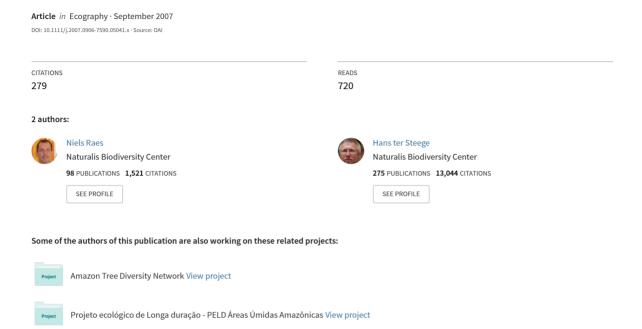
A null-model for significance testing of presence-only species distribution models





doi: 10.1111/j.2007.0906-7590.05041.x

© 2007 The Authors. Journal compilation © 2007 Ecography Subject Editor: Miguel Araújo. Accepted 3 August 2007

A null-model for significance testing of presence-only species distribution models

Niels Raes and Hans ter Steege

N. Raes (raes@nhn.leidenuniv.nl), National Herbarium of the Netherlands, Leiden Univ. branch, Einsteinweg 2, NL-2300 RA, Leiden, the Netherlands – H. ter Steege, Inst. of Environmental Biology, Section Plant Ecology and Biodiversity and the National Herbarium of the Netherlands, Utrecht Univ. branch, Utrecht Univ., Sorbonnelaan 14, NL-3584 CA, Utrecht, the Netherlands.

Species' distribution models (SDMs) attempt to predict the potential distribution of species by interpolating identified relationships between species' presence/ absence, or presence-only data on one hand, and environmental predictors on the other hand, to a geographical area of interest. Currently, they are widely applied in biogeography, conservation biology, ecology, palaeo-ecology, invasive species studies, and wildlife management (Guisan and Zimmermann 2000, Araújo and Pearson 2005, Thuiller et al. 2005, Peterson 2006, Araújo and Guisan 2006, Guisan et al. 2006). More recently, vast numbers of herbarium and natural history museum collections have become available (Graham et al. 2004) and techniques to apply this special type of presence-only data have been developed (Hirzel et al. 2002, Anderson et al. 2003, Pearce and Boyce 2006, Elith et al. 2006, Phillips et al. 2006). Despite the widespread use of SDMs, several high-priority research interests remain to be investigated (Guisan and Thuiller 2005, Araújo and Guisan 2006). One of these is the improvement of SDM validation, or the quantification of a model's predictive performance (Araújo and Guisan 2006). The fact that the standard validation procedures for an SDM are not sufficient to assess the applicability of an SDM in a predictive context, was first shown by Olden et al. (2002). They showed that after SDM validation it is critical to assess whether the SDM prediction differs from what would be expected on the basis of chance alone. SDMs producing random predictions are neither helpful nor useful (Olden et al. 2002). Thus, in this paper we introduce a null-model methodology that allows testing whether SDMs developed with presence-only data differ significantly from what would be expected by chance. We also

demonstrate that it is critical and possible to correct for collector-bias in specimen data in this test.

SDM validation and measures of accuracy

Validation of SDMs can be carried out with several different measures of model accuracy. The most widely applied measures of model accuracy include sensitivity, specificity, Cohen's kappa, and the area under the curve (AUC) of the receiver operating characteristic (ROC) plot (Fielding and Bell 1997, Manel et al. 2001, McPherson et al. 2004). Most measures of SDM accuracy, including the four mentioned above, are directly or indirectly derived from a confusion matrix (Fielding and Bell 1997). Sensitivity quantifies the proportion of observed presences correctly predicted as presence, the true positive fraction. Specificity quantifies the true negative fraction. Cohen's kappa quantifies overall agreement between predictions and observations, corrected for agreement expected to occur by chance. These three measures of accuracy require that probabilities of occurrence obtained with SDMs are transformed into discrete presences or absences, for which purpose a threshold of 0.5 is commonly used (McPherson et al. 2004, Liu et al. 2005, Jiménez-Valverde and Lobo 2007). The AUC value of the ROC plot is a method that does not require discrete presence/ absence predictions, and is therefore a measure of accuracy that is threshold independent (Pearce and Ferrier 2000, McPherson et al. 2004).

The ROC plot is obtained by plotting sensitivity as a function of the falsely-predicted positive fraction, or

commission error (1-specificity), for all possible thresholds of a probabilistic prediction of occurrence. The resulting area under the ROC curve provides a single measure of overall model accuracy, which is independent of a particular threshold. AUC values range from 0 to 1, with a value of 0.5 indicating model accuracy not better than random, and a value of 1.0 indicating perfect model fit (Fielding and Bell 1997). An AUC value can be interpreted as indicating the probability that, when a presence site (site where a species is recorded as present) and an absence site (site where a species is recorded as absent) are drawn at random from the population, the presence site has a higher predicted value than the absence site (Elith et al. 2006, Phillips et al. 2006).

All four measures of model accuracy were tested extensively for statistical artefacts, and the AUC value was the only measure of SDM accuracy that was invariable to the proportion of the data representing species' presence, known as prevalence (Pearce and Ferrier 2000, Manel et al. 2001, McPherson et al. 2004). Insensitivity to prevalence is of special relevance when the AUC values are used to assess model accuracy for SDMs that have been developed with presence-only data. When the required absences are lacking, they are replaced by pseudo-absences. Pseudo-absences are sites, randomly selected across the geographical area of interest, at localities where no species presence was recorded and for which species occurrence is set as absent (Ferrier et al. 2002, Anderson et al. 2003, Elith et al. 2006, Phillips et al. 2006). A sufficiently large sample of pseudo-absences is needed to provide a reasonable representation of the environmental variation exhibited by the geographical area of interest, typically 1000-10000 points (Stockwell and Peters 1999, Ferrier et al. 2002, Phillips et al. 2006). These large numbers of pseudo-absences automatically result in low prevalence values. The number of records by which a species is represented in herbaria and natural history museums range from 1 to 150-200 records (Stockwell and Peterson 2002). Even when a species is represented by 200 unique presence-only records and 1000 pseudo-absences are used, prevalence is only 16.7% (200/1200).

A major drawback of using pseudo-absences, however, is that the maximum achievable AUC value indicating perfect model fit, is no longer 1, but 1-a/2 (where a is the fraction of the geographical area of interest covered by a species' true distribution, which typically is not known (Phillips et al. 2004, 2006). Nevertheless, random prediction still corresponds to an AUC value of 0.5. Therefore, standard thresholds of AUC values indicating SDM accuracy (e.g. the threshold of AUC > 0.7 that is often used; Pearce and Ferrier 2000, Swets et al. 2000, Manel et al. 2001), do not apply.

A null-model approach for significance testing of presence-only SDMs

To test the significance of an SDM we propose to test the AUC value (of the SDM) against a null distribution of expected AUC values based on random collection data (sensu Olden et al. 2002). A null-distribution, or null-model, is a model that is based on randomizations of ecological data or random sampling from a known or imagined distribution (Jetz et al. 2004, Gotelli and McGill 2006). A null-model is straightforward in theory and closely resembles hypothesis testing in conventional statistical analysis. To build a null-model, first the AUC value of the real SDM is determined. Next, a null-model is generated by randomly drawing collection localities without replacement, from the geographical area for which the species distribution is modelled. The number of randomly drawn collection localities is equal to the actual number of collections for that species. This is repeated 999 times to generate a frequency histogram of AUC values, expected if the null hypothesis is true. The position of the observed AUC value in the null distribution of the "randomly" generated AUC values is then used to assign a probability value, just as in a conventional statistical analysis (Dolédec et al. 2000, Olden et al. 2002, Gotelli and McGill 2006). We use a one-sided 95% confidence interval (C.I.) since we are only interested in whether an SDM performs significantly better than expected by chance, rather than assessing whether it performs significantly worse. We interpret a significant model to indicate that the relations between species' presence localities and the predictor variable values at those locations are stronger than can be expected by chance.

An additional advantage of significance testing of an SDM with a null-model is that we can use all presence records to develop and test the SDM. Common practice in measuring an SDM's accuracy is the splitsample approach. This approach splits the available species records into a training and test sample (Fielding and Bell 1997). It is assumed that a randomly selected test sample from original data constitutes independent observations, which can be used for statistical testing (Araújo et al. 2005). However, such a test sample is not fully independent due to spatial autocorrelation (Araújo et al. 2005, McPherson and Jetz 2007). Moreover, dependent on the random split, different values of SDM accuracy may be obtained (Phillips et al. 2006). Phillips et al. (2006) showed that SDMs for a species represented by 128 records and 10 different random splits, yielded AUC values ranging from 0.819 until 0.903. More extremely, our unpublished results yielded AUC values for a species represented by 8 records ranging between 0.079 and 0.912 based on 100 random splits.

Testing an SDM against a null-model, however, could suffer from one more problem. When drawing random points from a geographical area one assumes that collectors visited all localities equally well. If this condition is not met, which is likely to be the case (Reddy and Davalos 2003, Romo et al. 2006, Hortal et al. 2007), the randomly drawn points, that are used to develop the null-model, might include ecological conditions that are not represented by the localities from where actual collections were gathered. This bias could results in a significant deviation from the null-model for species that are randomly distributed over the actual collection localities.

The impact of collection bias on significance testing

SDMs predict the presence and absence of a species for a given geographical area, based on the localities where the records were collected and the values of environmental predictors at those sites. SDMs are especially useful when only part of the entire geographical area has been sampled, as is generally the case. This works fine as long as the collection localities are randomly spread over the complete geographical area. Unfortunately, collectors tend to visit areas which are easily accessible, such as areas close to cities, roads, rivers, and nature reserves resulting in serious collection biases (Reddy and Davalos 2003, Parnell et al. 2003, Kadmon et al. 2004, Hortal et al. 2007). The influence of collection biases on the accuracy of SDMs largely depends on the range of values of each of the environmental variables covered by the collection localities, known as climatic, or environmental bias (Kadmon et al. 2003, 2004). Kadmon et al. (2003) showed that environmental biases, expressed as the degree of sampling bias with respect to the environmental conditions under which a species is known to occur, had a significant negative effect on the predictive accuracy of the SDM. Although this is a serious issue of concern (Araújo and Guisan 2006), it is not specific to any methodology used to develop SDMs. However, it is relevant when the accuracy of an SDM is tested against a null-model.

When collecting is environmentally biased, an SDM is more likely to deviate significantly from a random null-model that does not include such bias. When, for example, collection localities are biased for mean annual temperature, a significant part of the species' actual temperature range could remain unsampled. When these data are used in an SDM that is tested against a null-model, based on records that were randomly drawn from the entire study area, this species will possibly show a preferred mean annual temperature range

compared to the randomly drawn points. It will accordingly more likely deviate significantly from the null-model than its actual range would justify. Such collection bias might thus result in certain areas being systematically underpredicted by the SDM. It should be noted, however, that this is true for all distribution modelling methods and can only be solved by additional data collection.

Fortunately, the problem of having a higher chance of significantly deviating from a randomly drawn nullmodel if collections are biased, can be solved by restricting the randomly drawn points to all known collection localities. Thus, drawing the null-model from a biased distribution. To test for environmental bias in known collection localities a distribution model using all known collection localities is tested against a nullmodel developed by 100-1000 times drawing an equal number of random points from the entire study area. If the distribution model's accuracy of known collection localities deviates significantly from this "second" nullmodel, then we conclude that the collection localities are environmentally biased. If this is the case then the SDMs have to be tested against a null-model that is based on actual collection localities.

A case study based on Bornean plant collections

To illustrate the applicability of a null-model approach to select SDMs that deviate significantly from random expectation, we selected all occurrences of the genus *Shorea* (Dipterocarpaceae) on the Malesian island Borneo (ca 8°N–5°S, 108°–120°E; Fig. 3) from the BRAHMS database of plant collections present at the National Herbarium of the Netherlands, Leiden Univ., the Netherlands. *Shorea* was selected because this genus has been thoroughly taxonomically revised and species identifications are reliable (Ashton 1983). The database contained 4466 records of 147 *Shorea* species for Borneo. Out of these 147 species, 116 were represented by 5, or more, unique collection localities. For those species, we developed SDMs.

To model the species distributions we used environmental predictor variables with a 5 arc-minutes resolution (ca 10 km at the equator). We selected the digital elevation model (DEM) and the 19 bioclimatic variables of the current conditions (ca 1950–2000) from the WORLDCLIM dataset (<http://www.worldclim.org>) for Borneo (Hijmans et al. 2005). Additionally, we selected 15 FAO soil variables (Anon. 2002). We also included a measure of the effect of the El Niño Southern Oscillation Event (ENSO). This variable was expressed as the relative average annual difference in Normalized Difference Vegetation Index (NDVI)

between the months of an ENSO, and a non-ENSO year. To this dataset we added the Walsh's index (Walsh 1996, Leigh Jr 2004). This index integrates the effects of annual rainfall and its seasonality. Finally, the elevation range derived from the SRTM 90 m Digital Elevation Data (http://srtm.csi.cgiar.org/) was added. All data layers were scaled to 5 arc-minute resolution, and resampled to the geographical extent of the most restricted FAO soil variable data layers. This resulted in 8577 data cells for Borneo. All data layer manipulations were performed with Manifold GIS (Manifold Net).

To model Shorea species distributions of Borneo we used Maxent (ver. 2.3.0; http://www.cs.princeto- n.edu/~shapire/maxent/>) (Phillips et al. 2006). Maxent, or the maximum entropy method for species' distribution modelling, estimates the most uniform distribution ("maximum entropy") across the study area, given the constraint that the expected value of each environmental predictor variable under this estimated distribution matches its empirical average (average values for the set of species' presence records) (Hernandez et al. 2006, Phillips et al. 2006). Maxent was specifically developed to model species distributions with presence-only data and has outperformed most other modelling applications (Elith et al. 2006, Hernandez et al. 2006, Pearson et al. 2007). An added advantage of Maxent is that it also performs the ROC statistical analysis. Since we tested whether an SDM's AUC value deviates significantly from a null-model, the "random test percentage" was set to zero resulting in training data only. To avoid the inclusion of multiple presence records in one grid cell per species we set Maxent to "remove duplicate presence records". This reduced the total available presence records for the 116 Shorea species represented by at least five records to 2552. The modelling rules were set to "Auto Features" using only linear features when <10 records were available, adding quadratic features for SDMs developed with 10 or more and <15 records, and including hinge features for species with 15 or more records. Maxent adds product and threshold features for those species represented by 80, or more, records. However, we set Maxent to use linear, quadratic and hinge features for all species represented by at least 15 records, due to odd behaviour of Maxent when product and threshold features were added (explained in the discussion). For each of the 116 Shorea species we developed an SDM with Maxent using all presence records under the modelling rules as described above. The number of unique records per species ranged from 5 until 92 (Appendix, Table S1, "# records"). The AUC values of all Shorea SDMs are presented as dots in Fig. 1, and under "AUC" in Table S1 (Appendix).

Testing SDMs against a null-model

To test whether Shorea SDMs significantly differed from what would be expected by chance, we calculated the 95% C.I. AUC value for each number of records by which the Shorea species were represented. We developed frequency histograms of expected AUC values by randomly drawing points without replacement from all 8577 available cells of Borneo (999 times), and model these with Maxent under the same conditions as the Shorea species. We developed frequency histograms of expected AUC values for 5-30 records (26 distributions), for 35-50 records with intervals of 5 records (4 distributions), and for 60-100 records with intervals of 10 records (5 distributions). For each frequency histogram, we assessed the 95% C.I. upper limit AUC value, by ranking the 999 AUC values and selecting the 949th value $(0.95 \times 999 = 949; \text{ Fig. 1},$ triangles). For each of the three resulting sets of 95% C.I. AUC values we applied a curve-fit (Fig. 1, asterisks). The fitted 95% C.I. AUC values of the null-models for the number of records by which each Shorea species is represented, are given in Table S1, "95% C.I. All" (Appendix).

With the fitted 95% C.I. AUC values, it is now easy to assess which of the *Shorea* species has an accuracy of its SDM that is significantly higher than expected by chance alone (p < 0.05). This was the case for 105 of the 116 *Shorea* species (91%) which were modelled (Appendix, Table S1, "95% C.I. All").

Testing SDMs against a bias corrected null-model

In order to assess whether the known collection localities are environmentally biased, we selected all databased and georeferenced plant specimen records from Borneo that were present in the BRAHMS database of the National Herbarium of the Netherlands. In total the database contained 142 097 properly georeferenced records. These records could be assigned to 1837 of the total of 8577 grid cells of Borneo. This means that only 21.4% of the grid cells of Borneo have been visited by collectors who actually made any collections (Fig. 3). The collections are clearly geographically biased, as evident from the geographical distribution of the dark grey squares in Fig. 3. However, predicting species presences or absences in non-visited areas is one of the major applications of the use of SDMs, so this should not be a major problem. More importantly, it is to assess whether these localities are environmentally biased, or whether certain conditions are over- or under-represented with respect to the environmental conditions for the entire geographical area of Borneo. For this purpose, we first developed a distribution model of the 1837 collection localities and

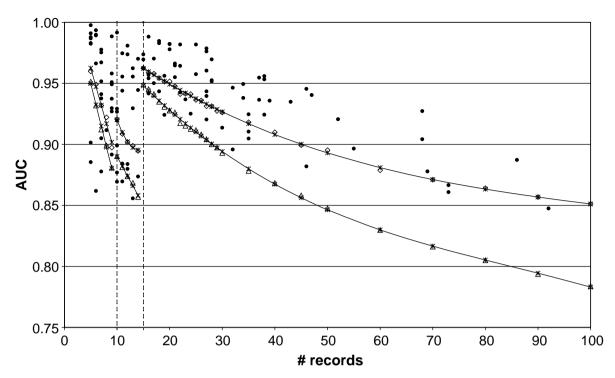


Fig. 1. Species' distribution model (SDM) AUC values (\bullet), the 95% confidence interval (C.I.) AUC values of the randomly drawn null-models (Δ), and the 95% C.I. AUC values of the environmentally bias corrected null-models (Δ). Asterisks give the fitted 95% C.I. AUC values for both series of null-models connected by a line. Vertical dotted lines indicated the consecutive addition to the initial linear modelling features, of quadratic, and hinge features by Maxent. SDM AUC values that are higher than their corresponding 95% C.I. AUC value of the fitted null-model, significantly deviate from what would be expected by random chance (p < 0.05).

assessed the model's AUC value. Then, we developed a frequency histogram of expected AUC values on basis of 1837 randomly drawn localities from the 8577 cells of Borneo (100 reps). Unfortunately the AUC value of the distribution model based on the collections localities, is significantly different from random expectation (p < 0.01; Fig. 2), hence, the collection localities are also environmentally biased.

The implication that collecting effort is environmentally biased for Borneo is that SDMs cannot be tested with null-models drawn randomly from all 8577 grid cells of Borneo. To overcome this problem we developed a second series of null-models, in the same way as described above, but now randomly drawing from the 1837 known collection locality cells. The resulting 95% C.I. AUC values of these null-models are presented as diamonds in Fig. 1. Again, we applied a fit through these values to establish the 95% C.I. AUC values against which the SDM AUC values were tested. These values are given in Table S1 (Appendix) under "95% C.I. Bias". Now only 80 of the 116 Shorea species (69%) have a SDM AUC value significantly different from a (bias corrected) null-model (Appendix, Table S1, "95% C.I. Bias"; Fig. 4A, C). This means that an additional 25 SDMs were rejected, compared to testing against environmentally unbiased null-models.

Discussion

By proposing the use of null-models in the field of presence-only species' distribution modelling, we introduce a novel methodology that allows for significance testing of SDMs. The new methodology makes use of all presence records to develop an SDM and to test its accuracy with the AUC procedure, a threshold-and prevalence-independent single measure of SDM accuracy. A significant SDM indicates that correlations between species' presence localities and the environmental predictor variables, as identified and interpolated by Maxent, deviate from random chance.

Secondly, we show the importance of correcting for environmental biases in data collection. Null models which incorporate the environmental bias within the collection data reject a significant fraction of SDMs which are significant based upon a randomly drawn null-model. If the collection localities are environmentally biased and a species is found throughout the subset

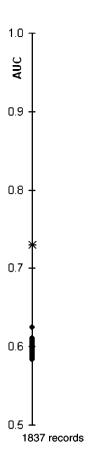


Fig. 2. The AUC value of the model based on the 1837 collection cells (*) and the 100 AUC values (\bullet) of models based on 1837 randomly drawn cells from the total 8577 cells of Borneo, indicating that the 1837 collection cells are significantly environmentally biased (p < 0.01).

of values represented by the collection localities, this species is likely to differ significantly from a null-model which is drawn from the total range of values. This results in an SDM that is an underestimation of the true geographical range of the species. This, because under these conditions the full range of values under which the species truly occurs is not incorporated in the SDM.

Although we introduce a null-model approach to the field of presence-only species' distribution modelling, the use of null-models for significance testing was successfully applied by Olden et al. (2002) for presence-absence SDM testing, and by Dolédec et al. (2000) in the field of community analysis. Our methodology differs from Olden et al. (2002) in that we adapted the null-model approach to make use of presence-only data, and test an SDM accuracy with the threshold- and prevalence independent AUC procedure (Swets 1988, Manel et al. 2001, McPherson et al. 2004, Guisan et al. 2006). This is important as in our case study the number of species presence records ranged from 5 to

92. Combined with 1000 pseudo-absences this resulted in prevalence values as low as 0.5–8.4%.

We interpret that species, for which the SDM AUC value significantly deviates from a null-model, have specific niche requirements that were met at the localities where they were collected. This agrees with the reasoning of Dolédec et al. (2000). They analysed community data with a new multivariate method they called OMI (for outlying mean index), to measure the distance between mean habitat conditions used by a species, and the mean habitat conditions of the sampling area (Dolédec et al. 2000). The OMI value (analogous to the SDM AUC value) of a species is tested against the null-distribution of "1000 random permutation values obtained under the null hypothesis that the species is indifferent to its environment". For species that significantly deviated from this, "theoretical ubiquitous species that tolerates the most general habitat conditions", it was concluded that the observed species position in habitat differed significantly from what would be expected by chance. This OMImethodology was later implemented in a species distribution modelling technique called ecologicalniche factor analysis (ENFA) (Hirzel et al. 2002), but testing against a null-distribution was never formalized.

The first to notice that accuracy assessment of presence-only SDMs alone was not sufficient, and SDMs should be tested against a random null hypothesis were Anderson et al. (2002). They used the split sample approach, dividing the available presence records of a species in a 75% training and 25% test dataset. After SDM development using the training data, they tested whether test points fell into areas predicted presence more often than expected at random, given the overall proportion of pixels predicted presence vs. predicted absence for that species (Anderson et al. 2002). The latest advances in this methodology were recently made, by introducing a jack-knife (or "leave-one-out") procedure for SDM accuracy assessment and a combined p-value significance test for significance testing of the presenceonly SDMs (for details see Pearson et al. 2007). However, this methodology does not take into account possible environmental bias in collection localities. If the full niche of a species is not represented by the collection localities, the species' predicted distribution will be smaller than its true distribution. Modelling applications, such as Maxent, are very well capable of predicting the species' distribution based on the available presence records without model under-fitting. A smaller predicted species' distribution automatically results in higher chance of significantly deviating from the random null hypothesis, the same way as in our case study more species significantly deviate from a randomly drawn nullmodel than from a null-model that is corrected for environmental bias (Fig. 1; Appendix, Table S1). Additionally, the jack-knife validation approach may

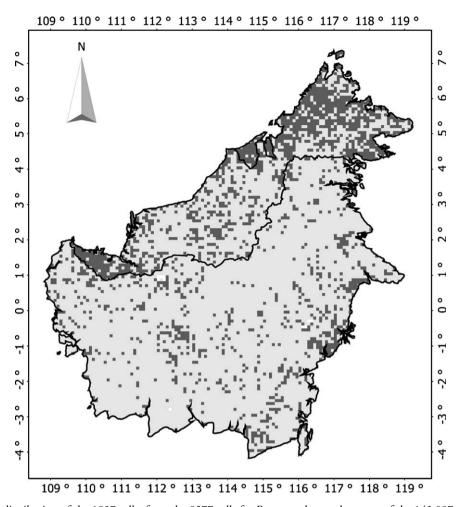


Fig. 3. Spatial distribution of the 1837 cells, from the 8577 cells for Borneo, where at least one of the 142 097 collections was made (indicated by dark grey squares). Light grey squares indicate the remaining 6740 unsampled cells. White cells indicate large lake areas for which no environmental data were available.

lead to overoptimistic estimates of the predictive power with larger sample sizes (Pearson et al. 2007).

Our results showed the importance of correcting for environmental bias in known collection localities when null-models are used for significance testing of presence-only SDMs. However, at the same time this requirement hampers the general applicability of the methodology. In our case study, we could make use of the full herbarium record database of the National Herbarium of the Netherlands, containing 142 097 georeferenced plant specimen records found in 1837 of the 8577 grid cells of Borneo. We recognize that this amount of data will not always be available. However, since the majority of collections has been made in close proximity to roads, rivers, cities, and nature reserves (Reddy and Davalos 2003, Kadmon et al. 2004, Hortal et al. 2007), an alternative could be to use a distance buffered road-river map, including cities and nature reserves, to select the grid cells and test these cells for environmental biases. If these cells are environmentally biased, the SDMs can then be tested against a nullmodel drawn from this pool of cells. However, this approach is less accurate and requires further testing.

Our results showed that for low prevalence values very high AUC values can be expected from randomly drawn points (Fig. 1; Appendix, Table S1). Olden et al. (2002) too reported such high accuracy values for low (and high) prevalence. The 95% C.I. AUC value of the bias corrected null-model for 15 records (prevalence = 1.48%) was as high as 0.9622 (Fig. 1; Appendix, Table S1). Nevertheless, 80 of the 116 *Shorea* species (69%) had an SDM AUC value higher than the 95% C.I. AUC value of the bias corrected null-model. Dolédec et al. (2000) reported that, for their application of a null-model for two case studies, 59 and 85% of their species respectively, had significant results. Pearson et al. (2006)

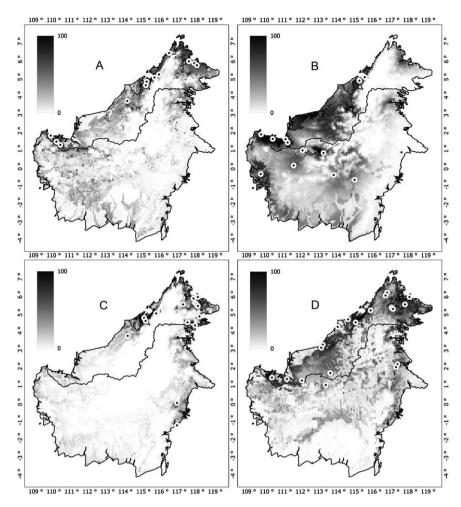


Fig. 4. A–D. Maxent predictions for two significant SDMs (A, C), and two non-significant SDMs (B, D). Collection localities are indicated by dots. (A) *Shorea isoptera* P.S. Ashton, (Appendix, Table S1, #45), (B) *S. platycarpa* Heim (Appendix, Table S1, #49), (C) *S. confusa* P.S. Ashton (Appendix, Table S1, #57), and (D) *S. macroptera* Dyer (Appendix, Table S1, #66).

report values from 62 to 100% depending on the modelling application and thresholds that were used. Our testing against a randomly drawn null model resulted in a comparably high percentage (91%) of significant SDM AUC values (p < 0.05). All these results are higher than the 50% reported by Olden et al. (2002).

Both the AUC values of the two null-models, and the SDMs, show a decreasing trend with increasing number of records (Fig. 1; Appendix, Table S1). This is most likely the result of applying ROC plots to SDMs, developed with presence-only data, reducing the maximum AUC value dependent on the species' true distribution (Phillips et al. 2006). Assuming that the predicted species' distributions are a good proxy for the species' true distributions, we assessed the area for which species were predicted to be present by converting the continuous probabilistic Maxent predictions of occurrence to discrete presence-absence values. We used the

maximized sensitivity-specificity sum threshold for this purpose (Liu et al. 2005). Regressing significant SDMs AUC values against the area for which they were predicted to be present (Appendix, Table S1; "Area (in%)") revealed a significant negative linear correlation $(AUC = 0.9913 - 0.0029 \times Area; p < 0.001; R^2 =$ 0.576). We consider this as a strong indication that it is not the accuracy of the models that is reduced but merely that the maximum achievable AUC value is reduced due to an increased true distribution of the species. We therefore do not support the statement that the predictive accuracy of the model decreases when the extent of a species distribution increases, as suggested by Hernandez et al. (2006). When an increased predicted distribution and related lower SDM AUC value is caused by a broad niche amplitude, however, as is the case for habitat generalists, an SDM accuracy is more likely not to deviate from a null-model and the SDM can therefore not be used. This is possibly the case for the SDMs presented in Fig. 4B, D (Appendix, Table S1; #49 and #66).

A consequence of implementing the proposed use of null-models for SDM evaluation, is that SDM accuracy is tested with the same data used to develop models, i.e., a form of model verification (Araújo and Guisan 2006). A problem with this approach is that SDMs may overfit the calibration, or training data (Araújo et al. 2005). Over-fitting, however, is not considered a problem if the goal is to describe a pattern and simultaneously reduce false negatives: i.e. true observations that are not predicted by the model (Araújo and Guisan 2006). An advantage is that all observations are used to develop the SDMs, making optimal use of all available information. If the modelled species' distributions are intended to be used for conservation planning, verification is an approved method to test whether an SDM performs as intended. However, if the models are used to predict range shifts under different climate change scenario's, or to assess the possible invasiveness of a species, an SDM's ability to correctly predict independent test data is preferred (Araújo and Guisan 2006). It should be kept in mind, however, that SDMs, as they are applied in this study, predict the potential distribution of a species and do not take into account competition, and historical or present geographical barriers (Soberón and Peterson 2005, Peterson 2006). Most studies addressing these issues use data partitioning methods to allocate records to training and test datasets. The most familiar technique is one-time data-splitting (Araújo et al. 2005). Our unpublished results indicated, however, that dependent on the spatial distribution and the random split of the records, SDM accuracies could be very different.

An advantage of Maxent is its ability to counteract the tendency of SDMs to over-fit when few presence records are available, due to its regularization procedure (Hernandez et al. 2006, Phillips et al. 2006). Therefore, we used the standard settings of Maxent. However, the null-models developed for 80, 90, and 100 records developed with the modelling rules set to "auto features" and the regularization multiplier set to 1, resulted in increasing 95% C.I. AUC values indicating over-fitting of the models (data not shown). For this reason, we set the modelling rules to use linear, quadratic and hinge functions to develop the null-distributions for those numbers of records and the SDMs developed with more than 79 records.

We are aware that spatial autocorrelation in the distribution of the species records and environmental variables may also influence SDM accuracy. Our intention was not to investigate the influence of spatial autocorrelation on SDM accuracy, however, but to provide a methodology for significance testing of presence-only SDMs. Simultaneously we showed that

the evaluation of presence-only SDM quality based on subjective ROC plot thresholds (e.g. $AUC \ge 0.7 =$ useful), cannot be applied. With this contribution, we hope to provide SDM users with a valuable tool to identify those species that can be accurately modelled, while providing an additional reason for being cautious about interpretations of SDMs that are not tested for significance.

Acknowledgements – We would like to thank Steven Phillips for adapting the Maxent application several times to allow us to perform the ROC analyses of the null-models. We also thank Pieter Baas, Chuck Cannon, Peter Hovenkamp, Marco Roos, Ferry Slik and four anonymous referees, all of whose comments allowed us to make useful improvements to the manuscript.

References

Anderson, R. P. et al. 2002. Geographical distributions of spiny pocket mice in South America: insights from predictive models. – Global Ecol. Biogeogr. 11: 131–141.

Anderson, R. P. et al. 2003. Evaluating predictive models of species' distributions: criteria for selecting optimal models.
 Ecol. Modell. 162: 211–232.

Anon. 2002. Terrastat; global land resources GIS models and databases for poverty and food insecurity mapping.
 FAO, Land and Water Digital Media Series # 20.

Araújo, M. B. and Pearson, R. G. 2005. Equilibrium of species' distributions with climate. – Ecography 28: 693–695.

Araújo, M. B. and Guisan, A. 2006. Five (or so) challenges for species distribution modelling. – J. Biogeogr. 33: 1677–1688.

Araújo, M. B. et al. 2005. Validation of species-climate impact models under climate change. – Global Change Biol. 11: 1504–1513.

Ashton, P. S. 1983. Dipterocarpaceae. – In: van Steenis, C. G. G. J. (ed.), Flora Malesiana I. Vol. 9, part 2. Martinus Nijhoff, The Hague, pp. 237–552.

Dolédec, S. et al. 2000. Niche separation in community analysis: a new method. – Ecology 81: 2914–2927.

Elith, J. et al. 2006. Novel methods improve prediction of species' distributions from occurrence data. – Ecography 29: 129–151.

Ferrier, S. et al. 2002. Extended statistical approaches to modelling spatial pattern in biodiversity in northeast New South Wales. I. Species-level modelling. – Biodiv. Conserv. 11: 2275–2307.

Fielding, A. H. and Bell, J. F. 1997. A review of methods for the assessment of prediction errors in conservation presence/absence models. – Environ. Conserv. 24: 38–49.

Gotelli, N. J. and McGill, B. J. 2006. Null versus neutral models: what's the difference? – Ecography 29: 793–800.

Graham, C. H. et al. 2004. New developments in museum-based informatics and applications in biodiversity analysis.Trends Ecol. Evol. 19: 497–503.

- Guisan, A. and Zimmermann, N. E. 2000. Predictive habitat distribution models in ecology. – Ecol. Modell 135: 147– 186.
- Guisan, A. and Thuiller, W. 2005. Predicting species distribution: offering more than simple habitat models.Ecol. Lett. 8: 993–1009.
- Guisan, A. et al. 2006. Making better biogeographical predictions of species' distributions. J. Appl. Ecol. 43: 386–392.
- Hernandez, P. A. et al. 2006. The effect of sample size and species characteristics on performance of different species distribution modeling methods. – Ecography 29: 773– 785.
- Hijmans, R. J. et al. 2005. Very high resolution interpolated climate surfaces for global land areas. Int. J. Climatol. 25: 1965–1978.
- Hirzel, A. H. et al. 2002. Ecological-niche factor analysis: how to compute habitat-suitability maps without absence data.
 Ecology 83: 2027–2036.
- Hortal, J. et al. 2007. Limitations of biodiversity databases: case study on seed-plant diversity in Tenerife, Canary Islands. – Conserv. Biol. 21: 853–863.
- Jetz, W. et al. 2004. The coincidence of rarity and richness and the potential signature of history in centres of endemism. – Ecol. Lett. 7: 1180–1191.
- Jiménez-Valverde, A. and Lobo, J. M. 2007. Threshold criteria for conversion of probability of species presence to either-or presence-absence. – Acta Oecol. 31: 361–369.
- Kadmon, R. et al. 2003. A systematic analysis of factors affecting the performance of climatic envelop models.Ecol. Appl. 13: 853–867.
- Kadmon, R. et al. 2004. Effect of roadside bias on the accuracy of predictive maps produced by bioclimatic models. – Ecol. Appl. 14: 401–413.
- Leigh Jr, E.G. 2004. How wet are the wet tropics? In: Losos, E. C. and Leigh Jr, E. G. (eds), Tropical forest diversity and dynamism. Univ. of Chicago Press, pp. 43–55.
- Liu, C. et al. 2005. Selecting thresholds of occurrence in the prediction of species distributions. – Ecography 28: 385– 393.
- Manel, S. et al. 2001. Evaluating presence-absence models in ecology: the need to account for prevalence. – J. Appl. Ecol. 38: 921–931.
- McPherson, J. M. and Jetz, W. 2007. Effects of species' ecology on the accuracy of distribution models. Ecography 30: 135–151.
- McPherson, J. M. et al. 2004. The effects of species' range sizes on the accuracy of distribution models: ecological phenomenon or statistical artefact? J. Appl. Ecol. 41: 811–823.

Download the appendix as file E5041 from www.oikos.ekol.lu.se/appendix>.

- Olden, J. D. et al. 2002. Predictive models of fish species distributions: a note on proper validation and chance predictions. Trans. Am. Fish. Soc. 131: 329–336.
- Parnell, J. A. N. et al. 2003. Plant collecting spread and densities: their potential impact on biogeographical studies in Thailand. – J. Biogeogr. 30: 193–209.
- Pearce, J. and Ferrier, S. 2000. Evaluating the predictive performance of habitat models developed using logistic regression. Ecol. Modell. 133: 225–245.
- Pearce, J. L. and Boyce, M. S. 2006. Modelling distribution and abundance with presence-only data. J. Appl. Ecol. 43: 405–412.
- Pearson, R. G. et al. 2007. Predicting species distributions from small numbers of occurrence records: a test case using cryptic geckos in Madagascar. – J. Biogeogr. 34: 102–117.
- Peterson, A. T. 2006. Uses and requirements of ecological niche models and related distributional models. Biodiv. Inform. 3: 59–72.
- Phillips, S. J. et al. 2006. Maximum entropy modeling of species geographic distributions. – Ecol. Modell. 190: 231–259.
- Phillips, S. J. et al. 2004. A maximum entropy approach to species distribution modeling. – Twenty-First International Conference on Machine Learning 2004, p. 9.
- Reddy, S. and Davalos, L. M. 2003. Geographical sampling bias and its implications for conservation priorities in Africa. – J. Biogeogr. 30: 1719–1727.
- Romo, H. et al. 2006. Identifying recorder-induced geographic bias in an Iberian butterfly database. – Ecography 29: 873–885.
- Soberón, J. and Peterson, A. T. 2005. Interpretation of models of fundamental ecological niches and species' distributional areas. – Biodiv. Inform. 2: 1–10.
- Stockwell, D. and Peters, D. 1999. The GARP modelling system: problems and solutions to automated spatial prediction. Int. J. Geogr. Inf. Sci. 13: 143–158.
- Stockwell, D. R. B. and Peterson, A. T. 2002. Effects of sample size on accuracy of species distribution models. – Ecol. Modell. 148: 1–13.
- Swets, J. A. 1988. Measuring the accuracy of diagnostic systems. Science 240: 1285–1293.
- Swets, J. A. et al. 2000. Better decisions through science.
 Sci. Am. 283: 82–87.
- Thuiller, W. et al. 2005. Niche-based modelling as a tool for predicting the risk of alien plant invasions at a global scale.
 Global Change Biol. 11: 2234–2250.
- Walsh, R. P. D. 1996. Climate. In: Richards, P. W. (ed.), The tropical rain forest. Cambridge Univ. Press, pp. 159– 205.