# RESEARCH



**Open Access** 

# PCTFPeval: a web tool for benchmarking newly developed algorithms for predicting cooperative transcription factor pairs in yeast

Fu-Jou Lai, Hong-Tsun Chang, Wei-Sheng Wu\*

*From* Joint 26th Genome Informatics Workshop and Asia Pacific Bioinformatics Network (APBioNet) 14th International Conference on Bioinformatics (GIW/InCoB2015) Tokyo, Japan. 9-11 September 2015

# Abstract

**Background:** Computational identification of cooperative transcription factor (TF) pairs helps understand the combinatorial regulation of gene expression in eukaryotic cells. Many advanced algorithms have been proposed to predict cooperative TF pairs in yeast. However, it is still difficult to conduct a comprehensive and objective performance comparison of different algorithms because of lacking sufficient performance indices and adequate overall performance scores. To solve this problem, in our previous study (published in BMC Systems Biology 2014), we adopted/proposed eight performance indices and designed two overall performance scores to compare the performance of 14 existing algorithms for predicting cooperative TF pairs in yeast. Most importantly, our performance comparison framework can be applied to comprehensively and objectively evaluate the performance of a newly developed algorithm. However, to use our framework, researchers have to put a lot of effort to construct it first. To save researchers time and effort, here we develop a web tool to implement our performance comparison framework, featuring fast data processing, a comprehensive performance comparison and an easy-to-use web interface.

**Results:** The developed tool is called PCTFPeval (Predicted Cooperative TF Pair evaluator), written in PHP and Python programming languages. The friendly web interface allows users to input a list of predicted cooperative TF pairs from their algorithm and select (i) the compared algorithms among the 15 existing algorithms, (ii) the performance indices among the eight existing indices, and (iii) the overall performance scores from two possible choices. The comprehensive performance comparison results are then generated in tens of seconds and shown as both bar charts and tables. The original comparison results of each compared algorithm and each selected performance index can be downloaded as text files for further analyses.

**Conclusions:** Allowing users to select eight existing performance indices and 15 existing algorithms for comparison, our web tool benefits researchers who are eager to comprehensively and objectively evaluate the performance of their newly developed algorithm. Thus, our tool greatly expedites the progress in the research of computational identification of cooperative TF pairs.

# Background

Understanding combinatorial or cooperative transcriptional regulation by two or more transcription factors (TFs) has become an important research topic in the recent decade. Researchers have studied and modelled

\* Correspondence: wessonwu@mail.ncku.edu.tw

various types of TF-TF interactions which contribute to positive or negative synergy in regulating genes [1-3]. Attributing to the availability of various kinds of genome-wide datasets (e.g. gene expression data, ChIP-chip data, TF binding site motifs, protein-protein interaction data and TF knockout data), researchers continued developing advanced algorithms to predict cooperative TF pairs. Some algorithms only utilized ChIP-chip data



© 2015 Lai et al. This is an Open Access article distributed under the terms of the Creative Commons Attribution License (http:// creativecommons.org/licenses/by/4.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited. The Creative Commons Public Domain Dedication waiver (http://creativecommons.org/publicdomain/ zero/1.0/) applies to the data made available in this article, unless otherwise stated.

Department of Electrical Engineering, National Cheng Kung University, Tainan, Taiwan

[3-6] or gene expression data [7], and the others integrated multiple data sources [8-17].

Since different algorithms integrated different data sources, used different rationales and predicted distinct lists of cooperative TF pairs, it is hard to tell which one is the best. Typically, researchers only compared their algorithm with a few existing algorithms using a few performance indices (see Table 1) and claimed their algorithm to be the best one. However, this kind of comparison is incomplete and subjective [18]. A comprehensive and objective performance comparison framework is urgently needed.

To meet this need, in our previous study [19], we proposed/adopted eight performance indices to compare the performance of 14 existing algorithms. Our results showed that the performance of an algorithm varies widely across different performance indices, implying that researchers may make a biased conclusion based on only a few performance indices. Therefore, in order to conduct a comprehensive and objective performance comparison, we designed two overall performance scores to summarize the comparison results of the eight performance indices.

Most importantly, our performance comparison framework can be applied to comprehensively and objectively evaluate the performance of a newly developed algorithm. Therefore, researchers who develop a new algorithm definitely would like to use our performance comparison framework to quickly evaluate the prediction performance in order for improvement when needed. However, to use our framework, researchers have to put a lot of effort to construct it first. Constructing our framework involves collecting and processing multiple genome-wide datasets from the public domain, collecting the lists of the predicted cooperative TF pairs from 15 existing algorithms in the literature, and writing a lot of

Table 1 The numbers of the compared algorithms, the performance indices, and the predicted cooperative TF pairs (PCTFPs) for each of the 15 existing algorithms.

Algorithm	# of existing algorithms used for performance comparison in their paper	# of indices used for performance evaluation in their paper	# of PCTFPs
Banerjee and Zhang (NAR 2003) [8]	0	1	31
Harbison et al. (Nature 2004) [4]	0	0	94
Nagamine et al. (NAR 2005) [9]	0	1	24
Tsai et al. (PNAS 2005) [10]	0	1	18
Chang et al. (Bioinformatics 2006) [11]	2	1	55
He et al. (IEEE GCCW 2006) [12]	2	1	30
Yu et al. (NAR 2006) [5]	0	1	300
Wang J (JBI 2007) [13]	3	1	14
Elati et al. (Bioinformatics 2007) [7]	4	1	20
Datta and Zhao (Bioinformatics 2008) [6]	3	1	25
Chuang et al. (BMC Bioinformatics 2009) [14]	4	2	13
Wang Y et al. (NAR 2009) [15]	5	2	159
Yang et al. (Cell Research 2010) [16]	3	1	186
Chen et al. (Bioinformatics 2012) [3]	2	2	221
Lai et al. (BMC Systems Biology 2014) [17]	11	3	27

codes to implement the eight performance indices. To save researchers time and effort, here we develop a web tool called PCTFPeval (Predicted Cooperative TF Pair evaluator) to implement our performance comparison framework, featuring fast data processing, a comprehensive performance comparison and an easy-to-use web interface. Constructing PCTFPeval is not a daunting task for us since we already have many experiences in developing databases and web tools [20-26].

# Implementation

# Fifteen existing algorithms used for performance comparison

Our tool provides 15 existing algorithms for users to conduct a performance comparison. As far as we know, this is the most comprehensive collection of the existing algorithms whose lists of the predicted cooperative TF pairs in yeast are available. The numbers of the predicted cooperative TF pairs from different algorithms vary widely, ranging from 13 to 300 (see Table 1).

# Eight existing performance indices used for performance evaluation

Our tool implements eight existing performance indices for users to evaluate the performance of an algorithm for predicting cooperative TF pairs in yeast. As far as we know, this is the most comprehensive collection of the existing performance indices. These eight performance indices can be divided into two types: TF-based indices and target gene based (TG-based) indices. Each type has four indices and different indices utilize different data sources and rationales (see Table 2).

# Two existing overall performance scores used for representing the comprehensive performance comparison results

Our tool implements two existing overall performance scores [19] to summarize the comparison results of the selected performance indices. The first one is called the comprehensive ranking score defined as the sum of the rankings in the selected performance indices [19]. The ranking of an algorithm in an index is k if its performance ranks #k among all the compared algorithms in that index. For example, the ranking of the best performing algorithm is 1. Therefore, the smaller the comprehensive ranking score, the better the overall performance of an algorithm.

The second overall performance score is called the comprehensive normalized score (*CNS*) defined as the sum of the normalized scores in the selected performance indices [19]. The *CNS* of the algorithm i is calculated as follows:

$$CNS(i) = \sum_{j=1}^{L} NS_{j}(i) = \sum_{j=1}^{L} \left( \frac{OS_{j}(i)}{\max(OS_{j}(1), OS_{j}(2), ..., OS_{j}(n))} \right)$$

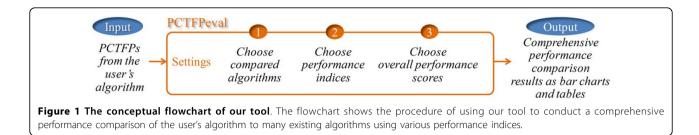
where  $NS_j(i)$  and  $OS_j(i)$  is the normalized score and the original score of the algorithm *i* calculated using the index *j*, respectively; *n* is the number of the algorithms being compared; *L* is the number of the selected indices. Note that  $0 \le NS_j(i) \le 1$  and  $NS_j(i) = 1$  if and only if the algorithm *i* is the best performing algorithm in the

Table 2 The eight performance indices implemented in our tool

Performance index type	Index	Data sources used	Rationale
	Index1	Yeast physical PPI data from BioGRID database [27]	Measure the overlap significance of the physical PPI partners of a PCTFP*
	Index2	Yeast physical PPI data from BioGRID database [27]	Measure the shortest path length of a PCTFP in the physical PPI network
	Index3	Yang et al.'s functional similarity scores of any two yeast genes [28]	Measure the functional similarity of a PCTFP
	Index4	Yang et al.'s high-quality benchmark set of 27 cooperative TF pairs in yeast [16]	Measure the overlap significance of the list of PCTFPs from an algorithm and the benchmark set of 27 cooperative TF pairs
TG-based	Index5	Balaji et al.'s co-regulatory coefficient dataset of 3459 TF pairs in yeast [29]	Measure the co-regulatory coefficient of a PCTFP
	Index6	Co-expression scores of any two yeast genes from SPELL database [30] and TF-gene documented regulation data from YEASTRACT database [31]	Measure the expression coherence of a PCTFP's common target genes
	Index7	Yang et al.'s functional similarity scores of any two yeast genes [28] and TF-gene documented regulation data from YEASTRACT database [31]	Measure the functional coherence of a PCTFP's common target genes
	Index8	Yeast physical PPI data from BioGRID database [27] and TF-gene documented regulation data from YEASTRACT database [31]	Measure the physical PPI coherence of a PCTFP's common target genes

\*PCTFP is the abbreviation for predicted cooperative TF pair.

Different indices utilizes different data sources and rationales. See our previous study [19] for the details about the mathematics of these eight performance indices.



index *j* (i.e. it has the highest original score calculated using the index *j*). The larger the *CNS*, the better the performance of an algorithm.

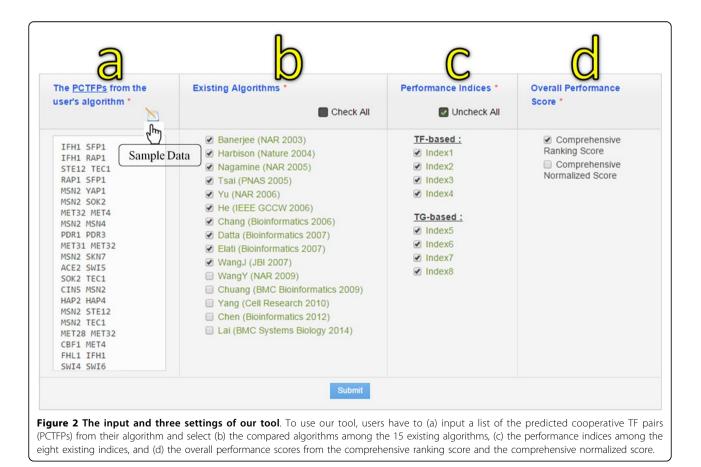
# **Results and discussion**

# Usage

The conceptual flowchart of our tool is shown in Figure 1. The friendly web interface allows users to input a list of the predicted cooperative TF pairs from their algorithm. Then three kinds of settings of our tool have to be specified. First, users have to choose the compared algorithms among the 15 existing algorithms. Second, users have to choose the performance indices among the eight existing indices. Finally, users have to choose the overall performance scores from the comprehensive ranking score and the comprehensive normalized score. After the submission, our tool conducts a comprehensive performance comparison of the user's algorithm to the compared algorithms using the selected performance indices. The comprehensive performance comparison results are then generated in tens of seconds and shown as both bar charts and tables.

# Case study

In our tool, a list of 40 TF pairs is provided as a sample data. For demonstration purpose, we regard the sample data as the list of the predicted cooperative TF pairs from a new algorithm and would like to conduct a comprehensive performance comparison of this new algorithm to the various existing algorithms using our tool. As shown in Figure 2, users input the sample data to our tool and select



#### Lai et al. BMC Bioinformatics 2015, **16**(Suppl 18):S2 http://www.biomedcentral.com/1471-2105/16/S18/S2



**Figure 3 The output of our tool**. Here we input the sample data (a list of 40 TF pairs) as a list of the predicted cooperative TF pairs (PCTFPs) from a user's algorithm and select 10 existing algorithms, eight performance indices, and the comprehensive ranking score as the overall performance score. (a) The comprehensive performance comparison results are shown as a bar chart and a table. It can be seen that the overall performance of the user's algorithm ranks three among all the 11 algorithms being compared. (b) When clicking the hyperlink of "Index5", users will get the performance comparison results (shown as both a bar chart and a table) using only the index 5. It can be seen that the user's algorithm is the best performing algorithm in the index 5. (c) When clicking the hyperlink of "Details of the score of Index5 for each compared algorithm", users will get a text file containing the original scores (calculated using the index 5) of all PCTFPs of each algorithm being compared.

(i) 10 existing algorithms for comparison, (ii) eight performance indices for evaluation, and (iii) the comprehensive ranking score as the overall performance score. After the submission, the comprehensive comparison results are generated and shown as both bar charts and tables (see Figure 3). It can be seen that the new algorithm performs well in the first five performance indices but performs worse in the last three performance indices. The overall performance of the new algorithm ranks three among all the 11 algorithms being compared. Getting the comprehensive comparison results from our tool, researchers immediately know that there is still room to improve the performance of their new algorithm.

# Conclusions

Knowing the cooperative TFs is crucial for understanding the combinatorial regulation of gene expression in eukarvotic cells. This is why the computational identification of cooperative TF pairs has become a hot research topic. Researchers will keep developing new algorithms. Using our tool, researchers can quickly conduct a comprehensive and objective performance comparison of their new algorithm to the various existing algorithms. If the performance of their new algorithm is not satisfactory, researchers can modify their algorithm and use our tool again to see if the performance is improved. Therefore, having our tool in hand, researchers can now totally focus on designing new algorithms and need not worry about how to comprehensively and objectively evaluate the performance of their new algorithms. In conclusion, our tool can greatly expedite the progress in this research topic.

# Availability and requirements

#### Project name: PCTFPeval

Project home page: http://cosbi.ee.ncku.edu.tw/ PCTFPeval/

Operating system(s): platform independent.

Programming language: PHP, Python and Javascript. Other requirements: Internet connection.

License: none required.

Any restrictions to use by non-academics: no restriction.

#### **Competing interests**

The authors declare that they have no competing interests.

#### Authors' contributions

WSW conceived the research topic and provided essential guidance. WSW and FJL developed the method and wrote the manuscript. FJL collected and processed all the genome-wide datasets used in this tool. HTC constructed the web interface of this tool. All authors read, edited and approved the final manuscript.

#### Acknowledgements

This study was supported by National Cheng Kung University and Ministry of Science and Technology of Taiwan MOST-103-2221-E-006 -174 -MY2.

#### Declarations

The publication of this paper was funded by National Cheng Kung University and Ministry of Science and Technology of Taiwan MOST-103-2221-E-006 -174 -MY2.

This article has been published as part of *BMC Bioinformatics* Volume 16 Supplement 18, 2015: Joint 26th Genome Informatics Workshop and 14th International Conference on Bioinformatics: Bioinformatics. The full contents of the supplement are available online at http://www.biomedcentral.com/ bmcbioinformatics/supplements/16/S18.

#### Published: 9 December 2015

#### References

- 1. Miller JA, Widom J: Collaborative competition mechanism for gene activation in vivo. *Mol Biol Cell* 2003, 23(5):1623-1632.
- Tanay A: Extensive low-affinity transcriptional interactions in the yeast genome. Genome Res 2006, 16:962-972.
- Chen MJ, Chou LC, Hsieh TT, Lee DD, Liu KW, Yu CY, Oyang YJ, Tsai HK, Chen CY: De novo motif discovery facilitates identification of interactions between transcription factors in Saccharomyces cerevisiae. *Bioinformatics* 2012, 28(5):701-708.
- Harbison CT, Gordon DB, Lee TI, Rinaldi NJ, Macisaac KD, Danford TW, Hannett NM, Tagne JB, Reynolds DB, Yoo J, Jennings EG, Zeitlinger J, Pokholok DK, Kellis M, Rolfe PA, Takusagawa KT, Lander ES, Gifford DK, Fraenkel E, Young RA: Transcriptional regulatory code of a eukaryotic genome. *Nature* 2004, 431(7004):99-104.
- Yu X, Lin J, Masuda T, Esumi N, Zack DJ, Qian J: Genome-wide prediction and characterization of interactions between transcription factors in Saccharomyces cerevisiae. *Nucleic Acids Res* 2006, 34(17):917-927.
- Datta D, Zhao H: Statistical methods to infer cooperative binding among transcription factors in Saccharomyces cerevisiae. *Bioinformatics* 2008, 24:545-552.
- Elati M, Neuvial P, Bolotin-Fukuhara M, Barillot E, Radvanyi F, Rouveirol C: LICORN: learning cooperative regulation networks from gene expression data. *Bioinformatics* 2007, 23(18):2407-2414.
- Banerjee N, Zhang MQ: Identifying cooperativity among transcription factors controlling the cell cycle in yeast. Nucleic Acids Res 2003, 31:7024-7031.
- Nagamine N, Kawada Y, Sakakibara Y: Identifying cooperative transcriptional regulations using protein-protein interactions. *Nucleic Acids Res* 2005, 33:4828-4837.
- Tsai HK, Lu HHS, Li WH: Statistical methods for identifying yeast cell cycle transcription factors. Proc Natl Acad Sci USA 2005, 102:13532-13537.
- Chang YH, Wang YC, Chen BS: Identification of transcription factor cooperativity via stochastic system model. *Bioinformatics* 2006, 22(18):2276-2282.
- He D, Zhou D, Zhou Y: Identifying synergistic transcriptional factors involved in the yeast cell cycle using Microarray and ChIP-chip data. In Proceedings of the Fifth International Conference on Grid and Cooperative Computing Workshops:21-23 October 2006; Hunan. Los Alamitos: IEEE Computer Society;Xiao N, Buyya R, Liu Y, Yang G 2006:357-360.
- 13. Wang J: A new framework for identifying combinatorial regulation of transcription factors: a case study of the yeast cell cycle. J Biomedical Informatics 2007, 40(6):707-725.
- Chuang CL, Hung K, Chen CM, Shieh GS: Uncovering transcriptional interactions via an adaptive fuzzy logic approach. *BMC Bioinformatics* 2009, 10:400.
- Wang Y, Zhang XS, Xia Y: Predicting eukaryotic transcriptional cooperativity by Bayesian network integration of genome-wide data. *Nucleic Acids Res* 2009, 37(18):5943-5958.
- Yang Y, Zhang Z, Li Y, Zhu XG, Liu Q: Identifying cooperative transcription factors by combining ChIP-chip data and knockout data. *Cell Res* 2010, 20(11):1276-1278.
- Lai FJ, Jhu MH, Chiu CC, Huang YM, Wu WS: Identifying cooperative transcription factors in yeast using multiple data sources. *BMC Systems Biology* 2014, 8 Suppl 5:S2.
- Norel R, Rice JJ, Stolovitzky G: The self-assessment trap: can we all be better than average? *Mol Syst Biol* 2011, 7:537.
- Lai FJ, Chang HT, Huang YM, Wu WS: A comprehensive performance evaluation on the prediction results of existing cooperative transcription factors identification algorithms. *BMC Systems Biology* 2014, 8 Suppl 4:S9.

- Chang DTH, Huang CY, Wu CY, Wu WS: YPA: an integrated repository of promoter features in Saccharomyces cerevisiae. *Nucleic Acids Res* 2011, 39(1):D647-D652.
- 21. Chang DTH, Li WS, Bai YH, Wu WS: YGA: identifying distinct biological features between yeast gene sets. *Gene* 2012, 518(1):26-34.
- Chiu CC, Chan SY, Wang CC, Wu WS: Missing value imputation for microarray data: a comprehensive comparison study and a web tool. BMC Syst Biol 2013, 7 Suppl 6:S12.
- 23. Yang TH, Wang CC, Wang YC, Wu WS: **YTRP: a repository for yeast** transcriptional regulatory pathways. *Database* 2014, bau014.
- Yang TH, Chang HT, Hsiao ESL, Sun JL, Wang CC, Wu HY, Liao PC, Wu WS: iPhos: toolkit to streamline the alkaline phosphatase assisted comprehensive LC-MS phosphorproteome investigation. BMC Bioinformatics 2014, 15(Suppl 16):S10.
- Yang TH, Wang CC, Hung PC, Wu WS: cisMEP: an integrated repository of genomic epigenetic profiles and cis-regulatory modules in Drosophila. BMC Syst Biol 2014, 8(Suppl 4):58.
- Hung PC, Yang TH, Liaw HJ, Wu WS: YNA: an integrative gene mining platform for studying chromatin structure and its regulation in Yeast. BMC Genomics 2014, 15(Suppl 9):S5.
- Stark C, Breitkreutz BJ, Chatr-Aryamontri A, Boucher L, Oughtred R, Livstone MS, Nixon J, Van Auken K, Wang X, Shi X, Reguly T, Rust JM, Winter A, Dolinski K, Tyers M: The BioGRID Interaction Database: 2011 update. Nucleic Acids Res 2011, 39(Database issue):D698-D704.
- Yang H, Nepusz T, Paccanaro A: Improving GO semantic similarity measures using download random walks. *Bioinformatics* 2012, 28(10):1383-1389.
- Balaji S, Babu MM, Iyer LM, Luscombe NM, Aravind L: Comprehensive analysis of combinatorial regulation using the transcriptional regulatory network of yeast. J Mol Biol 2006, 360(1):213-227.
- Hibbs MA, Hess DC, Myers CL, Huttenhower C, Li K, Troyanskaya OG: Exploring the functional landscape of gene expression: directed search of large microarray compendia. *Bioinformatics* 2007, 23(20):2692-2699.
- Abdulrehman D, Monteiro PT, Teixeira MC, Mira NP, Lourenço AB, dos Santos SC, Cabrito TR, Francisco AP, Madeira SC, Aires RS, Oliveira AL, Sá-Correia I, Freitas AT: YEASTRACT: providing a programmatic access to curated transcriptional regulatory associations in Saccharomyces cerevisiae through a web services interface. *Nucleic Acids Res* 2011, 39(Database issue):D136-D140.

#### doi:10.1186/1471-2105-16-S18-S2

**Cite this article as:** Lai *et al.*: PCTFPeval: a web tool for benchmarking newly developed algorithms for predicting cooperative transcription factor pairs in yeast. *BMC Bioinformatics* 2015 16(Suppl 18):S2.

# Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

BioMed Central

Submit your manuscript at www.biomedcentral.com/submit