In *Silico* prediction and Comparative Modeling of proteins in Creeping Fig (*Ficus pumila*) plant

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ABSTRACT

In the present study, the *Ficus pumila* have taken to analyze the proteins by their preliminary characters from the database and predicted vital role of different sequences. The *F. pumila* (Creeping fig) is a prostrate/climbing shrub, experiments proved various active phytochemicals and antioxidant, antimicrobial, antinutagenic, analgesic, anti-inflammatory, antihyperglycemic, hypolipidemic, anti-hyperprolactinemic, anticholinesterase, nephroprotective properties. In addition to all metabolites, it also constitutes specific proteins that were evaluated through *insilico* homology modeling. Though it is considered as a poisonous weed, the protein present in this plant is evaluated by physicochemical, phylogeny and amino acid proportions by protparam, Swiss model, SOPMA, Clustal omega tools to describe its structural features and to understand molecular function. The computed theoretical isoelectric point (pI) found to be more than 7 indicates basic nature of proteins. The aliphatic index ranges 67-113 indicates thermal stability of proteins. The predicted Grand average hydropathy(GRAVY) shows possibilities of enhanced interaction of these proteins with water by lowest value.

Functional analysis of these proteins was performed by SOSUI server which predicted transmembrane helix and solubility. Secondary structure analysis was carried out by SOPMA revealed that Alpha helix and random coil dominated followed by extended strand, and beta turns among secondary structure elements. The modelling of three-dimensional structure of proteins was performed by Swiss model. The model was validated using protein structure checking tool- VADAR. Particularly, NAD(P)H – quinone oxidoreductase and Glyceraldehyde-3-phosphate dehydrogenase structures were analysed by phylogenetic analysis to trace relationship and reported. The results suggesting its possible role in cellular and metabolic functions.

Keywords: NAD(P)H –quinone oxidoreductase; Glyceraldehyde-3-phosphate dehydrogenase; *Ficus pumila*; phylogeny; Plant protein; Phylogenetic analysis

INTRODUCTION

Recent advances in sequence analysis methods and software have revolutionized the characterization and phylogenetic studies [1]. The availability of structural models of proteins is the key to understanding biological processes at a molecular level. The lot of protein sequences that can be modeled, as well as the exactness of the prediction, is growing gradually because of the growth and number of known protein sequences and structures as well as advances in the modeling software [2].

Sequence databases are of special importance for different fields of biological research because they are comprehensive sources of information on nucleotide sequences and proteins primary, secondary and tertiary characters. The tools for the computational analysis of the data and retrieved sequences are necessary resources for biological and medical research [3]. Plants are an abundant source of natural components especially proteins. In this study, proteins of *F.
Ficus pumila have been validated and expressed as preliminary characters of the same for the betterment of plant proteomic researches. Thus understanding its structural details would be a great revolution for engineering new herbicides, developing resistant crops and antimicrobial drugs.[4].

The Ficus species are used as food and as medication in traditional practices like Ayurveda, Siddha and Chinese Medicine. The F. pumila (Creeping fig) is a prostrate or climbing shrub, experiments proved various active phytochemicals like alkaloids, carbohydrates, tannins, flavonoids, saponin, glycosides, and steroid/triterpenoids.[1] The biological activities have proved through various experiments predominantly antioxidant, antimicrobial, antimitogenic, analgesic, anti-inflammatory, antiproliferative, hypoglycemic, hypolipidemic, anti-hyperprolactinemic, anticholinesterase, nephroprotective analytic methods. In addition to all metabolites, it also constitutes specific proteins that were evaluated in this attempt through insilico homology modeling. Traditionally this plant is used for malaria, fever, shivering, and vomiting in New Guinea. In China and Japan, it has been using for rheumatism, arthritis and pains due to sprains and even to treat diabetes and high blood pressure.[5]. The biologically active plant proteins need to get attention to acquaintance science and to make potential and sustainable alternative sources for various applications and essentials. This proteome is analyzable according to distribution of domains and protein families, and secondary and tertiary structures of proteins and can be made comparable to other proteomes.[9].

Despite progress in protein structure prediction, qualified modeling remains the only process that can constantly predict the 3-D structure of a protein with accuracy comparable to a low-resolution experimentally determined structure. Even models with mistakes may be useful because some features of function can be predictable from only coarse structural features[6]. The structures were taken to account along with phylogenetic analysis to trace the relationships.

MATERIALS AND METHODS

Sequence Retrieval
The FASTA sequences of the proteins were retrieved by using Genbank database hosted by the NCBI[7].

Primary Structure Prediction
The Exapy Propurm server has been used to get the physiochemical properties of proteins such as theorehtical Isoelectric Point (pl), molecular weight, total number of positive and negative residues, extinction coefficient, instability index(Ii), aliphatic index (Ai) and grand average of hydropathy (GRAVY)[8].

Structure Prediction
SOPMA (Self Optimized Prediction Method with Alignment) was used for the secondary structure prediction. Swiss model and Phyre2 were used for tertiary structure prediction of the proteins.

Functional Characterization
SOSUI tool used to describe whether the protein is soluble or transmembrane in nature. InterPro is a combined resource for protein families, domains and functional sites. Inter Pro incorporates the major protein signature databases into a single resource. Superfamily and molecular function were predicted by Inter Proprotein sequencing and classification. Motif Search from Genome Net was used for predicting the functional domains.

Sequence alignment was performed using pairwise sequence alignment tool (NCBI- BLAST) (http://blast.ncbi.nlm.nih.gov/Blast.cgi) and multiple sequence alignment was done using the EBI-CLUSTAL OMEGA (http://www.ebi.ac.uk/Tools/msa/clustalo/) tool. This Clustal Omega has features for adding sequences to and exploiting information in existing alignments, making use of the vast amount of precomputed information in public databases like Pfam[9]. The importance of this work was to novelty the regions of sequence comparison, which in other words allows us to yield functional and evolutionary relationships among the proteins considered in this study.

The phylogenetic analysis of ten to twenty proteins with most similarity percentage was completed to determine the number of proteins that part common structural and functional features. As an input to Clustal Omega, all sequences in FASTA formats were supplied with default options. The output was analyzed for sequences and the phylogenetic tree was constructed based on the bootstrap Neighbour Joining (NJ) method[10]. The steadiness of the internal nodes was assessed by bootstrap analysis with 1000 replicates.

RESULTS AND DISCUSSIONS
To classify and trace out the evolutionary path among related species, it is important to investigate them at the molecular level for better understanding of the evolution of plants.[11].

Parameters computed using Exapy’s ProfParam tool revealed that the highest molecular weight has observed as 45498 with the length of 424 in ATP synthase. It is also indicated that the pl of 5 proteins were less than 7, indicates those proteins were acidic (Table 1), from which pl of 2 proteins was nearly 7, indicates those proteins are near neutral and another 5 have basic character. The proteins are found to be compact and stable at their pl[12]. The computed isoelectric point (pl) will be useful for developing a buffer system for purification by the isoelectric focusing method[13]. Except 3, all the proteins of Ficus pumila showed instability index smaller than 40, which indicates the proteins are stable. The aliphatic index ranges from 67.41-112.02 indicates the thermal stability of the proteins. Grand average hydropathy (GRAVY) was predicted to be lower value shows the possibility of better interaction of these proteins with water[14]. The range of GRAVY of Ficus pumila proteins was –0.419 to 0.265.
Table 1: Accession numbers of the *F. pumila* proteins and physicochemical characters:

| Protein name                                      | Accession No. | Length | Mol.Wt   | pI  | R- | R+ | EC | li | AI  | Gravy |
|---------------------------------------------------|---------------|--------|----------|-----|----|----|----|----|-----|-------|
| NAD(P)H-quinoneoxidoreductase subunit             | AAN63314.1    | 388    | 44707.60 | 9.59| 20 | 35 | 68425 | 28.14| 100.28 | 0.265 |
| Chitinase                                         | ABB86300.1    | 301    | 33233.09 | 5.74| 31 | 25 | 48330 | 40.13| 67.41  | -0.367 |
| Oleosin                                           | ABQ57397.1    | 153    | 16314.90 | 9.77| 8  | 12 | 8480  | 44.58| 108.43 | 0.159 |
| Glyceraldehyde-3-phosphate dehydrogenase          | AEX08243.1    | 114    | 12039.58 | 6.48| 14 | 14 | 8480  | 32.26| 84.56  | -0.237 |
| Cystatin                                          | ACY30464.1    | 114    | 12821.06 | 9.67| 9  | 15 | 24980 | 13.58| 112.02 | 0.078 |
| Ribulose bisphosphate carboxylase large chain     | AFJ76695.1    | 233    | 26036.51 | 6.93| 29 | 29 | 39100 | 31.78| 76.18  | -0.394 |
| Maturase K                                        | AFJ76529.1    | 265    | 31615.58 | 9.43| 18 | 29 | 60070 | 39.02| 87.51  | -0.075 |
| ATP synthase subunit alpha                        | AAW33097.1    | 424    | 45498.31 | 7.78| 45 | 46 | 21110 | 34.86| 102.85 | 0.024 |
| 1-aminocyclopropane-1-carboxylate oxidase         | AEX08288.1    | 194    | 22293.55 | 5.26| 33 | 24 | 24075 | 22.28| 89.90  | -0.419 |
| Caleosin                                          | ABV72237.1    | 239    | 26975.52 | 5.37| 30 | 22 | 48360 | 47.02| 82.09  | -0.265 |

The computed protein concentration and existence coefficients help in the quantitative study of protein-protein and protein-ligand interactions in solution\(^{10}\) as well as in the further bioresearch and related product developments. Secondary structure prediction of *Ficus pumila* proteins by SOPMA revealed that α – helix, random coil, β – turn and extended strand were more prevalent. In Oleosin, Glyceraldehyde 3-phosphate dehydrogenase and 1-aminocyclopropane-1-carboxylate oxidase, α – helix, which plays a vital role in protein structure and function determination, predominate; whereas in Maturase K, ATP synthase subunit alpha extended strand dominates followed by Beta helix (Table 2). The random coil has dominated in Chitinase and Caleosin proteins of *Ficus pumila*.

Table 2: SOPMA prediction of *Ficus pumila* proteins

| Proteins                                           | Alpha helix (Hh) | Extended strand (Ee) | Beta turn (Tt) | Random coil (Cc) |
|----------------------------------------------------|------------------|----------------------|----------------|------------------|
| NAD(P)H-quinoneoxidoreductase subunit (ndhF)       | 33.25            | 21.13                | 4.90           | 40.72            |
| Chitinase                                          | 21.93            | 14.95                | 5.32           | 57.81            |
| Oleosin                                            | 45.75            | 15.03                | 9.15           | 30.07            |
| Glyceraldehyde 3-phosphate dehydrogenase (G3pdh)  | 40.35            | 21.93                | 4.39           | 33.33            |
| Cystatin                                           | 39.47            | 16.67                | 4.39           | 39.47            |
| Ribulose bisphosphate carboxylase (rbc)            | 33.91            | 20.17                | 5.15           | 40.77            |
| Maturase K                                         | 37.74            | 23.02                | 5.28           | 33.96            |
| ATP synthase subunit alpha (atp1)                  | 32.31            | 22.64                | 10.61          | 34.43            |
| 1-aminocyclopropane-1-carboxylate oxidase (aco1)    | 41.24            | 18.04                | 7.22           | 33.51            |
| Caleosin                                           | 33.05            | 9.21                 | 7.11           | 50.63            |
The SOSUI predicted the NAD(P)H-quinone oxido reductase subunit amino acid sequence (length 461 AA) is of a membrane protein which has 6 transmembrane helices which show average hydrophobicity is 0.142082. The TMHMM v.2.0 structure of is shown in Fig. 1. In the case of Glyceraldehyde-3-phosphate dehydrogenase, the amino acid sequence (length 189 AA) is of a soluble protein with average hydrophobicity is 0.479365.

Fig. 1: TMHMM result showing transmembrane region for NAD(P)H-quinone oxidoreductase subunit of Ficus pumila.

Domains are evolutionary units, frequently known as repeated sequences or 3D structures Fig. 2. In our results shows the distribution of the main chain bond lengths and bond angles were found to be inside the limits for these proteins. Such figures assigned by the Ramachandran plot exemplify good quality of the predicted models (Fig. 3). It can determine angles permitted and obtained the insight into the structure of peptides.

Fig. 2: 3D structure prediction through Swiss Models
a) Chitinase; b) Glyceraldehyde 3-phosphate dehydrogenase; c) Oleosin; d) Cystatin; e) Ribulose bisphosphate carboxylase large chain; f) Maturase K; g) 1-aminocyclopropane-1-carboxylate oxidase; h) ATP synthase subunit alpha; i) Caleosin
These proteins of *Ficus pumila* was subjected to BLASTp analysis to find the other plant species having the same query protein. The evolutionary relationship was done with NAD(P)H-quinone oxidoreductase subunit and glyceraldehyde-3-phosphophate dehydrogenase proteins. NADPH is an essential cofactor in many metabolic reactions. In plant peroxisomes, several enzyme activities are strictly dependent on the presence of NADPH. NADPH, whose regeneration is critical for reductive biosynthesis and detoxification pathways, is an essential component in cell redox homeostasis and other physiological function of amino acid-producing strains [16].

Role of Predicted pfam domains of the proteins in plants:

Pfam is a database of such protein domain families, with each family represented by multiple sequence alignments and profile hidden Markov models (HMMs). In addition, each family has associated annotation, literature references and links to other databases. The entries in Pfam are available via the worldwide web and in flat file format. One of the main challenges when constructing such a database is to simultaneously satisfy the conflicting demands of completeness on the one hand and quality of alignment and domain definitions on the other. We have also identified many novel family memberships in known proteins, including new kazal, Fibronectin type III, and response regulator receiver domains. Pfam-A families have permanent accession numbers and form a library of HMMs available for searching and automatic annotation of new protein sequences [17]. In the present study, the Pfam domain have located in different position those were tabulated with their E-value (Table 3) and structure (Fig. 4).

Table 3: Predicted Pfam domains of *F. pumila* proteins

| Sl. No | Protein | Pfam Domain | Position | E-value | Description |
|-------|---------|-------------|----------|---------|-------------|
| 1     | Oleosin | Oleosin     | 33…142  | PF01277, Oleosin |
| 2     | Caleosin | Oleosin     | 60…227  | PF05042, Caleosin related protein |
| 3     | Cystatin | SQAPI       | 32…111  | PF16845, Aspartic acid proteinase inhibitor |
|       |         | Cystatin    | 34…111  | PF00031, Cystatin domain |
|       |         | PP1         | 39…104  | PF07430, Phloem filament protein PPcystatin-like domain |
| 4     | Chitinase | Glyco_hydro_19 | 89…301 | 4.1e-69 | PF00182, Chitinase class I |
|       |         | Chitin_bind_1 | 28…66  | 7.4e-05 | PF00187, Chitin recognition protein |
| 5     | NdhF    | Proton_antipo_C | 124…376 | 2.8e-100 | PF01010, NADH-dehydrogenase subunit F, TMs (complex 1) |
|       |         | Proton_antipo_M | 1…118  | 9.7e-31 | PF00361, Proton-conducting membrane transporter |
| 6     | Rbc1    | RuBisCO_large | 143…451 | 4.3e-128 | PF00016, Ribulosebisphosphate carboxylase large chain, catalytic domain |
|       |         | RuBisCO_large_N | 13…133 | 3.4e-45 | PF02788, Ribulosebisphosphate carboxylase large chain, N-terminal domain |
| 7     | G3pdh   | Gp_dh_C     | 1…120   | 4.4e-55 | PF02800, Glyceraldehyde-3-phosphate dehydrogenase, C-terminal domain |
| 8     | atp1    | ATP-synt_ab | 121…345 | 2.3e-71 | PF00006, ATP synthase alpha/beta family, nucleotide-binding domain |
|       |         | ATP-synt_ab_C | 352…424 | 2.3e-33 | PF00366, ATP synthase alpha/beta family, C terminal domain |
|       |         | ATP-synt_ab_N | 1…64   | 2.5e-18 | PF02874, ATP synthase alpha/beta family, beta-barrel domain |
| 9     | Aco1    | 2OG-FeII_oxy | 131…193 | 2.3e-15 | PF03171, 2OG-Fe(II) oxygenase superfamily |
|       |         | DIOX_N      | 1…39    | 6.5e-06 | PF14226, non-ahaemdioxygenase in morphine synthesis N-terminal |
The Glyceraldehyde-3-phosphate dehydrogenase involving plant metabolism and evidence that the complex interactions existing between metabolism and development. Glyceraldehyde-3-phosphate dehydrogenase (NAD-GAPDH) is involved in a critical energetic step of glycolysis and also has several significant functions moreover its enzymatic activity \cite{18, 19}. It showed resemblances with 21 plants from 13 families where 6 plants from Brassicaceae which possess 94-96% identity shows the true relationship with *Ficus pumila*. The whole tree indices the identity ranges from 93.9 to 98.2%. It was also traced with higher identity (95.6%) with families such as Solanaceae, Rosaceae and Lamiaceae. The plant *Arachis hypogaea* show 98% identity and *Vigna angularis* shows 94.7%, therefore two plants are in the nearest relationship in Leguminosae family. This finding suggests that *Ficus pumila* and closely related plants of different families listed here may evolved from a common ancestor. Merging and separation are two distinguished phylogenetic properties, which can be valuable to determine uniqueness, closely as well as distantly related group of protein sequences \cite{20}. Glyceraldehyde-3-phosphate dehydrogenase protein sequences of *Diplocyclos palmatus* are almost coincidences with the known plant protein sequences of Euphorbiaceae, and Araliaceae with the identity of more than 95% \cite{21}.

It showed resemblances with ten families of twelve plant species ranges from 74 to 94%. It was traced in families such as Rosaceae, Solanaceae and Moraceae (own family). It indicates the phylogeny of the plant with the families mostly up to above 90% identity followed by Rutaceae, Asteraceae and Brassicaceae shows the identity of 87-89%. It shows 90.8% identity with the own family member *Morus indica* proved the true relationship with *Ficus pumila*. The phylogenetic tree of NAD(P)H-quinoneoxidoreductase subunit of the plants is presented in Table 4 & Fig. 5. In present study the Glyceraldehyde-3-phosphate dehydrogenase protein sequences of *F. pumila* shows the resemblance with Leguminosae, Brassicaceae, Solanaceae and Rosaceae families ranges from 95-98% predominantly (Table 5 & Fig.6).
Table 4: Phylogeny of *Ficus pumila* traced through NAD(P)H-quinone oxidoreductase subunit

| Accession No | Plant species containing NAD(P)H-quinone oxidoreductase subunit protein | Family | Identity % |
|--------------|---------------------------------------------------------------------------|--------|------------|
| ABB21003.1   | *Morus indica*                                                            | Moraceae | 90.8       |
| ESR41467.1   | *Citrus clementina*                                                       | Rutaceae | 87.2       |
| PRQ15668.1   | *Rosa chinensis*                                                          | Rosaceae | 93.1       |
| OTG25873.1   | *Helianthus annuus*                                                       | Asteraceae | 87.8     |
| EFH44954.1   | *Arabidopsis lyrata*                                                      | Brassicaceae | 88.4     |
| TYJ27490.1   | *Gossypium hirsutum*                                                      | Malvaceae | 86.8       |
| OIT27308.1   | *Nicotiana attenuata*                                                    | Solanaceae | 92.3       |
| RYQ84036.1   | *Arachis hypogaea*                                                        | Leguminosae | 86.9       |
| RX09703.1    | *Malus domestica*                                                         | Rosaceae | 92.2       |
| PUZ60784.1   | *Panicum hallii var. hallii*                                              | Poaceae | 79.5       |
| ADO65021.1   | *Prunus persica*                                                          | Rosaceae | 74.1       |
| AAT98557.1   | *Panax ginseng*                                                           | Araliaceae | 74.0       |

Fig. 5: Phylogenetic tree of *Ficus pumila* traced through NAD(P)H-quinone oxidoreductase subunit

Table 5: Phylogeny of *Ficus pumila* traced through glyceraldehyde-3-phosphate dehydrogenase protein

| Accession No | Plant species containing glyceraldehyde-3-phosphate dehydrogenase protein | Family | Similarity % |
|--------------|---------------------------------------------------------------------------|--------|--------------|
| RYR67635.1 | *Arachis hypogaea*                                                        | Leguminosae | 98.2        |
| CAA7039284.1 | *Microtus pierrica*                                                       | Brassicaceae | 96.5        |
| PHU23654.1 | *Capsicum chinense*                                                       | Solanaceae | 95.6        |
| CDY42906.1 | *Brassica napus*                                                           | Brassicaceae | 95.6        |
| ESQ35475.1 | *Eutremaasalusugineum*                                                    | Brassicaceae | 95.6        |
| EFH69008.1 | *Arabidopsis lyrata*                                                      | Brassicaceae | 95.6        |
| ONI08955.1 | *Prunus persica*                                                           | Rosaceae | 95.6        |
| TEY91672.1 | *Salvia splendens*                                                         | Lamiaceae | 95.6        |
| KF73847.1 | *Arabisalpina*                                                             | Brassicaceae | 95.6        |
| KZM93169.1 | *Daucus carota*                                                            | Apiaceae | 94.7        |
| PKJ77035.1 | *Panicum gramineum*                                                        | Lythraceae | 94.7        |
| KAB2624871.1 | *Pyrus ussuriensis*                                                        | Rosaceae | 94.7        |
| KGW61990.1 | *Eucalyptus grandis*                                                       | Myrtaceae | 94.7        |
| BAT79321.1 | *Vigna angularis*                                                          | Leguminosae | 94.7        |
| EOA38185.1 | *Capsella rubella*                                                         | Brassicaceae | 94.7        |
| PL735941.1 | *Lactuca sativa*                                                           | Asteraceae | 94.7        |
| RXW96054.1 | *Malus domestica*                                                          | Rosaceae | 94.7        |
| PSS30165.1 | *Actinidichinae*                                                           | Actinidiaceae | 94.7     |
| KNA16142.1 | *Spinacia oleracea*                                                        | Amaranthaceae | 94.7     |
| PIA51356.1 | *Aquilegia coerulea*                                                       | Ranunculaceae | 94.7     |
| KDO82023.1 | *Citrus sinensis*                                                          | Rutaceae | 93.9        |
CONCLUSION

Mapping the physicochemical character of plant proteins and their relationship with other families will provide an important guideline for the future study of plant biology and drug development. Physicochemical characters of various proteins of Ficus pumila have been analysed and their three dimensional structure were predicted. Multiple alignments are a central feature of Pfam. The web sites provide access to the seed and full alignments in a variety of formats, allowing users to input Pfam data into their own software. Alignments are best viewed with specialised programs that can highlight similar regions and carry out manipulations of the alignment. The phylogeny of the plant proteins NAD(P)H-quinone oxidoreductase and Glyceraldehyde-3-phosphate dehydrogenase were traced by their relationships with other plants. The results suggested that the selected proteins having unique characters as well as the phylogeny indicate the convergence and divergence with different plants of different families even through the long history of evolution. Further, phytochemicals and the peptides of the proteins will be used for further studies to assess their potential biological activity in targeting various diseases. This team will carry these biomaterials to explore the many therapeutic properties in future for the drug target identification.

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