



Review

Machine-learning techniques for the prediction of protein–protein interactions

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Protein–protein interactions (PPIs) are important for the study of protein functions and pathways involved in different biological processes, as well as for understanding the cause and progression of diseases. Several high-throughput experimental techniques have been employed for the identification of PPIs in a few model organisms, but still, there is a huge gap in identifying all possible binary PPIs in an organism. Therefore, PPI prediction using machine-learning algorithms has been used in conjunction with experimental methods for discovery of novel protein interactions. The two most popular supervised machine-learning techniques used in the prediction of PPIs are support vector machines and random forest classifiers. Bayesian-probabilistic inference has also been used but mainly for the scoring of high-throughput PPI dataset confidence measures. Recently, deep-learning algorithms have been used for sequence-based prediction of PPIs. Several clustering methods such as hierarchical and *k*-means are useful as unsupervised machine-learning algorithms for the prediction of interacting protein pairs without explicit data labelling. In summary, machine-learning techniques have been widely used for the prediction of PPIs thus allowing experimental researchers to study cellular PPI networks.

Keywords. Clustering; deep learning; decision tree; machine-learning techniques; protein–protein interaction; support vector machine

1. Introduction

Protein–protein interaction (PPI) networks play fundamental roles in regulating nearly all biological processes and form the basis for the cellular structure and functions. Deciphering the interaction networks of proteins, therefore, helps in improving our knowledge of functions of proteins, and is also crucial for understanding cellular pathways, and developing effective therapies for the treatment of human diseases. Several high-throughput experimental techniques such as yeast 2-hybrid (Y2H), affinity purification-mass spectrometry (AP-MS) and protein microarrays have been employed for PPI discovery, but their accuracy is questionable due to the occurrence of a large number of false positives (FPs) as well as false negatives (FNs) (Mrowka *et al.* 2001). In this context, it has been noted that fast and scalable machine-learning algorithms have proven extremely useful for the prediction of novel PPIs, and can improve the efficacy of experimental identification of PPIs, when used in conjunction. Furthermore, it has been found that computational predictions of PPIs demonstrated almost similar accuracy levels to those of

large-scale experimental PPI datasets (von Mering *et al.* 2002).

So far, it has been a daunting task for the proteomics researchers to generate a comprehensive picture of entire interactomes, especially for complex eukaryotic organisms such as human beings. The entire human interactome has been estimated to comprise ~130,000 binary PPIs (Venkatesan *et al.* 2009), of which only ~60,000 PPIs have been currently compiled from various published and unpublished experimental results through the Human Reference Protein Interactome (HuRI) project (Bader *et al.* 2017). Experimental studies have also been conducted to unravel the interactomes of other model organisms such as *Escherichia coli*, *Helicobacter pylori*, *Saccharomyces cerevisiae* (yeast), *Caenorhabditis elegans* (roundworm), *Drosophila melanogaster* (fruit fly), *Mus musculus* (mouse) and *Rattus norvegicus* (rat) (Walhout *et al.* 2000; Rain *et al.* 2001; Xenarios *et al.* 2002). However, as in humans, most of these interactomes have also been incomplete to date. For example, the interactome involving only 37.1% of mouse proteins and 20.2% of *C. elegans* proteins is known with a varying degree of certainty (Alonso-López *et al.* 2016).

Nevertheless, several experimental datasets, especially for the yeast interactome (Uetz *et al.* 2000; Gavin *et al.* 2002; Krogan *et al.* 2006), as well as, some PPI databases such as DIP (Salwinski *et al.* 2004), IntAct (Orchard *et al.* 2014) and BioGRID (Chatr-Aryamontri *et al.* 2017), are available in the public domain. In addition, different experimental methods may be able to detect different types of interactions, thereby reporting different subsets of the actual interactome. For example, interaction networks derived from Y2H and AP-MS experiments have different topological and biological properties, because Y2H is better adapted to capture transient interactions between signalling molecules, whereas, AP-MS data are enriched in stable protein complexes (Saha *et al.* 2010). Since PPI networks are inherently dynamic in nature, the lack of tissue-specific or condition-specific (e.g. healthy vs disease state) experimental PPI data further complicates the problem.

Therefore, computational methods can be highly beneficial for the study of interactomes, as they can extrapolate the experimental PPI data for elucidating the complete interactome of an organism, and even predict the interactome for homologous organisms. Furthermore, since experimental strategies for PPI identification are expensive in terms of both time and money, *in silico* prediction of PPIs provides a complementary strategy for query proteome/interactome annotation at a low cost and less time. Although protein–protein docking methods are available, their usefulness in PPI prediction is limited, since there is difficulty in docking proteins undergoing large conformational changes, as well as due to the unavailability of experimentally derived 3D structures for a majority of proteins. In this review article, we have provided an overview of different types of machine-learning strategies that have been utilized for predicting PPIs, and discussed their respective advantages and disadvantages. We have also summarized the datasets used and the organisms studied in the prediction methods developed under each of these categories.

2. Machine-learning techniques

The use of machine-learning techniques in the prediction of PPIs began in 2001 through the independent efforts of a few research groups (Bock and Gough 2001; Sprinzak and Margalit 2001; Zhou and Shan 2001). The prediction of binary PPIs generally involves the sequence or structure of both the interacting proteins as the input and the probability of these two proteins to interact with each other as the output. The observable quantities given as input to a machine-learning algorithm for creating the statistical prediction models are called ‘features’. Thus, for PPI-prediction methods, the typically used features include amino acid composition, domain/motif composition or hydrophobicity profiles of input protein sequences, interface properties of protein 3D structures, genomic features such as gene neighbouring or phylogenetic relationship, and network

topology-based features such as degree distribution or clustering coefficient.

Machine-learning techniques used for predicting PPIs can be broadly divided into two main categories: supervised and unsupervised, based on whether the input variables need to be labelled according to the expected outcome or not (figure 1A). Supervised learning infers a mapping function from a set example input–output pairs, which could be used for predicting the outcome for other inputs, whereas, unsupervised learning discovers the hidden structure within unlabelled training data for drawing meaningful conclusions. Examples of supervised machine-learning algorithms used for PPI prediction include artificial neural networks (ANNs), Bayesian inference, support vector machines (SVMs) and decision tree-based methods such as random forest (RF). Clustering techniques such as *k*-means, single-linkage and spectral clustering are associated with unsupervised machine-learning methods used for PPI prediction. Supervised machine learning is implemented for classification problems, i.e. mapping of input data into specific classes, where a set of quantitative or categorical features are analysed for identified features that can discriminate the input variables into specified classes. Thus, percentage composition of the 20 standard amino acid residues may be computed using sequences of pairs of interacting proteins. Since defining a high-confidence negative dataset for PPI prediction is difficult, negative datasets have been created by considering randomly selected protein pairs not known to interact with each other (Ben-Hur and Noble 2005), or belonging to different sub-cellular locations (Xia *et al.* 2010).

In binary classification problems such as PPI prediction, there are only two categories: the ‘positive’ (p) category, containing proteins that interact with each other, and the ‘negative’ (n) category, containing proteins that do not interact. If the class prediction for each instance is made based on a score that can be represented as a continuous random variable (X), then at a given threshold value (T), the instance is classified as ‘positive’ if $X > T$, and ‘negative’ otherwise. There are four possible prediction outcomes from algorithms undertaking such binary classification problems. If two proteins that have been experimentally verified to interact with each other are correctly classified by the algorithm as an interacting protein pair, then it is called a true-positive (TP) prediction; but if wrongly classified as a non-interacting pair, then this is said to be a FN prediction. Conversely, when two proteins not known to interact are classified by the algorithm as non-interactors, a true-negative (TN) prediction occurs, while if they are classified as interactors then it is a FP prediction.

The prediction efficiency of all classification algorithms can be measured using several threshold-dependent parameters such as precision, sensitivity or recall, specificity, accuracy, *F1*-score and Mathew’s correlation coefficient (MCC), using the following formulae:

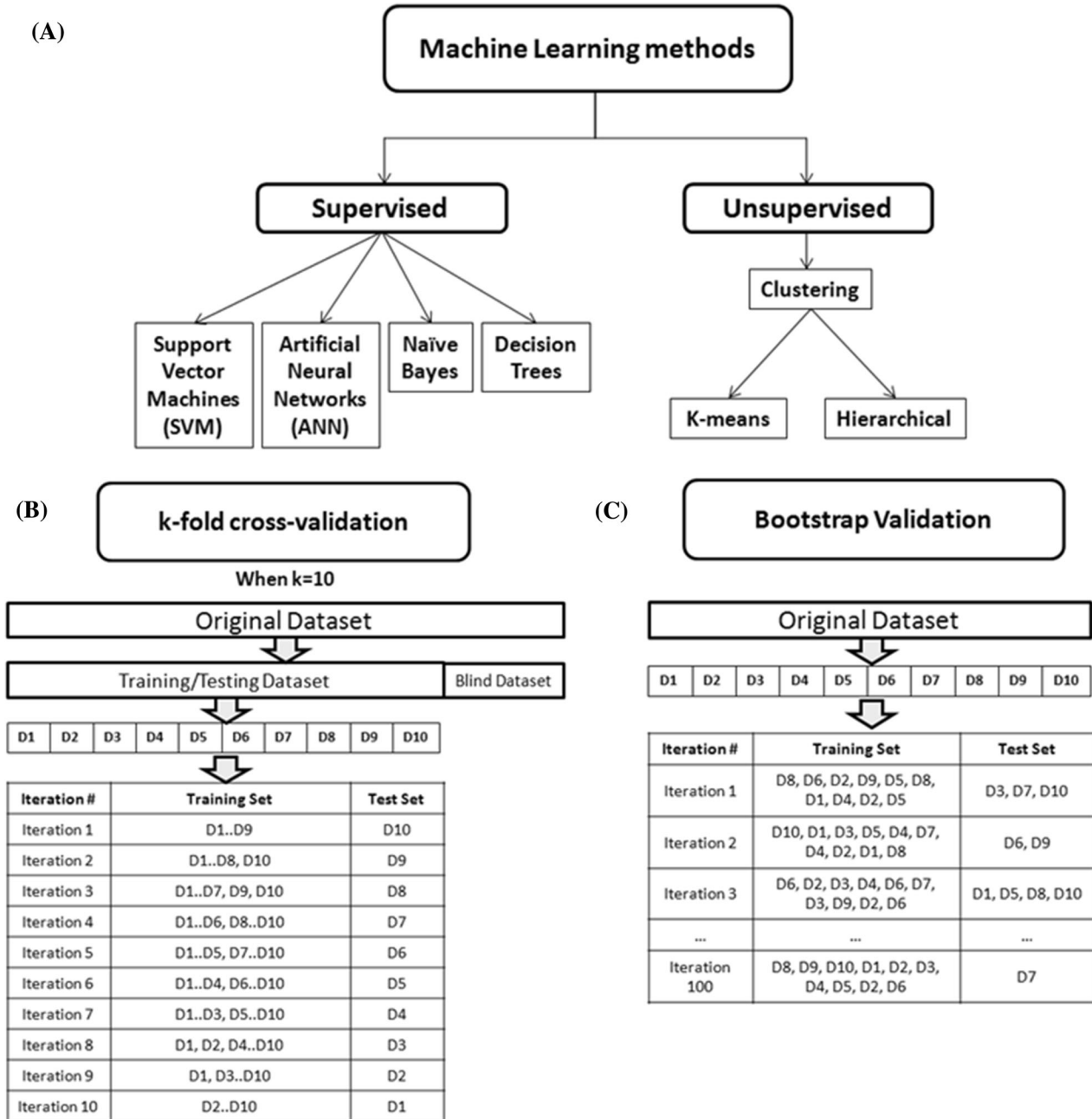


Figure 1. Diagrammatic representation of machine-learning methods, datasets and methods of validation used in prediction of PPIs. (A) Classification of different machine-learning methods into supervised and unsupervised approaches. (B) Training, testing and blind datasets for *k*-fold cross-validation. (C) Training and testing datasets for bootstrap validation.

$$\text{Precision} = \frac{TP}{TP + FP}; \quad \text{Recall/Sensitivity} = \frac{TP}{TP + FN};$$

$$\text{Specificity} = \frac{TN}{TN + FP}$$

$$\text{Accuracy} = \frac{TP + TN}{TP + FP + TN + FN} \times 100;$$

$$F1 \text{ score} = \frac{2TP}{2TP + FP + FN}$$

$$MCC = \frac{(TP)(TN) - (FP)(FN)}{\sqrt{[TP + FP][TP + FN][TN + FP][TN + FN]}} \times 100$$

MCC is generally believed to be a more accurate criterion for evaluating the prediction performance of binary classifiers than accuracy and *F1* score, because it takes into account the balanced ratios of all the four prediction outcomes, viz., TP, FP, TN and FN.

For a threshold-independent overall representation of the prediction performance, a receiver operating characteristics (ROC) curve or a precision-recall (PR) curve is plotted. The area under curve (AUC) statistic for either of these curves, therefore, becomes a threshold-independent measure of

prediction performance. The ROC curve is created by plotting the true-positive rate (TPR) against the false-positive rate (FPR) at various thresholds, where $\text{TPR} = \frac{\text{TP}}{\text{TP} + \text{FN}}$ and $\text{FPR} = \frac{\text{FP}}{\text{FP} + \text{TN}}$. The random variable X that represents the prediction score follows a probability density $f_1(x)$ if the instance actually belongs to the ‘positive’ category, and $f_0(x)$ if it originates from the ‘negative’ category. Therefore, $\text{TPR}(T) = \int_T^\infty f_1(x)dx$ and $\text{FPR}(T) = \int_T^\infty f_0(x)dx$, and when $\text{TPR}(T)$ is plotted parametrically vs $\text{FPR}(T)$ with threshold T as the varying parameter, then

$$\begin{aligned} \text{AUC} &= \int_{-\infty}^{\infty} \text{TPR}(T) \text{FPR}'(T) dT \\ &= \int_{-\infty}^{\infty} \int_{-\infty}^{\infty} I(T' > T) f_1(T') f_0(T) dT' dT \\ &= P(X_1 > X_0) \end{aligned}$$

where X_1 is the score for a positive instance and X_0 is the score for a negative instance.

The aim of PPI predicting machine-learning algorithms is to predict whether a new protein pair will interact with each other or not, after being trained with known PPI pairs. This property is known as generalization and it depends on how well the complexity of the prediction model (hypothesis) generated by the algorithm matches with the complexity of the function underlying the input dataset. If the hypothesis is not complex enough to model the samples, it is called underfitting, whereas, if the hypothesis is too complex and the training samples are not enough to constrain it, this is called overfitting, which leads to problems in generalization. Hence, the generalization ability of a model must be evaluated by dividing the training samples into a larger training dataset and a smaller testing set, for determining the model complexity, as shown in figure 1B. Finally, an independent or blind dataset, containing new input variables not used in training or testing, is also required for evaluating the expected performance of the trained model. For example, Barman *et al.* (2015) have used *E. coli* PPI data downloaded from the Bacterial Protein Interaction Database (*Bacterio.me.org*) for training and testing of the prediction models, and *E. coli* PPIs observed in solved X-ray crystal structures obtained from the PDB as a blind set. In ANNs, the training dataset is used to initially fit the parameters during model construction, and the fitted model is successively evaluated using a separate validation dataset to prevent overfitting and minimize errors. Another test dataset is finally used for an unbiased evaluation of whether the model fits the training dataset. For example, Sun *et al.* (2017) have used 33,052 positive and 32,816 negative samples from their initial PPI dataset as a training dataset, 3493 positive and 3507 negative samples as a hold-out validation dataset and several external test sets for final model evaluation.

The datasets for model training, testing and validation can be prepared using techniques such as randomization, cross-validation and bootstrapping (figure 1B and C). If the dataset

is large enough, then it can be randomly divided into K parts, and each of these parts can again be randomly divided into training and testing sets, and this process can be repeated K times. This is called randomization, which ensures the random sampling of training and testing sets from the data so that the learning process is independent of the selection of training data. However, since datasets large enough for proper randomization are rarely available, the same dataset is repeatedly split into training and testing sets in different ways by a technique called cross-validation or rotation estimation, which can again be exhaustive or non-exhaustive. Exhaustive cross-validation may involve either leave- p -out cross-validation (LpOCV), where p observations are set aside as the test set and the remaining observations are taken as the training set, or leave-one-out cross-validation (LOOCV), where $p = 1$. The most commonly used form of non-exhaustive cross-validation is k -fold cross-validation, where the original sample is randomly partitioned into k equal-sized subsamples, from which a single subsample is retained as the test set and the remaining subsamples are used as the training set. The process is then repeated k times, with each of the k subsamples used exactly once in the test set, as shown in figure 1B. The most commonly used is the 5-fold cross-validation, where the training dataset is divided into five subsets of which four subsets are used in training the model, and the remaining one is used for testing it, and the process is repeated five times, using a different subset in each iteration.

Furthermore, in the real-world biological classification problems, the number of TN PPI pairs far exceeds the number of TP ones. It is therefore, advisable to use both balanced (equal number of TP and TN) and unbalanced ($\text{TN} \gg \text{TP}$) training datasets in such cases (Blagus and Lusa 2010; Yu *et al.* 2010; Barman *et al.* 2015).

2.1 Support vector machines (SVMs)

SVM is a statistical learning algorithm developed by Vapnik (1999). SVM-mediated pattern classification is well known for generalization ability, and thus has been widely used for binary classification problems involving biological data, such as classifying interacting and non-interacting protein pairs. The goal of the SVM algorithm is to find an optimal hyperplane that separates the training samples by a maximal margin, with all positive samples lying on one side and all negative samples lying on the other side. Suppose that we are given a training dataset of N instance-labelled pairs $X = [(x_1, y_1), (x_2, y_2), \dots, (x_N, y_N)]$ with input data $x_i \in R^n$ and labelled output data $y_i \in [+1, -1]$, for $i = 1, 2, 3, \dots, N$. The SVM algorithm solves the quadratic optimization problem by minimizing the function as below:

$$\min_{w, b, \xi} \frac{1}{2} \|w\|^2 + C \sum_{i=1}^N \xi_i \quad (1)$$

$$\text{subject to } y_i(\langle w, x_i \rangle + b) \geq 1 - \xi_i, \quad \xi_i \geq 0, \quad (2)$$

where w is the weight vector, b is the bias term, C is the penalty factor and ξ_i is the slack variable. Since $\|w\|^2$ is convex, minimizing equation (1) under the linear constraints in (2) can be solved with Lagrange multipliers. The complexity parameter denoted by C is a trade-off between training error and margin, which determines the penalty that should be imposed on training data points that end up on the wrong side of the decision boundary.

Generally, the training examples x_i are mapped into a high-dimensional feature space through some nonlinear function ϕ . This transformation helps us to make the class data distributions linearly separable. The feature vectors in SVM thus appear in the form of dot products of two data points as $K(x_i, x_j) = x_i^T \cdot x_j$, which is called a ‘kernel’ and represents a measure of similarity between the two data points x_i and x_j . Hence, even though $\phi(x)$ maybe complex and very high dimensional, after the transformation the dot product, $K(x_i, x_j) = \phi(x_i)^T \cdot \phi(x_j)$, also known as the ‘kernel function’, can have a simple form. Typical kernel functions used in SVM can be linear, polynomial, sigmoid or radial basis function (RBF), as shown below:

- Linear: $K(x_i, x_j) = x_i^T x_j$;
- Polynomial: $K(x_i, x_j) = (\gamma x_i^T x_j + \gamma)^D, \gamma > 0$;
- Sigmoid: $K(x_i, x_j) = \tanh(\gamma x_i^T x_j + B)$;
- RBF: $K(x_i, x_j) = e^{-(\|x_i - x_j\|^2 / 2\sigma^2)}$, or $K(x_i, x_j) = e^{-\gamma \|x_i - x_j\|^2}, \gamma > 0$.

It has been observed that generally the RBF kernel performs best for complex biological classification problems such as predicting PPI pairs (Barman *et al.* 2014).

The majority of SVM-based PPI-prediction methods depend on protein-primary sequences as input (Bock and Gough 2001; Ben-Hur and Noble 2005; Martin *et al.* 2005; Shen *et al.* 2007; Guo *et al.* 2008; Barman *et al.* 2015; Srinivasulu *et al.* 2015; Sriwastava *et al.* 2015; You *et al.* 2015b), since sequence information is available for most proteins. SVM-prediction methods have also been trained with input features such as PDB structures (Zhu *et al.* 2006), domain compositions (Chatterjee *et al.* 2011; Hou *et al.* 2012) and gene ontology (GO) annotations (Mei 2013) of interacting proteins. SVM-based methods for the prediction of specific domain-binding linear motif peptides that may mediate PPIs have also been proposed (Kundu *et al.* 2014; Sarkar *et al.* 2018). Several examples of SVM-based PPI-prediction methods are summarized in table 1.

Although SVM is very efficient in classifying datasets with unspecified complexity, one of its major drawbacks is the requirement of a considerable amount of computational resources for training and parameter optimization, since the number of support vectors grows linearly with the scale of the training data. In addition, the outputs are not probabilistic, and the relationship between training data and output may be complex and not clearly comprehensible.

2.2 Decision tree-based methods

Decision-tree algorithms (Breiman *et al.* 1984) comprise recursive partitioning of the input space by selecting the best attribute, and expanding the leaf nodes of the tree until a predefined stopping criterion, such as a minimum number of training instances assigned to each leaf node of the tree, is met. The best test condition for splitting is determined by different algorithms using different metrics such as Gini impurity and information gain. Gini impurity is a measure of the misclassification to denote the probability of a randomly chosen element from the set to be incorrectly labelled according to the distribution of labels in the subset. Gini impurity for a set of items with J classes, with $i \in [1, 2, \dots, J]$, and p_i being the fraction of items labelled with class i in the set, can be computed as

$$\begin{aligned} I_G(p) &= \sum_{i=1}^J p_i \sum_{k \neq i} p_k = \sum_{i=1}^J p_i (1 - p_i) = \sum_{i=1}^J (p_i - p_i^2) \\ &= \sum_{i=1}^J p_i - \sum_{i=1}^J p_i^2 = 1 - \sum_{i=1}^J p_i^2 \end{aligned}$$

Information gain or entropy is another metric used to determine the best feature to be considered for splitting at each step of tree-building. Entropy is defined as $H(T) = I_E(p_1, p_2, \dots, p_J) = -\sum_{i=1}^J p_i \log_2 p_i$, with p_1, p_2, \dots, p_J being fractions that add up to 1 and represent the percentage of each class present in the child node that results from a split in the tree:

$$\begin{aligned} I_G(T, \alpha) &= H(T) - H(T|\alpha) \\ &= -\sum_{i=1}^J p_i \log_2 p_i - \sum_a p(a) \\ &\quad - \sum_{i=1}^J -Pr(i|a) \log_2 Pr(i|a) \end{aligned}$$

In an RF classification algorithm, many decision trees are constructed based on random feature vectors sampled independently from a training dataset. For a new input, the feature vector is passed through each of the trees and classification is performed by majority voting among the independent trees. The main advantage of this method is that for a large dataset having a large number of features, no separate feature selection algorithm is needed as the algorithm itself can rank features according to their significance in classification.

Various decision tree-based PPI-prediction methods have been developed using several input features such as protein sequence (Chen and Jeong 2009; Xia *et al.* 2010; Zahiri *et al.* 2013; Wei *et al.* 2015; You *et al.* 2015a; Liu *et al.* 2016; Sze-To *et al.* 2016; Zhou *et al.* 2017) and 3D structure (Li *et al.* 2012; Maheshwari and Brylinski 2017). Some methods also use genomic features (Wang *et al.* 2009) and domain composition (Chen and Liu 2005; Rodgers-Melnick *et al.* 2013) as listed in table 2.

Table 1. SVM-based methods for PPI prediction

Type of feature	Dataset(s) used for model building	Organism(s)	Reference(s)
Sequence-based	Database of interacting proteins (DIPs)	<i>E. coli</i> , yeast, <i>Drosophila</i> , human, mouse, rat, bovine	Bock and Gough (2001)
		<i>E. coli</i> , <i>H. pylori</i> , yeast, human, mouse	Martin <i>et al.</i> (2005)
		<i>E. coli</i> , yeast, human	Sriwastava <i>et al.</i> (2015)
		Yeast	Guo <i>et al.</i> (2008)
			You <i>et al.</i> (2015b)
			Ben-Hur and Noble (2005)
			Ben-Hur and Noble (2005)
			Bandyopadhyay and Mallick (2017)
			Shen <i>et al.</i> (2007)
	Biomolecular Interaction Network Database (BIND)	Yeast	
		Human	
	Human Protein References Database (HPRD)	<i>E. coli</i>	Barman <i>et al.</i> (2015)
		Yeast	Ruan <i>et al.</i> (2018)
		–	Srinivasulu <i>et al.</i> (2015)
		–	
	Su <i>et al.</i> (2008)	–	
		–	
	Kiemer <i>et al.</i> (2007)	–	
		–	
	Chen <i>et al.</i> (2013), Yugandhar and Gromiha (2014)	–	
		–	
Structure-based	Protein Data Bank (PDB)	–	Bradford and Westhead (2005)
		–	Zhu <i>et al.</i> (2006)
GO-based	NCBI HIV-1 Human Interaction Database	Human with HIV-1	Mei (2013)
		Yeast	Bandyopadhyay and Mallick (2017)
Domain/motif-based	Database of interacting proteins (DIPs)	<i>E. coli</i> , yeast, <i>Drosophila</i> , human, mouse, rat, bovine	Chatterjee <i>et al.</i> (2011)
		Human with viruses such as HIV-1/SV-40/HBV/HCV, etc.	Barman <i>et al.</i> (2014)
		Yeast	Hou <i>et al.</i> (2012)
		–	Kundu <i>et al.</i> (2014)
		–	
		–	
		–	
		–	
		–	
		–	
		–	Sarkar <i>et al.</i> (2018)

Decision tree-based classification methods have been found to show superior performance in the prediction of PPIs as compared to other types of machine-learning algorithms (Xia *et al.* 2010; You *et al.* 2015a; Zhou *et al.* 2017). In particular, the RF classifier consistently emerged as one of the top performing algorithms across features and datasets (Qi *et al.* 2006; Zhou *et al.* 2017), as illustrated in table 3. However, decision tree-based classification methods, especially RFs, are sensitive to high noise in the data, are prone to overfitting, and highly correlated features can affect their prediction performance.

2.3 Probabilistic/Bayesian classification

Bayesian-probabilistic classifiers are generally preferred by biologists over other machine-learning techniques, because their functionalities are easier to understand as compared to ‘black-box’ predictors such as SVMs and neural networks, and they can be used with both numerical and categorical data.

Probabilistic classifiers denote algorithms that classify the input data points using conditional probability distributions

to model the relationship between the features of the training samples and the class which they belong to. Hence, if the features are denoted by $x_i (i = 1, \dots, M)$, then the feature vector for each data point can be represented as $x = [x_1, x_2, \dots, x_M]$, and the probability of the data point belonging to each of the N classes ($c = c_1, c_2, \dots, c_N$) as $P(C = c_1|x), P(C = c_2|x), \dots, P(C = c_N|x)$. The probability for each class can be computed using the Bayes’ theorem as

$$P(C|x) = \frac{P(x|C)P(C)}{P(x)} = \frac{P(x|C)P(C)}{\sum_{c_i \in C} P(x|C = c_i)P(C = c_i)}$$

$P(x|C)$ and $P(C)$ can be estimated from the training data for each class. Thus, a new data point is classified as the class with the maximum probability $c^* = \arg \max_{c_j} P(C = c_j|x)$ and $j = 1, \dots, N$.

After modelling the class conditional probabilities, the probabilistic approach seeks to classify the input data points to the class with the maximum probability. In case of a binary classification problem this amounts to computing the ratio $\Upsilon = \frac{P(x|C=c_1)}{P(x|C=c_2)}$, and then choosing c_1 if $\Upsilon > 1$, and c_2 otherwise, because the decision boundary is formed by the region of the feature space where $\Upsilon = 1$.

Table 2. Decision tree-based methods for PPI prediction

Type of feature	Dataset(s) used for model building	Organism(s)	Reference(s)
Sequence-based	Database of interacting proteins (DIPs)	Yeast	You <i>et al.</i> (2015a)
		Yeast and <i>H. pylori</i>	Xia <i>et al.</i> (2010)
	Human Protein References Database (HPRD)	Human	Zahiri <i>et al.</i> (2013)
	Protein Data Bank (PDB)	–	Chen and Jeong (2009)
	Murakami and Mizuguchi (2010)	–	Wei <i>et al.</i> (2015)
	Murakami and Mizuguchi (2010)	–	Liu <i>et al.</i> (2016)
	Wu <i>et al.</i> (2009)	Yeast	Sze-To <i>et al.</i> (2016)
Structure-based	Martin <i>et al.</i> (2005), You <i>et al.</i> (2014), Huang <i>et al.</i> (2015)	<i>H. pylori</i> , yeast and human	Zhou <i>et al.</i> (2017)
	Protein Data Bank (PDB)	–	Maheshwari and Brylinski (2017)
Genomic feature-based	Database of Three-dimensional Interacting Domains (3did)	–	Li <i>et al.</i> (2012)
	Munich Information Center for Protein Sequences (MIPS)	Yeast	Wang <i>et al.</i> (2009)
Domain/motif-based	Database of interacting proteins (DIPs)	Yeast Yeast, <i>Arabidopsis</i> , <i>Drosophila</i> , human, mouse	Chen and Liu (2005) Rodgers-Melnick <i>et al.</i> (2013)

Table 3. Comparison of SVM- and RF-based PPI-prediction methods on PPI datasets from different organisms (Zhou *et al.* 2017)

Organism	Classifier	Accuracy (%)	F-score (%)	MCC (%)
Yeast	SVM	93.25	92.97	86.51
	RF	94.61	94.44	89.37
<i>H. pylori</i>	SVM	85.94	85.90	71.91
	RF	86.28	86.27	72.58
Human	SVM	96.45	96.20	92.96
	RF	97.57	97.44	95.15

Probabilistic methods based on log-odds scoring schemes have been widely used in PPI prediction, as well as for filtering high-throughput experimental datasets that can potentially include several FPs (Gavin *et al.* 2006; Krogan *et al.* 2006; Saha *et al.* 2010; Choi *et al.* 2011). Bayesian-probabilistic frameworks have also been proposed for PPI prediction using genomic features such as co-expression values, essentiality and co-localization (Jansen *et al.* 2003), structural features (Murakami and Mizuguchi 2010; Zhang *et al.* 2012) and sequence signatures (Sprinzak and Margalit 2001). Some prominent examples of PPI-prediction methods based on Bayesian conditional probability relationships are summarized in table 4.

The major disadvantage of Bayesian classification is the forced assumption of independence among the features, which is difficult to ensure in real-world problems. In addition, specifying a prior probability and computing a posterior probability for each data point can be extremely difficult and computationally infeasible in some cases.

2.4 Artificial neural networks

Artificial neural network (ANN) is one of the oldest machine-learning algorithms that can be used to perform

nonlinear statistical modelling and develop binary classification models, and has now evolved into state-of-the-art deep-learning algorithms such as stacked autoencoders. ANNs require less formal statistical training, and are able to implicitly detect complex nonlinear relationships between dependent and independent variables, as well as, detect all possible interactions between predictor variables.

ANNs consist of a network of connections, each of which transfers the output of a neuron i to the input of a neuron j . Thus, i is the predecessor of j and j is the successor of i . Each connection is assigned a weight w_{ij} . The propagation function $p_j(t) = \sum_i o_i(t)w_{ij}$, computes the input $p_j(t)$ to the neuron j from the outputs $o_i(t)$ of predecessor neurons. Thus, a neuron with label j receives an input $p_j(t)$ from predecessor neurons, and consists of the following components:

- an activation $a_j(t)$, depending on a discrete time parameter,
- a threshold θ_j , which stays fixed unless changed by a learning function,
- an activation function f that computes the new activation at a given time $t + 1$ from $a_j(t)$, θ_j and the net input $p_j(t)$ giving $a_j(t + 1) = f(a_j(t), p_j(t), \theta_j)$ and
- an output function f_{out} for computing the output from the activation as

$$o_j(t) = f_{\text{out}}(a_j(t))$$

PPI-prediction methods based on ANNs and deep learning have been developed using several sequence (You *et al.* 2013; Yousef and Moghadam Charkari 2013; Sun *et al.* 2017; Wang *et al.* 2017b; Huang *et al.* 2018) and structural (Zhou and Shan 2001; Fariselli *et al.* 2002; Ofra and Rost 2007; Wang *et al.* 2010; Du *et al.* 2016) features as listed in table 5.

Table 4. Probabilistic/Bayesian methods for PPI prediction

Type of feature	Dataset(s) used for model building	Organism(s)	Reference(s)
PPI data analysis	Gavin <i>et al.</i> (2006) Krogan <i>et al.</i> (2006) Sardiú <i>et al.</i> (2008), Sowa <i>et al.</i> (2009) Krogan <i>et al.</i> (2006), Yu <i>et al.</i> (2008) Collins <i>et al.</i> (2013), Teo <i>et al.</i> (2015)	Yeast –	Gavin <i>et al.</i> (2006) Krogan <i>et al.</i> (2006) Choi <i>et al.</i> (2011) Saha <i>et al.</i> (2010) Teo <i>et al.</i> (2016)
Sequence-based	Database of interacting proteins (DIPs)	Yeast Yeast and <i>H. pylori</i>	Sprinzak and Margalit (2001) An <i>et al.</i> (2016) Li <i>et al.</i> (2017) Wang <i>et al.</i> (2017a)
Structure-based	Protein Data Bank (PDB)	Yeast and human –	Zhang <i>et al.</i> (2012) Murakami and Mizuguchi (2010)
Genomic feature-based	Munich Information Center for Protein Sequences (MIPS)	–	Jansen <i>et al.</i> (2003)
Domain/motif-based	Human Protein References Database (HPRD) Munich Information Center for Protein Sequences (MIPS) Human Protein References Database (HPRD) Landgraf <i>et al.</i> (2004), Tonikian <i>et al.</i> (2009)	Human Yeast Human Yeast	Scott and Barton (2007) Sprinzak and Margalit (2001) Scott and Barton (2007) Jain and Bader (2016)

Table 5. ANN-based methods for PPI prediction

Type of feature	Dataset(s) used for model building	Organism(s)	Reference(s)
Sequence-based	Database of interacting proteins (DIPs)	Yeast Yeast and <i>H. pylori</i>	You <i>et al.</i> (2013) Huang <i>et al.</i> (2018) Yousef and Moghadam Charkari (2013) Wang <i>et al.</i> (2017b)
Structure-based	Human Protein References Database (HPRD) Protein Data Bank (PDB) Database of Three-dimensional Interacting Domains (3did)	Human – – – –	Sun <i>et al.</i> (2017) Huang <i>et al.</i> (2018) Zhou and Shan (2001) Fariselli <i>et al.</i> (2002) Ofra and Rost (2007) Wang <i>et al.</i> (2010) Du <i>et al.</i> (2016)

The disadvantages of using ANN methods include their ‘black box’ nature, greater computational burden, proneness to overfitting and the empirical nature of model development.

2.5 Clustering

Clustering is the major form of unsupervised machine-learning technique applied to classification problems, which tries to segregate data points into groups such that data points placed in the same group are more similar to each

other than to those in other groups. Clustering is useful in exploratory pattern analysis, pattern classification, decision making and also for outlier detection. The main advantage of clustering is its ability to determine the intrinsic classification within a set of unlabelled data, hence not requiring a separate training stage. Hence, clustering is generally used in cases where the class labels are not known in advance.

The different distance metrics used by clustering algorithms include: (a) Euclidean distance metric, (b) Euclidean squared distance metric, (c) Manhattan (city-block) distance, (d) Chebyshev distance, (e) Pearson’s correlation coefficient,

Table 6. Clustering-based methods for PPI prediction

Type of feature	Dataset(s) used for model building	Organism(s)	Reference(s)
Network topology-based	Database of interacting proteins (DIPs) Munich Information Center for Protein Sequences (MIPS) Others	Yeast Yeast Yeast	Liu <i>et al.</i> (2015) Bader and Hogue (2003) Spirin and Mirny (2003) Xu and Guan (2014)
Structure-based	Protein Data Bank (PDB)	–	Fukuhara and Kawabata (2008)

(f) squared Pearson's correlation coefficient and (g) Spearman's rank correlation coefficient.

Clustering techniques have been used for PPI prediction using mainly the overall network topology of known interaction networks (Bader and Hogue 2003; Spirin and Mirny 2003; Xu and Guan 2014; Liu *et al.* 2015), except for one method that uses structural features (Fukuhara and Kawabata 2008), as shown in table 6.

The disadvantages of clustering include the inability to process high-dimensional datasets, getting stuck into local optima, ambiguity regarding cluster descriptors and the inherent randomness of the method, which causes problems with reproducibility. In addition, the final output shows the similarity of the object to a single cluster only, whereas, in reality, the object might have similarities to other clusters also, and this information is lost.

3. Discussion

The use of machine-learning methods for prediction of PPIs can help in discovery of novel PPIs by filtering out the large proportion of FPs and FNs reported by high-throughput experimental procedures and improving interactome coverage. Predicted PPIs from machine-learning methods can also be used to prioritize pairs of proteins for experimental assays, and may provide further insight into the specific context for the PPI such as tissue or phenotype. PPI networks discovered through computational prediction and subsequent experimental validation can be used for prediction of gene function (Mostafavi and Morris 2012), identification of disease genes (Navlakha and Kingsford 2010) and the discovery of novel therapeutics (Barabasi *et al.* 2011). Computational analyses of various physico-chemical, structural and functional attributes of interacting protein pairs also provide a better understanding of the molecular basis of PPIs. Nevertheless, the sets of features that have been used to predict PPIs are still not able to fully capture the dynamic and intricate specifications that can identify the true PPIs unambiguously.

The prediction of interacting protein pairs is a complex problem, since it is dependent not only on the sequence or structure, but also on other parameters such as cellular concentration and localization. Initially, various sequence- and structure-based features were used for training of SVMs, ANNs and Bayesian models for prediction of novel PPIs. It was observed that the prediction performance of RF-based methods is better than that of the more popular SVM-based methods. Several studies described in tables 1, 2, 4–6 have used a combination of more than one feature in a single method for prediction of PPIs with higher accuracy. However, the evaluation and incorporation of an optimum set of features encompassing all biochemical, contextual and structural information associated with proper identification of PPIs still remains elusive. Hence, the field of prediction of PPIs using machine-learning techniques is still open in terms of selecting new features, developing new algorithms and parameter optimization.

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