# Guidelines for the prediction of species interactions through binary classification

Timothée Poisot 1,2

<sup>1</sup> Université de Montréal <sup>2</sup> Québec Centre for Biodiversity Sciences

## **Correspondance to:**

 $Timoth\'{e}e \ Poisot - \ timothee.poisot@umontreal.ca$ 

This work is released by its authors under a CC-BY 4.0 license

Last revision: June 10, 2022

- 1. The prediction of species interactions is gaining momentum as a way to circumvent limitations in data volume. Yet, ecological networks are challenging to predict because they are typically small and sparse. Dealing with extreme class imbalance is a challenge for most binary classifiers, and there are currently no guidelines as to how predictive models can be trained for this specific problem.
- 2. Using simple mathematical arguments and numerical experiments in which a variety of classifiers (for supervised learning) are trained on simulated networks, we develop a series of guidelines related to the choice of measures to use for model selection, and the degree of unbiasing to apply to the training dataset.
- 3. Neither classifier accuracy nor the ROC-AUC are informative measures for the performance of interaction prediction. PR-AUC is a fairer assessment of performance. In some cases, even standard measures can lead to selecting a more biased classifier because the effect of connectance is strong. The amount of correction to apply to the training dataset depends on network connectance, on the measure to be optimized, and only weakly on the classifier.
- 4. These results reveal that training machines to predict networks is a challenging task, and that in virtually all cases, the composition of the training set needs to be fine-tuned before performing the actual training. We discuss these consequences in the context of the low volume of data.

- Ecological networks are a backbone for key ecological and evolutionary processes; yet enumerating all of
- the interactions between S species is a daunting task, as it scales with  $S^2$ , i.e. the squared species richness
- Martinez, 1992). Recent contributions to the field of ecological network prediction (Becker et al., 2022;
- 4 Pichler et al., 2020; Strydom et al., 2021) highlight that although interactions can be predicted by adding
- <sup>5</sup> ecologically relevant information (in the form of, e.g. traits), we do not have robust guidelines as to how
- 6 the predictive ability of these models should be evaluated, nor about how the models should be trained.
- 7 Here, by relying on simple derivations and a series of simulations, we formulate a number of such
- 8 guidelines, specifically for the case of binary classifiers derived from thresholded values. Specifically, we
- 9 conduct an investigation of the models in terms of their skill (ability to make the right prediction), bias
- 10 (trends towards systematically over-predicting one class), class imbalance (the relative number of cases
- 11 representing interactions), and show how these effects interact. We conclude on the fact that models with
- the best interaction-scale predictive score do not necessarily result in the most accurate representation of
- 13 the network.
- 14 The prediction of ecological interactions shares conceptual and methodological issues with two fields in
- biology: species distribution models (SDMs), and genomics. SDMs suffers from issues affecting
- 16 interactions prediction, namely low prevalence (due to sparsity of observations/interactions) and data
- aggregation (due to bias in sampling some locations/species). In previous work, Allouche et al. (2006)
- suggested that  $\kappa$  was a better test of model performance than the True Skill Statistic (TSS; which we refer
- to as Youden's informedness); these conclusions were later criticized by Somodi et al. (2017), who
- emphasized that informedness' is affected both by prevalence and bias. Although this work offers
- 21 recommendations about the comparison of models, it doesn't establishes baselines or good practices for
- training on imbalanced ecological data, or ways to remedy the imbalance. Steen et al. (2021) show that,
- when applying spatial thinning (a process that has no analogue in networks), the best approach to train
- ML-based SDMs varies according to the balancing of the dataset, and the evaluation measures used. This
- suggests that there is no single "recipe" that is guaranteed to give the best model. By contrast to networks,
- SDMs have the advantage of being able to both thin datasets to remove some of the sampling bias (e.g.
- 27 Inman et al., 2021), but also to create pseudo-absences to inflate the number of supposed negatives in the
- <sup>28</sup> dataset (e.g. Iturbide et al., 2015).
- 29 An immense body of research on machine learning application to life sciences is focused on genomics
- 30 (which has very specific challenges, see a recent discussion by Whalen et al., 2021); this sub-field has

- 31 generated recommendations that do not necessarily match the current best-practices for SDMs, and
- therefore hint at the importance of domain-specific guidelines. Chicco & Jurman (2020) suggest using
- Matthews correlation coefficient (MCC) over  $F_1$ , as a protection against over-inflation of predicted results;
- Delgado & Tibau (2019) advocate against the use of Cohen's  $\kappa$ , again in favor of MCC, as the relative
- nature of  $\kappa$  means that a worse classifier can be picked over a better one; similarly, Boughorbel et al.
- 36 (2017) recommend MCC over other measures of performance for imbalanced data, as it has more
- desirable statistical properties. More recently, Chicco et al. (2021) temper the apparent supremacy of the
- 38 MCC, by suggesting it should be replaced by Youden's informedness (also known as J, bookmaker's
- 39 accuracy, and the True-Skill Statistic) when the imbalance in the dataset may not be representative of the
- 40 actual imbalance.
- Species interaction networks are often under-sampled (Jordano, 2016a, 2016b), and this under-sampling is
- 42 structured taxonomically (Beauchesne et al., 2016), structurally (de Aguiar et al., 2019) and spatially
- (Poisot, Bergeron, et al., 2021; Wood et al., 2015). As a consequence, networks suffer from data
- deficiencies both within and between datasets. This implies that the comparison of classifiers across
- space, when undersampling varies locally (see e.g. McLeod et al., 2021) is non-trivial. Furthermore, the
- baseline value of classifiers performance measures under various conditions of skill, bias, and prevalence,
- 47 has to be identified to allow researchers to evaluate whether their interaction prediction model is indeed
- 48 learning. Taken together, these considerations highlight three specific issues for ecological networks.
- 49 First, what values of performance measures are indicative of a classifier with no skill? This is particularly
- 50 important as it can evaluate whether low prevalence can lull us into a false sense of predictive accuracy.
- 51 Second, independently of the question of model evaluation, is low prevalence an issue for *training* or
- testing, and can we remedy it? Finally, because the low amount of data on interaction makes a lot of
- imbalance correction methods (see e.g. Branco et al., 2015) hard to apply, which indicators can be
- optimized by sacrificing least amount of positive interaction data?
- 55 It may sound counter-intuitive to care so deeply about how good a classifier with no-skill is, as by
- definition, is has no skill. The necessity of this exercise has its roots in the paradox of accuracy: when the
- desired class ("two species interact") is rare, a model that gets less ecologically performant by only
- predicting the opposite class ("these two species do not interact") sees its accuracy increase; because most
- of the guesses have "these two species do not interact" as a correct answer, a model that never predicts
- 60 interactions would be right an overwhelming majority of the time; it would also be utterly useless. Herein

- 61 lies the core challenge of predicting species interactions: the extreme imbalance between classes makes
- the training of predictive models difficult, and their validation even more so as we do not reliably know
- 63 which negatives are true. The connectance (the proportion of realized interactions, usually the number of
- 64 interactions divided by the number of species pairs) of empirical networks is usually well under 20%, with
- larger networks having a lower connectance (MacDonald et al., 2020), and therefore being increasingly
- 66 difficult to predict.

# A primer on binary classifier evaluation

- Binary classifiers, which it to say, machine learning algorithms whose answer is a categorical value, are
- 69 usually assessed by measuring properties of their confusion matrix, i.e. the contingency table reporting
- 70 true/false positive/negative hits. A confusion matrix is laid out as

$$\begin{pmatrix} tp & fp \\ fn & tn \end{pmatrix}$$
.

- In this matrix, tp is the number of times the model predicts an interaction that exists in the network (true
- 72 positive), fp is the number of times the model predicts an interaction that does not exist in the network
- <sub>73</sub> (false positive), fn is the number of times the model fails to predict an interaction that actually exists in the
- network (false negatives), and to is the number of times the model correctly predicts that an interaction
- does not exist (true negatives). From these values, we can derive a number of measures of model
- performance (see Strydom et al., 2021 for a review of their interpretation in the context of networks). At a
- coarse scale, a classifier is accurate when the trace of the matrix divided by the sum of the matrix is close
- to 1, with other measures informing us on how the predictions fail.
- 79 There is an immense diversity of measures to evaluate the performance of classification tasks (Ferri et al.,
- 2009). Here we will focus on five of them with high relevance for imbalanced learning (He & Ma, 2013).
- The choice of metrics with relevance to class-imbalanced problems is fundamental, because as Japkowicz
- 82 (2013) unambiguously concluded, "relatively robust procedures used for unskewed data can break down
- miserably when the data is skewed." Following Japkowicz (2013), we focus on two ranking metrics (the
- areas under the Receiver Operating Characteristic and Precision Recall curves), and three threshold

- metrics ( $\kappa$ , informedness, and MCC; we will briefly discuss  $F_1$  but show early on that it has undesirable
- 86 properties).
- The  $\kappa$  measure (Landis & Koch, 1977) establishes the extent to which two observers (the network and the
- prediction) agree, and is measured as

$$2\frac{tp\times tn-fn\times fp}{(tp+fp)\times (fp+tn)+(tn+fp)\times (tn+fn)}\,.$$

- 189 Informedness (Youden, 1950) (also known as bookmaker informedness or the True Skill Statistic) is
- TPR + TNR 1, where TPR = tp/(tp + fn) and TNR = tn/(tn + fp). Informedness can be used to find
- 91 the optimal cutpoint in thresholding analyses (Schisterman et al., 2005); indeed, the maximal
- 92 informedness corresponds to the point on the ROC curve that is closest to the perfect classifier point. The
- 93 formula for informedness is

$$\frac{tp}{tp+fn} + \frac{tn}{tn+fp} - 1.$$

94 The MCC is defined as

$$\frac{tp \times tn - fn \times fp}{\sqrt{(tp + fp) \times (tp + fn) \times (tn + fp) \times (tn + fn)}}.$$

- Finally,  $F_1$  is the harmonic mean of precision (the chance that interaction was correctly detected as such)
- and sensitivity (the ability to correctly classify interactions), and is defined as

$$2\frac{tp}{2\times tp + fp + fn}.$$

- 97 A lot of binary classifiers are built by using a regressor (whose task is to guess the value of the interaction,
- 98 and can therefore return a value considered to be a pseudo-probability); in this case, the optimal value
- below which predictions are assumed to be negative (i.e. the interaction does not exist) can be determined
- by picking a threshold maximizing some value on the ROC or the PR curve. The area under these curves
- (ROC-AUC and PR-AUC henceforth) give ideas on the overall goodness of the classifier, and the ideal
- threshold is the point on these curves that minimizes the tradeoff represented in these curves. Saito &

Rehmsmeier (2015) established that the ROC-AUC is biased towards over-estimating performance for imbalanced data; on the contrary, the PR-AUC is able to identify classifiers that are less able to detect 104 positive interactions correctly, with the additional advantage of having a baseline value equal to 105 prevalence. Therefore, it is important to assess whether these two measures return different results when applied to ecological network prediction. The ROC curve is defined by the false positive rate on the x axis, 107 and the true positive rate on the y axis, and the PR curve is defined by the true positive rate on the x axis, 108 and the positive predictive value on the y axis. By comparison with the previous paragraph, it is obvious 109 that  $F_1$  and MCC have ties to the PR curve (being close to the expected PR-AUC), and that informedness 110 has ties to the ROC curve (whereby the threshold maximizing informedness is also the point of maximal 111 inflection on the ROC curve). One important difference between ROC and PR is that the later does not 112 prominently account for the size of the true negative compartments: in short, it is more sensitive to the correct positive predictions. In a context of strong imbalance, PR-AUC is therefore a more stringent test of 114 model performance. 115

#### Baseline values for the threshold metrics

In this section, we will assume a network of connectance  $\rho$ , *i.e.* having  $\rho S^2$  interactions (where S is the species richness), and  $(1-\rho)S^2$  non-interactions. Therefore, the vector describing the *true* state of the network (assumed to be an unweighted, directed network) is a column vector  $\mathbf{o}^T = [\rho, (1-\rho)]$  (we can safely drop the  $S^2$  terms, as we will work on the confusion matrix, which ends up expressing *relative* values). We will apply skill and bias to this matrix, and measure how a selection of performance metrics respond to changes in these values, in order to assess their suitability for model evaluation.

#### Confusion matrix with skill and bias

In order to write the values of the confusion matrix for a hypothetical classifier, we need to define two characteristics: its skill, and its bias. Skill, here, refers to the propensity of the classifier to get the correct answer (*i.e.* to assign interactions where they are, and to not assign them where they are not). A no-skill classifier guesses at random, *i.e.* it will guess interactions with a probability  $\rho$ . The predictions of a no-skill classifier can be expressed as a row vector  $\mathbf{p}^T = [\rho, (1 - \rho)]$ . The confusion matrix  $\mathbf{M}$  for a no-skill classifier is given by the element-wise (Hadamard, outer) product of these vectors  $\mathbf{o} \odot \mathbf{p}$ , *i.e.* 

$$\mathbf{M} = \begin{pmatrix} \rho^2 & \rho(1-\rho) \\ (1-\rho)\rho & (1-\rho)^2 \end{pmatrix}.$$

In order to regulate the skill of this classifier, we can define a skill matrix **S** with diagonal elements equal to s, and off-diagonal elements equal to (1-s), which allows to regulate how many predictions are wrong, under the assumption that the bias is the same (*i.e.* the classifier is as likely to make a false positive or a false negative). The skill-adjusted confusion matrix is  $\mathbf{M} \odot \mathbf{S}$ , *i.e.* 

$$\begin{pmatrix} \rho^2 & \rho(1-\rho) \\ (1-\rho)\rho & (1-\rho)^2 \end{pmatrix} \odot \begin{pmatrix} s & (1-s) \\ (1-s) & s \end{pmatrix}.$$

When s = 0,  $Tr(\mathbf{M}) = 0$  (the classifier is *always* wrong), when s = 0.5, the classifier is no-skill and guesses at random, and when s = 1, the classifier is perfect.

The second element we can adjust in this hypothetical classifier is its bias, specifically its tendency to over-predict interactions. Like above, we can do so by defining a bias matrix  $\mathbf{B}$ , where interactions are over-predicted with probability b, and express the final classifier confusion matrix as  $\mathbf{M} \odot \mathbf{S} \odot \mathbf{B}$ , *i.e.* 

$$\begin{pmatrix} \rho^2 & \rho(1-\rho) \\ (1-\rho)\rho & (1-\rho)^2 \end{pmatrix} \odot \begin{pmatrix} s & (1-s) \\ (1-s) & s \end{pmatrix} \odot \begin{pmatrix} b & b \\ (1-b) & (1-b) \end{pmatrix}.$$

39 The final expression for the confusion matrix in which we can regulate the skill and the bias is

$$\mathbf{C} = \begin{pmatrix} s \times b \times \rho^2 & (1-s) \times b \times \rho(1-\rho) \\ (1-s) \times (1-b) \times (1-\rho)\rho & s \times (1-b) \times (1-\rho)^2 \end{pmatrix}.$$

In all further simulations, the confusion matrix **C** is transformed so that it sums to unity, *i.e.* the entries are the *proportions* of guesses.

#### What are the baseline values of performance measures?

In this section, we will change the values of b, s, and  $\rho$ , and report how the main measures discussed in 143 the introduction (MCC,  $F_1$ ,  $\kappa$ , and informedness) respond. Before we do so, it is important to explain why we will not focus on accuracy too much. Accuracy is the number of correct predictions  $(Tr(\mathbf{C}))$  divided by the sum of the confusion matrix. For a no-skill, no-bias classifier, accuracy is equal to  $\rho^2 + (1 - \rho)^2$ ; for 146  $\rho = 0.05$ , this is  $\approx 0.90$ , and for  $\rho = 0.01$ , this is equal to  $\approx 0.98$ . In other words, the values of accuracy are 147 high enough to be uninformative (for  $\rho$  small,  $\rho^2 \ll (1-\rho)^2$ ). More concerning is the fact that introducing bias changes the response of accuracy in unexpected ways. Assuming a no-skill classifier, the numerator 149 of accuracy becomes  $b\rho^2 + (1-b)(1-\rho)^2$ , which increases when b is low, which specifically means that at 150 equal skill, a classifier that under-predicts interactions will have higher accuracy than an un-biased 151 classifier (because the value of accuracy is dominated by the size of tn, which will increase). These issues 152 are absent from balanced accuracy, but should nevertheless lead us to not report accuracy as the primary 153 measure of network prediction success; moving forward, we will focus on other measures. In order to examine how MCC,  $F_1$ ,  $\kappa$ , and informedness change w.r.t. the imbalance, skill, and bias, we 155 performed a grid exploration of the values of logit(s) and logit(b) linearly from -10 to 10; logit(x) = -10156 means that x is essentially 0, and logit(x) = 10 means it is essentially 1 – this choice was motivated by the 157 fact that most responses are non-linear with regards to bias and skill. The values or  $\rho$  were taken linearly 158 in [0, 0.5], which is within the range of connectance for species interaction networks. Note that at this 159 point, there is no network model to speak of; the confusion matrix we discuss can be obtained for any classification task. Based on the previous discussion, the desirable properties for a measure of classifier 161 success should be: an increase with classifier skill, especially at low bias; a hump-shaped response to bias, 162 especially at high skill, and ideally centered around logit(b) = 0; an increase with prevalence up until 163 equiprevalence is reached.

#### [Figure 1 about here.]

165

In fig. 1, we show that none of the four measures satisfy all the considerations at once:  $F_1$  increases with skill, and increases monotonously with bias; this is because  $F_1$  does not account for true negatives, and the increase in positive detection masks the over-prediction of interactions. Informedness varies with skill, reaching 0 for a no-skill classifier, but is entirely unsensitive to bias. Both MCC and  $\kappa$  have the same

behavior, whereby they increase with skill.  $\kappa$  peaks at increasing values of bias for increasing skill, *i.e.* is likely to lead to the selection of a classifier that over-predicts interactions. By contract, MCC peaks at the same value, regardless of skill, but this value is not logit(b) = 0: unless at very high classifier skill, MCC risks leading to a model that over-predicts interactions. In fig. 2, we show that all measures except  $F_1$  give a value of 0 for a no-skill classifier, and are forced towars their correct maximal value when skill changes (*i.e.* a more connected networks will have higher values for a skilled classifierd, and lower values for a classifier making mostly mistakes).

#### [Figure 2 about here.]

These two analyses point to the following recommendations: MCC is indeed more appropriate than  $\kappa$ , as although sensitive to bias, it is sensitive in a consistent way. Informedness is appropriate at discriminating 179 between different skills, but confounded by bias. As both of these measures bring valuable information on 180 the model behavior, we will retain them for future analyses.  $F_1$  is increasing with bias, and should not be prioritized to evalue the performance of the model. The discussion of sensitivity to bias should come with 182 a domain-specific caveat: although it is likely that interactions documented in ecological networks are 183 correct, a lot of non-interactions are simply unobserved; as predictive models are used for data-inflation (i.e. the prediction of new interactions), it is not necessarily a bad thing in practice to select models that 185 predict more interactions than the original dataset, because the original dataset misses some interactions. 186 Furthermore, the weight of positive interactions could be adjusted if some information about the extent of 187 undersampling exists (e.g. Branco et al., 2015). In a recent large-scale imputation of interactions in the 188 mammal-virus networks, Poisot, Ouellet, et al. (2021) for example estimated that 93% of interactions are 189 yet to be documented. 190

# 91 Numerical experiments on training strategy

177

In the following section, we will generate random bipartite networks, and train four binary classifiers (as well as an ensemble model using the sum of ranged outputs from the component models) on 50% of the interaction data. In practice, testing usually uses 70% of the total data; for ecological networks, where interactions are sparse *and* the number of species is low, this may not be the best solution, as the testing set becomes constrained not by the *proportion* of interactions, but by their *number*. Preliminary

experiments using different splits revealed no qualitative change in the results. Networks are generated by 197 picking a random infectiousness trait  $v_i$  for 100 species (from a beta distribution  $B(\alpha = 6, \beta = 8)$ 198 distribution), and a resistance trait  $h_i$  for 100 species (from  $B(\alpha = 2, \beta = 8)$  distribution). There is an 199 interaction between i and j when  $v_i - \xi/2 \le h_j \le v_i + \xi/2$ , where  $\xi$  is a constant regulating the 200 connectance of the network (visual exploration of the parameters show that there is an almost 1:1 201 relationship between  $\xi$  and connectance), and varies uniformly in [0.05, 0.35]. This model gives fully 202 interval networks that are close analogues to the bacteria-phage model of Weitz et al. (2005), with both a 203 modular structure and a non-uniform degree distribution. This dataset is easy for almost any algorithm to 204 learn: when trained with features  $[v_i, h_j, abs(v_i, h_j)]^T$  to predict the interactions between i and j, all four 205 models presented below were able to reach almost perfect predictions all the time (data not presented 206 here) - this is in part because the rule (there is maximum value of the distance between traits for which there is an interaction) is fixed for all interactions, and any method able to learn non-linear relationships 208 should infer it without issues. In order to make the problem more difficult to solve, we use  $[v_i, h_j]$  as a 209 feature vector (i.e. the traits on which the models are trained), and therefore the models will have to uncover that the rule for interaction is  $abs(v_i, h_i) \le \xi$ . The models therefore all have the following form, 211 where  $i_{i,j}$  is an interaction from species i to species j: 212

$$\begin{bmatrix} i_{1,1} \\ i_{1,2} \\ \vdots \\ i_{m,n-1} \\ i_{m,n} \end{bmatrix} \propto \begin{bmatrix} v_1 & h_1 \\ v_1 & h_2 \\ \vdots & \vdots \\ v_m & h_{n-1} \\ v_m & h_n \end{bmatrix}$$

The training sample is composed of 50% of the  $10^4$  possible entries in the network, *i.e.* n = 5000. Out of these interactions, we pick a proportion  $\nu$  (the training set balance) to be positive, so that the training set has  $\nu n$  interactions, and  $(1 - \nu)n$  non-interactions. We vary  $\nu$  uniformly in ]0, 1[. This allows to evaluate how the measures of binary classification performance respond to artificially rebalanced dataset for a given network connectance. The rest of the dataset is used as a testing set, on which all further measures are calculated. Note that although the training set is balanced arbitrarily, the testing set is assembled so that it has the exact connectance of the entire network; this ensures that the model is evaluated under the class imbalance where the predictions will be made, which represents a more meaningful evaluation.

Note also that although the simulated networks are bipartite, the algorithms have no "knowledge" of the network structure, and simply look at pairs of species; therefore, the approach outlined here would also 222 work for unipartite networks. 223 The dataset used for numerical experiments is composed of a grid of 35 values of connectance (from 0.011 224 to 0.5) and 35 values of  $\nu$  (from 0.02 to 0.98); for each pair of values, 500 networks are generated and 225 predicted. For each network, we train four machines: a trait-based k-NN (e.g. Desjardins-Proulx et al., 226 2017), a regression tree, a regression random forest, and a boosted regression tree. Following results from Pichler et al. (2020), linear models have not been considered (in any way, the relationship in the simulated 228 networks is non-linear). The point of these numerical experiments is *not* to recommend the best model 229 (this is likely problem-specific), but to highlight a series of recommendations that would work for supervised learning tasks. All models were taken from the MLJ. jl package (Blaom et al., 2020; Blaom & 231 Vollmer, 2020) in Julia 1.7 (Bezanson et al., 2017). All machines use the default parameterization; this is 232 an obvious deviation from best practices, as the hyperparameters of any machine require training before 233 its application on a real dataset. As we use 612500 such datasets, this would require over 2 millions unique 234 instances of tweaking the hyperparameters, which is prohibitive from a computing time point of view. An 235 important thing to keep in mind is that the problem we simulate has been designed to be simple to solve: 236 we expect all machines with sensible default parameters to fare well — the results presented in the later 237 sections show that this assumption is warranted, and we further checked that the models do not overfit by 238 ensuring that there is never more than 5% of difference between the accuracy on the training and testing 239 sets. All machines return a quantitative prediction, usually (but not necessarily) in [0, 1], which is proportional (but not necessarily linearly) to the probability of an interaction between i and j. The 241 ROC-AUC and PR-AUC (and therefore the thresholds) can be measured by integrating over the domain of 242 the values return by each machine, but in order to make the average-based ensemble model more meaningful, all predictions are expressed in [0, 1]. 244 In order to pick the best confusion matrix for a given trained machine, we performed a thresholding 245 approach using 500 steps on predictions from the testing set, and picking the threshold that maximized 246 Youden's informedness. During the thresholding step, we measured the area under the receiver operating 247 characteristic (ROC-AUC) and precision-recall (PR-AUC) curves, as measures of overall performance over the range of returned values. We report the ROC-AUC and PR-AUC, as well as a suite of other measures as introduced in the next section, for the best threshold. The ensemble model was generated by summing the

predictions of all component models on the testing set (ranged in [0, 1]), then put through the same thresholding process. The complete code to run the simulations is available at 10.17605/0SF.IO/JKEWD.

After the simulations were completed, we removed all runs (*i.e.* triples of model,  $\xi$ , and  $\nu$ ) for which at least one of the following conditions was met: the accuracy was 0, the true positive or true negative rates were 0, the connectance was larger than 0.25. This removes both the obviously failed model runs, and the networks that are more densely connected compared to the connectance of empirical food webs (and are therefore less difficult to predict, being less imbalanced; preliminary analyses of data with a connectance larger than 0.3 revealed that all machines reached consistently high performance).

#### 259 Effect of training set balance on performance

275

In fig. 3, we present the response of two thresholding measures (PR-AUC and ROC-AUC) and two ranking 260 measures (Informedness and MCC) to a grid of 35 values of training set balance, and 35 values of 261 connectance, for the four component models as well as the ensemble. ROC-AUC is always high, and does 262 not vary with training set balance. On the other hand, PR-AUC shows very strong responses, increasing with training set balance. It is notable here that two classifiers that seemed to be performing well (Decision 264 Tree and Random Forest) based on their MCC are not able to reach a high PR-AUC even at higher 265 connectances. All models reached a higher performance on more connected networks, and using more 266 balanced training sets. In all cases, informedness was extremely high, which is an expected consequence 267 of the fact that this is the value we optimized to determine the cutoff. MCC increased with training set 268 balance, although this increase became less steep with increasing connectance. Three of the models (kNN, 269 decision tree, and random forest) only increased their PR-AUC sharply when the training set was heavily 270 imbalanced towards more interactions. Interestingly, the ensemble almost always outclassed its 271 component models. For larger connectances (less difficult networks to predict, as they are more balanced), 272 MCC and informedness stared decreasing when the training set bias got too close to one, suggesting that a 273 training set balance of 0.5 may often be appropriate if these measures are the one to optimize. 274

#### [Figure 3 about here.]

Based on the results presented in fig. 3, it seems that informedness and ROC-AUC are not necessarily able to discriminate between good and bad classifiers (although this result may be an artifact for informedness,

as it has been optimized when thresholding). On the other hand, MCC and PR-AUC show a strong response to training set balance, and may therefore be more useful at model comparison.

#### 280 Required amount of positives to get the best performance

289

299

The previous results revealed that the measure of classification performance responds both to the bias in
the training set *and* to the connectance of the network; from a practical point of view, assembling a
training set requires one to withhold positive information, which in ecological networks are very scarce
(and typically more valuable than negatives, on which there is a doubt). For this reason, across all values
of connectance, we measured the training set balance that maximized a series of performance measures.
When this value is high, the training set needs to skew more positive in order to get a performant model;
when this value is about 0.5, the training set needs to be artificially balanced to optimize the model
performance. These results are presented in fig. 4.

#### [Figure 4 about here.]

The more "optimistic" measures (ROC-AUC and informedness) required a biasing of the dataset from 290 about 0.4 to 0.75 to be maximized, with the amount of bias required decreasing only slightly with the 291 connectance of the original network. MCC and PR-AUC required values of training set balance from 0.75 292 to almost 1 to be optimized, which is in line with the results of the previous section, i.e. they are more stringent tests of model performance. These results suggest that learning from a dataset with very low 294 connectance can be a different task than for more connected networks: it becomes increasingly important 295 to capture the mechanisms that make an interaction exist, and therefore having a slightly more biased training dataset might be beneficial. As connectance increases, the need for biased training sets is less 297 prominent, as learning the rules for which interactions do not exist starts gaining importance. 298

#### [Figure 5 about here.]

When trained at their optimal training set balance, connectance still had a significant impact on the
performance of some machines (fig. 5). Notably, Decision Tree, and k-NN, as well as Random forest to a
lower extent, had low values of PR-AUC. In all cases, the Boosted Regression Tree was reaching very good
predictions (especially for connectances larger than 0.1), and the ensemble was almost always scoring

perfectly. This suggests that all the models are biased in different ways, and that the averaging in the
ensemble is able to correct these biases. We do not expect this last result to have any generality, and
provide a discussion of a recent example in which the ensemble was performing worse than its
components models.

### Do better classification accuracy result in more realistic networks?

In this last section, we generate a network using the same model as before, with  $S_1$ ,  $S_2 = 50$ , 80 species, a connectance of  $\approx 0.16$  ( $\xi = 0.19$ ), and a training set balance of 0.5, as fig. 4 suggests this is the optimal training set balance for this range of connectance. The prediction made on the complete dataset is presented in fig. 6.

#### [Figure 6 about here.]

313

The trained models were then thresholded (again by optimising informedness), and their predictions transformed back into networks for analysis; specifically, we measured the connectance, nestedness ( $\eta$ ; 315 Bastolla et al., 2009), modularity (Q; Barber, 2007), asymmetry (A; Delmas et al., 2018), and Jaccard 316 network dissimilarity (Canard et al., 2014). This process was repeated 250 times, and the results are 317 presented in tbl. 1. The k-NN model is an interesting instance here: it produces the network that looks the 318 most like the original dataset, despite having the lowest PR-AUC, suggesting it hits high recall at the cost 319 of low precision. The ensemble was able to reach a very high PR-AUC (and a very high ROC-AUC), which 320 translated into more accurate reconstructions of the structure of the network (with the exception of 321 modulairty, which is underestimated by 0.03). This result bears elaborating. Measures of model 322 performance capture how much of the interactions and non-interactions are correctly identified. As long 323 as these predictions are not perfect, some interactions will be predicted at the "wrong" position in the 324 network; these measures cannot describe the structural effect of these mistakes. On the other hand, 325 measures of network structure can have the same value with interactions that fall at drastically different 326 positions; this is in part because a lot of these measures covary with connectance, and in part because as 327 long as these values are not 0 or their respective maximum, there is a large number of network 328 configurations that can have the same value. That ROC-AUC is consistently larger than PR-AUC may be a 329 case of this measure masking models that are not, individually, strong predictors (Jeni et al., 2013). In this

specif example, the combination of individually "adequate" models resulted in an extremely strong
ensemble, suggesting that the correct prediction of interactions (as measured by MCC, Inf., ROC-AUC,
and PR-AUC) and network properties is indeed a feasible task under appropriately hyper-parameterized
models.

Table 1: Values of four performance metrics, and five network structure metrics, for 500 independent predictions similar to the ones presented in fig. 6. The values in **bold** indicate the best value for each column (including ties). Because the values have been rounded, values of 1.0 for the ROC-AUC column indicate an average  $\geq 0.99$ .

Model	MCC	Inf.	ROC-AUC	PR-AUC	Conn.	η	Q	A	Jaccard
Decision tree	0.59	0.94	0.97	0.04	0.17	0.64	0.37	0.42	0.1
BRT	0.46	0.91	0.97	0.36	0.2	0.78	0.29	0.41	0.19
Random Forest	0.72	0.98	0.99	0.1	0.16	0.61	0.38	0.42	0.06
k-NN	0.71	0.98	0.99	0.02	0.16	0.61	0.39	0.42	0.06
Ensemble	0.74	0.98	1.0	0.79	0.16	0.61	0.38	0.42	0.06
Data					0.16	0.56	0.41	0.42	0.0

# Guidelines for the assessment of network predictive models

We establish that due to the low prevalence of interactions, even poor classifiers applied to food web data 336 will reach a high accuracy; this is because the measure is dominated by the accidentally correct 337 predictions of negatives. On simulated confusion matrices with ranges of imbalance that are credible for 338 ecological networks, MCC had the most desirable behavior, and informedness is a linear measure of 339 classifier skill. By performing simulations with four models and an ensemble, we show that informedness 340 and ROC-AUC are consistently high on network data, whereas MCC and PR-AUC are more accurate 341 measures of the effective performance of the classifier. Finally, by measuring the structure of predicted 342 networks, we highlight an interesting paradox: the models with the best performance measures are not necessarilly the models with the closest reconstructed network structure. We discuss these results in the 344 context of establishing guidelines for the prediction of ecological interactions. 345

It is noteworthy that the ensemble model was systematically better than the component models. We do

not expect that ensembles will always be better than single models. Networks with different structures

simple rule we used here. In a recent multi-model comparison involving supervised and unsupervised 349 learning, Becker et al. (2022) found that the ensemble was not the best model, and was specifically 350 under-performing compared to models using biological traits. This may be because the dataset of Becker et al. (2022) was known to be under-sampled, and so the network itself contained less information than 352 the network and species traits. There is no general conclusion to draw from either these results or ours, 353 besides reinforcing the need to be pragmatic about which models should be included in the ensemble, and 354 whether to use an ensemble at all. In a sense, the surprising performance of the ensemble model should 355 form the basis of the first broad recommendation: optimal training set balance and its interaction with 356 connectance and the specific binary classifier used is, in a sense, an hyperparameter that should be 357 assessed. The distribution of results in fig. 4 and fig. 5 show that there are variations around the trend, and 358 multiple models should probably be trained on their "optimal" training/testing set, as opposed to the same 359 ones. 360 The results presented here highlight an interesting paradox: although the k-NN model was ultimately able 361 to get a correct estimate of network structure (see tbl. 1 and fig. 6), it ultimately remains a poor classifier, 362 as evidenced by its low PR-AUC. This suggests that the goal of predicting interactions and predicting 363 networks may not always be solvable in the same way – of course a perfect classifier of interactions would 364 make a perfect network prediction; indeed, the best scoring predictor of interactions (the ensemble model) 365 had the best prediction of network structure. The tasks of predicting networks structure and of predicting 366 interactions within networks are essentially two different ones. For some applications (.e.g. comparison of network structure across gradients), one may care more about a robust estimate of the structure, at the 368 cost at putting some interactions at the wrong place. For other applications (e.g. identifying pairs of 369 interacting species), one may conversely care more about getting as many pairs right, even though the mistakes accumulate in the form of a slightly worse estimate of network structure. How these two 371 approaches can be reconciled is something to evaluate on a case-by-case basis, especially since there is no 372 guarantee that an esemble model will always be the most precise one. Despite this apparent tension at the 373 heart of the predictive exercise, we can use the results presented here to suggest a number of guidelines. 374 First, because we have more trust in reported interactions than in reported absences of interactions (which are overwhelmingly pseudo-absences), we can draw on previous literature to recommend informedness as a measure to decide on a threshold for binary classification (Chicco et al., 2021); this being said, because 377

than the one we simulated here may respond in different ways, especially if the rules are fuzzier than the

informedness is insensitive to bias (although it is a linear measure of skill), the overall model performance is better evaluated through the use of MCC (figs. 4, 5). Because  $F_1$  is monotonously sensitive to classifier 379 bias fig. 1 and network connectance fig. 2, MCC should be preferred as a measure of model evaluation and 380 comparison. When dealing with multiple models, we therefore suggest to find the optimal threshold using informedness, and to pick the best model using MCC (assuming one does not want to use an ensemble 382 model). 383 Second, accuracy alone should not be the main measure of model performance, but rather an expectation of how well the model should behave given the class balance in the set on which predictions are made; 385 this is because, as derived earlier, the expected accuracy for a no-skill no-bias classifier is  $\rho^2 + (1 - \rho)^2$ 386 (where  $\rho$  is the class balance), which will most often be large. This pitfall is notably illustrated in a recent food-web model (Caron et al., 2022) wherein the authors, using a training set of  $n = 10^4$  with only 100 388 positive interactions (representing 0.1% of the total interactions), reached a good accuracy. Reporting a 389 good accuracy is not informative, especially when accuracy isn't (i) compared to the baseline expected value under the given class balance, and (ii) interpreted in the context of a measure that is not sensitive to 391 the chance prediction of many negatives (like MCC). 392 Third, because the PR-AUC responds more to network connectance (fig. 5) and training set imbalance (fig. 4) than ROC-AUC, it should be used as a measure of model performance over the ROC-AUC. This is 394 not to say that ROC-AUC should be discarded (in fact, a low ROC-AUC is undoubtedly a sign of an issue 395 with the model), but that its interpretation should be guided by the PR-AUC value. Specifically, a high ROC-AUC is not informative, as it can be associated to a low PR-AUC (see e.g. Random Forest in tbl. 1) 397 This again echoes recommendations from other fields (Jeni et al., 2013; Saito & Rehmsmeier, 2015). We 398 therefore expect to see high ROC-AUC values, and then to pick the model that maximizes the PR-AUC 399 value. Taken together with the previous two guidelines, we strongly encourage to (i) ensure that accuracy 400 and ROC-AUC are high (in the case of accuracy, higher than expected under no-skill no-bias situation), 401 and (ii) to discuss the performance of the model in terms of the most discriminant measures, i.e. PR-AUC 402 and MCC. 403 Finally, network connectance (i.e. the empirical class imbalance) should inform the composition of the 404 training and testing set, because it is an ecologically relevant value. In the approach outlined here, we treat the class imbalance of the training set as an hyper-parameter, but test the model on a set that has the same class imbalance as the actual dataset. This is an important distinction, as it ensure that the prediction

environment matches the testing environment (as we cannot manipulate the connectance of the empirical dataset on which the predictions will be made), and so the values measured on the testing set (or validation 409 set if the data volume allows one to exists) can be directly compared to the values for the actual prediction. 410 A striking result from fig. 4 is that Informedness was almost always maximal at 50/50 balance (regardless of connectance), whereas MCC required more positives to be maximized when connectance increases, 412 matching the idea that it is a more stringent measure of performance. This has an important consequence 413 in ecological networks, for which the pool of positive cases (interactions) to draw from is typically small: 414 the most parsimonious measure (i.e. the one requiring to discard the least amount of interactions to train 415 the model) will give the best validation potential, and in this light is very likely informedness (maximizing 416 informedness is, in fact, the generally accepted default for imbalanced classification regardless of the 417 problem domain; Schisterman et al., 2005). This last result further strengthens the assumption that the amount of bias is an hyper-parameter that must be fine-tuned, as using the wrong bias can lead to models 419 with lower performance; for this reason, it makes sense to not train all models on the same 420 training/testing set, but rather to optimize the set composition for each of them. One key element for real-life data that can make the prediction exercise more tractable is that some 422 interactions can safely be assumed to be impossible; indeed, a lot of networks admit a stochastic block 423 model as a good approximation (e.g. Xie et al., 2017). In ecological networks, this can be due to spatial 424 constrains (Valdovinos, 2019), or to the long-standing knowledge that some links are "forbidden" due to 425 traits (Olesen et al., 2011) or abundances (Canard et al., 2014). The matching rules (Olito & Fox, 2015; 426 Strona & Veech, 2017) can be incorporated in the model either by adding compatibility traits, or by only training the model on pairs of species that are not likely to be forbidden links; having this information 428 would allow to assemble training/testing sets that have true negatives, and in this situation, it may be 429 possible to use the more usual 70/30 split. Besides forbidden links, a real-life case that may arise is multi-interaction or multi-layer networks (Pilosof et al., 2017). These can be studied using the same 431 general approach outlined here, either by assuming that pairs of species can interact in more than one way 432 (wherein one would train a model for each type of interaction, based on the relevant predictors), or by 433 assuming that pairs of species can only have one type of interaction (wherein this becomes a multi-label 434 classification problem). 435 Acknowledgements: We acknowledge that this study was conducted on land within the traditional

unceded territory of the Saint Lawrence Iroquoian, Anishinabewaki, Mohawk, Huron-Wendat, and

- Omàmiwininiwak nations. We thank Colin J. Carlson, Michael D. Catchen, Giulio Valentino Dalla Riva,
- and Tanya Strydom for inputs on earlier versions of this manuscript. This research was enabled in part by
- support provided by Calcul Québec (www.calculquebec.ca) through the Narval general purpose cluster.
- TP is supported by the Fondation Courtois, a NSERC Discovery Grant and Discovery Acceleration
- 442 Supplement, by funding to the Viral Emergence Research Initiative (VERENA) consortium including NSF
- 443 BII 2021909, and by a grant from the Institut de Valorisation des Données (IVADO).

#### References

- 445 Allouche, O., Tsoar, A., & Kadmon, R. (2006). Assessing the accuracy of species distribution models:
- Prevalence, kappa and the true skill statistic (TSS). *Journal of Applied Ecology*, 43(6), 1223–1232.
- https://doi.org/10.1111/j.1365-2664.2006.01214.x
- Barber, M. J. (2007). Modularity and community detection in bipartite networks. *Physical Review E*, 76(6),
- 449 066102. https://doi.org/10.1103/PhysRevE.76.066102
- Bastolla, U., Fortuna, M. A., Pascual-García, A., Ferrera, A., Luque, B., & Bascompte, J. (2009). The
- architecture of mutualistic networks minimizes competition and increases biodiversity. *Nature*,
- 452 458(7241), 1018–1020. https://doi.org/10.1038/nature07950
- Beauchesne, D., Desjardins-Proulx, Archambault, P., & Gravel, D. (2016). Thinking Outside the
- Box-predicting Biotic Interactions in Data-poor Environments. Vie Et Milieu-Life and enVironment,
- 455 *66*(3-4), 333–342.
- Becker, D. J., Albery, G. F., Sjodin, A. R., Poisot, T., Bergner, L. M., Chen, B., Cohen, L. E., Dallas, T. A.,
- Eskew, E. A., Fagre, A. C., Farrell, M. J., Guth, S., Han, B. A., Simmons, N. B., Stock, M., Teeling, E. C.,
- & Carlson, C. J. (2022). Optimising predictive models to prioritise viral discovery in zoonotic
- reservoirs. The Lancet Microbe. https://doi.org/10.1016/S2666-5247(21)00245-7
- 460 Bezanson, J., Edelman, A., Karpinski, S., & Shah, V. (2017). Julia: A Fresh Approach to Numerical
- 461 Computing. SIAM Review, 59(1), 65–98. https://doi.org/10.1137/141000671
- Blaom, A. D., Kiraly, F., Lienart, T., Simillides, Y., Arenas, D., & Vollmer, S. J. (2020). MLJ: A Julia package
- for composable machine learning. *Journal of Open Source Software*, 5(55), 2704.
- https://doi.org/10.21105/joss.02704

```
Blaom, A. D., & Vollmer, S. J. (2020). Flexible model composition in machine learning and its
       implementation in MLJ. http://arxiv.org/abs/2012.15505
466
    Boughorbel, S., Jarray, F., & El-Anbari, M. (2017). Optimal classifier for imbalanced data using Matthews
467
       Correlation Coefficient metric. PloS One, 12(6), e0177678.
468
       https://doi.org/10.1371/journal.pone.0177678
469
    Branco, P., Torgo, L., & Ribeiro, R. (2015). A Survey of Predictive Modelling under Imbalanced Distributions.
       http://arxiv.org/abs/1505.01658
471
    Canard, E. F., Mouquet, N., Mouillot, D., Stanko, M., Miklisova, D., & Gravel, D. (2014). Empirical
       evaluation of neutral interactions in host-parasite networks. The American Naturalist, 183(4), 468-479.
473
       https://doi.org/10.1086/675363
474
    Caron, D., Maiorano, L., Thuiller, W., & Pollock, L. J. (2022). Addressing the Eltonian shortfall with
       trait-based interaction models. Ecology Letters, 25(4), 889–899. https://doi.org/10.1111/ele.13966
476
    Chicco, D., & Jurman, G. (2020). The advantages of the Matthews correlation coefficient (MCC) over F1
       score and accuracy in binary classification evaluation. BMC Genomics, 21(1), 6.
478
       https://doi.org/10.1186/s12864-019-6413-7
479
    Chicco, D., Tötsch, N., & Jurman, G. (2021). The Matthews correlation coefficient (MCC) is more reliable
480
        than balanced accuracy, bookmaker informedness, and markedness in two-class confusion matrix
481
       evaluation. BioData Mining, 14, 13. https://doi.org/10.1186/s13040-021-00244-z
482
    de Aguiar, M. A. M., Newman, E. A., Pires, M. M., Yeakel, J. D., Boettiger, C., Burkle, L. A., Gravel, D.,
483
       Guimarães, P. R., O'Donnell, J. L., Poisot, T., Fortin, M.-J., & Hembry, D. H. (2019). Revealing biases in
484
       the sampling of ecological interaction networks. PeerJ, 7, e7566.
485
       https://doi.org/10.7717/peerj.7566
486
    Delgado, R., & Tibau, X.-A. (2019). Why Cohen's Kappa should be avoided as performance measure in
487
       classification. PloS One, 14(9), e0222916. https://doi.org/10.1371/journal.pone.0222916
488
    Delmas, E., Besson, M., Brice, M.-H., Burkle, L. A., Dalla Riva, G. V., Fortin, M.-J., Gravel, D., Guimarães,
489
       P. R., Hembry, D. H., Newman, E. A., Olesen, J. M., Pires, M. M., Yeakel, J. D., & Poisot, T. (2018).
490
```

Analysing ecological networks of species interactions. Biological Reviews, 112540.

https://doi.org/10.1111/brv.12433

491

492

```
Desjardins-Proulx, P., Laigle, I., Poisot, T., & Gravel, D. (2017). Ecological interactions and the Netflix
       problem. PeerJ, 5(e3644). https://doi.org/10.7717/peerj.3644
494
    Ferri, C., Hernández-Orallo, J., & Modroiu, R. (2009). An experimental comparison of performance
495
       measures for classification. Pattern Recognition Letters, 30(1), 27–38.
496
       https://doi.org/10.1016/j.patrec.2008.08.010
497
    He, H., & Ma, Y. (Eds.). (2013). Imbalanced Learning: Foundations, Algorithms, and Applications (1st
       edition). Wiley-IEEE Press.
499
    Inman, R., Franklin, J., Esque, T., & Nussear, K. (2021). Comparing sample bias correction methods for
500
       species distribution modeling using virtual species. Ecosphere, 12(3), e03422.
501
       https://doi.org/10.1002/ecs2.3422
502
    Iturbide, M., Bedia, J., Herrera, S., del Hierro, O., Pinto, M., & Gutiérrez, J. M. (2015). A framework for
       species distribution modelling with improved pseudo-absence generation. Ecological Modelling, 312,
504
       166-174. https://doi.org/10.1016/j.ecolmodel.2015.05.018
505
    Japkowicz, N. (2013). Assessment Metrics for Imbalanced Learning. In Imbalanced Learning (pp.
506
       187-206). John Wiley & Sons, Ltd. https://doi.org/10.1002/9781118646106.ch8
507
    Jeni, L. A., Cohn, J. F., & De La Torre, F. (2013). Facing Imbalanced Data-Recommendations for the Use
508
       of Performance Metrics. 2013 Humaine Association Conference on Affective Computing and Intelligent
509
       Interaction, 245–251. https://doi.org/10.1109/ACII.2013.47
510
    Jordano, P. (2016a). Chasing Ecological Interactions. PLOS Biol, 14(9), e1002559.
511
       https://doi.org/10.1371/journal.pbio.1002559
512
    Jordano, P. (2016b). Sampling networks of ecological interactions. Functional Ecology.
       https://doi.org/10.1111/1365-2435.12763
514
    Landis, J. R., & Koch, G. G. (1977). The Measurement of Observer Agreement for Categorical Data.
515
       Biometrics, 33(1), 159-174. https://doi.org/10.2307/2529310
516
    MacDonald, A. A. M., Banville, F., & Poisot, T. (2020). Revisiting the Links-Species Scaling Relationship in
517
       Food Webs. Patterns, 1(0). https://doi.org/10.1016/j.patter.2020.100079
518
    Martinez, N. D. (1992). Constant Connectance in Community Food Webs. The American Naturalist,
519
       139(6), 1208-1218. http://www.jstor.org/stable/2462337
520
```

```
McLeod, A., Leroux, S. J., Gravel, D., Chu, C., Cirtwill, A. R., Fortin, M.-J., Galiana, N., Poisot, T., & Wood,
```

- S. A. (2021). Sampling and asymptotic network properties of spatial multi-trophic networks. *Oikos*,
- n/a(n/a). https://doi.org/10.1111/oik.08650
- Olesen, J. M., Bascompte, J., Dupont, Y. L., Elberling, H., Rasmussen, C., & Jordano, P. (2011). Missing
- and forbidden links in mutualistic networks. *Proc. R. Soc. B*, 278(1706), 725–732.
- 526 https://doi.org/10.1098/rspb.2010.1371
- Olito, C., & Fox, J. W. (2015). Species traits and abundances predict metrics of plant–pollinator network
- structure, but not pairwise interactions. *Oikos*, *124*, 428–436.
- Pichler, M., Boreux, V., Klein, A., Schleuning, M., & Hartig, F. (2020). Machine learning algorithms to
- infer trait-matching and predict species interactions in ecological networks. *Methods in Ecology and*
- Evolution, 11(2), 281-293. https://doi.org/10.1111/2041-210X.13329
- Pilosof, S., Porter, M. A., Pascual, M., & Kéfi, S. (2017). The multilayer nature of ecological networks.
- Nature Ecology & Evolution, 1, 0101. https://doi.org/10.1038/s41559-017-0101
- Poisot, T., Bergeron, G., Cazelles, K., Dallas, T., Gravel, D., MacDonald, A., Mercier, B., Violet, C., &
- Vissault, S. (2021). Global knowledge gaps in species interaction networks data. *Journal of*
- Biogeography, jbi.14127. https://doi.org/10.1111/jbi.14127
- Poisot, T., Ouellet, M.-A., Mollentze, N., Farrell, M. J., Becker, D. J., Albery, G. F., Gibb, R. J., Seifert, S. N.,
- & Carlson, C. J. (2021). Imputing the mammalian virome with linear filtering and singular value
- decomposition. http://arxiv.org/abs/2105.14973
- Saito, T., & Rehmsmeier, M. (2015). The Precision-Recall Plot Is More Informative than the ROC Plot
- When Evaluating Binary Classifiers on Imbalanced Datasets. *PLOS ONE*, 10(3), e0118432.
- https://doi.org/10.1371/journal.pone.0118432
- 543 Schisterman, E. F., Perkins, N. J., Liu, A., & Bondell, H. (2005). Optimal Cut-point and Its Corresponding
- Youden Index to Discriminate Individuals Using Pooled Blood Samples. *Epidemiology*, 16(1), 73–81.
- https://doi.org/10.1097/01.ede.0000147512.81966.ba
- Somodi, I., Lepesi, N., & Botta-Dukát, Z. (2017). Prevalence dependence in model goodness measures with
- special emphasis on true skill statistics. *Ecology and Evolution*, 7(3), 863–872.
- https://doi.org/10.1002/ece3.2654

```
Steen, V. A., Tingley, M. W., Paton, P. W. C., & Elphick, C. S. (2021). Spatial thinning and class balancing:
       Key choices lead to variation in the performance of species distribution models with citizen science
550
       data. Methods in Ecology and Evolution, 12(2), 216–226. https://doi.org/10.1111/2041-210X.13525
551
    Strona, G., & Veech, J. A. (2017). Forbidden versus permitted interactions: Disentangling processes from
552
       patterns in ecological network analysis. Ecology and Evolution, 7(14), 5476–5481.
553
       https://doi.org/10.1002/ece3.3102
554
    Strydom, T., Catchen, M. D., Banville, F., Caron, D., Dansereau, G., Desjardins-Proulx, P., Forero-Muñoz,
555
       N. R., Higino, G., Mercier, B., Gonzalez, A., Gravel, D., Pollock, L., & Poisot, T. (2021). A roadmap
556
       towards predicting species interaction networks (across space and time). Philosophical Transactions of
557
       the Royal Society B: Biological Sciences, 376(1837), 20210063.
558
       https://doi.org/10.1098/rstb.2021.0063
559
    Valdovinos, F. S. (2019). Mutualistic networks: Moving closer to a predictive theory. Ecology Letters, 0(0).
       https://doi.org/10.1111/ele.13279
561
    Weitz, J. S., Hartman, H., & Levin, S. A. (2005). Coevolutionary arms races between bacteria and
       bacteriophage. Proceedings of the National Academy of Sciences of the United States of America, 102(27),
563
       9535-9540. https://doi.org/10.1073/pnas.0504062102
564
    Whalen, S., Schreiber, J., Noble, W. S., & Pollard, K. S. (2021). Navigating the pitfalls of applying machine
565
       learning in genomics. Nature Reviews Genetics, 1–13.
566
       https://doi.org/10.1038/s41576-021-00434-9
567
    Wood, S. A., Russell, R., Hanson, D., Williams, R. J., & Dunne, J. A. (2015). Effects of spatial scale of
568
       sampling on food web structure. Ecology and Evolution, 5(17), 3769–3782.
569
       https://doi.org/10.1002/ece3.1640
570
    Xie, J.-R., Zhang, P., Zhang, H.-F., & Wang, B.-H. (2017). Completeness of Community Structure in
571
       Networks. Scientific Reports, 7(1), 5269. https://doi.org/10.1038/s41598-017-05585-6
572
    Youden, W. J. (1950). Index for rating diagnostic tests. Cancer, 3(1), 32–35.
573
```

https://doi.org/10.1002/1097-0142(1950)3:1%3C32::AID-CNCR2820030106%3E3.0.CO;2-3

574

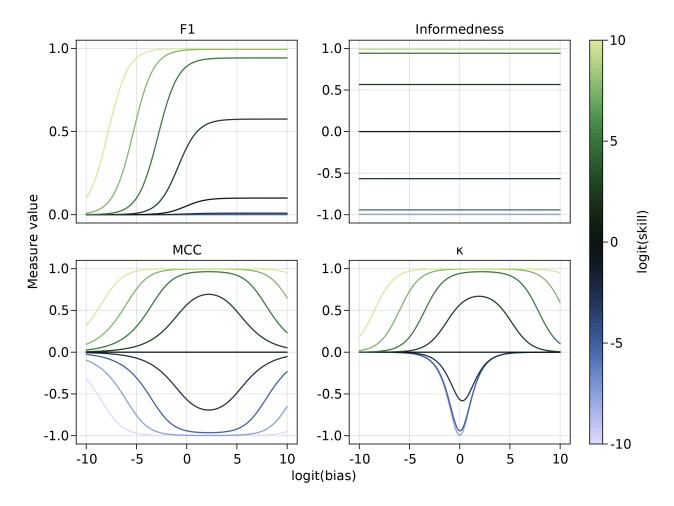


Figure 1: Consequences of changing the classifier skills (s) and bias (s) for a connectance  $\rho=0.15$ , on  $F_1$ , informedness, MCC, and  $\kappa$ . Accuracy increases with skill, but also increases when the bias tends towards estimating *fewer* interactions (this follows from the derivations in the text, not shown in the figure). Interestingly,  $\kappa$  responds as expected to skill (being negative whenever s<0.5), and peaks for values of  $b\approx0.5$ ; nevertheless, the value of bias for which  $\kappa$  is maximized in *not* b=0.5, but instead increases with classifier skill. In other words, at equal skill, maximizing  $\kappa$  would lead to select a *more* biased classifier.

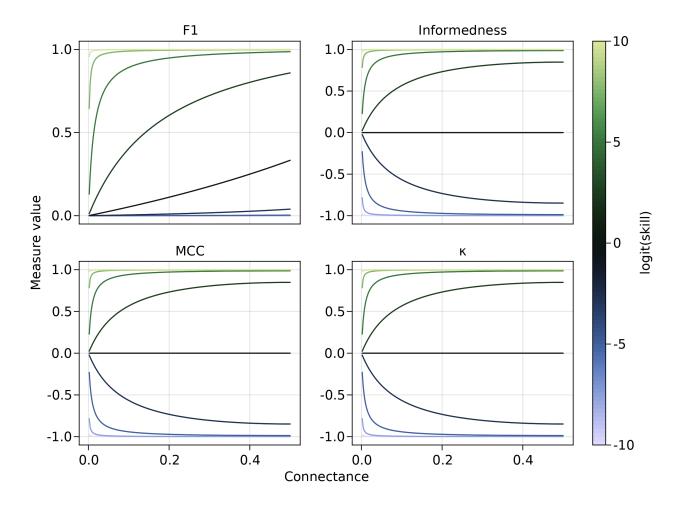


Figure 2: As in fig. 1, consequences of changing connectance for different levels of classifier skill, assuming no classifier bias. Informedness,  $\kappa$ , and MCC do increase with connectance, but only when the classifier is not no-skill; by way of contrast, a more connected network will give a higher  $F_1$  value even with a no-skill classifier.

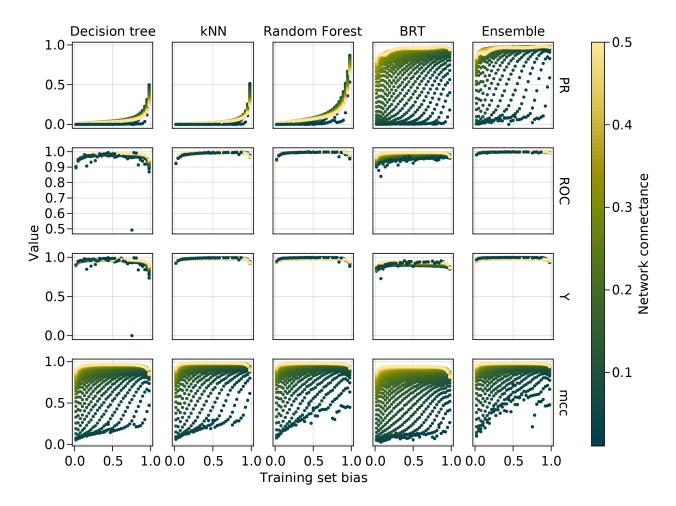


Figure 3: Response of MCC, Informedness, ROC-AUC, and PR-AUC to changes in the training set balance (on the *x* axis) for a series of increasing connectances (color). All of these values approach 1 for a good model, but should be lower when the prediction is more difficult. Informedness is consistently high, and by contrast, MCC increases with additional training set balance. Across all models, training on a more connected network is easier. ROC-AUC is consistently high, and therefore not properly able to separate good from poor classifiers. On the other hand, PR-AUC responds to changes in the training set.

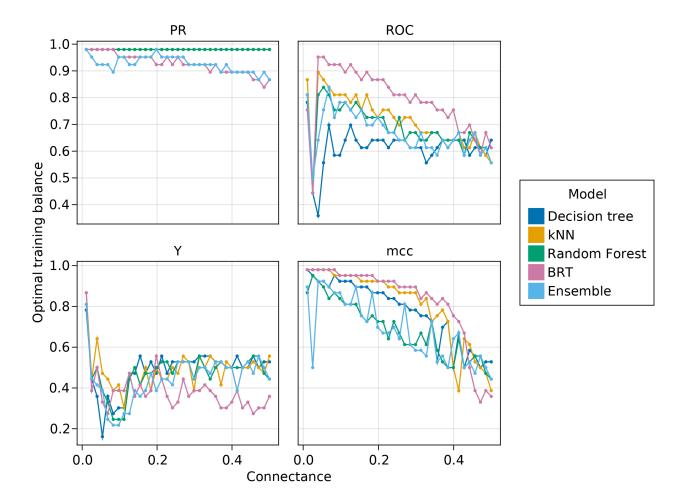


Figure 4: Value of the optimal training set balance for the different models and measures evaluated here, over a range of connectances. Informedness was reliably maximized for balanced training sets, and kept this behavior across models. For other measures, larger connectances in the true network allowed lower biases in the training set. In a large number of cases, "over-correcting" by having training sets with more than half instances representing interactions would maximize the values of the model performance measures.

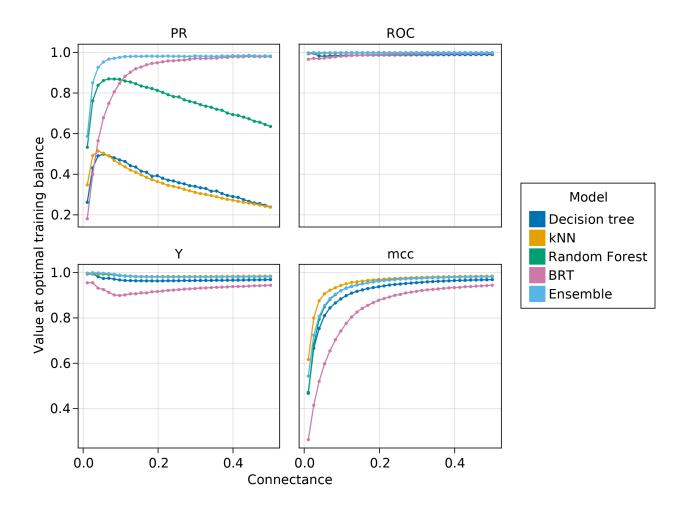


Figure 5: When trained on their optimally biased training set, most models were able to maximize their performance; this is not true when measuring PR-AUC for decision tree, k-NN, and to a lower extent RF. The ensemble had a consistently high performance despite incorporating low-performing models.

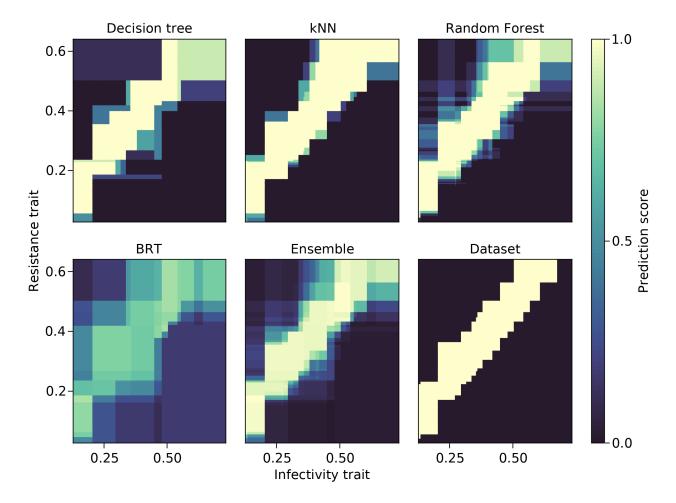


Figure 6: Visualisation of the raw (un-thresholded) models predictions for one instance of a network prediction problem (shown in the "Dataset" panel). Increasing the value of the  $\xi$  parameter would make the diagonal structure "broader," leading to more interactions. A visual inspection of the results is important, as it highlights how some models can "miss" parts of the network; by combining them in an ensemble, these gaps compensate one another, and lead (in this case) to a better prediction.