**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/dRHP-GreyFun.

KEYWORDS: Remote homology prediction; Function domains; Grey incidence analysis; Grey-PSSM

**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](#_ENREF_1), [2](#_ENREF_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](#_ENREF_3)]), 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](#_ENREF_8)]). Unfortunately, such principle cannot cover the cases of remote homology proteins. In view of this, considerable efforts [[9-17](#_ENREF_9)] 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](#_ENREF_9), [12](#_ENREF_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. This also happened in the benchmark datasets in [[15](#_ENREF_15)]. Secondly, the ranking algorithm used in [[12](#_ENREF_12), [17](#_ENREF_17)] 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 N2 proteins pair samples. Thirdly, the ProDec-BLSTM[[15](#_ENREF_15)] predictor, which onverted the protein sequence into a representing matrix by PSSM ( position specific scoring matrix) and fed it into the net input layer, got the best performance, but not all proteins could be formulated by PSSM.

The present study was initiated to address the three 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 [[18](#_ENREF_18)] that have been widely and increasingly used by many investigators, 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](#_ENREF_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[[19](#_ENREF_19)] 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[[20](#_ENREF_20)].

**2.2 Grey Incidence Analysis (GIA) of proteins formulated by Grey-PseAAC**

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

|  |  |
| --- | --- |
|  | (1) |

where is the *i*-th residue in the protein. Since all the existing machine-learning algorithms can only handle vector but not sequence samples [[21](#_ENREF_21)], 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)[[22](#_ENREF_22), [23](#_ENREF_23)] 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., [[24-31](#_ENREF_24)] as well as a long list of references cited in [[32](#_ENREF_32), [33](#_ENREF_33)]). According to the general PseAAC[[18](#_ENREF_18)], the protein of Eq.1 can be formulated as

|  |  |
| --- | --- |
|  | (2) |

where **T** is the transposing operator, the subscript is an integer, and its value and the components will depend on how to extract the desired features and properties from the protein sequence.

In this study, the model “Grey-PseAAC” proposed by Lin[[34](#_ENREF_34)] is adopted. It fristly codes the amino acid by using the five physical-chemical properties: hydrophobicity, pk1 (, pk2 (NH3), PI () and molecular weight. Then the sequential intrinsic information are extracted by the grey model GM(1,1)[[35](#_ENREF_35)]. For the concrete procedures, refer to the original papers[[34](#_ENREF_34)].

After the Grey-PseAAC treatment, we have finally got a 30-D PseAAC vector for Eq.2; i.e., its subscript parameter and each of the 30 components therein has been uniquely defined.

Assume

(3)

are the set of protein samples, and is the protein. According to Equals [6]~[11] in Ref. [[36](#_ENREF_36)], the distance is defined as the grey incidence degree between and . The larger the value of , the more similar they are.

**2.3 Domain Similarity Analysis**

There are also other models used to formulating proteins except for the PseAAC. Here, we propose a novel mothed to describe the proteins. For a protein , we discribe its functional domains set by the following steps.

Step 1,  is the homology set of protein and it is extracted by searching against UniProt release 2018\_08 Swiss-Prot FASTA format flatfile by HMMER [[37-39](#_ENREF_37)]. We just use the top 10 sequences if the search results have more than 10 sequences. Therefore there are at most 10 proteins in .

Step 2, for a protein in ,, 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 and as follwing.

(3)

where denote the i-th functional domain in and the i-th clan in . 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 , the k-th homology protein of protein , is expressed as a set

{} (4)

It is mean that all functional domains of constitute the set .

Step 3, the protein is expressed as a domains set

(5)

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

Now a protein is expressed as 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, is discribed as follows.

1) If,

2) Else If

We define to denote the clans set of . includes the clans of each element in

2.1) If , . F is a constant and in this study F is equal to 0.2.

2.2) Else,

Where, |•| means the count of set, ∩ is the intersection operator of two sets, and ∪is the union operator of two sets. From above description, we have and the larger the , the more similarity they are.

**2.4 Operation Engine or Algorithm**

In this study, the Grey Relational Analysis [[35](#_ENREF_35), [40](#_ENREF_40)] 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. Results 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 [[41](#_ENREF_41)]. Because the LambdaMART ranking algorithm used in preview studies [[9](#_ENREF_9), [12](#_ENREF_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,

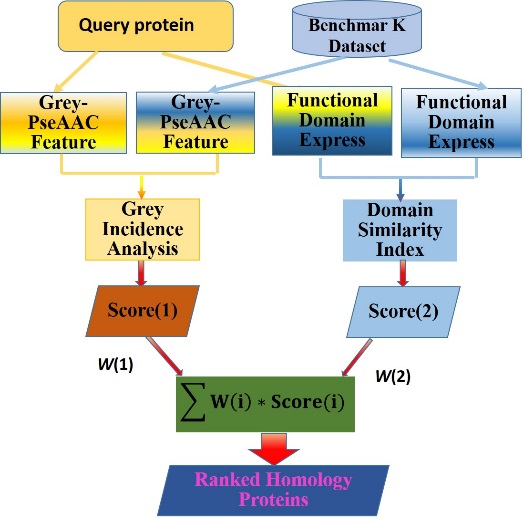


Figure 1. A flowchart to illustrate how the proposed predictor is working

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.

**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 |
| GIA (Grey-PSSM) | 0.8937 | 0.7149 |
| Jaccard Index | 0.8196 | 0.8070 |
| Domain Similarity Index (DSI) | 0.9053 | 0.8454 |
| GIA and Jaccard Index | 0.9301 | 0.8533 |
| PSI-BLAST+DSI | 0.9277 | 0.8791 |
| **PHom-GRADSI (GIA+DSI)** | **0.9620** | **0.8861** |

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:

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 domains have same clan.

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

Moreover, for the 23 proteins whose functional domains sets are empty, we predicted their remote homology families by BLAST and GIA of Grey-PseAAC. The **table 2** showed that GIA got better performance than BLAST. We can draw that the grey incidence analysis of Grey-PseAAC can compensate for the worst failure of DSI.

**Table 2**

A comparison of the test result for 23 proteins whose functional domains set are empty again the benchmark dataset

|  |  |  |
| --- | --- | --- |
| **Methods** | **ROC1** | **ROC50** |
| PSI-BLAST | 0.7131 | 0.5065 |
| GIA (Grey-PseAAC) | 0.8895 | 0.5486 |

**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 novel method DSI is proposed. It describes 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 novel method in https://github.com/jcilwz/dRHP-GreyFun. Everyone can download programs from this website and the usage is described in ReadMe.txt.

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