

The Algorithms in Disarray Hashing Technique That Preserves Gene Ontology Hierarchy Application of the Phish Methodology

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Abstract—Gene Ontology (GO) is a systematic taxonomy that employs organized languages to explain the biochemical activities, functional activities, including tissue placements underlying transcription factors. Gene Ontology (GO) markers link proteins to GO concepts as well as show which genetic variants perform the physiological activity specified by the words. Forecasting proper GO descriptors for individuals from a large number of GO words specified by GO, on the other hand, would be a challenging task. To address these issues, researchers present a conceptual approach for functional genomics predictions involved in Gene Ontology Hierarchical Preserving Hashing (Phish). Phish begins by comparing the taxonomy homology of GO keywords. It all then optimizes a succession of hashed algorithms to store huge GO concepts via a small binary format, using a hierarchical structure scrambling method to maintain the hierarchical system between GO words. Following that, Phish uses those scrambling techniques to transfer the gene-term connection matrices into a cheaper region, where it conducts lexical resemblance gene function estimation. Phish outperforms existing similar methods as well as being resistant to the number of encryption schemes, according to experimental observations on 3 model organisms for intercultural gene function forecasting. In particular, researchers use Phish as a BLAST-based gene function forecast component. According to the findings of the experiments, Phish greatly enhances forecasting accuracy.

Keywords: Gene Ontology; Hierarchy Preserving Hashing; gene function; Phish Technology

I. INTRODUCTION

Most biological activities, including those of metabolic, hormone control, including cell signaling, rely on genetic variants, including such enzymes including RNAs [1]. Medication research, illness research, gene collection expression values, as well as other disciplines, could benefit from detailed information on all these genetic variants. Moreover, because detecting the gene functions using wet-lab approaches was labor-intensive, costly, as well as limited bandwidth, operational classifications of genetic variants lagged substantially while behind the speed of acquired genetic information [2-4]. Moreover, the operational classifications of genomes were influenced by biologists' research goals, including experimentation morals

including people and animals. As a result, developing mathematical algorithms to effectively as well as accurately identify genetic activities was highly necessary. GO was created as a collaborative attempt to enable computerized gene function forecasting as well as to organize biological material of genes as well as their metabolites in a formalized as well as uniform manner among genomic information [5]. The Gene Ontology (GO) would be a collection of concepts as well as a foundation for describing the activities of genetic variants across all species. The GO research was made up of two parts: the Ontology, which uses a directed acyclic graph to organize GO words, hierarchical structures, and goes markers, which link proteins but rather gene products to GO concepts [6].

A GO word in the DAG denotes higher specific operational information than its progenitor words. If a GO word was associated with a gene, its ancestor phrases also were associated with that gene, but not another way around. On the other side, if a word must not be attributed to a gene, then neither would its descendent words [7]. In the field of gene function, structural similarity-based techniques have been intensively researched. Those approaches were predicated on the fact that semantic relatedness produced from gene GO classifications correlates with familiarity estimated from a variety of biological information, including such sequencing, gene expression patterns, as well as posttranslational modifications [8]. These primarily aim to fill up gaps in GO classifications imperfectly identified genes [9]. The structures of GO descriptors could be designed to estimate gene function using a tree structure or perhaps a Bayes classifier coupled to the data. However, to effectively estimate the likelihood function between GO words, such a template method requires enough annotations [10].

II. RELATED WORKS

As a result, it is incompatible with sparse words assigned to less than 30 genes. In practice, the majority of GO words were sparse words; with various GO words having varying amounts of associated genes, as well as the amounts are quite unbalanced. Because sparse words convey

a higher precise basic understanding than their parent phrases, which have been linked to many more genes, it's preferable to forecast the future relationship among sparse words as well as genetic mutations. To consider explicitly, researchers employed the GO structure to calculate the relationship between GO words as well as put the results into a gene function predictions algorithm [11]. Their research shows that integrating the relationship would increase forecasting accuracy greatly, particularly for sparse GO keywords. Many gene GO classifications were imperfect, and fresh GO classifications were added to those same proteins on a constant schedule. To predict empty gene GO annotations, a randomized walking systematic model has been used [12]. It must be motivated by the finding that the GO words refilled to a gene usually match the grandchildren of the words previously ascribed to the genes. That approach also generates additional labels for a gene depending on the characteristics of its conceptual surrounding genes as well as the predicted identifiers [13] to take the benefit of annotation from other genes. Just GO evaluations of genes from the very same genus have been used to evaluate those linguistic resemblance alternatives. Because God would be the genus and therefore can tag genetic variation, species with the same GO concepts, the semantic relationship among genes across various organisms may also be assessed [14].

These investigations reveal that GO comments from separate species were complementing to be together, but also that the forecast performance improvement was clear for two varieties with strong homology; however, the increase was insignificant for two varieties having poor homologous recombination. The abovementioned conceptual techniques can't effectively quantify the semantic relatedness among genes having regard to huge GO keywords based on partial as well as sparse GO classifications [15]. Large GO keywords also are a problem for gene function forecasting models that combine heterogeneous biological information. The GO hierarchy could be utilized to make reliable forecasts as well as increase forecasting accuracy dramatically. These GO words reduction approaches essentially use the GO hierarchy largely represented by the gene-term interaction matrices explicitly as well as convert the GO DAG into an undirected one to make management sets. As a result, those who do not follow the GO structure well enough, negatively affected [16]. Humans present a computer methodology based on GO Phish for functional genomics estimation to overcome the difficulties with existing GO words pressure methods. The fundamental goal of Phish would be to solve the problem of accurately guessing gene-to-massive-GO-term correlations. Phish could not only retain the hierarchy system areas And provide words in a cheap environment but rather the taxonomic homology between GO words, increasing the accuracy of gene function forecasting in that storage.

III. PROPOSED METHODS

To evaluate taxonomic resemblance among words as well as preserve the GO hierarchy, Phish initially utilizes a meaningful biological matching score. Then it uses a ranked team scrambling method to optimize a set of scrambling

algorithms that could keep GO words' similarities as well as their structures of power. The gene-term correlation matrices were subsequently compressed into a low-dimensional one using these cryptographic techniques. Following that, Phish uses the compressing gene-term interaction matrix to quantify semantic relatedness among genes as well as guesses activities of a gene depending on descriptions of its semantical neighbors. Those forecasts then were projected return to the initial GO words universe, resulting in the formation of gene-to-massive GO term relationships.

The goal of scrambling would be to learn information as well as task-specific hashing algorithm that produce compressed digital signals while maintaining the original information's comparable associations. Phish, on the other hand, seeks to represent huge GO keywords using compressed binary digits that maintain both the taxonomic identity and the structures of power among those. Researchers employ Lin's resemblance throughout this research, which would be a sample taxonomic measurement extensively used during hierarchical Ontology. To accomplish a purpose, researchers utilize a well-known metric called Standardized Discount Cumulative Benefit to assess the integrity of the hierarchical ordering leaderboard:

$$NDCG = \frac{1}{L_x} \int_{y=1}^n + \frac{2^{k_{xy}} - 1}{\log_2(1 + y)} = \frac{1}{L_y} \int_{y=1}^n + \frac{2^{k_{xy}} - 1}{\log_2(1 + s_x^y)} \quad (1)$$

$$s_x^y = 1 + \int_{v=1}^n + sgn(Ham(t_x, t_y) > Ham(t_x, t_v)) \\ = 1 + \int_{v=1}^n + sgn(t_x^M(t_v, t_y) > 0) \quad (2)$$

Humans could see from the approximations in Equations (1) & (2) That NDCG favors phrases having smaller ranking commands over words with big rating values. As a result, rather than include all my words, NDCG was frequently cut at a specific order of popularity. Because the deepest point of branch words in the GO structure was currently 15, q could essentially be set to 15. The bigger the NDCG number, the further the scrambling algorithms coincide with the priority order, according to the specification of NDCG. Whenever the Distance measure between GO words generated respective individual hashes was entirely consistent with hierarchy structure as well as mutual information among tea as well as other words, the maximum NDCG score was reached. By maximizing the NDCG measurement, the desired scrambling algorithms could guarantee that the scrambling algorithms were compatible with the taxonomy resemblance of GO words, as well as preserving the relative ranking of GO aspects. The NDCG across m GO periods was maximized as follows, depending on the above order to prepare:

$$B(T) = \int_{x=1}^n + \frac{1}{L_x} \int_{y=1}^n + \frac{2^{k_{xy}} - 1}{\log_2(1 + s_x^y)} \quad (3)$$

$$s_x^y = 1 + \int_{v=1}^n + sgn(t_x^T(t_v, t_y) > 0) \quad (4)$$

IV. RESULTS AND DISCUSSIONS

To assess the effectiveness of Phish-based cross-species gene function forecasting, researchers performance management process in a medieval to current manner. The previous comments seem to be the GO annotations of 3 model organisms completed on 2020-05-07, while the latest comments seem to be the GO evaluations completed on 2021-09-01. Researchers use previous identifiers to educate Phish as well as create gene function predictions, as well as then we could use contemporary identifiers to confirm the predictions. Humans obtained the historical and current GO classifications from Gene Ontology simultaneously, as well as the current ontology documents. 2 GO adheres to the standard of annotating genetic instructions at every genus database to some of the most relevant and comprehensive levels in the ontologism that accurately reflects the biology of the gene product; straight comments were interpretations that have been available immediately in the reference documents. Humans additionally annotation all the parent words of direct connections of a gene to the very same gene using the True Path Rule. Humans exclusively utilize words accessible, including both preserved Ontology data during trials to minimize the influence of GO modification, as performed in CAFA.

V. COMPARATIVE EXAMINATIONS

Baseline as well as Baseline explicitly calculates the semantic relatedness among genes in the same GO words field using the Based feature selection as well as a Cosine measurement approach, and afterward, utilize a comparable KNN classifier as Hash GO as well as InterGFP for functional genomics estimation. ClusDCA seems to be a variation of Cloudscape that employs SVD to condense the Gene Ontology mapping function into a cheap environment before doing gene function estimation in the reduced area, similar to HPHash as well as Hash GO.hash seems to be a variation of Phish that seeks cryptographic techniques immediately using a clustering algorithm of GO instead of using Lin's resemblance. Because the correlation between a nongenetic as well as other genes was 0, those approaches cannot identify activities for genes which annotation was wholly unexplained.

As a result, researchers confirm Plash's efficacy in sequencing information functional genomic forecasting, which may also generate predictions for genes that seem to be essentially annotated. Researchers use Phish to condense the wide gene-term correlation matrices into a cheap version, as well as BLAST to calculate sequence identity across genes, before estimating the model depending on the collapsed relationship matrices as well as pattern similarities. Humans utilize Rank Loss rather than RankingLoss to keep these things comparable to the rest of the assessment criteria. Throughout this approach, the greater the efficiency, the greater the value of all of the other evaluation criteria. Humans select the most suitable q conditions with the biggest probabilities as the appropriate words of a gene, as well as q has been equivalent to the expected amount of captions of a gene, as completed in an earlier study, as well as humans recognize the top q

conditions with the biggest probabilities as the appropriate words of a gene, as completed in a past analysis.

Some measures appear to be related simply to the projected numerical connection matrices. Because all these measures assess gene function predictive performance from either a variety of perspectives, it's challenging for one technique to regularly beat others throughout all measures. To compare as well as quantify the efficacy of refilling GO descriptions of largely characterized genes, researchers perform research on GO classifications of 3 model organisms stored in various decades.

TABLE 1. PREDICTION ON ARCHIVED GO ANNOTATION

Empty Cell	Empty Cell	(Ham)	(Cos)	(BMA)	(G)	Empty Cell	(G)	Empty Cell
MicroAvgF1	BP	1.326	5.8278	1.3267	1.322	1.3306	1.3178	1.3571
MF		1.326	1.3782	5.869	1.3642	1.3683	1.3615	1.4065
CC		1.3696	1.3496	5.8251	1.3399	1.3487	1.3209	1.3651
MacroAvgF1	BP	1.3252	5.8479	1.3341	1.3174	1.3486	1.3275	1.3653
MF		1.332	1.362	5.8455	1.3433	1.3593	1.3389	1.3891
CC		1.2798	1.3324	5.8063	1.3155	1.303	1.2909	1.3501
AvgAUC	BP	1.4324	5.9209	1.4267	1.4319	1.433	1.4276	1.4384
MF		1.4326	1.4181	5.9277	1.4325	1.4299	1.426	1.4516
CC		1.4243	1.4078	5.9156	1.4234	1.4253	1.4236	1.4345
Fmax	BP	1.33	5.8558	1.3569	1.3607	1.372	1.3634	1.4049
MF		1.4082	1.4288	5.8679	1.4208	1.4223	1.4223	1.4463
CC		1.3831	1.4018	5.8639	1.3808	1.389	1.3891	1.4194
1 - RankLoss	BP	1.3451	5.8851	1.3537	1.3824	1.3906	1.3534	1.4558
MF		1.3892	1.4308	5.8906	1.4136	1.4173	1.4059	1.4754
CC		1.3499	1.4054	5.842	1.3967	1.4001	1.3827	1.4555
AvgPrecision	BP	1.2902	5.7972	1.2912	1.283	1.2941	1.2791	1.3542
MF		1.3677	1.3945	5.8562	1.3624	1.3662	1.3607	1.4256
CC		1.3208	1.3608	5.8147	1.3295	1.3417	1.3092	1.3788
MCC	BP	1.3365	5.8592	1.3446	1.3308	1.3596	1.3405	1.375
MF		1.3426	1.3726	5.8557	1.3546	1.3699	1.3515	1.3977
CC		1.2903	1.3422	5.8158	1.3149	1.3262	1.3044	1.36
Smin ↓	BP	20.79	24.3039	14.0866	8.1719	7.7425	12.9943	5.8751
MF		2.3713	2.4523	7.2884	2.2527	2.2134	2.2134	1.8398
CC		3.0564	3.0403	7.6051	3.0542	3.1048	3.0254	2.693

The revised GO classifications of every species stored in 2021 have been used to verify the predictions, and go classifications of these creatures stored in 2020 were utilized to build gene function, forecasting models.

The neighborhood size k is set to 300; the password bits duration d was adjusted to 60, the parameters were set to 0.1, as well as the variable was adjusted to 0.01. Those settings have been set as indicated by the researchers in the primary documents. The neighborhood, as well as dimension parameters for the methods based, are much like before. Table 1 of the Support and contribution contains the documented experimental data among those contrasting approaches. The boldface information within every column among those figures seems to be the best possible result from multiple methodologies of comparative. Across all 3 species, including across practically all assessment methods,

Phish beats alternative comparative approaches in identifying gene function, as shown in the following figure. Hash GO, similar to Phish, utilizes network cryptographic algorithms to compress the gene-term connection matrices into a low-dimensional another before doing semantically resemblance gene function forecasting; however, it often falls short of Phish.

These findings demonstrate that the selected GO hierarchal structure scrambling method outperforms Hash GO as well as cloudscape in respect of research including using hierarchical structural linkages among huge GO words. Another intriguing discovery would be that Phish usually fails to Phish, even though they might equally preserve the GO words' hierarchical structures. The reason seems to be that Phish employs the GO dense mapping function, which really only stores the family link areas and provides words and does not properly store additional genetic heritage connections. S, on the other hand, has been used by Phish to store not just too genetic heritage links areas and providing words, as well as other types of associations. Although those who utilize a comparable KNN classifier like Phish to forecast genes involved, two control techniques yield similar outcomes with everyone, plus those who frequently maintain similar findings using InterGFP. This seems to be because those same three techniques employ the high dimension dense GO words universe to quantify the semantic relatedness of genes as well as forecast genes involved. This finding implies that the strength of semantic interaction calculated using huge GO words suffers from superficial as well as inadequate annotating, but the semantic relatedness generated from condensed Y suffers less. The findings for Fax, Smin, and MCC on Early humans were seen in Fig.1 (a), (b), (c) of the Resulted. Phish was extremely resistant to hashing bits sized, or perhaps the complexity of a compacted gene-term correlation matrix. Such consists of results, on the other hand, were susceptible to do and then have oscillations inside the examined d ranges. Furthermore, researchers find that Phish facilitates movement, steady sometimes at $d = 5$ as well as $d = 10$, and that this is untouched by password mismatch. These findings show that high numbers of GO words could be represented using simple binary frequent patterns, and that certain GO word seems semantically related enough to be combined. In reality, researchers investigated the responsiveness of d in two additional organisms; the experimental findings reveal identical interpretations of the results, and they've been deleted from this paper to conserve content. Humans could deduce from either of these results that specifying an important performance value of d using HPHash was simple.

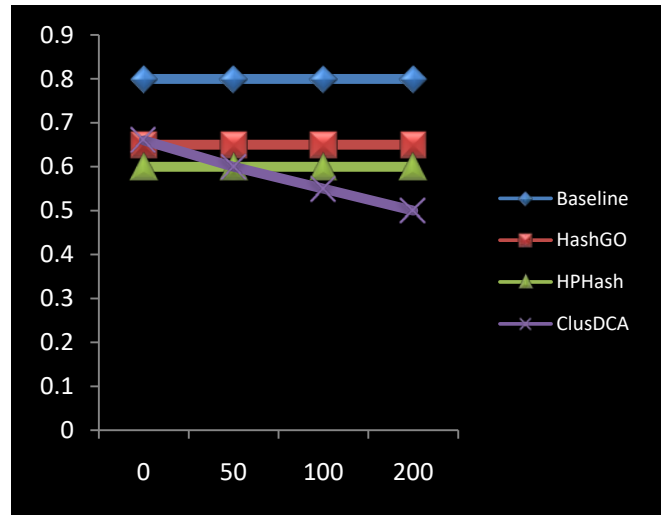


Fig.1(a). Sensitivity analysis of hash codes length

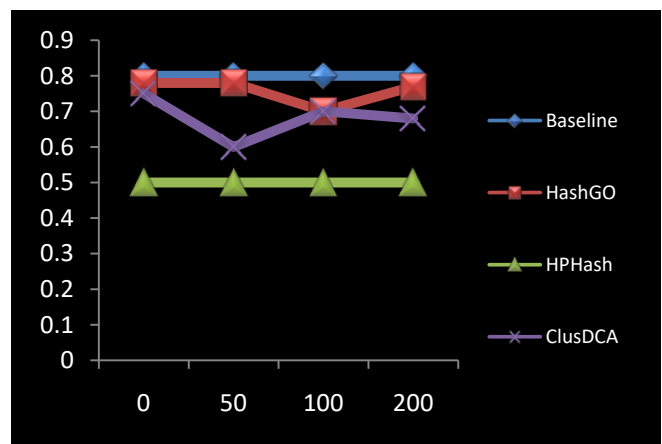


Fig.1(b). Sensitivity analysis of hash codes length

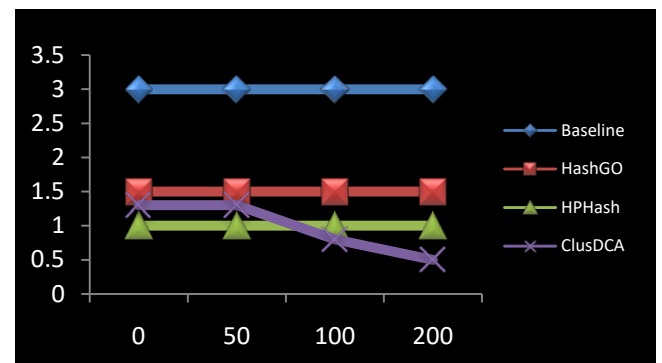


Fig.1(c). Sensitivity analysis of hash codes length

The outcomes on Homo sapiens for various k input parameters were seen in Fig.2 of the Supplemental Material. Humans could see that the initial value of k seems to have little effect on Phish. Many comparative approaches, from the other side, were susceptible to k . Moreover, selecting an efficient k for most of these comparison techniques was difficult since there is no definite trend for selecting an appropriate k that produces superior Fax, Sin, and MCC. Phish, on the other hand, was essentially unaffected by k . The excellence of Phish could be credited to the GO power structure conserving hash functions, which could also effectively start exploring as well as implement the

relationship existing between GO aspects, allowing for a more precise measurement of similarity measures among genes, including the replenishment of GO captions as well as the reduction of the influence of imperfect captions. Humans could deduce from all these findings that choosing an efficient k for Phish was simple.

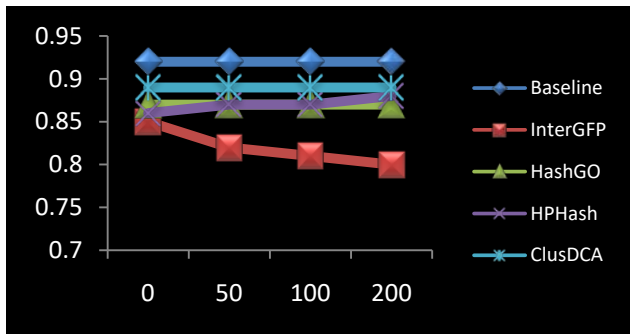


Fig.2(a) . Sensitivity analysis of neighbor size k.

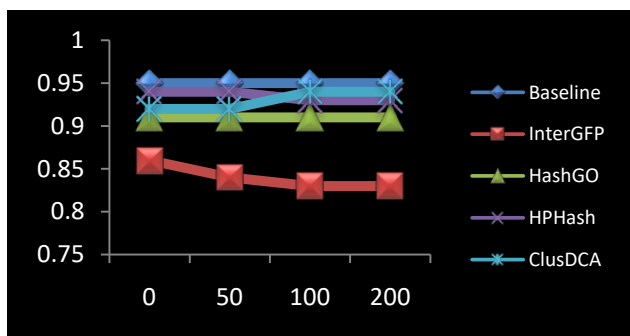


Fig.2(b) . Sensitivity analysis of neighbor size k.

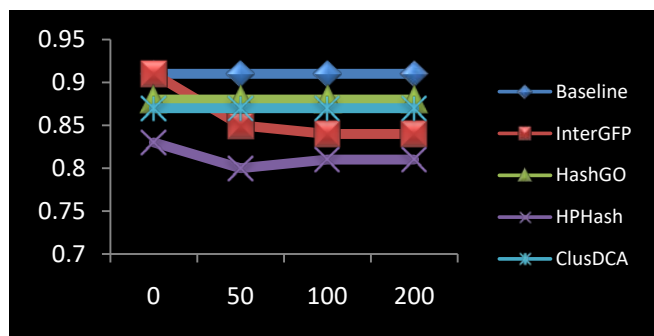


Fig.2(c). Sensitivity analysis of neighbor size k.

Fig.2 (a), (b), (c) displays that BLAST + Phish outperforms BLAST regularly. Since the previous gene-term affiliation structure would be a sparse matrix as well as the compact somebody may not be as sparse, the BLAST predicted gene-term affiliation structure seems to be sparse as well, while the affiliation framework of BLAST + Phish has been retained by cryptographic hash features H and therefore also contains the most nonzero records. Because all these nonzero values affect Aargau, BLAST + Phish occasionally do have a smaller Aargau than BLAST. BLAST + Phish, on the other hand, consistently outperforms BLAST in terms of Fax as well as Sin. The variance among BLAST as well as BLAST + Phish across one of these 3 species as well as assessment criteria also was assessed using signed-rank testing, with a p-value of less below 0.001. Based on this approach, researchers could

infer that Phish may be utilized as a component for sequencing information gene function forecast, and it can increase forecasting accuracy greatly.

VI. CONCLUSIONS

Among the most important responsibilities in the specific biological age was predicting genetic mutations. Lexical similarity-based techniques were also used to deal with the challenge that has shown considerable success, however, the majority of methods struggle from scant as well as inadequate genes GO classifications. To enhance the assessment of semantic relatedness among genes as well as for functional genomics forecasting, various GO keyword condensing methodologies have been developed. Even during the process of compression, though, individuals might not always obey the Gene Ontology structure, which may impact forecasting accuracy. Researchers offer a Phish based on Gene Ontology Hierarchy Preserving Pattern matching to resolve this challenge as well as quickly predict the relationships among genes as well as huge GO keywords. During the optimization of hashes, Phish may preserve the taxonomic resemblance of GO words as well as the structures of power among those. The hashes reduce strong gene-term connection matrices to a cheap version, making it easier to accurately measure semantic relatedness among families as well as predict gene function.

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