



HAL
open science

Species specific amino acid sequence-protein local structure relationships: An analysis in the light of a structural alphabet.

Alexandre de Brevern, Agnel Praveen Joseph

► To cite this version:

Alexandre de Brevern, Agnel Praveen Joseph. Species specific amino acid sequence-protein local structure relationships: An analysis in the light of a structural alphabet.: species sequence - structure relationship. *Journal of Theoretical Biology*, 2011, 276 (1), pp.209-17. 10.1016/j.jtbi.2011.01.047 . inserm-00646218

HAL Id: inserm-00646218

<https://www.hal.inserm.fr/inserm-00646218>

Submitted on 29 Nov 2011

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Species Specific Amino Acid Sequence – Protein Local Structure Relationships: an analysis in the light of a structural alphabet.

Alexandre G. de Brevern^{1,2,3,§}, Agnel Praveen Joseph^{1,2,3}

¹ Dynamique des Structures et Interactions des Macromolécules Biologiques (DSIMB), INSERM, U665, Paris, 6, rue Alexandre Cabanel, F-75015, France.

² Université Paris Diderot - Paris 7, UMR-S665, Paris, F-75015, France.

³ Institut National de la Transfusion Sanguine (INTS), Paris, F-75015, France.

[§] Corresponding author: Alexandre G. de Brevern, INSERM UMR-S 665, DSIMB, Université Paris Diderot – Paris 7, Institut National de Transfusion Sanguine (INTS), 6, rue Alexandre Cabanel, 75739 Paris cedex 15, France

Email addresses : alexandre.debrevern@univ-paris-diderot.fr, agnel.praveen@univ-paris-diderot.fr

Short title: species' sequence – structure relationship

Abstract

Protein structure analysis and prediction methods are based on non-redundant data extracted from the available protein structures, regardless of the species from which the protein originates. Hence, these datasets represent the global knowledge on protein folds, which constitutes a generic distribution of amino acid sequence – protein structure (AAS-PS) relationships. In this study, we try to elucidate whether the AAS-PS relationship could possess specificities depending on the specie.

For this purpose, we have chosen three different species: *Saccharomyces cerevisiae*, *Plasmodium falciparum* and *Arabidopsis thaliana*. We analyzed the AAS-PS behaviors of the proteins from these three species and compared it to the “expected” distribution of a classical non-redundant databank. With the classical secondary structure description, only slight differences in amino acid preferences could be observed. With a more precise description of local protein structures (Protein Blocks), significant changes could be highlighted.

Saccharomyces cerevisiae's AAS-PS relationship is close to the general distribution, while striking differences are observed in the case of *Arabidopsis thaliana*. *Plasmodium falciparum* is the most distant one.

This study presents some interesting view-points on AAS-PS relationship. Certain species exhibit unique preferences for amino acids to be associated with protein local structural elements. Thus, AAS-PS relationships are species dependant. These results can give useful insights for improving prediction methodologies which take the species specific information into account.

Key-words: amino acids, protein folds, secondary structure, structural alphabet, Protein Blocks, malaria.

1. Introduction

The knowledge of three dimensional structures of proteins gives valuable insights into their functions. Prediction of different features of protein structure like secondary structures (Jones, 1999; Madera et al., 2010; Pollastri et al., 2002), protein disorder (Madera et al., 2010; Xue et al., 2010), transmembrane regions (Illergard et al., 2010; Pylouster et al., 2010), phosphorylation sites (Biswas et al., 2010), protein flexibility (Bornot et al., 2011) or the generation of structural models (de Brevern, 2010; Kelley and Sternberg, 2009), are mainly based on machine learning algorithms (Brylinski and Skolnick, 2008; Rangwala et al., 2009; Xu et al., 2008). Protein structure analyses and prediction methods derive information from non-redundant databanks that represent the state-of-the-art of available data (*i.e.*, solved protein structures). In a way, they reflect a generic ('expected') distribution of amino acid sequence – protein structure (AAS-PS) relationship. Though the term “protein sequence-structure relationships” have been used extensively in literature, to avoid confusion with a few discussions on genomic data and to add clarity, we use the abbreviation AAS-PS here.

The number of protein sequences available is enormous and rapidly growing. Indeed, the number of completely sequenced genomes currently available is more than 1,250 (Kyrpides, 1999; Liolios et al., 2008). On the other hand, due to technical difficulties, the number of protein structures available is highly limited (Berman et al., 2003; Berman et al., 2000; Lattman, 2004; Service, 2008) and these prediction approaches can help in narrowing down the gap between the sequence and structure space.

Though the AAS-PS relationships observed across different proteins in a databank follow similar patterns, it is not known whether there could be variations based on the species from which the proteins are derived. Analyses of genomic data have given some interesting insights in this direction. Considerable variation in the composition of nucleotides has been observed

in some genomes, *e.g.*, *Plasmodium falciparum* (Gardner et al., 2002; Singer and Hickey, 2000). This compositional bias can be attributed to species-specific and/or environmental specific preferences (Dumontier et al., 2002; Wang and Lercher, 2009). Nucleotide bias is also reported to have an effect on the codon usage and hence, the amino acid composition of encoded proteins (Paila et al., 2008; Singer and Hickey, 2000). An interesting work has been carried out on the comparison of protein structural properties in *Bacillus subtilis* and *Escherichia coli* (Marashi et al., 2007). The main focus was on the amino acid solvent accessibilities and helix propensities and they could identify specificities in the properties observed in each species. Considering these observations, we base our work on the hypothesis that differences in the AAS-PS behaviors could be observed across species.

Indeed, a few research studies on similar grounds have been reported recently. Out of the sequences available for *P. falciparum* proteins, the functions of 5,460 are predicted and 3,200 are of unknown function (www.plasmodb.org). Sequence annotation methods work mainly by analogy, *i.e.*, by aligning the protein sequences. Improvement in the quality of sequence alignment could be achieved with the help of information on the amino acid compositional bias or with the use of composition adjusted substitution matrices (Brick and Pizzi, 2008; Paila et al., 2008; Yu et al., 2003). Asymmetric substitution matrices were also derived from proteomes with biased amino acid distributions (Bastien et al., 2005).

These studies reflect the fact that different species are marked with variations in their genomes and this is reflected at the level of amino acids that build proteins. Nevertheless, structural researches rely equally on the information from all the proteins, without taking into account the species they come from. Considering these points, we have analyzed whether the proteins originating from different species exhibit different AAS-PS relationships. For this purpose, we have analyzed protein structures with two main view points: (i) classical secondary structure and (ii) a finer description of local fold based on a structural alphabet

named Protein Blocks. The secondary structure description based on regular structural elements, *i.e.*, α -helix and β -sheet (Pauling and Corey, 1951; Pauling et al., 1951), is widely used to describe and visualize 3D structures. Nevertheless, these regular structures do not provide a complete description of the local protein structure; more than 50% of the structure is described as coil which is defined to represent highly variable backbone conformations. However, regular local folds could be traced in the segments assigned as coils.

Hence, local protein structure libraries that are able to give a complete approximation of protein structures, without any *a priori* knowledge of secondary structure assignments, have been developed. The complete set of local structure prototypes defines a structural alphabet (Karchin et al., 2003; Offmann et al., 2007). The number of prototypes in a library is important for characterizing the local fold (Hunter and Subramaniam, 2003; Martin et al., 2008; Sander et al., 2006; Tung et al., 2007; Unger et al., 1989). The structural alphabet used in this study, is composed of 16 local structure prototypes that are 5 residues long, called Protein Blocks (PBs, see Figure 1) (de Brevern et al., 2000; Joseph et al., 2010a; Joseph et al., 2010b). They efficiently approximate every part of protein structures (de Brevern, 2005)

From different eukaryotic kingdoms, three species namely *Arabidopsis thaliana* (*At*), *Plasmodium falciparum* (*Pf*) and *Saccharomyces cerevisiae* (*Sc*) were chosen for the study. The genomes of *At* and *Sc* have similar AT content (~62-64%) while *Pf* is highly diverged (82%). With the availability of complete genome sequences, sufficient data on the proteome could be also extracted for analyzing the compositions. For these three species, we characterize the amino acid preferences associated with different local folds. These behaviors were then compared with the general or 'expected' AAS-PS distribution. This will help to put into evidence the interest and importance of considering the species of origin to refine the knowledge on AAS-PS relationship. Moreover, from a methodological point of view, this

study highlights the interest of using a more detailed local structure description with a structural alphabet, to improve and refine the analysis of sequence-structure relationships.

2. Materials and Methods

2.1. Data sets.

In the same way, we have generated three species-specific non-redundant databanks. The Protein Data Bank was searched for structures from three species namely *Arabidopsis thaliana*, *Plasmodium falciparum* and *Saccharomyces cerevisiae*. Only X-ray crystallography structures were selected and the sequence identity cutoff of 90% was chosen. In September 2008, 68 crystallographic structures of *P. falciparum* (*i.e.*, 36,175 amino acids), 123 of *A. thaliana* (*i.e.*, 173,450 amino acids) and 427 of *S. cerevisiae* (*i.e.* 202,410 amino acids), were available (see the Supplementary material 1 for the list of identifiers). These three databases are referred as *Pf*, *At* and *Sc* respectively in the text. To analyze the effect of redundancy on the results, new non-redundant databanks have been generated for the three species at 50 and 25% sequence identity cutoffs.

To compare the composition of the proteome and sequences of structures available in the PDB, proteome sequences were obtained from publically available databases. The *A. thaliana* proteome was downloaded from the National Center for Biotechnology Information database (ftp://ftp.ncbi.nih.gov/genomes/Arabidopsis_thaliana/). About 33,711 annotated sequences were available (version of January 2008). The *P. falciparum* proteome was obtained from the PlasmoDB database (<http://plasmodb.org/common/downloads/release-5.4/Pfalciparum/>). From the database release 5.4, 5,460 automatically annotated sequences were downloaded.

2.2. Secondary structure assignments

Secondary structure assignment has been carried out using DSSP (Kabsch and Sander, 1983) (CMBI version 2000). DSSP assignment is based on the hydrogen bond lengths between backbone atoms of amino acids describing the 3D structure. As the assignment of more than 3 basic states is given by DSSP, we have grouped the results in three states: the α -helix contains α , 3_{10} and π - helices, the β -strand contains only the β -sheet assignment and the coil includes the other assignments (β -bridges, turns, bends and coil), as done in earlier studies (Fourrier et al., 2004; Tyagi et al., 2009a). Default parameters have been used.

2.3. Protein Blocks

Protein Blocks (PBs) correspond to a set of 16 local prototypes, labeled from *a* to *p* (cf. Figure 1), described as a series of (Φ , Ψ) dihedral angles corresponding to 5 residues (Joseph et al., 2010a; Joseph et al., 2010b). They were obtained using an unsupervised classifier similar to Kohonen Maps (Kohonen, 2001) and Hidden Markov Models (Rabiner, 1989). The PBs *m* and *d* can be roughly described as prototypes for the central region of α -helix and β -strand, respectively. PBs *a* through *c* primarily represent the N-cap of β -strand while *e* and *f* correspond to C-caps; PBs *g* through *j* are specific to coils, PBs *k* and *l* correspond to N cap of α -helix while PBs *n* through *p* to C-caps. This structural alphabet allows a reasonable approximation of local protein 3D structures (de Brevern et al., 2000) with a root mean square deviation (*rmsd*) of about 0.42 Å (de Brevern, 2005). PB assignment has been made using an in-house tool (available at <http://www.dsimb.inserm.fr/DOWN/LECT/>), similar assignment is also made by PBE web server (<http://bioinformatics.univ-reunion.fr/PBE/>) (Tyagi et al., 2006). They have been used to describe the 3D protein backbones (de Brevern, 2005) and to perform local structure prediction (Benros et al., 2009; de Brevern et al., 2000; de Brevern et

al., 2004; Dong et al., 2008; Etchebest et al., 2005; Rangwala et al., 2009; Zimmermann and Hansmann, 2008). PB based approaches have been proved to be effective in prediction based on protein sequence-structure relationships (de Brevern et al., 2007; Li et al., 2009; Tyagi et al., 2009b). Several other applications based on the PB structural alphabet have also given successful results (de Brevern et al., 2009; Dudev and Lim, 2007; Faure et al., 2009; Thomas et al., 2006; Tyagi et al., 2008). The characteristics of our structural alphabet have been compared with those of 8 other structural alphabets. This comparative study clearly shows that our PB alphabet is highly informative, with the best predictive ability of those tested (Karchin, 2003; Karchin et al., 2003).

2.4. Z-score

The Z-score quantifies the preference for each amino acid to occur in a local structural element (secondary structures and PBs). The amino acid occurrences in each secondary structure and PB have been normalized into a Z-score:

$$Z(n_{i,j}) = \frac{n_{i,j}^{obs} - n_{i,j}^{th}}{\sqrt{n_{i,j}^{th}}}$$

with $n_{i,j}^{obs}$ the observed occurrence number of amino acid i in position j for a given secondary structure or PB and n_{ij}^{th} , the expected number. The product of the occurrences in position j with the frequency of amino acid i in the entire databank equals $n_{i,j}^{th}$. Positive Z-scores correspond to over-represented amino acids and vice versa; threshold values of 4.42 and 1.96 were chosen to indicate the level of significance (probability less than 10^{-5} and $5 \cdot 10^{-2}$ respectively). On the Figures, a color code is used to highlight and easily compare the behavior in different databanks; only the preferences corresponding to the central positions are presented. This approach was earlier used to analyze the amino acid representativeness in

Protein Blocks in our earlier works (Benros et al., 2009; Benros et al., 2006; de Brevern and Hazout, 2001; de Brevern and Hazout, 2003; de Brevern et al., 2000; de Brevern et al., 2002; de Brevern et al., 2007; de Brevern et al., 2004; Etchebest et al., 2005; Etchebest et al., 2007; Ghozlane et al., 2009).

3. Results

3.1. Analysis of the Databanks

The species specific databanks contain only structures from the species: *Saccharomyces cerevisiae* (noted as *Sc* in the text), *Arabidopsis thaliana* (*At*) and *Plasmodium falciparum* (*Pf*). A classical non-redundant databank (*NR*) was also obtained from the PDB (Berman et al., 2000). The three specialized databanks (*Sc*, *At* and *Pf*) represents 202,410, 173,450 and 36,175 amino acid residues respectively. With regards to our previous studies (Bornot et al., 2009; de Brevern, 2005; de Brevern et al., 2000; de Brevern et al., 2002; de Brevern et al., 2004; Etchebest et al., 2005; Fourrier et al., 2004), the number of amino acids in each databank is sufficient to analyze the sequence – structure relationships in terms of secondary structure and Protein Blocks (Tyagi et al., 2009b).

Amino acid frequencies observed in each databank are reported in Table 1. As explained earlier, the *NR* is considered as the reference dataset. It corresponds to the “expected” distribution of amino acids. The amino acid frequencies of the *At* databank are most similar to those of the *NR* databank. Only one amino acid, *i.e.*, Methionine, is under-represented by about 40%, when compared to its frequency in the *NR* databank. For the other 19 amino acids, the variation is below 10%. For the *Sc* databank, only 3 out of the 20 amino acids exhibit notable changes in their relative occurrence frequencies. Isoleucine and Lysine are over-represented, with an increase of about 30% in the occurrence frequency, while Alanine is

under-represented with a decline in the frequency by 30%. For the other 17 amino acids, the variations are below 10%. The amino acid frequencies associated with the *Pf* databank has profound differences. Half of the amino acids exhibit variations of more than 30%. Five amino acids namely (Isoleucine, Tyrosine, Cysteine, Asparagine and Lysine) have significantly higher frequencies, and on the other hand, four others (Alanine, Tryptophan, Glutamine and Arginine) have lower frequencies compared with those observed for the *NR* databank.

3.2. Comparison with the proteome sequence dataset

As a first step, we performed a precise analysis of the protein structure databanks to ensure that they reflect the amino acid distribution observed in the proteome of the respective species. These differences observed with respect to the *NR* databank could be due to the amino acid composition of each species. Figure 2 shows the distribution of amino acids observed within the proteomes of *Arabidopsis thaliana* and *Plasmodium falciparum* (indicated as *At_p* and *Pf_p*) and also in the structural databanks *At* and *Pf*. Thus, the amino acid composition of protein structure databanks mainly reflects the proteomic composition of the organisms. Nevertheless, particular behaviors can be observed for two amino acids while comparing the compositions of *Pf* and *Pf_p*. The frequency of Asparagine (N) and Lysine (K) in *Pf_p* is significant, *i.e.*, 14.4% and 11.7% respectively, and these are higher than those observed in the structural *Pf* databank. The increase is about two times for N and 25% for K. These amino acids mainly characterize the low-complexity regions which are largely absent in the solved structures (see Supplementary material 2 for more details and also refer to (Aravind et al., 2003; DePristo et al., 2006; Joubert and Joubert, 2008)).

The amino acid distributions differ between species (and this is remarkable for *Pf*) and our structure datasets present a good reflection of the proteome content. Hence they can be used as representative datasets. Moreover, even if the amino acid frequencies vary between species (see Table 1), we can observe that the frequencies of regular secondary structures and Protein Blocks (PBs), are quite similar, as indicated in Table 2. Hence, all the datasets can be compared in terms of local protein structures.

3.3. Secondary structure contents

In order to analyze the variation in AAS-PS relationship between species in terms of secondary structures, the Z-scores distribution of amino acids associated with α -helix, β -strand and coil have been computed for the 4 databanks. As Z-score follows a χ -square distribution, a statistical threshold can be used to quantify the significance of the under or over-representation of an amino acid in the secondary structural element. Z-scores were computed on a sequence window of 5 residues (corresponding to PB length). Figures 3a to 3c show amino acid frequency (in terms of Z-score) associated with α -helix, β -strand and coil for *At*, *Pf*, *Sc* and *NR* databanks. We perform a precise analysis *per* state to explore *specific* behaviors within species. Methionine is seen to have a high preference to be associated with α -helices, this over-representation is not observed in *Pf* alone (see Figure 3a). The distribution of Histidine and Cysteine in *At* and *Pf* databanks, show significant differences. The amino acid distribution in β -strands of *Pf* shows striking variation when compared to the other datasets, especially the preference for Tryptophan is not seen (see Figure 3b). For the coil state, the amino acid distribution in *At* shows some differences, marked with a higher preference for Phenylalanine, Tyrosine, Tryptophan, Glutamine, Glutamate and Arginine while in the case of *Pf*, only the over-representation for Tryptophan is absent (see Figure 3c).

Figure 3d shows a hierarchical clustering of different species based on the amino acid distributions. As expected, the secondary structures contents show similarities across the 4 databanks; the 3 states (α -helix, β -strand and coil) cluster separately, each state localized on a specific branch. We can notice that the distribution in *Sc* is close to *NR* for all states, while amino acid preferences in *At* has similarity with *Pf*. Though a few variations are noted, a detailed picture of species specific behavior in terms of amino acid distribution cannot be obtained using the secondary structure description alone.

3.4. Protein Block contents in terms of species

The 16-state description of local structures in terms of Protein Blocks (PBs) is expected to highlight finer differences in the amino acid preferences in each species (Joseph et al., 2010b). We can categorize the amino acid preferences for different PBs into 3 classes while comparing *At*, *Pf* and *Sc* with *NR*: (i) *new* over- or under-representation, *i.e.*, a higher or lower amino acid preference not seen in *NR*, but found in the other datasets, (ii) *absence of* over- or under-representation, *i.e.*, an over- or under-representation seen in *NR* but *missing* in other datasets, and (iii) the most important variation, *inversion in the preference*, *i.e.*, an amino acid which has high preference for association with a PB in *NR*, is poorly represented in one or more of the other datasets and *vice – versa* (see the Supplementary material 3).

For the 16 different PBs, *Sc* has 32 *new* and significant amino acid over- or under-representations, only 4 amino acid preferences are *missing* and no *inversion* is observed. This roughly corresponds to a change affecting about 2.25 amino acids per PB (11.25%). *Pf* has a more complex behavior as it has 16 *new* significant associations, but 27 preferences are *missing* and 3 have been *inverted*. These modifications affect an average of 2.87 amino acids per PB (14.35%). Finally, *At* shows 52 *new* significant over- or under- amino acid

representations, 15 are *missing* and 6 *inversions* are observed, this corresponds to a change of about 4.5 amino acids per PB (22.50%). However, the extent of variation depends strongly of the kind of PB. Some have none or only a few differences when compared to *NR*. For instance, the distribution of PB *f*'s amino acid preferences is similar for all the datasets; PB *m* has only one *new* over or under-representation in *Sc* and two in *Pf* and *At*.

3.5. Some characteristic Protein Blocks

Figure 4 gives a visual representation of the amino acid preferences for 4 representatives PBs. PBs *j* and *k* exhibit few variations with respect to *NR*, while PB *e* shows more changes and PB *a* is the most variant (see Table 3). PB *j* (see Figure 4c) is quite determinant as it shows a strong over-representation of Glycine and all the other amino acids are under-represented. So, the over-representation of Glutamine observed for *At* is a surprising inversion. For *Sc*, the strong under-representation of Asparagine is not observed, Alanine and Arginine are highly under-represented in *Pf* and other significant variations can be observed. For PB *k* (see Figure 4d), the distribution is more complex with 4 over-represented and 13 under-represented amino acids. An *inversion* is observed for Arginine in the case of *Pf* and for Serine in *At*. A new over-representation is seen in *Sc* and *At*, two preferences are *missing* for *At* and three are absent in the case of *Pf*.

For PB *e* (see Figure 4b), 14 differences were observed: (i) two *inversions* (both in the case of *At*), (ii) three new over-representation for *Sc*, five for *At* and one for *Pf*, and (iii) one *missing* amino acid preference for *At* and 2 for *Pf*. PB *a* (see Figure 4a) exhibits the most striking variations that are species specific. Out of the 25 major changes observed, 4 are found in *Sc*, 12 in *At* and 9 in *Pf*. Importantly, PB *a* is characterized by 5 over- and 4 amino acid under-representations unique for the species. Interestingly, even if PB *a* connects repetitive

structures (PBs *m* to PB *d* see (de Brevern et al., 2002)), it has weak preferences for Proline and Glycine. New preferences dominate the majority of changes; 4 are observed in *Sc*, 10 in *At* and 5 in *Pf*. These specificities are further strengthened with 2 *missing* amino acids observed in *At* and *Pf* and also the *inversion* of 2 amino acid preferences in *Pf*.

3.6. Global comparison in terms of PBs

As mentioned earlier, slight differences can be observed while analyzing secondary structures, but still same states are clustered together. With PB description, strong differences have been detected across species. The hierarchical clustering of the associated amino acid distributions (see Figure 5) shows that AAS-PS relationship of PBs from different species could be really different. Indeed, three kinds of behaviors could be observed (see the Supplementary material 4). Firstly, the amino acid distributions associated with a PB are clustered together, for the four different databanks, *e.g.*, PBs *g*, *n* and *o*. Secondly, three amino acid distributions (for different databanks) are clustered and the amino acid distribution of *At* is associated with another cluster. This behavior is observed for PBs *e* and *h*. Thirdly, the most common case (11 out of 16 BPs), three amino acid distributions are clustered and the amino acid distribution corresponding to *Pf* is not associated with them. The latter behaviors are due to two major factors: (i) the difference in informativity of AAS-PS relationship and (ii) the specific amino acid preferences which are species -dependent.

The informativity can be quantified as the sum of absolute values of Z-scores. *At* has an additional informativity of 16% compared to *NR*. This value decreases to +4% for *Sc*. For *Pf*, a very weak informativity is observed, *i.e.*, 45% lower than that of *NR*. Thus, *Pf* exhibits weak AAS-PS relationships compared to other datasets. It must be also noted that some PBs

of *At* have more informative distributions, *e.g.* PB *a*: +72%, PB: *e* +60%, PB: *h* +39% and PB: *g* +20%, but still the clustering show variations.

4. Discussion

4.1. Global view

In this study, we have analyzed protein AAS-PS relationships existing in *A. thaliana*, *Plasmodium falciparum* and *S. cerevisiae*. We have shown that non-redundant protein structure databanks specific for these species can be compared without systematic bias, even if the *P. falciparum* specific databank has compositional disequilibrium in terms of certain amino acids.

For the four databanks, the analysis of AAS-PS relationships associated with the secondary structures highlights only a few differences. At the light of Protein Block description, striking specie specific behaviors are observed. The AAS-PS relationship of *Sc* is quite close to that of *NR* (*i.e.*, expected distribution). Only slight differences are seen, *e.g.*, no *inversion* is observed. *A. thaliana* has both an AT-content and proteome composition similar to *S. cerevisiae*. Nevertheless, *At* exhibits very distinct AAS-PS relationships. Though the informativity associated with the distributions in *Sc* can be compared to that of *NR*, *At* is clearly more informative, having an average variation of 5 amino acid preferences *per* PB which is quite striking. As seen in Table 3, these differences could result in very different amino acid signatures when compared to *NR*. These precise details on amino acid preferences were not highlighted in their study of Marashi and co-workers (Marashi et al., 2007). In the same way, it is expected that *P. falciparum* with 82% AT content and strong biases in amino acid composition, shows strong divergence in terms of amino acid usage. Its AAS-PS preferences are quite surprising. Clustering of these amino acid signatures (see Figure 5)

shows that *Pf* is quite far from the other species, and also from the expected view of AAS-PS (*NR*, see Supplementary material 5).

The last remark is promising indeed. The extensive genetic diversity of blood stage antigens is one of the key challenges for vaccine development against malaria. A recent approach proposed for antigen discovery is based on the bioinformatic selection of heptad repeat motifs corresponding to α -helical coiled coil structures (Kulangara et al., 2009). Considering the fact that the AAS-PS seen in *Pf* is diverged when compared to the expected / observed values (*NR*), such approaches could be refined or improved. For *At*, the differences with respect to the expected amino acid distributions are mainly due to *new* amino acids preferences (52), coupled with the *absence* of certain amino acid preferences observed in *NR* (15), some *inversions* of preferences are also observed (6). Indeed, this can be considered as a specialization unique for the species. This ratio of significant *new* and *missing* amino acid preferences is about 3.47 for *At*, whereas this is not the case for *Pf* where the ratio is only 0.59.

4.2. Specific case of *P. falciparum*

The sequence – structure relationship of *Pf* is weaker than that of the other databanks. In the hierarchical clustering shown in Figure 5, the PBs of *Sc*, *NR* and *At* are spread all over the dendrogram, while the PBs of *Pf* are seen to cluster in group located in the central part of the dendrogram. For proteomic approaches also, the results of our study could be of interest. The variation in the amino acid preferences associated with the local folds can have an implication in the conservation or substitutions of amino acids observed in the sequences of the species. 58% of predicted proteins of *P. falciparum* available in PlasmoDB have no known function (Aurrecochea et al., 2009). As *P. falciparum* proteins have diverged extensively during the

course of evolution, it is difficult to find homologues for many of these proteins *e.g.*, protein kinases (Anamika et al., 2005). Dedicated amino acid substitution matrix that is biased towards the amino acid distribution observed in *P. falciparum*, could help in better homology detection and functional annotation of *P. falciparum* proteins (Bastien et al., 2005; Brick and Pizzi, 2008; Nidhi et al., 2009; Paila et al., 2008).

Indeed, it is possible to propose equivalent (easily exchangeable) amino acids, for the species under consideration in order to obtain a reduced amino acid alphabet (Etchebest et al., 2007). This can be followed by simpler pairwise alignments, with a limited number of combinations. The detection of putative or/and new homologues would be easier and particularly interesting for *Plasmodium falciparum*, as its proteins represent potential therapeutic drug targets to combat malaria (Salzemann et al., 2009).

4.3. Pilot study

A pilot study has been carried out to assess the possible influence of species AAS-PS in local structure prediction. A classical Bayesian approach (Altschul et al., 1997) has been used to predict Protein Blocks from sequence (Joseph et al., 2010a). The principle was to train and validate one species specific databank and assess the prediction on the other databanks. It gives an idea of the influence of species specific preferences on prediction efficiency. 100 independent simulations have been carried out, as in one of our earlier works (Tyagi et al., 2009b). Figure 6 summarizes the results of this analysis.

For the training based on the amino acid preferences observed in the *NR* databank, the average prediction rate was higher for the non-redundant set (~34%) when compared to the species specific validation sets. The least prediction rate was obtained in the case of *Pf* (~30%) for which highly diverged sequence-structure relationships were observed, as

explained above. Predictions for *Sc* and *At* are slightly lower than the *NR* prediction rate (~32%).

When species specific training sets were used for the prediction, better prediction rates were obtained for the test sets from the same species (see supplementary material 6). The prediction rates were highly sensitive to the random choice of protein sequences, underlining the need for more protein structures for *Plasmodium falciparum* or *Arabidopsis thaliana*. Even though these preliminary results need to be assessed in-depth, they give good insights on the idea that the knowledge of species specific sequence-structure relationship can be used to improve the efficiency of prediction algorithms.

5. Conclusion

Our work sheds light into the fact that the sequence-structure relationship observed within a species can have considerable degree of variation when compared to the global preferences. This idea can have a positive implication in prediction and threading approaches, many of which make use of sequence alignments (Yang et al., 2008; Zhang, 2008). Here, we have shown the importance of specie dependence on protein sequence-structure relationship. Adjustments could be made in regards to the species under consideration, our future works will mainly focus in this direction. As the knowledge of the three dimensional structures of proteins has the potential to accelerate drug discovery (Congreve et al., 2005; Doppelt-Azeroual et al., 2009), recent developments in bioinformatics that mainly aim in filling the gap between sequence and structure space, have greatly transformed this research area (Doppelt et al., 2007; Hajduk, 2006; Moriaud et al., 2009). Thus, our work could be supportive in such approaches.

Hence this study puts emphasis on the fact that variations in protein sequence-structure relationships could exist between species with biased or unbiased amino acid compositions, demonstrated by the comparative analysis of *A. thaliana*, *S. cerevisiae* and *P. falciparum*. The efficiency of using a precise local structure description with the Protein Block approach is also underlined, and this approach appears to be more informative in the particular case of the *P. falciparum*. In addition, our preliminary results enable the consideration of other interesting tracks either in optimizing prediction methods or in homology modeling (Birkholtz et al., 2008).

Acknowledgments

The authors are deeply indebted to the excellent work done by H el ene Valadi e on this project. This work was supported by grants from the Atomic Energy Commission (CEA), Minist ere de la Recherche, Universit e Paris Diderot – Paris 7, the Institute of Blood Transfusion (INTS) and the Institute for Health and Medical Research (INSERM). APJ has a grant from CEFIPRA number 3903-E. AdB acknowledges to CEFIPRA for collaborative grant (number 3903-E).

Figures

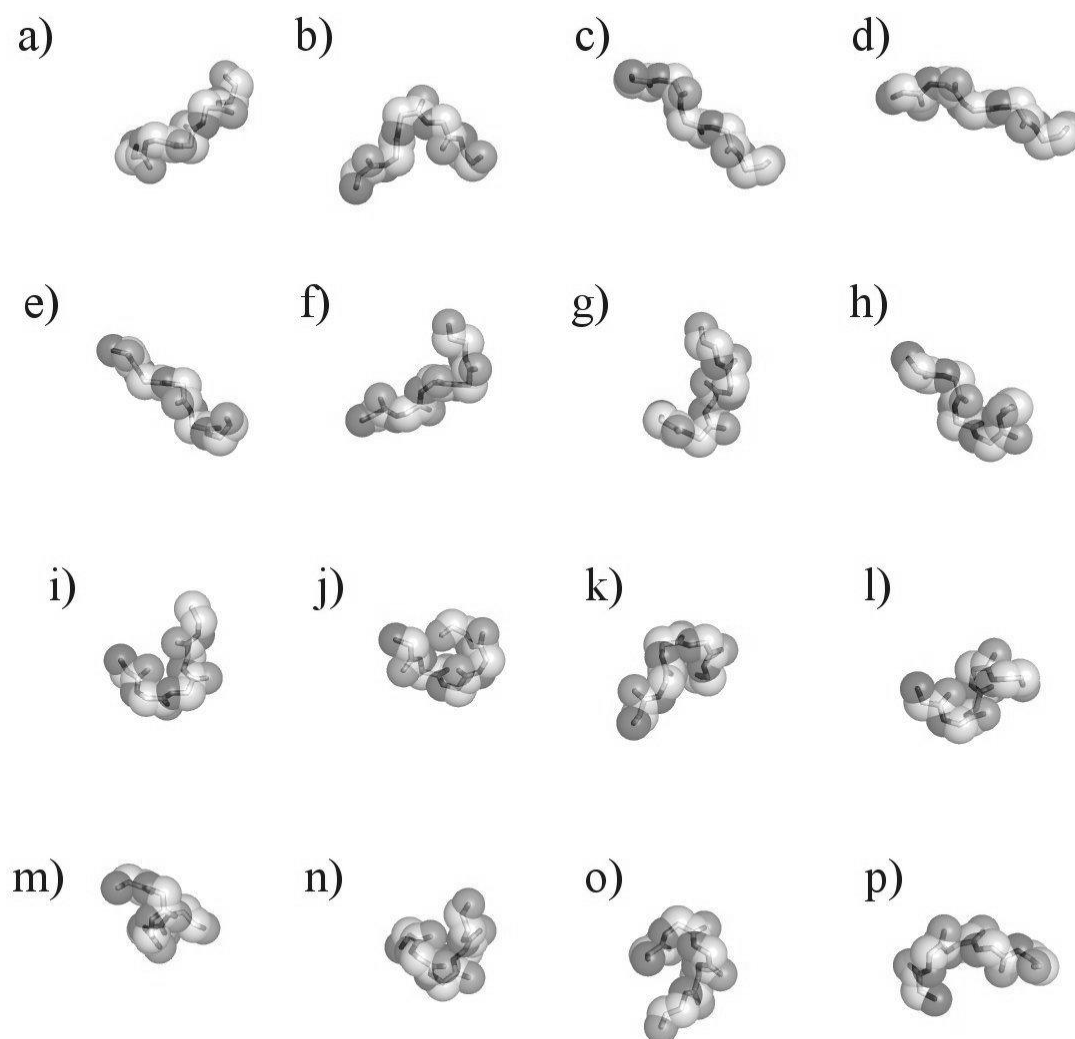


Figure 1 – *Protein Blocks*. From left to right and top to bottom, PyMol images (DeLano, 2002) of the 16 Protein Blocks of the structural alphabet (de Brevern et al., 2000). Each prototype is five residues long and corresponds to a series of eight dihedral angles (ϕ, ψ). The PBs *m* and *d* can be roughly described as prototypes for the central α -helix and the central β -strand, respectively. For each PB, the preferred N terminal conformation is on the left and the C-terminal on the right.

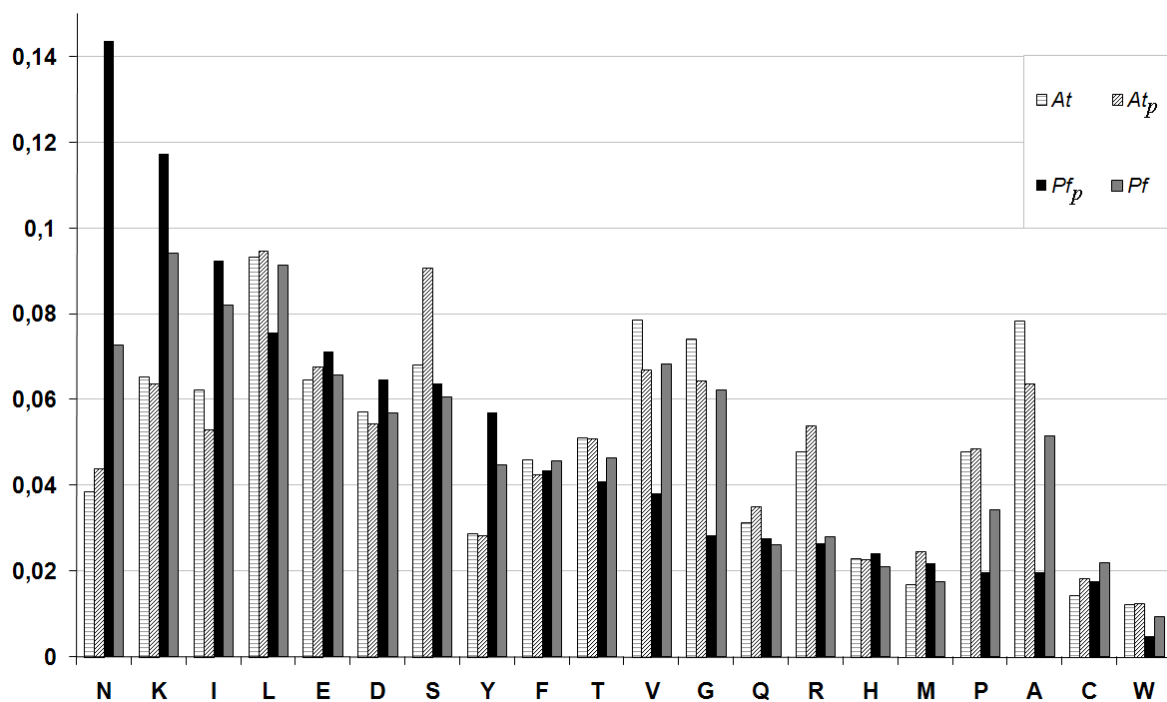


Figure 2 - Comparison of amino acid distributions. Amino acid distributions observed for *P. falciparum* and *A. thaliana* proteomes (noted Pf_p and At_p) and also in the available structures of *P. falciparum* and *A. thaliana* (noted Pf and At). Amino acids are ranked on the basis of their frequencies in the *P. falciparum* proteome.

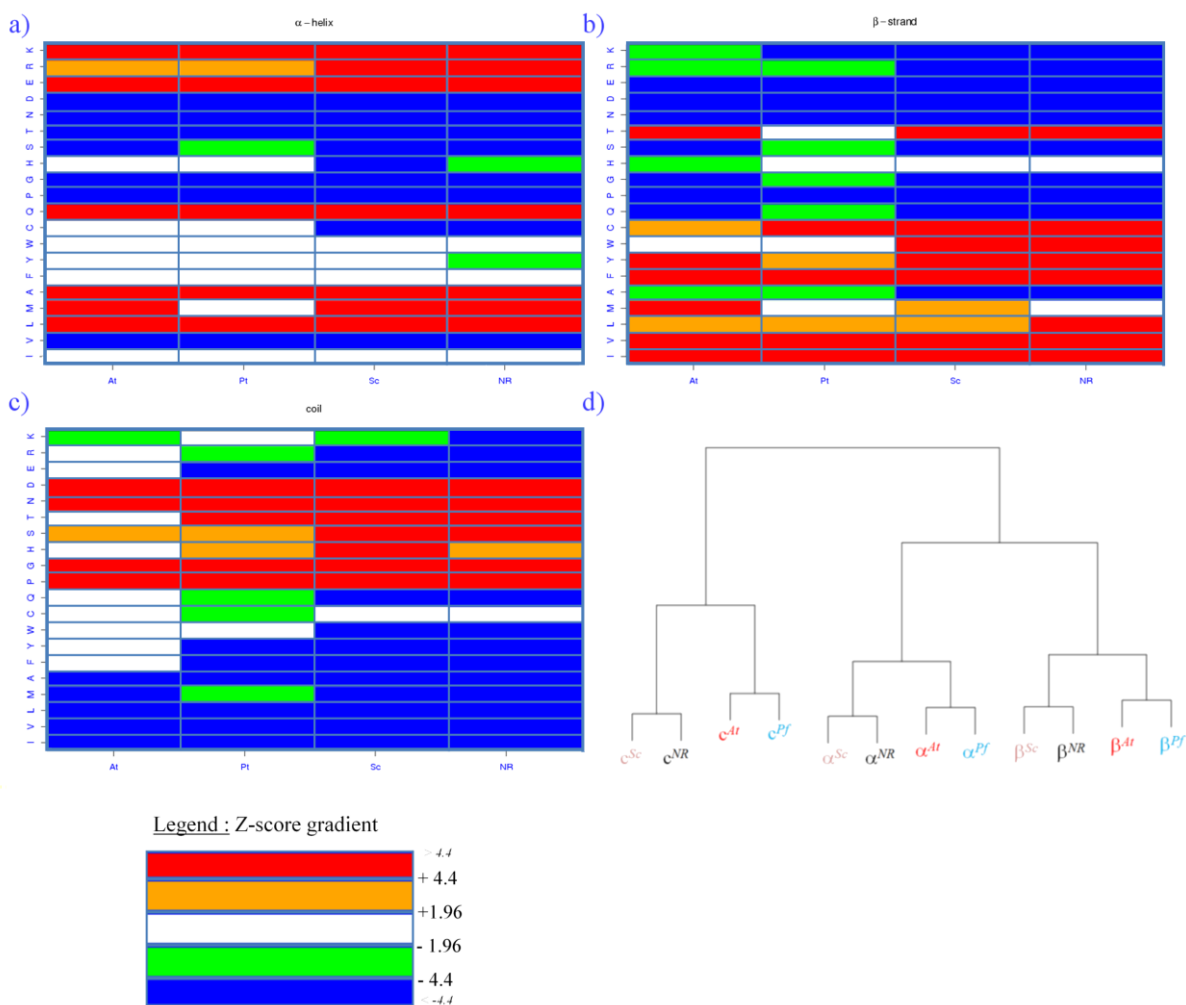


Figure 3 – Amino acid preferences of classical secondary structures. (a-c) Z-scores associated with the central positions of α -helix, coil and β -strand as assigned by DSSP, for the *At*, *Pf*, *Sc* and *NR* databanks. Each color indicates a Z-Score range; (blue): Z-scores $< (-4.4)$, (green): $(-4.4) < Z$ -scores $< (-1.96)$, (white): $(-1.96) < Z$ -scores $< (1.96)$, (orange): $(1.96) < Z$ -scores $< (4.4)$, and (red): Z-scores $> (+4.4)$. A legend for color scale is also provided. The amino acids are labeled on the vertical axis in the following order: IVLMAFYWCQPGHSTNDEK. (d) Hierarchical clustering of the preferences (3a-3c) associated with the secondary structures in *At*, *Pf*, *Sc* and *NR* databanks .

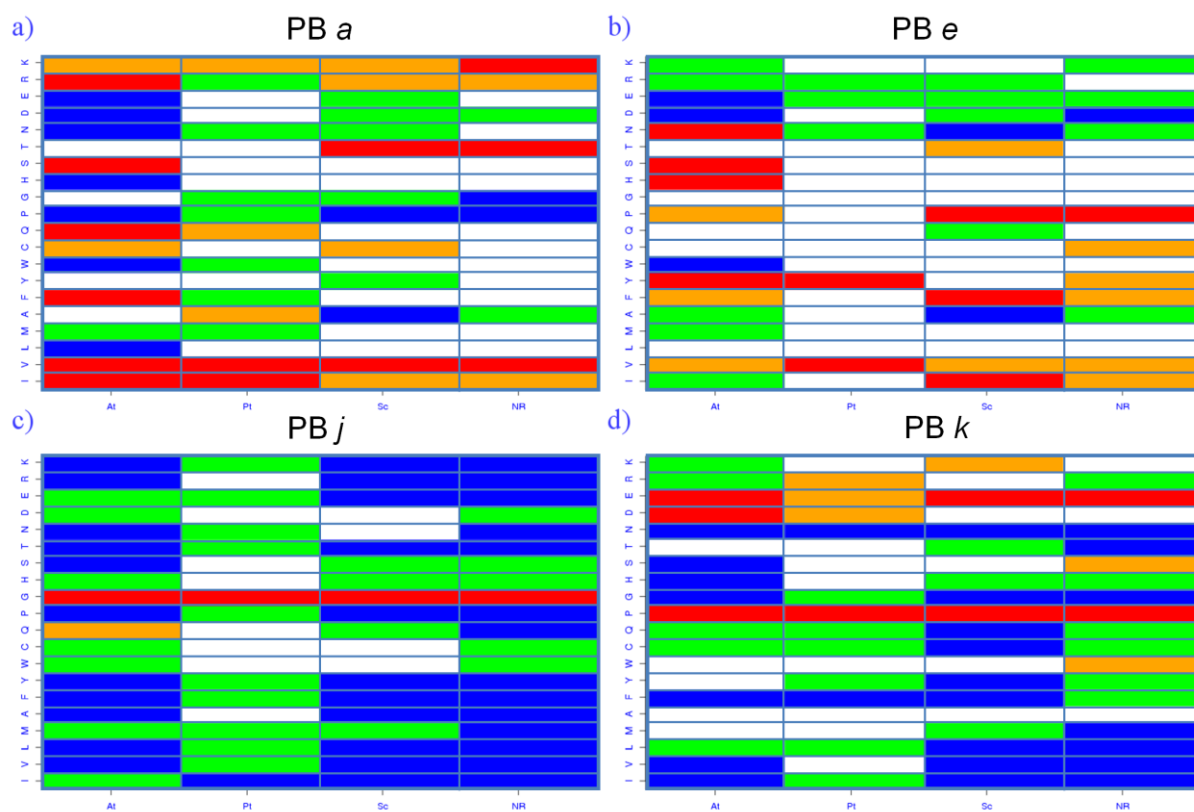


Figure 4 – Amino acid preferences of PBs. (a-d) Z-scores associated with the central positions of PB *a*, PB *e*, PB *j* and PB *k* respectively, for *At*, *Pf*, *Sc* and NR (see Figure 3 legend for details).

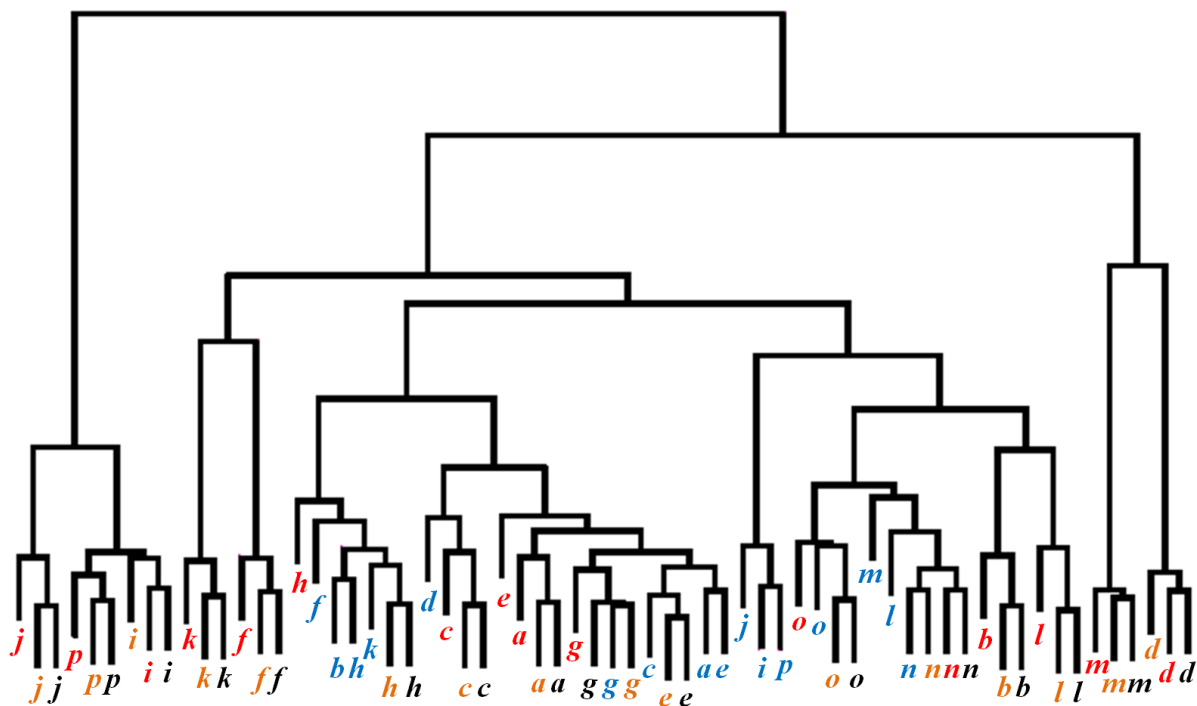


Figure 5 – Clustering of the amino acid sequence – protein structure relationships associated with PBs. Hierarchical clustering of preferences of PB central positions, seen in *At* (red), *Pf* (blue), *Sc* (pink) and *NR* (black) databanks. The PBs are noted by their respective alphabets (from *a* to *p*).

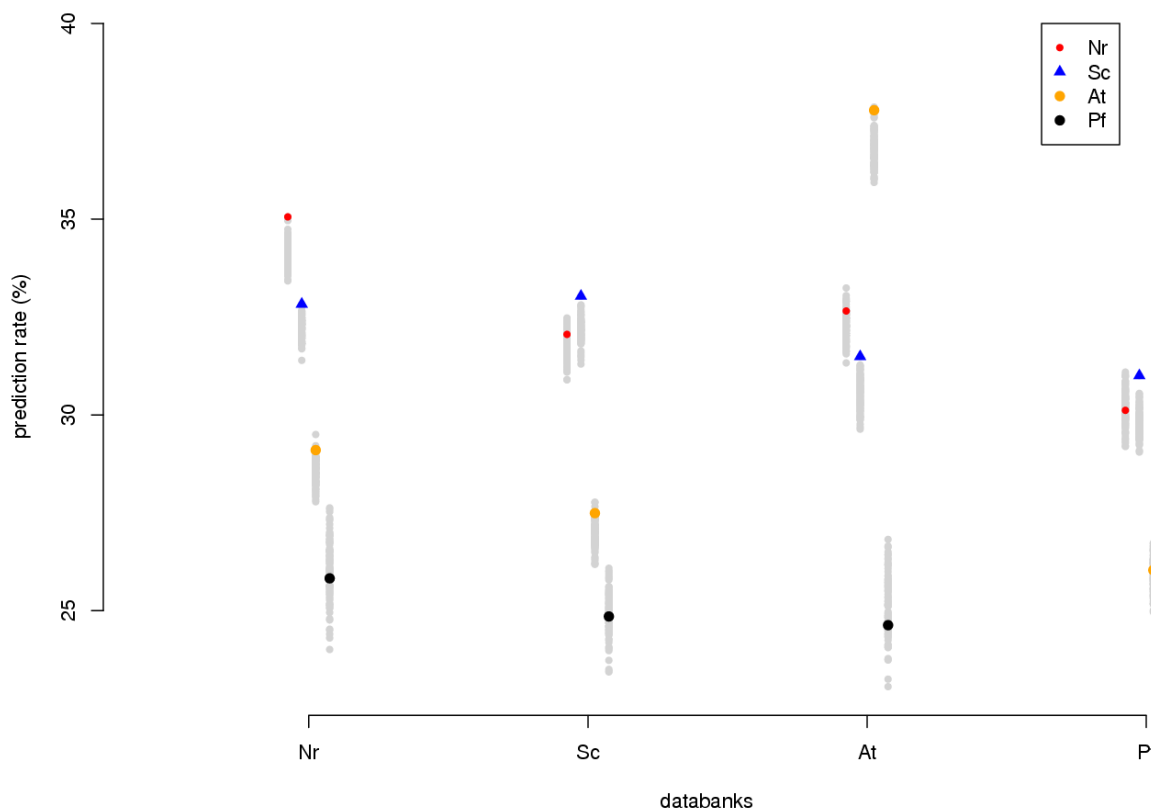


Figure 6 – *PBs prediction and influence of specie context*. 100 independent simulations have been carried out to predict PBs using a Bayesian prediction procedure (de Brevern et al., 2000; Etchebest et al., 2005; Tyagi et al., 2009b). From left to right (x axis), each label (*Nr*, *Sc*, *At* and *Pf*) indicates databank on which the training is carried out. For each of the trainings, the prediction rates on on all databanks are plotted. The 100 prediction rates are shown in grey while a selected prediction for *Nr* is in red, for *Sc* in blue, for *At* in orange and for *Pf* in black (see Supplementary data 6).

Tables

	<i>NR</i>	<i>Sc</i>	<i>At</i>	<i>Pf</i>
I	5.50	6.99	6.18	8.26
V	7.04	6.58	8.11	6.77
L	8.87	9.81	<i>9.40</i>	9.22
M	2.05	1.58	1.28	1.73
A	8.71	6.39	7.50	5.03
F	4.09	<i>4.46</i>	4.55	4.50
Y	3.54	<i>3.77</i>	2.97	4.46
W	1.52	1.18	<i>1.42</i>	0.94
C	1.35	<i>1.28</i>	<i>1.43</i>	2.23
Q	3.81	<i>3.54</i>	3.16	2.63
P	4.68	4.16	<i>4.86</i>	3.53
G	7.75	5.88	7.65	6.13
H	2.38	<i>2.19</i>	2.50	2.08
S	5.76	6.86	6.94	6.02
T	5.60	<i>5.48</i>	4.91	4.70
N	4.33	5.20	3.85	7.23
D	5.94	<i>6.11</i>	<i>5.71</i>	5.80
E	6.52	6.80	6.62	6.59
R	4.97	4.25	<i>4.81</i>	2.84
K	5.59	7.41	6.15	9.33

Table 1 – *Amino acid frequencies of the four databanks.* In bold red are highlighted the amino acids which have a variation in frequency, greater than 25%, when compared to the *NR* databank. Variations between 10-25% are shown in italics.

		<i>NR</i>	<i>Sc</i>	<i>At</i>	<i>Pf</i>
secondary structures	α -helix	37.37	40.77	36.37	39.48
	coil	42.72	40.96	39.96	41.16
	β -sheet	21.54	19.85	23.67	20.91
Protein Blocks	<i>a</i>	3.90	3.28	3.42	3.39
	<i>b</i>	4.15	3.84	4.19	4.23
	<i>c</i>	7.91	7.53	7.93	7.70
	<i>d</i>	18.23	17.37	17.69	16.93
	<i>e</i>	2.34	1.64	1.88	2.07
	<i>f</i>	6.50	6.48	6.55	6.76
	<i>g</i>	1.10	1.02	1.17	1.10
	<i>h</i>	2.29	1.71	1.94	1.97
	<i>i</i>	1.79	1.30	1.50	1.57
	<i>j</i>	0.79	0.57	0.56	0.76
	<i>k</i>	5.40	5.41	5.38	5.47
	<i>l</i>	5.38	5.63	5.79	5.75
	<i>m</i>	31.63	36.03	33.61	34.18
	<i>n</i>	2.16	2.05	2.15	2.23
	<i>o</i>	2.88	2.73	2.68	2.59
	<i>p</i>	3.54	3.40	3.55	3.31

Table 2 – *Secondary structures and Protein Block frequencies of the four databanks.* Secondary structure and Protein Block frequencies were calculated for each of the 3 databases specific for *S. cerevisiae*, *A. thaliana* and *P. falciparum*, noted as *Sc*, *At* and *Pf* respectively; and for the non-redundant databank, noted as *NR*. The secondary structures were defined by DSSP, and classified in 3 states -corresponding to α -helix, β -sheet and coil (see material and methods).

		<i>central</i>	<i>miss</i>
PB <i>a</i> (+)	<i>NR</i>	iVTrK	
	<i>Sc</i>	iVcTrk	
	<i>At</i>	IVFcQSRk	<i>T</i>
	<i>Pf</i>	IVaqk	<i>TR</i>
(-)	<i>NR</i>	aPGd	
	<i>Sc</i>	AyPgnde	
	<i>At</i>	LmWPHNDE	<i>G</i>
	<i>Pf</i>	mfwpgnr	

		<i>central</i>	<i>miss</i>
PB <i>e</i> (+)	<i>NR</i>	ivfycP	
	<i>Sc</i>	IvFPt	
	<i>At</i>	vfYpHSN	<i>P</i>
	<i>Pf</i>	VY	<i>P</i>
(-)	<i>NR</i>	anDek	
	<i>Sc</i>	AqNder	
	<i>At</i>	imaWDERk	
	<i>Pf</i>	ner	<i>D</i>

		<i>central</i>	<i>miss</i>
PB <i>j</i> (+)	<i>NR</i>	G	
	<i>Sc</i>	G	
	<i>At</i>	qG	
	<i>Pf</i>	G	
(-)	<i>NR</i>	IVLMAFYwcQPhsTNdERK	
	<i>Sc</i>	IVLmAFYqPhsTERK	<i>N</i>
	<i>At</i>	iVLmAFYwcPhSTNdeRK	<i>Q</i>
	<i>Pf</i>	Ivlmfyqptnek	<i>AR</i>

		<i>central</i>	<i>miss</i>
PB <i>k</i> (+)	<i>NR</i>	wPsE	
	<i>Sc</i>	PEk	
	<i>At</i>	PDE	
	<i>Pf</i>	Pder	
(-)	<i>NR</i>	IVLMfycqGhTnr	
	<i>Sc</i>	IVLmFYCQGhtN	
	<i>At</i>	IVIFcqGHSNrk	<i>MT</i>
	<i>Pf</i>	ilFycqgN	<i>VMT</i>

Table 3 – Significant amino acid preferences in Protein Blocks. Amino acid preferences of central positions of (a) PB *a*, (b) PB *e*, (c) PB *j* and (d) PB *k*, respectively, for the *NR*, *Sc*, *At* and *Pf* databanks. Under “miss” column is given the list of amino acids whose preference is absent, compared to the *NR* databank. (+) and (-) correspond to over-representations (Z-score > 1.96) and under-representation (Z-score < -1.96) of the amino acids. If the amino acid is in uppercase, the Z-score is higher than 4.4 (or lower than -4.4). Amino acids in blue are over-represented (or under-represented) and not found over-represented (or under-represented) in the *NR* databank. Amino acid in red indicates inversion, *i.e.*, the amino acid that is over-represented in the *NR* databank is under-represented in the specialized databank or vice-versa.

Supplementary Material

Supplementary material 1 – *PDB identifiers for the 3 databanks*. List of identifiers for the three databanks, for *A. thaliana*, *P. falciparum* and *S. cerevisiae*. These identifiers correspond to X-ray crystallographic structures with less than 90% pairwise sequence identity.

Supplementary material 2 – *The high propensities of amino acids N and K in the Pf genome*.

Supplementary material 3 – *Significant amino acids in Protein Blocks*. (central) Amino acid preferences of five positions of all the 16 PBs, for the *NR*, *Sc*, *At* and *Pf* databanks – see legend of Table 3 for details.

Supplementary material 4 – *Clustering of the sequence – structure relationship of PBs*. Each of the 16 PBs (from PB *a* to PB *p*) is highlighted in the hierarchical clustering based on the amino acid preferences of PB central positions, of the 4 databanks, as depicted on the Figure 5.

Supplementary material 5 – *Hierarchical clustering of preferences at the central positions of the PBs for At, Pf, Sc and NR*.

References

- Altschul, S. F., Madden, T. L., Schaffer, A. A., Zhang, J., Zhang, Z., Miller, W., Lipman, D. J., 1997. Gapped BLAST and PSI-BLAST: a new generation of protein database search programs. *Nucleic Acids Res* 25, 3389-402.
- Anamika, Srinivasan, N., Krupa, A., 2005. A genomic perspective of protein kinases in *Plasmodium falciparum*. *Proteins* 58, 180-9, doi:10.1002/prot.20278.
- Aravind, L., Iyer, L. M., Wellems, T. E., Miller, L. H., 2003. *Plasmodium* biology: genomic gleanings. *Cell* 115, 771-85, doi:S0092867403010237 [pii].
- Aurrecochea, C., Brestelli, J., Brunk, B. P., Dommer, J., Fischer, S., Gajria, B., Gao, X., Gingle, A., Grant, G., Harb, O. S., Heiges, M., Innamorato, F., Iodice, J., Kissinger, J. C., Kraemer, E., Li, W., Miller, J. A., Nayak, V., Pennington, C., Pinney, D. F., Roos, D. S., Ross, C., Stoeckert, C. J., Jr., Treatman, C., Wang, H., 2009. PlasmoDB: a functional genomic database for malaria parasites. *Nucleic Acids Res* 37, D539-43, doi:gkn814 [pii]10.1093/nar/gkn814.
- Bastien, O., Roy, S., Marechal, E., 2005. Construction of non-symmetric substitution matrices derived from proteomes with biased amino acid distributions. *C R Biol* 328, 445-53.
- Benros, C., de Brevern, A. G., Hazout, S., 2009. Analyzing the sequence-structure relationship of a library of local structural prototypes. *J Theor Biol* 256, 215-26, doi:S0022-5193(08)00465-7 [pii]10.1016/j.jtbi.2008.08.032.
- Benros, C., de Brevern, A. G., Etchebest, C., Hazout, S., 2006. Assessing a novel approach for predicting local 3D protein structures from sequence. *Proteins* 62, 865-80.
- Berman, H., Henrick, K., Nakamura, H., 2003. Announcing the worldwide Protein Data Bank. *Nat Struct Biol* 10, 980, doi:10.1038/nsb1203-980 nsb1203-980 [pii].
- Berman, H. M., Westbrook, J., Feng, Z., Gilliland, G., Bhat, T. N., Weissig, H., Shindyalov, I. N., Bourne, P. E., 2000. The Protein Data Bank. *Nucleic Acids Res* 28, 235-42.
- Birkholtz, L., van Brummelen, A. C., Clark, K., Niemand, J., Marechal, E., Llinas, M., Louw, A. I., 2008. Exploring functional genomics for drug target and therapeutics discovery in *Plasmodia*. *Acta Trop* 105, 113-23, doi:S0001-706X(07)00262-8 [pii]10.1016/j.actatropica.2007.10.013.
- Biswas, A. K., Noman, N., Sikder, A. R., 2010. Machine learning approach to predict protein phosphorylation sites by incorporating evolutionary information. *BMC Bioinformatics* 11, 273, doi:1471-2105-11-273 [pii]10.1186/1471-2105-11-273.
- Bornot, A., Etchebest, C., de Brevern, A. G., 2009. A new prediction strategy for long local protein structures using an original description. *Proteins* 76, 570-87, doi:10.1002/prot.22370.
- Bornot, A., Etchebest, C., de Brevern, A. G., 2011. Predicting protein flexibility through the prediction of local structures. *Proteins*, in press, doi:10.1002/prot.22922.
- Brick, K., Pizzi, E., 2008. A novel series of compositionally biased substitution matrices for comparing *Plasmodium* proteins. *BMC Bioinformatics* 9, 236, doi:1471-2105-9-236 [pii]10.1186/1471-2105-9-236.
- Brylinski, M., Skolnick, J., 2008. A threading-based method (FINDSITE) for ligand-binding site prediction and functional annotation. *Proc Natl Acad Sci U S A* 105, 129-34, doi:0707684105 [pii]10.1073/pnas.0707684105.
- Congreve, M., Murray, C. W., Blundell, T. L., 2005. Structural biology and drug discovery. *Drug Discov Today* 10, 895-907, doi:S1359-6446(05)03484-7 [pii]10.1016/S1359-6446(05)03484-7.

- de Brevern, A. G., 2005. New assessment of a structural alphabet. *In Silico Biol* 5, 283-9.
- de Brevern, A. G., 2010. 3D structural models of transmembrane proteins. *Methods Mol Biol* 654, 387-401, doi:10.1007/978-1-60761-762-4_20.
- de Brevern, A. G., Hazout, S., 2001. Compacting local protein folds with a "hybrid protein model". *Theo Chem Acc* 106, 36-47.
- de Brevern, A. G., Hazout, S., 2003. 'Hybrid protein model' for optimally defining 3D protein structure fragments. *Bioinformatics* 19, 345-53.
- de Brevern, A. G., Etchebest, C., Hazout, S., 2000. Bayesian probabilistic approach for predicting backbone structures in terms of protein blocks. *Proteins* 41, 271-87.
- de Brevern, A. G., Valadie, H., Hazout, S., Etchebest, C., 2002. Extension of a local backbone description using a structural alphabet: a new approach to the sequence-structure relationship. *Protein Sci* 11, 2871-86.
- de Brevern, A. G., Etchebest, C., Benros, C., Hazout, S., 2007. "Pinning strategy": a novel approach for predicting the backbone structure in terms of protein blocks from sequence. *J Biosci* 32, 51-70.
- de Brevern, A. G., Autin, L., Colin, Y., Bertrand, O., Etchebest, C., 2009. In silico studies on DARC. *Infect Disord Drug Targets* 9, 289-303.
- de Brevern, A. G., Benros, C., Gautier, R., Valadie, H., Hazout, S., Etchebest, C., 2004. Local backbone structure prediction of proteins. *In Silico Biol* 4, 381-6.
- DeLano, W. L. T., 2002. The PyMOL Molecular Graphics System DeLano Scientific, San Carlos, CA, USA. <http://www.pymol.org>.
- DePristo, M. A., Zilversmit, M. M., Hartl, D. L., 2006. On the abundance, amino acid composition, and evolutionary dynamics of low-complexity regions in proteins. *Gene* 378, 19-30, doi:S0378-1119(06)00253-8 [pii] 10.1016/j.gene.2006.03.023.
- Dong, Q., Wang, X., Lin, L., 2008. Prediction of protein local structures and folding fragments based on building-block library. *Proteins* 72, 353-66, doi:10.1002/prot.21931.
- Doppelt-Azeroual, O., Moriaud, F., Delfaud, F., de Brevern, A. G., 2009. Analysis of HSP90 related folds with MED-SuMo classification approach. *Drug Design, Development and Therapy* 3, 59-72.
- Doppelt, O., Moriaud, F., Bornot, A., de Brevern, A. G., 2007. Functional annotation strategy for protein structures. *Bioinformation* 1, 357-9.
- Dudev, M., Lim, C., 2007. Discovering structural motifs using a structural alphabet: application to magnesium-binding sites. *BMC Bioinformatics* 8, 106.
- Dumontier, M., Michalickova, K., Hogue, C. W., 2002. Species-specific protein sequence and fold optimizations. *BMC Bioinformatics* 3, 39.
- Etchebest, C., Benros, C., Hazout, S., de Brevern, A. G., 2005. A structural alphabet for local protein structures: improved prediction methods. *Proteins* 59, 810-27.
- Etchebest, C., Benros, C., Bornot, A., Camproux, A. C., de Brevern, A. G., 2007. A reduced amino acid alphabet for understanding and designing protein adaptation to mutation. *Eur Biophys J* 36, 1059-69.
- Faure, G., Bornot, A., de Brevern, A. G., 2009. Analysis of protein contacts into Protein Units. *Biochimie* 91, 876-87, doi:S0300-9084(09)00106-0 [pii] 10.1016/j.biochi.2009.04.008.
- Fourrier, L., Benros, C., de Brevern, A. G., 2004. Use of a structural alphabet for analysis of short loops connecting repetitive structures. *BMC Bioinformatics* 5, 58.
- Gardner, M. J., Hall, N., Fung, E., White, O., Berriman, M., Hyman, R. W., Carlton, J. M., Pain, A., Nelson, K. E., Bowman, S., Paulsen, I. T., James, K., Eisen, J. A., Rutherford, K., Salzberg, S. L., Craig, A., Kyes, S., Chan, M. S., Nene, V., Shallom,

- S. J., Suh, B., Peterson, J., Angiuoli, S., Pertea, M., Allen, J., Selengut, J., Haft, D., Mather, M. W., Vaidya, A. B., Martin, D. M., Fairlamb, A. H., Fraunholz, M. J., Roos, D. S., Ralph, S. A., McFadden, G. I., Cummings, L. M., Subramanian, G. M., Mungall, C., Venter, J. C., Carucci, D. J., Hoffman, S. L., Newbold, C., Davis, R. W., Fraser, C. M., Barrell, B., 2002. Genome sequence of the human malaria parasite *Plasmodium falciparum*. *Nature* 419, 498-511, doi:10.1038/nature01097 nature01097 [pii].
- Ghozlane, A., Joseph, A. P., Bornot, A., de Brevern, A. G., 2009. Analysis of protein chameleon sequence characteristics. *Bioinformatics* 8.
- Hajduk, P. J., 2006. Puzzling through fragment-based drug design. *Nat Chem Biol* 2, 658-9, doi:nchembio1206-658 [pii] 10.1038/nchembio1206-658.
- Hunter, C. G., Subramaniam, S., 2003. Protein local structure prediction from sequence. *Proteins* 50, 572-9.
- Illergard, K., Callegari, S., Elofsson, A., 2010. MPRAP: an accessibility predictor for α -helical transmembrane proteins that performs well inside and outside the membrane. *BMC Bioinformatics* 11, 333, doi:1471-2105-11-333 [pii] 10.1186/1471-2105-11-333.
- Jones, D. T., 1999. Protein secondary structure prediction based on position-specific scoring matrices. *J Mol Biol* 292, 195-202.
- Joseph, A. P., Bornot, A., de Brevern, A. G., 2010a. Local Structure Alphabets. In: Rangwala, H., Karypis, G., Eds.), *Protein Structure Prediction* wiley, pp. in press.
- Joseph, A. P., Agarwal, G., Mahajan, S., Gelly, J.-C., Swapna, L. S., Offmann, B., Cadet, F., Bornot, A., Tyagi, M., Valadié, H., Schneider, B., Cadet, F., Srinivasan, N., de Brevern, A. G., 2010b. A short survey on Protein Blocks. *Biophysical Reviews* 2, 137-145.
- Joubert, Y., Joubert, F., 2008. A structural annotation resource for the selection of putative target proteins in the malaria parasite. *Malar J* 7, 90, doi:1475-2875-7-90 [pii] 10.1186/1475-2875-7-90.
- Kabsch, W., Sander, C., 1983. Dictionary of protein secondary structure: pattern recognition of hydrogen-bonded and geometrical features. *Biopolymers* 22, 2577-637.
- Karchin, R., 2003. Evaluating local structure alphabets for protein structure prediction (PhD). Vol. PhD, pp. 301.
- Karchin, R., Cline, M., Mandel-Gutfreund, Y., Karplus, K., 2003. Hidden Markov models that use predicted local structure for fold recognition: alphabets of backbone geometry. *Proteins* 51, 504-14.
- Kelley, L. A., Sternberg, M. J., 2009. Protein structure prediction on the Web: a case study using the Phyre server. *Nat Protoc* 4, 363-71, doi:nprot.2009.2 [pii] 10.1038/nprot.2009.2.
- Kohonen, T., 2001. *Self-Organizing Maps* (3rd edition). Springer.
- Kulangara, C., Kajava, A. V., Corradin, G., Felger, I., 2009. Sequence conservation in *Plasmodium falciparum* α -helical coiled coil domains proposed for vaccine development. *PLoS One* 4, e5419, doi:10.1371/journal.pone.0005419.
- Kyrpides, N. C., 1999. Genomes OnLine Database (GOLD 1.0): a monitor of complete and ongoing genome projects world-wide. *Bioinformatics* 15, 773-4.
- Lattman, E., 2004. The state of the Protein Structure Initiative. *Proteins* 54, 611-5, doi:10.1002/prot.20000.
- Li, Q., Zhou, C., Liu, H., 2009. Fragment-based local statistical potentials derived by combining an alphabet of protein local structures with secondary structures and solvent accessibilities. *Proteins* 74, 820-36, doi:10.1002/prot.22191.

- Liolios, K., Mavromatis, K., Tavernarakis, N., Kyripides, N. C., 2008. The Genomes On Line Database (GOLD) in 2007: status of genomic and metagenomic projects and their associated metadata. *Nucleic Acids Res* 36, D475-9.
- Madera, M., Calmus, R., Thiltgen, G., Karplus, K., Gough, J., 2010. Improving protein secondary structure prediction using a simple k-mer model. *Bioinformatics* 26, 596-602, doi:btq020 [pii] 10.1093/bioinformatics/btq020.
- Marashi, S. A., Behrouzi, R., Pezeshk, H., 2007. Adaptation of proteins to different environments: a comparison of proteome structural properties in *Bacillus subtilis* and *Escherichia coli*. *J Theor Biol* 244, 127-32, doi:10.1016/j.jtbi.2006.07.021.
- Martin, J., de Brevern, A. G., Camproux, A. C., 2008. In silico local structure approach: a case study on outer membrane proteins. *Proteins* 71, 92-109, doi:10.1002/prot.21659.
- Moriaud, F., Doppelt-Azeroual, O., Martin, L., Oguievetskaia, K., Koch, K., Vorotyntsev, A., Adcock, S. A., Delfaud, F., 2009. Computational Fragment-Based Approach at PDB Scale by Protein Local Similarity. *J Chem Inf Model*, doi:10.1021/ci8003094 [pii].
- Nidhi, T., Swapna, L. S., Mohanty, S., Agarwal, G., Gowri, V. S., Anamika, K., Priya, M. L., Krishnadev, O., Srinivasan, N., 2009. Evolutionary divergence of *Plasmodium falciparum*: Sequences, protein-protein interactions, pathways and processes. *Infectious Diseases - Drug Targets* 3.
- Offmann, B., Tyagi, M., de Brevern, A. G., 2007. Local Protein Structures. *Current Bioinformatics* 3, 165-202.
- Paila, U., Kondam, R., Ranjan, A., 2008. Genome bias influences amino acid choices: analysis of amino acid substitution and re-compilation of substitution matrices exclusive to an AT-biased genome. *Nucleic Acids Res* 36, 6664-75, doi:gkn635 [pii] 10.1093/nar/gkn635.
- Pauling, L., Corey, R. B., 1951. The pleated sheet, a new layer configuration of polypeptide chains. *Proc Natl Acad Sci U S A* 37, 251-6.
- Pauling, L., Corey, R. B., Branson, H. R., 1951. The structure of proteins; two hydrogen-bonded helical configurations of the polypeptide chain. *Proc Natl Acad Sci U S A* 37, 205-11.
- Pollastri, G., Przybylski, D., Rost, B., Baldi, P., 2002. Improving the prediction of protein secondary structure in three and eight classes using recurrent neural networks and profiles. *Proteins* 47, 228-35.
- Pylouster, J., Bornot, A., Etchebest, C., de Brevern, A. G., 2010. Influence of assignment on the prediction of transmembrane helices in protein structures. *Amino Acids* 39, 1241-54, doi:10.1007/s00726-010-0559-6.
- Rabiner, L. R., 1989. A tutorial on hidden Markov models and selected application in speech recognition. *Proceedings of the IEEE* 77, 257-286.
- Rangwala, H., Kauffman, C., Karypis, G., 2009. svmPRAT: SVM-based protein residue annotation toolkit. *BMC Bioinformatics* 10, 439, doi:1471-2105-10-439 [pii] 10.1186/1471-2105-10-439.
- Salzemann, J., Botha, M., Dacosta, A., Degliesposti, G., Isea, R., Kim, D., Maass, A., Kenyon, C., Rastelli, G., Hofmann-Apitius, M., Breton, V., 2009. WISDOM-II: Screening against multiple targets implicated in malaria using computational grid infrastructures. *Malar J* 8, 88.
- Sander, O., Sommer, I., Lengauer, T., 2006. Local protein structure prediction using discriminative models. *BMC Bioinformatics* 7, 14.
- Service, R. F., 2008. Structural biology. Protein structure initiative: phase 3 or phase out. *Science* 319, 1610-3, doi:319/5870/1610 [pii] 10.1126/science.319.5870.1610.

- Singer, G. A., Hickey, D. A., 2000. Nucleotide bias causes a genomewide bias in the amino acid composition of proteins. *Mol Biol Evol* 17, 1581-8.
- Thomas, A., Deshayes, S., Decaffmeyer, M., Van Eyck, M. H., Charlotiaux, B., Brasseur, R., 2006. Prediction of peptide structure: how far are we? *Proteins* 65, 889-97.
- Tung, C. H., Huang, J. W., Yang, J. M., 2007. Kappa-alpha plot derived structural alphabet and BLOSUM-like substitution matrix for rapid search of protein structure database. *Genome Biol* 8, R31.
- Tyagi, M., de Brevern, A. G., Srinivasan, N., Offmann, B., 2008. Protein structure mining using a structural alphabet. *Proteins* 71, 920-37, doi:10.1002/prot.21776.
- Tyagi, M., Bornot, A., Offmann, B., de Brevern, A. G., 2009a. Analysis of loop boundaries using different local structure assignment methods. *Protein Sci* 18, 1869-81, doi:10.1002/pro.198.
- Tyagi, M., Bornot, A., Offmann, B., de Brevern, A. G., 2009b. Protein short loop prediction in terms of a structural alphabet. *Comput Biol Chem* 33, 329-33, doi:S1476-9271(09)00051-6 [pii] 10.1016/j.compbiolchem.2009.06.002.
- Tyagi, M., Sharma, P., Swamy, C. S., Cadet, F., Srinivasan, N., de Brevern, A. G., Offmann, B., 2006. Protein Block Expert (PBE): a web-based protein structure analysis server using a structural alphabet. *Nucleic Acids Res* 34, W119-23, doi:34/suppl_2/W119 [pii] 10.1093/nar/gkl199.
- Unger, R., Harel, D., Wherland, S., Sussman, J. L., 1989. A 3D building blocks approach to analyzing and predicting structure of proteins. *Proteins* 5, 355-73.
- Wang, G.-Z., Lercher, m. J., 2009. Biased amino acid composition in warm-blooded animals. *Nature Proceedings*.
- Xu, J., Jiao, F., Yu, L., 2008. Protein structure prediction using threading. *Methods Mol Biol* 413, 91-121, doi:1-59745-574-1:91 [pii].
- Xue, B., Dunbrack, R. L., Williams, R. W., Dunker, A. K., Uversky, V. N., 2010. PONDR-FIT: a meta-predictor of intrinsically disordered amino acids. *Biochim Biophys Acta* 1804, 996-1010, doi:S1570-9639(10)00013-0 [pii] 10.1016/j.bbapap.2010.01.011.
- Yang, Y. D., Park, C., Kihara, D., 2008. Threading without optimizing weighting factors for scoring function. *Proteins* 73, 581-96, doi:10.1002/prot.22082.
- Yu, Y. K., Wootton, J. C., Altschul, S. F., 2003. The compositional adjustment of amino acid substitution matrices. *Proc Natl Acad Sci U S A* 100, 15688-93, doi:10.1073/pnas.2533904100 2533904100 [pii].
- Zhang, Y., 2008. Progress and challenges in protein structure prediction. *Curr Opin Struct Biol* 18, 342-8, doi:S0959-440X(08)00034-1 [pii] 10.1016/j.sbi.2008.02.004.
- Zimmermann, O., Hansmann, U. H., 2008. LOCUSTRA: accurate prediction of local protein structure using a two-layer support vector machine approach. *J Chem Inf Model* 48, 1903-8, doi:10.1021/ci800178a.