and reproduction of the olive fruit fly *Bactocera* (*Dacus*) *oleae*. *Journal of Applied Entomology*, **135**, 359–3366.

Pearson, R.G., & Dawson, T.P. (2003). Predicting the impacts of climate change on the distribution of species: are bioclimate envelope models useful? *Global Ecology and Biogeography*, **12**, 361–371.

Peel, M.C., Grieser, J., Beck, C., Rudolf, B., & Rubel, F. (2007). Updated world map of the Koppen-Geiger climate classification. *Meteorologische Zeitschrift*, **15**, 259–263.

This article is protected by copyright. All rights reserved.

Pereira, J.A., Alves, M.R., Casal, S., & Oliveira, M.B.P.P. (2004). Effect of olive fruit fly infestation on the quality of olive oil from cultivars Cobrancosa, Madural and Verdeal Transmontana. *Italian Journal of Food Science*, **16**(3), 355–365.

Peterson, A.T., Soberón, J., & Sánchez-Cordero, V. (1999). Conservatism of ecological niches in evolutionary time. *Science*, **285**, 1265–1267.

Phillips, S.J., Anderson, R.P., & Schapire, R.E. (2006). Maximum entropy modeling of species geographic distributions. *Ecological Modelling*, **190**, 231–259.

Phillips, S.J., & Dudík, M. (2008). Modeling of species distributions with Maxent: new extensions and a comprehensive evaluation. *Ecography*, **31**, 161–175.

Phillips, S.J., Anderson, R.P., Dudík, M., Schapire, R.E., & Blair, M.E. (2017). Opening the black box: an open-source release of Maxent. *Ecography*, **40**, 887–893.

Phillips, S.J., Dudík, M. & Schapire, R.E. (2019). Maxent software for modeling species niches and distributions (Version 3.4.1). Available at: http://biodiversityinformatics.amnh. org/open source/maxent/. Accessed: 9 Feb 2019.

Podgomik, M., Vuk, I., Arbeiter, A., Hladnik, M., & Bandelj, D. (2013). Population fluctuation of adult males of the olive fruit fly *Bactrocera oleae* (Rossi) analysis in olive orchards in relation to abiotic factors. *Entomology News*, **123**, 15–25.

Ponti, L., Gutierrez, A.P., Ruti, P.M., & Dell'Aquila, A. (2014). Fine-scale ecological and economic assessment of climate change on olive in the Mediterranean Basin reveals winners and losers. *Proceedings of the National Academy of Sciences, USA*, **111**(15), 5598–5603.

R Core Team. (2018). *R: a Language and Environment for Statistical Computing*. Vienna, Austria: R Foundation for Statistical Computing. Available at: http://www.R-project.org. Accessed: 01 Feb 2018.

This article is protected by copyright. All rights reserved.

Reboredo-Rodríguez, P., González-Barreiro, C., Cancho-Grande, B., Simal-Gándara, J., & Trujillo, I. (2018). Genotypic and phenotypic identification of olive cultivars from northwestern Spain and characterization of their extra virgin olive oils in terms of fatty acid composition and minor compounds. *Scientia Horticulturae*, **232**, 269–279.

Reuter, H.I., Nelson, A., & Jarvis, A. (2007). An evaluation of void filling interpolation methods for SRTM data. *International Journal of Geographic Information Science*, **21**, 9, 983–1008.

Sutherst, R.W. (2013). Pest species distribution modelling: origins and lessons from history. *Biological Invasions*, **16**, 239–256.

Steven, J., & Phillips, S.J. (2009). A Brief Tutorial on Maxent. Network of Conservation Educators and Practitioners, Center for Biodiversity and Conservation, American Museum of Natural History. *Lessons in Conservation*, **3**, 108–135.

Swets, J.A. (1988). Measuring the accuracy of diagnostic systems. *Science*, **240**, 1285–1293.

Tanasijevic, L., Todorovic, M., Pereira, L.S., Pizzigalli, C., & Lionello, P. (2014). Impacts of climate change on olive crop evapotranspiration and region requirements in the Mediterranean region. *Agricultural Water Management*, **144**, 54–68.

Tobias Sing, Oliver Sander, Niko Beerenwinkel, Thomas Lengauer. (2005). ROCR: visualizing classifier performance in R. *Bioinformatics*, **21**(20), 3940–3941.

Thuiller, W., Lavorel, S., Sykes, M.T., & Araujo, M.B. (2006). Using niche-based modelling to assess the impact of climate change on tree functional diversity in Europe. *Diversity and Distributions*, **12**, 49–60.

Veloz, S.D., Williams, J.W., Blois, J.L., He, F., Otto-Bliesner, B., & Liu, Z. (2012). No-analog climates and shifting realized niches during the late quaternary: implications for 21st-century predictions by species distribution models. *Global Change Biology*, **18**, 1698–1713.

Wang, X.G., Levy, K., Nadel, H., Johnson, M.W., Blanchet, A., Argov, Y., Pickett, C.H., & Daane, K.M. (2013). Overwintering survival of olive fruit fly (Diptera: Tephritidae) and two introduced parasitoids in California. *Environmental Entomology*, **42**(3), 467–76.

Wang, X.G., Johnson, M.W., Daane, K.M., & Nadel, H. (2009). High summer temperatures affect the survival and reproduction of olive fruit fly (Diptera: Tephritidae). *Environmental Entomology*, **38**(5), 1496–1504.

Yackulic, C.B., Chandler, R., Zipkin, E.F., Royle, J.A., Nichols, J. D., Campbell Grant, E.H., & Veran, S. (2013). Presence-only modelling using MAXENT: when can we trust the inferences? *Methods in Ecology and Evolution*, **4**, 236–243.

Yokoyama, V.Y., & Miller, G.T. (2007). Olive fruit fly biology and cultural control practices in California. *International Organization for Biological Control Western Palearctic Section Bulletin*, **30**, 263–269.

Code in database	Bioclimatic variable
bio1	Annual mean temperature
bio2	Mean diurnal range (mean of monthly (max temp - min temp))
bio3	Isothermality (bio2/bio7) (× 100)
bio4	Temperature seasonality (standard deviation × 100)
bio5	Max temperature of warmest month
bio6	Min temperature of coldest month
bio7	Temperature annual range (bio5-bio6)
bio8	Mean temperature of wettest quarter
bio9	Mean temperature of driest quarter
bio10	Mean temperature of warmest quarter
bio11	Mean temperature of coldest quarter
bio12	Annual precipitation
bio13	Precipitation of wettest month
bio14	Precipitation of driest month
bio15	Precipitation seasonality (coefficient of variation)
bio16	Precipitation of wettest quarter
bio17	Precipitation of driest quarter
bio18	Precipitation of warmest quarter
bio19	Precipitation of coldest quarter

Table 1. Description of bioclimatic variables provided by the WorldClim database.

Table 2. Percent of contribution of each selected bioclimatic driver used to model the climatic suitability of *Bactrocera oleae* in the Iberian Peninsula.

Bioclimatic variable	Percent contribution
Precipitation of coldest quarter	22.20
Precipitation of driest month	20.00
Mean diurnal range (Mean of monthly (max temp - min temp))	14.90
Min temperature of coldest month	13.80
Temperature seasonality (Standard deviation ×100)	12.30
Precipitation seasonality (Coefficient of variation)	9.10
Temperature annual range (Max temperature of warmest month - Min temperature of coldest month)	5.40
Mean temperature of wettest quarter	1.60
Mean temperature of driest quarter	0.50
Isothermality (Mean diurnal range / Temperature annual range) × 100	0.20

Figure captions

Figure 1. Occurrence-based distribution map of *Bactrocera oleae* throughout the Iberian Peninsula. Blue dots represent occurrence records projected on a digital elevation model. Altitude is expressed in meters (m).

Figure 2. Multicollinearity analysis of the pool of 19 bioclimatic variables available from WorldClim for the selected geographical extent of analysis (the Mediterranean basin: 15°W, 40°E, 27°S, 50°N). Pies represent the amount of Pearson correlation within each pair of variables. The description of bioclimatic variables is available at Table 1.

Figure 3. Mean of 20 Maxent prediction replicates using the linear and quadratic (LQ) model for the climatic suitability of *Bactrocera oleae* in the Iberian Peninsula. Colors have been chosen to allow risk assessment in terms of pest potential presence (i.e. reddish areas indicate higher climatic suitability).

Figure 4. Response curve of each selected bioclimatic driver used to model the climatic suitability of *Bactrocera oleae* in the Iberian Peninsula. Asterisks indicate the three most contributing variables to the model: precipitation of coldest quarter (bio19), precipitation of driest month (bio14), and mean diurnal range (bio2). Codes of variables correspond to Table 1.

Figure 5. Extended distribution (mean of 20 Maxent prediction replicates using the linear and quadratic (LQ) model) of the climatic suitability (p) of *Bactrocera oleae* in the Iberian Peninsula. Black thick lines represent country administrative limits. Black thin lines represent administrative limits of districts and autonomous communities for Portugal and Spain respectively. Orange dots represent occurrence records of *B. oleae*, green dots represent occurrence records of *Olea europaea*, and the green area represents the current

chorological map of *O. europaea*. The red, purple, and blue circles encompass the autonomous community of Galicia, the central area of Castilla y León, and the Spanish side of the International Douro Natural Park. These areas of new olive cultivation with no previous capturing of the pest have been suggested by the model as suitable for pest establishment. Reddish areas indicate higher climatic suitability.



















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