Data integration through heterogeneous ensembles for protein function prediction

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Introduction

An exciting aspect of the data revolution in biomedical sciences has been the collection of multiple data modalities reflecting diverse aspects of entities of interest. For example, multiple aspects of proteins, such as amino acid sequences, three-dimensional structures, and protein-protein interactions, are used to study their function¹−³ and other properties. Several data integration strategies have been developed to leverage the inherent diversity and complementarity of these modalities to build comprehensive predictive model(s) for properties of biomedical entities⁴−⁶. However, most of these strategies focus on building uniform integrated representations⁷, such as networks², ⁸−⁹ kernels¹⁰ and tensors¹¹, an approach sometimes referred to as early or intermediate integration⁵, ¹². While such uniform integration is expected to reinforce the consensus among the modalities, this approach may not be ideal when different classes of prediction methods are effective for individual data types, such as deep learning¹³ for medical images¹⁴ and clinical text¹⁵, XGBoost¹⁶ for structured data¹⁷, ¹⁸ and label propagation for networks¹⁹. We refer to dataset- or data type-specific models inferred using these types of methods as local models.

To better utilize the properties of local models, we propose a potentially more effective approach to data integration and predictive modeling by assimilating these models into heterogeneous ensembles²⁰. These ensembles can incorporate a large number and variety of base predictors, including local models, and can benefit from both the consensus and diversity among these predictors. Due to these properties, heterogeneous ensembles have been effective at improving predictive performance for protein function prediction (PFP)²⁰−²², DREAM Challenges²³, and other applications²⁴−²⁶.

Here, we present a novel approach named Ensemble-based data Integration (EI) for predictive modeling. EI leverages established heterogeneous ensemble methods, specifically stacking²⁷ and ensemble selection (ES)²⁸, which are generally applied to base predictors derived from the same data set²⁰. In the data integration scenario, EI uses these methods to aggregate local models inferred from different data sets to develop an ensemble predictor for the target problem (Figure 1). Note that this approach has been referred to as late integration in the literature, although we deploy it more systematically and at a larger scale than in previous work⁴−⁶. We tested EI’s effectiveness for the challenging problem of protein function prediction (PFP)¹−³ using diverse networks from the STRING database²⁹, and compared its performance with data integration algorithms developed for networks³⁰, ³¹.

Materials and Methods
**Ensemble Integration.** Figure 1 shows an overview of the ensemble integration (EI) approach. First, we trained one local model each using ten Weka-based standard binary classification algorithms, such as Support Vector Machine (SVM), Logistic Regression (LR) and Decision Tree (DT), from base predictor training subsets of each of the individual datasets being integrated. We used random under-sampling to balance the positive and negative classes before training these base predictors. Next, to construct the integrated ensemble, we executed a second set of algorithms, namely ES developed by Caruana et al. and Stacking, on the predictions generated by these local models on separate ensemble training subsets of the individual datasets. We used eight standard classification algorithms for Stacking, as in previous work. This whole procedure was conducted within a five-fold nested cross-validation setup typically used for heterogeneous ensemble learning (described in detail in ref. 20), which reduces adverse effects of overfitting while ensuring fair evaluation of the ensemble methods being tested. The source code implementing this EI framework is publicly available.

**Baseline data/predictor integration methods.** We compared EI’s performance with those of the Mashup and deepNF early network integration algorithms. Specifically, we tested the same ten types of classifiers on the Mashup- and deepNF-integrated versions of the individual networks. This baseline’s design enabled a fair comparison with EI, since each approach represent a single level of integration, i.e., early versus late, respectively.

As an alternate baseline, we also tested the integration of base predictors inferred from individual datasets into dataset-specific heterogeneous ensembles. This baseline enabled us to compare the above data integration strategies with the individual datasets being integrated.

In both the above baselines, as in EI, random under-sampling was applied to balance the positive and negative classes before training the base/individual predictors.

**Experimental data.** We tested all the above integration approaches for PFP, specifically GO term annotation prediction from six human STRING datasets/networks. These included protein-protein interactions (PPI), interactions in curated databases, co-expression networks, and genomic neighborhood, co-occurrence and fusion interactions. For consistency, these datasets were exactly the same as those used to evaluate Mashup and deepNF in their respective publications. For all the datasets, both individual and integrated, we used each protein’s adjacency vector as its feature values vector for training and evaluating predictive models. This feature encoding has recently been shown to be effective for network-based gene classification.

Our PFP experiments focused on 112 GO Molecular Function and Biology Process terms that were assigned to at least a thousand human genes each in September 2017 with a non-IEA evidence code. For each GO term, we defined proteins annotated to it as positive examples, and any that were neither annotated to the term, nor its ancestors or descendants, as negative examples. For each term, only proteins that were either positive or negative examples were used in all the experiments. If any of these proteins were not covered by an individual data set, the protein was assigned a corresponding feature vector consisting of all zeros.

We also calculated specific properties of the GO terms considered using the GOATools package (version 0.8.4). The depth of term was defined as the length of the shortest path from the root of the corresponding ontology to , while its information content (IC) is defined as $-\log_{10}(p(\ell))$, where $p(\ell)$ is the probability of a human protein being annotated with $\ell$.

**Evaluation methodology.** In all our assessments, we used the recommended $F_{\text{max}}$ evaluation measure, i.e., the maximum value of the F-measure across all prediction score thresholds. For each GO term, we compared the representative predictors in the following three categories:

- The heterogeneous ensemble algorithm yielding the best-performing EI predictor.
- The best-performing base predictor on the Mashup- and deepNF-integrated datasets.
- The best-performing heterogeneous ensemble algorithm on each individual dataset.
This methodology enables a fair comparison among the representative predictors of the EI and baseline approaches. Finally, we statistically compared the performances of these approaches across all the GO terms tested using the recommended Friedman and Nemenyi tests\(^{41}\).

**Results**

Figure 2 shows the distributions of the \(F_{\text{max}}\) scores obtained by the best implementations of the different integration approaches for the each of GO terms tested. EI achieved the highest median \(F_{\text{max}}\) score overall. It significantly outperformed the best individual predictors on the integrated Mashup (\(p=1.55\times10^{-15}\)) and deepNF (\(p=2.12\times10^{-12}\)) networks, as well the best-performing heterogeneous ensembles on every individual dataset (\(p \leq 9.14\times10^{-8}\)). This improvement is likely due to the ability of EI to encapsulate local information in the individual datasets into the most effective base predictors before their assimilation into heterogeneous ensembles. It is also notable that the heterogeneous ensembles on the individual PPI datasets performed slightly better than predictors from the Mashup and deepNF networks (\(p=0.008\) and 0.092, respectively), reaffirming the utility of these ensembles for predictive modeling.

It is well-known that the performance of PFP algorithms can vary substantially across GO terms depending on their information content (IC) (inversely related to its size, i.e., the number of proteins annotated to a term) and depth in the respective GO hierarchy (specificity of the term)\(^{40,42}\). Thus, we also compared EI, deepNF and Mashup, the main data integration strategies evaluated in this study, with regard to how their performance varied with these properties of GO terms. In terms of both these properties, EI consistently performed better than both deepNF and Mashup at increasing depths (Figure 3(A)) and levels of IC (Figure 3(B)). These trends indicate
that EI can perform well even for GO terms that are deeper or have fewer annotated genes than terms near the root of the hierarchy. This is again likely due to its ability to capture and aggregate information local to individual datasets, which is not enabled sufficiently by early integration.

**Discussion**

Here, we proposed a novel approach to late data integration for predictive modeling, namely Ensemble Integration (EI), where local models derived from individual datasets are integrated into heterogeneous ensembles to develop predictive models. Through the problem of protein function prediction, we demonstrated that EI performs better than early or no data integration approaches.

This proof of principle study has limitations, which offer several avenues for future work. Foremost, our assessments only included a relatively small number of GO terms with at least a thousand human genes annotated to each of them. Developing EI methods for more specific GO terms and other sparse labels, which have many fewer positive examples, is an important challenge. Moreover, even though the individual STRING datasets considered in this study were natively structured as networks, we used their adjacency matrices as sources of feature vectors for traditional classification algorithms such as SVM, LR and DT. Although this feature-based representation has been recently shown to be more effective for (gene) classification\(^ {37}\), this assessment may need to be re-evaluated in the context of network integration. Additionally, in the current implementation of EI, we assigned all-zero feature vectors to proteins disconnected from an individual network. In future work, it will be important to investigate other strategies for handling such missing values. It is also important to repeat our assessments for different subsets of individual STRING networks and GO evidence codes to eliminate non-obvious circularities between the two data sources. Finally, it will also be useful to study the mathematical properties of EI that led to its improved performance over other data integration algorithms in our experiments, as well as the methodologies to interpret the ensembles EI yields.

**Acknowledgements**

This work was supported by NIH grant R01GM114434, an IBM faculty award, and IARPA, under Cooperative Agreement Number [W911NF-17-2-0105]. It was also enabled by Scientific Computing resources at Mount Sinai. We thank colleagues in the FunGCAT/IGACAT team and at Mount Sinai for discussions and suggestions. The views and conclusions herein do not represent those of the U.S. Government, which is authorized to reproduce and distribute reprints notwithstanding any copyright annotation thereon.
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