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Predict Remote Homology Proteins by Merging Grey Incidence Analysis and Domain Similarity Analysis

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ABSTRACT

Protein remote homology detection is a challenging problem for drug development. Although there are a couple of methods to deal with this problem, the benchmark datasets based on which the existing models were trained and tested contained many high homologous samples due to the fact that the cutoff threshold was set at 95%. In this study, we reconstructed the benchmark dataset by setting the threshold at 40%, meaning none of the proteins included has more than 40% pairwise sequence identity with any other. Using the new benchmark dataset, we proposed a new method called PHom-GRADSI to detect the remote homologous proteins by integrating various ranking approaches via grey incidence analysis and function domain similarity index. Rigorous cross-validations have indicated that the new predictor is superior to its counterparts in both enhancing successes rates and reducing computational cost. The predictor can be download from https://github.com/jcilwz/RemoteHomology/tree/master/program.

1. INTRODUCTION

Detecting remote homology relationship among proteins plays one of the fundamental and central roles in computational proteomics. It is particularly useful for drug development (see, e.g.,[1, 2]). With the development of sequencing techniques, the protein sequence data rapidly raise. To find those proteins structure and function is more and more urgent. Although X-ray crystallography is a powerful tool in determining protein 3D structures, it is time-consuming and expensive. Particularly, not all proteins can be successfully crystallized, particularly for membrane proteins. The NMR technique is indeed a very powerful tool in determining the 3D structures for membrane proteins as indicated by a series of recent publications (see, e.g., [3-7]), it is time-consuming and costly. To acquire the structural information in a timely manner, one has to resort to various structural bioinformatics tools based on the sequence similarity principle (see, e.g., [8]). Unfortunately, such principle cannot cover the cases of remote homology proteins. In view of this, considerable efforts [9-14] have been made to detect remote homology proteins.

Although these methods each had their own merits and did play stimulating role in this area, further work is needed. Firstly, the benchmark datasets used in their studies had high similarity. For instance, the benchmark dataset in [9, 12] contains 7329 proteins from 1070 different super families, with pairwise sequence identity cutoff set at 95%. In other words, it would allow those proteins with higher than 80% similarity in the data set. Secondly, the ranking algorithm used in those studies would spend a lot of time to training the learning model. For example, if the training dataset has N proteins, the LambdaMART need to deal with N^2 proteins pair samples.

The present study was initiated to address the two problems with the aim to develop a more powerful method in this regard.

2. MATERIALS AND METHOD

2.1 Benchmark Dataset

According to Chou's 5-step rules [15] that have been widely and increasingly used by many investigators (see, e.g., [16-32]), the first prerequisite in establishing a new predictor is to construct or select an effective benchmark dataset.

In this study, the benchmark dataset was taken from Liu et al. [12]. It included 7329 proteins from 1070 different super families and 1824 families derived from SCOP database. To reduce the redundancy and homology bias, the program CD-HIT[33] was adopted to cut down those proteins that had ≥40% pairwise sequence identity to any other in the dataset. Furthermore we removed those families that just had one protein sequence. Finally, we obtained 3128 proteins from 540 super-families and 777 families.

2.2 Sample Similarity Analysis

2.2.1 Grey Incidence Analysis of proteins formulated by Grey-PSSM

Given a protein with L amino acid residues, it is usually expressed by

$$\mathbf{P} = \mathrm{R}_1\mathrm{R}_2\mathrm{R}_3\cdots\mathrm{R}_i\cdots\mathrm{R}_L \tag{1}$$

where R_i $(i = 1, 2, \cdots, L)$ is the i-th residue in the protein. Since all the existing machine-learning algorithms can only handle vector but not sequence samples [34], one has to convert Eq.1 into a vector model. But a biological sequence expressed as a vector in the discrete framework may lose all the sequence-order or pattern information.

To avoid completely losing this kind of information for proteins, the pseudo amino acid composition (PseAAC)[35, 36] was proposed. Ever since the concept of Chou's PseAAC was proposed, it has been widely used in nearly all the areas of computational proteomics (see, e.g., [37-44] as well as a long list of references cited in[45, 46]). According to the general PseAAC[15], the protein of Eq.1 can be formulated as

$$\mathbf{P} = [\Psi_1 \ \ \Psi_2 \ \ \cdots \ \ \Psi_u \ \ \cdots \ \ \Psi_\Omega]^{\mathbf{T}} \tag{2}$$

where $\mathbf{T}$ is the transposing operator, the subscript Ω is an integer, and its value and the components Ψ_u $(u = 1, 2, \cdots)$ will depend on how to extract the desired features and properties from the protein sequence.

In this study, the model, Grey-PSSM proposed by Lin[47, 48], is adopted. It extracted the sequential evolution information by the Position Specific Scoring Matrix (PSSM). For the concrete procedures, refer to the original papers [47, 48].

After the Grey-PSSM treatment, we have finally got a 60-D PseKNC vector for Eq.2; i.e., its subscript parameter $\Omega = 60$ and each of the 60 components therein has been uniquely defined.

Assume

$$\mathbb{S} = \{P_1, P_2, \cdots, P_N\} \tag{3}$$

are the set of protein samples, and $P_i (1 \le i \le N)$ is the i^{th} protein. According to Equals [6]~[11] in Ref. [49], the distance $\Gamma(P_i, P_j)$ is defined as the grey incidence degree between P_i and P_j. The larger the value of $\Gamma(P_i, P_j)$, the more similar they are.

2.2.2 Domain Similarity Analysis

There are also other models used to formulating proteins excepte for the PseAAC. Here, we propose a noval mothed to discribe the proteins. For a protein $P_i \in \mathbb{S}$, we discribe its functional domains set by the following steps.

Step 1, $\mathbb{S}_i^{homo}$ is the homology set of protein P_i and it is extracted by searching against UniProt release 2018_08 Swiss-Prot FASTA format flatfile by HMMER[50-52]. We just use the top 10 sequences if the search results have

more than 10 sequences. Therefore there are at most 10 proteins in $\mathbb{S}_i^{homo}$.

Step 2, for a protein in $\mathbb{S}_i^{homo}$, $h_k^i \in \mathbb{S}_i^{homo}$ $(1 \le k \le 10)$, we annotate its functional domains by running hmmscan program against Pfam-A database (Pfam release 32.0). The Pfam-A includes 17,929 functional domains and 688 clans. We define the sets $\mathbb{F}$ and $\mathbb{C}$ as follwing.

$$\mathbb{F} = \{f_1, f_2, \cdots, f_{17929}\}$$
$$\mathbb{C} = \{c_1, c_2, \cdots, c_{688}\} \quad (3)$$

where f_i $(1 \le \mathrm{i} \le 17929)$ denote the i-th functional domain in $\mathbb{F}$ and c_i $(1 \le i \le 688)$ the i-th clan in $\mathbb{C}$. Some functional domains have same clan. For example, the domains of "PF15884" and "PF17050" have the same clan "CL0683". Therefore, the functional domains set of protein h_k^i, the k-th homology protein of protein P_i, is expressed as a set

$$D_k^i = \{f_i | f_i \in \mathbb{F}\} \quad (4)$$

It is mean that all functional domains of h_k^i constitute the set D_k^i.

Step 3, the protein P_i is expressed as a domains set

$$D_i = \bigcup_{k=1}^{10} D_k^i \quad (5)$$

D_i is a set which is unioned together the functional domain set of each homology protein of the protein P_i. We define D_i as the functional domain set formulation of P_i.

As aboved steps, a protein is expressed a set including some functional domains from Pfam-A. For the proteins in same family or clan have similar functional domains, new distance between two proteins, named as Domain Similarity Index (DSI), can be defined based on the functional domains.

The algorithm of distance between P_i, P_j is discribed as follows.

1) If $D_i \cap D_j \neq \emptyset$, $\mathrm{DSI}(P_i, P_j) = \frac{|D_i \cap D_j|}{|D_i \cup D_j|}$

2) Else If $D_i \cap D_j = \emptyset$

We define $Clan_i$ to denote the clans set of P_i. $Clan_i$ includes the clans of each element in D_i

2.1) If $Clan_i \cap Clan_j \neq \emptyset$, $\mathrm{DSI}(P_i, P_j) = F$. F is a constant and in this study F is equal to 0.2.

2.2) Else, $\mathrm{DSI}(P_i, P_j) = 0$

Where, $|\bullet|$ means the count of set, $\cap$ is the intersection opertator of two sets, and $\cup$ is the union operator of two sets. From above discription, we have $0 \le \mathrm{DSI}(P_i, P_j) \le 1$ and the larger the $\mathrm{DSI}(P_i, P_j)$, the more similarity they are.

2.3 Operation Engine or Algorithm

In this study, the Grey Relational Analysis [53, 54] and the Domain Similarity Index was utilized to rank the relationship of proteins. Given a query protein, the system will search it against the benchmark dataset and return the top ranked proteins. The predictor thus formed is called "PHom-GRADSI". Illustrated in **Figure 1** is a flowchart to show how the proposed predictor is working. In this paper, w(1) and w(2) are equal to 0.5.

3. RESULT AND DISCUSSION

The jackknife test is deemed the least arbitrary and most objective among three cross-validation methods: independent dataset test, K-fold cross-validation test and jackknife test [55]. Because the LambdaMART ranking algorithm used in preview studies [9, 12] consumed more training time and computer memory, as a compromise the 5-fold cross-validation test was adopted there. Now, we employed GRA and DSI to compute the relationship score between the query protein and benchmark dataset proteins, significantly reducing the computing time and memory. Therefore it would be feasible to use the most rigorous jackknife test to examine the prediction quality. The outcome thus obtained are given in **Table 1**, where we can see that PHom-GRADSI achieved the best performance in both the score of ROC1 and the score of ROC50.

In the same time, we used the Jaccard Index (JI) to calculate the similarity of the proteins. The JI is used to measure the similarity of two sets A and B. It is defined as following:

$$\mathrm{JI(A,B)} = \frac{|A \cap B|}{|A \cup B|}$$

The proteins were formulated as the functional domain set as Eq.(4). In the method of Jaccard Index, the score of ROC and ROC50 were 0.8196 and 0.8070 respectively. In contrast to this, the score of ROC and ROC50 achieved by DSI were 0.9053 and 0.8454 respectively. It can be concluded that DSI preforms better than Jaccard Index. The main reason is that some different domains have same clan. If there are not same elements in two proteins' functional domain set, the distance of Jaccard Index is zero, but the distance of DSI is greater zero. It is evident that the non zero distance of these proteins is more reasonable because their some functional domain have same clan.

Because not all proteins can be finded their homology protins, these proteins' functional domain set formulation is empty set (see Eq.(4)). For example, there are 23 protiens who can not be formulated as functional domain set in the **Benchmark Dataset**. The distences between these proteins and other proteins are zero according to the definition of JI and DSI . So we can not distinguish the similarity of these proteins who have no homology protiens. This situation is the worst failure DSI. In order to overcome the failure we merged GIA and Jaccard Index and merging GIA and DSI in predicting, respectively. From the

Tabel 1, it is showed that the values of ROC1 and ROC50 are improved at least 0.04 after merging GIA into Jaccard Index or DSI. We can draw the grey incidence analysis can compensate for the worst failure of DSI.

4. CONCLUSION

Protein remote homology detection is vital for studying protein structures and functions. It is anticipated that the proposed method may become a useful high throughput toll for both basic research and drug design. In this study, a noval method DSI is proposed. It discribes a protein as a functional domain set and measure the distance of two proteins by comparing two proteins' functional domain set similarity. The work testifies the DSI method is effective. This method formulating proteins and calculating distance between proteins may be used in other fields of predicting protein function or structure. We deliver programs of this noval method in https://github.com/jcilwz/RemoteHomology/tree/master/program. Everyone can download programs from this website and the usage is described in ReadMe.txt.

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FIGURE LEGENDS

Figure 1. A flowchart to illustrate how the proposed predictor is working.

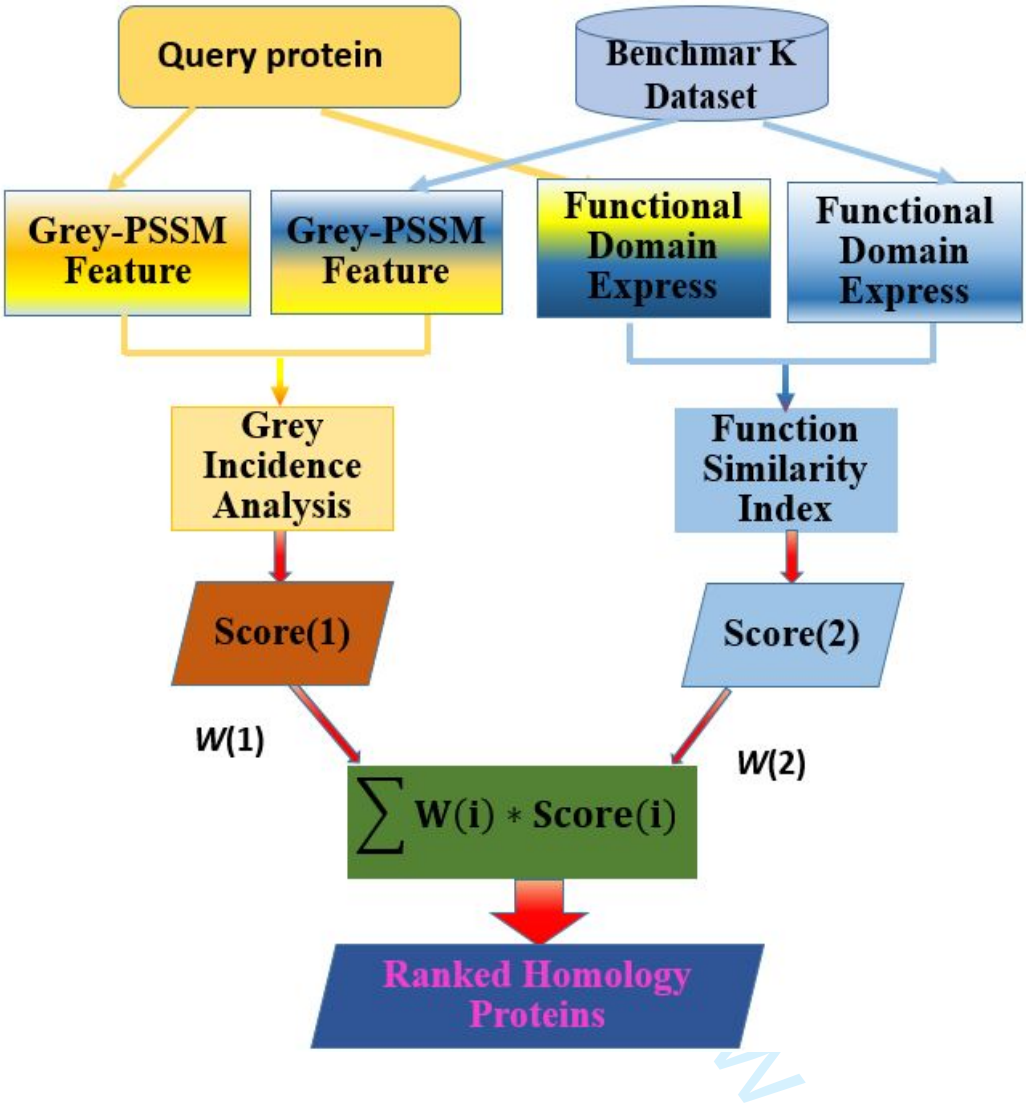

Table 1. A comparison of the jackknife test results for protein remote homology detection on the benchmark dataset

Methods	ROC1	ROC50
PSI-BLAST	0.7113	0.7647
GRA (Grey-PSSM)	0.8937	0.7149
Jaccard Index	0.8196	0.8070
Domain Similarity Index (DSI)	0.9053	0.8454
GRA and Jaccard Index	0.9301	0.8533
PHom-GRADSI	**0.9620**	**0.8861**

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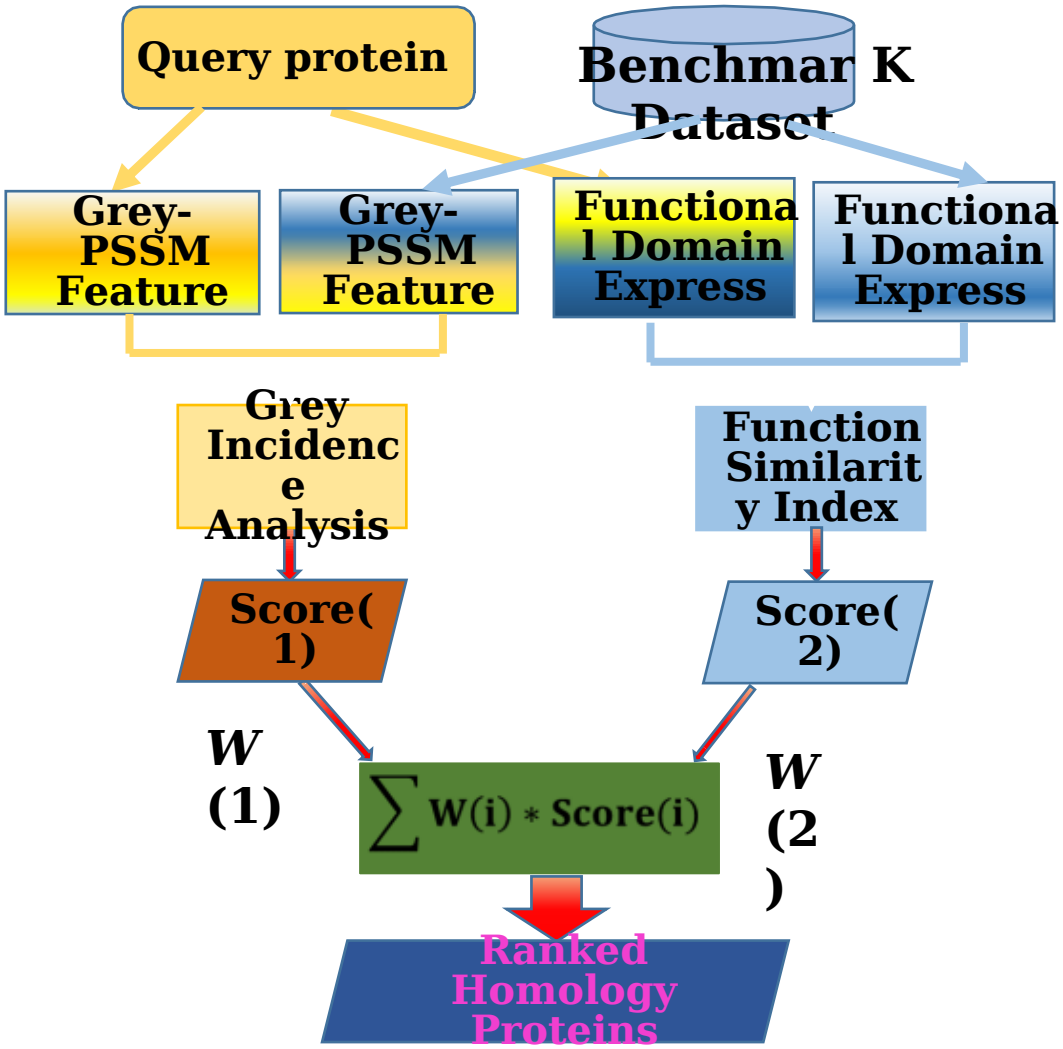
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Benchmar K Dataset
Grey-PSSM Feature
Grey-PSSM Feature
Functiona l Domain Express
Functiona l Domain Express
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Function Similarit y Index
Score(1)
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W (1)
∑ W(i) * Score(i)
W (2)
Ranked Homology Proteins