Discerning Systematic Bias in S. Cervisiae Pathways Using Novel Bayesian Statistics Problem Structuring Methods

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BACKGROUND

One of the major problems in computational biology is developing algorithms that cope with multifunctionality of proteins due to the reuse of biological pathways and components in different contexts. Because of this inherent complexity, new work coping with genome scale problems is potentially pressured to conform to existing biases or limitations present in the literature.

DESCRIPTION OF ALGORITHM

We propose an algorithm that utilizes a novel form of Bayesian problem structuring to correlate overfitting with these aforementioned problems in an ensemble of classifiers attempting to predict gene function based on incomplete or inherently biased expert knowledge. We tested this novel form of Bayesian problem structuring through iteratively removing random genes from the well-studied MAPK pheromone response pathway in S. cerevisiae. We then subsequently used this modified pathway as a training set in our classifier ensemble that is applied to features from freely available heterogeneous data sources. We then performed cross validation, creating features from the normalized results and the amount of overfitting observed in the first round of classification. Classification outputs from this initial ensemble are converted into features and fed into a second “Black Box” classifier that outputs the probability of expert knowledge being inherently biased.

CONCLUSION

Early results indicate that this problem structuring method has potential, based on a small subset of very well studied S. cerevisiae pathways, to predict pathways that may significantly suffer from study biases or lack of complete knowledge.

REFERENCES


Jordan, Michael I. What Are The Open Problems In Bayesian Statistics. ISBA Bulletin Vol 18, Number 1. 2011

Mavrovouniotis ML. Describing Multiple Levels of Abstraction in the Metabolism. ISMB. 1994.


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