
Water-mediated ionic interactions in protein structures

R SABARINATHAN, K AISHWARYA, R SARANI, M KIRTI VAISHNAVI and K SEKAR*

Bioinformatics Centre, Centre of Excellence in Structural Biology and Bio-computing, Indian Institute of Science, Bangalore 560 012, India

**Corresponding author (Fax, +91-080-23600683/23600551; Email, sekar@physics.iisc.ernet.in)*

It is well known that water molecules play an indispensable role in the structure and function of biological macromolecules. The water-mediated ionic interactions between the charged residues provide stability and plasticity and in turn address the function of the protein structures. Thus, this study specifically addresses the number of possible water-mediated ionic interactions, their occurrence, distribution and nature found in 90% non-redundant protein chains. Further, it provides a statistical report of different charged residue pairs that are mediated by surface or buried water molecules to form the interactions. Also, it discusses its contributions in stabilizing various secondary structural elements of the protein. Thus, the present study shows the ubiquitous nature of the interactions that imparts plasticity and flexibility to a protein molecule.

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1. Introduction

Water molecules play a vital role in governing biological macromolecules aiding in the stabilization of the three-dimensional architecture (Franks 2002; Chaplin 2006; Raschke 2006), dynamics (Otting *et al.* 1991; Halle 2004; Raschke 2006) and function (Eisenmesser *et al.* 2005; Smolin *et al.* 2005; Zhang *et al.* 2007). Further, the formation of water bridges between the protein complexes enhances water-mediated protein–protein interactions (Langhorst *et al.* 2000; Davey *et al.* 2002), protein–DNA interactions (Morton and Ladbury 1996; Grove 2003) and protein–ligand interactions (Poornima and Dean 1995). In addition, these interactions are involved in many biological functions like biomolecular recognition (Papoian *et al.* 2003), stabilization of interfaces and protein folding and unfolding (Papoian *et al.* 2004; Levy and Onuchic 2006). In a protein molecule, an ionic interaction is defined as an electrostatic interaction between the polar atoms of the positively (Arg, Lys and His) and negatively (Glu and Asp) charged residues. Ionic interactions play an important role in oligomerization, molecular recognition, domain motion,

thermostability and α -helix capping (Perutz 1970; Fersht 1972; Barlow and Thornton 1983; Musafia *et al.* 1995; Xu *et al.* 1997; Kumar *et al.* 2000; Kumar and Nussinov 2002; Shankar *et al.* 2007). Thus, a water-mediated ionic interaction is defined as when one or more water molecules mediate an interaction between a pair of charged residues. For example, disruption of surface salt bridges (a class of ionic interactions) by water molecules in proteins permits protein–DNA interactions (Grove 2003) because it creates the cationic surface complementary to the anionic DNA phosphate in the wrapped complex interface (Saecker and Record 2002).

Hydration of ionic interactions may lower the energy of the molecule (Pratt *et al.* 1994) as seen in the case of the enzyme cytochrome P450 (Opera *et al.* 1997). When three-dimensional protein structures were compared both at room and cryo-temperatures, it was found that the room temperature structure contained a bidentate salt bridge, whereas the cryo-temperature structure had a salt bridge mediated by a water molecule (Natesh *et al.* 2003). This implies that the mediation of water molecule plays a crucial role in the formation of some alternate arrangements in proteins. Recent studies have emphasized on the

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inclusion of water molecules when modelling protein systems (Papoian *et al.* 2003; Jiang *et al.* 2005; Van Dijk and Bonvin 2006) and in hydration-based ligand design (Poornima and Dean 1995). Some studies on the interactions formed by buried (Park and Saven 2005) and surface water molecules have also been attempted (Teyra and Pisabarro 2007; Rodier *et al.* 2005). These studies take into account not only the water-mediated ionic interactions but also long-range hydrophobic interactions and the interactions between neutral moieties. Although there has been a specific study on the contribution of water-mediated salt bridges that enhances thermal stability of the 'tail' region of GrpE protein from *E. coli* (Mehl *et al.* 2007) and thermophilic mutants (Bogin *et al.* 2002), there is not much information available about the nature and distribution of various types of water-mediated ionic interactions over a large dataset of protein structures available in its archive, Protein Data Bank, or PDB (Berman *et al.* 2000). Therefore, the present study aims at analysing the frequency of occurrence and distribution of water-mediated ionic interactions in secondary structures and the possible role it plays within and between the subunits. Such a study would contribute to the understanding of biomacromolecular interactions as well as reveals some valuable insights into protein stability and flexibility.

2. Materials and methods

2.1 Dataset

In order to reduce the bias caused by redundancy, a statistical analysis of protein structures required the use of non-redundant data. Thus, in this study, a total of 9577 unique protein chains with low mutual sequence identity of less than 90% were obtained from the non-redundant data (Hobohm and Sander 1994). While constructing the dataset, only one chain was included from a multimeric protein structure (for example, only one chain was considered in the case of a protein molecule composed of two identical chains in the asymmetric unit with sequence identity of more than 90%). Only those X-ray-analysed crystal structures whose resolution was better than or equal to 1.5 Å and R-factor better than or equal to 20% were selected and included in the dataset. After applying these quality cut-off criteria, the dataset reduced to 1021 chains corresponding to 979 proteins, of which 37 proteins had multiple chains. On the basis of Structural Classification of Proteins (SCOP) classification (Murzin *et al.* 1995), the protein chains in the dataset represent all protein classes (figure 1). The three-dimensional atomic coordinates of the protein chains were downloaded from the locally available PDB FTP anonymous server, Bioinformatics centre, Indian Institute of Science, India.

2.2 Methods

For the purpose of this study, the following simplification was made regarding the water-mediated ionic interactions. Only the crystallographic water molecules present in the PDB files are included for computation. In basic residues, N^{δ1} and N^{ε2} atoms of histidine, N^ε atom of lysine and the distal nitrogen atoms (NH1, NH2 and N^ε) of arginine were considered as potential hydrogen bond donors. On the other hand O^{δ1} and O^{δ2} of aspartate and O^{ε1} and O^{ε2} of glutamate are considered as potential hydrogen-bond acceptors among the acidic residues. In addition, free nitrogen and carbonyl oxygen atoms present in the N- and C-terminals of the protein chains were also taken into account for this study. Using hydrogen bond distance (2.6 Å to 3.5 Å) criteria, the charged residues and the water molecules involved in mediating ionic interactions were extracted from the above dataset. Then, the resulting data was validated using the program HBPLUS (McDonald and Thornton 1999) to check for hydrogen bonds between the charged atoms and water molecules by using the two following geometric criteria: (1) the donors and acceptors were seen to exist within the distance of 3.5 Å and the angle between them was greater than 90° and (2) the solvent accessible area of water molecules involved in the ionic interactions were computed using the program NACCESS (Hubbard and Thornton 1993) with a probe radius 1.4 Å. Water molecules with accessible surface area less than or equal to 2.5 Å² were considered as internal or buried water molecules, whereas the other water molecules were considered as surface water molecules. The mobility or positional spread of each water molecule was calculated using atomic displacement parameter (often referred to as B-factor). The normalized B-factor was calculated using the formula $B'_i = (B_i - \langle B \rangle) / B$, where B_i is the B-factor of each atom, $\langle B \rangle$ is the mean B-factor of protein molecule and B is the standard deviation of the B-factors (Smith *et al.* 2003). Water molecules with a low normalized B-factor hold high stability. Secondary structural assignments were predicted using the program STRIDE (Frishman and Argos 1995) and the necessary PERL scripts were developed locally to perform the above calculations.

3. Results and discussion

3.1 Distribution of charged residues

It is interesting to note that out of 1021 protein chains in the dataset, water-mediated ionic interactions are observed in 821 chains based on the two defined geometric criteria. The average percentage of charged residues (positive and negative) present in the dataset corresponds to 12% (table 1). The number of positively charged residues is approximately equal to the number of negatively charged residues in the

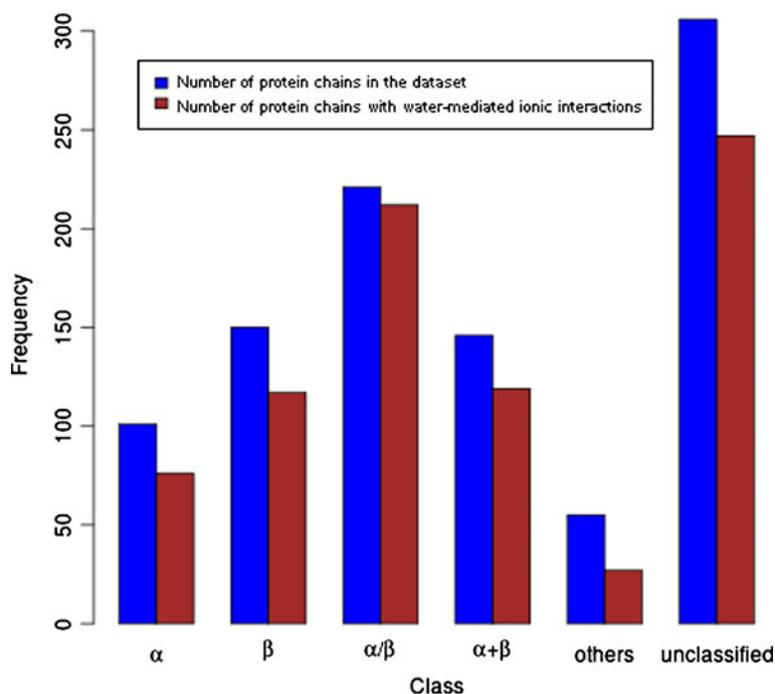


Figure 1. Distribution of different classes (based on SCOP classification) of protein chains used in the present study and the number of protein structures containing water-mediated ionic interactions.

dataset. However, the percentage of positively (32.8%) charged residues involved in the water-mediated ionic interaction are higher than the participation of negatively (24.6%) charged residues (table 1). Even though the reason for this difference is solely dependent on the hydrogen bond formation with the mediating water molecule, the positively charged residues have the highest propensity (see following section for details).

3.2 Distribution of water-mediated ionic interactions

A total of 3876 water-mediated ionic interactions were observed in 821 protein chains (interactions present within

Table 1. Number of charged residues present in the 90% non-redundant protein chains and their involvement in the water-mediated ionic interaction

Residue type	Number of charged residues in the dataset	Number of residues involved in water-mediated ionic interactions
Histidine	5125 (2.37%)	371 (7.23%)
Arginine	10434 (4.84%)	1604 (15.37%)
Lysine	11778 (5.46%)	1197 (10.16%)
Aspartate	12699 (5.89%)	1666 (13.12%)
Glutamate	13358 (6.19%)	1537 (11.51%)

and between the subunits). Water-mediated ionic interaction is represented as: a-HOH-b, where 'a' and 'b' are the positively and negatively charged residues, respectively, and HOH is the water molecule that mediates the interaction. The combined pair of residues ('a-b') is termed as a residue pair that contains water-mediated ionic interaction. Out of these, 2020 water-mediated ionic interactions were formed between basic residues (His/Lys/Arg) and an acidic aspartate residue. The remaining 1856 interactions were found with glutamate (table 2). Very few water-mediated ionic interactions were observed between the residue pairs His-Asp (248) and His-Glu (190). This might be due to the fact that the imidazole ring of histidine makes it difficult to expose both nitrogen atoms to the solvent at the same time; however, it allows the formation of a single hydrogen bond and is more significant when it is buried. Comparatively more interactions were observed between Lys-Asp and Lys-Glu (table 2). This is because lysine residue has sp^3 -hybridized N^ζ atom that can form three hydrogen bonds due to its side-chain flexibility. Finally, the highest number of interactions was observed between Arg-Asp and Arg-Glu (table 2). Three nitrogen atoms (NH1, NH2 and N^ϵ) of arginine contribute to these interactions, in which both NH1 and NH2 are often involved in a single or double hydrogen bond formation whereas N^ϵ is involved in the formation of a single hydrogen bond. In addition, the hydrogen acceptor present at the other end of aspartate and glutamate residues has two sp^2 -hybridized carboxyl oxygen atoms. Each oxygen atom can accept

Table 2. Distribution of water-mediated ionic interactions between different charged residues

	Aspartate		Glutamate		Total
	Same subunit	Different subunits	Same subunit	Different subunits	
Histidine	243	5	189	1	438
Arginine	1088	8	953	9	2058
Lysine	671	5	696	8	1380
Total	2020		1856		3876

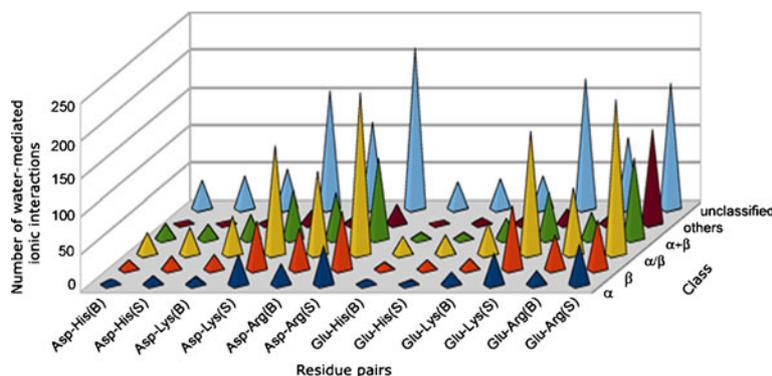
hydrogen in the form of two hydrogen bonds. Many residues make at least one hydrogen bond and a significant number of residues make two. Buried residues display an increased propensity towards the formation of two hydrogen bonds with aspartate. This leads to the distribution of different combinations of ionic interactions between charged residues. The water-mediated ionic interactions between the N-terminal nitrogen and C-terminal oxygen atoms were identified in two protein subunits.

The water molecules mediating ionic interactions were identified in both buried and surface regions of the protein structures based on its surface accessible area (Supplementary figure 1). Further, the stability of these water molecules was high, which corresponded to their low normalized B-factor. These interactions are commonly found in all classes of protein structures (figure 1). The distribution of water molecules mediating the ionic interactions in different classes of proteins is shown in figure 2. In all classes of proteins, the number of ionic interactions (figure 2) mediated by the buried and surface water molecules (between the residues pair Asp-His and Glu-His) are almost equal. However, the ionic interactions between Asp-Lys, Glu-Lys, Asp-Arg and Glu-Arg are mediated more (60%) by surface water molecules than buried water molecules (figure 2). It shows that the

distribution of charged residues such as aspartate, glutamate, lysine and arginine on the surface of the protein molecules are highly exposed for water-mediated ionic interactions and leads to the participation in biomolecular interactions.

3.3 Types of water-mediated ionic interactions

Water-mediated ionic interactions can be divided into two main types, namely, complete and incomplete, based on the number of atoms involved in the formation ionic interaction (Kumar *et al.* 2000; Shankar *et al.* 2007). As mentioned earlier, a water-mediated ionic pair contains a residue pair (an acidic and a basic residue) and a water molecule. In an ionic interaction, either all the side-chain polar atoms from both acidic and basic residues or three atoms (one from basic and two from acidic residues and vice versa) participate in the formation of a water-mediated ionic interaction. In a complete water-mediated ionic interaction, one of the hydrogen bonds formed between water molecule and charged residue is a bifurcated bond. Figure 3A depicts the complete water-mediated ionic interaction between the residue pair Arg 173–Asp 48 via HOH 215 in the three-dimensional structure of alcohol dehydrogenase [PDB-id: 1O2D] (Schwarzenbacher *et al.* 2004). When only one atom from the acidic residue and one atom from the basic residue are involved, it is called as an incomplete water-mediated ionic interaction (figure 3B). In the 90% non-redundant dataset, the percentages of complete and incomplete water-mediated ionic interactions are 11.3% and 88.7%, with respect to the total number of water-mediated ionic interactions observed (supplementary tables 1 and 2). In complete water-mediated ionic interactions, 35% of the interactions are mediated by the buried water molecules. This helps to stabilize the mobility of water molecules in the core region by forming more than one hydrogen bond with the respective charged residues. In addition, the buried charged atoms get solvated by this interaction and provide

**Figure 2.** Distribution of the water-mediated ionic interactions mediated by buried (B) and surface (S) water molecules present in different classes of proteins.

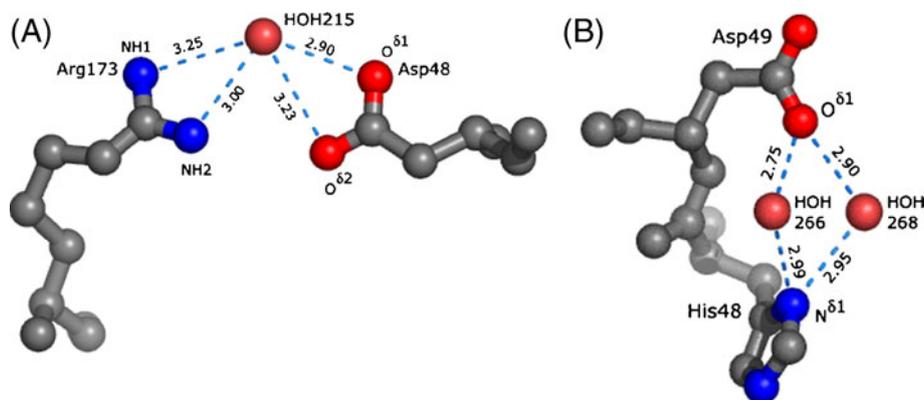


Figure 3. Types of water-mediated ionic interactions: (A) Complete water-mediated ionic interaction observed in the three-dimensional structure of alcohol dehydrogenase [PDB-id: 1O2D], between the residues Arg 173 and Asp 48 (chain A) mediated by the water molecule HOH 215. (B) Incomplete water-mediated ionic interaction found in the crystal structure of bovine pancreatic phospholipase A₂ [PDB-id: 1UNE] is formed between the catalytically important residues (His 48 and Asp 49) via two water molecules, HOH 266 and HOH 268. The distances are in Å units.

stability to the inner hydrophobic core of the protein molecule. In incomplete water-mediated ionic interactions, the total number of interactions mediated by surface water molecules is slightly higher when compared with the complete water-mediated ionic interactions. The high distribution of incomplete water-mediated ionic interactions in the surface region helps in biomolecular recognition, which in turn aids in the formation of different protein complexes. It is to be noted that the corresponding percentages of complete and incomplete water-mediated ionic interactions in the 90% non-redundant dataset is in complete reversal to the direct ionic interaction results published by our group and the values are 58% and 41%, respectively (Shankar *et al.* 2007).

Incomplete water-mediated ionic interactions possess significant functionalities as observed in the crystal structure of bovine pancreatic phospholipase A₂ (Sekar and Sundaralingam 1999). The ionic interaction is formed between two catalytically important residues His48 and Asp49 (figure 3B). Thus, it can be concluded that, due to the mediation of surface water molecules and the high flexibility, the charged residues on the surface of protein molecules get hydrated. Therefore, the great conformational or steric freedom of surface residues allows the formation of an incomplete water-mediated ionic interaction with great ease.

3.4 Networks of water-mediated ionic interactions

Instead of a single residue pair, when three or more charged residues form a water-mediated ionic interaction with at least one residue in common, it is known as a network of water-mediated ionic interactions (figure 4A–B). In the present dataset, a total of 20% of water-mediated

ionic interactions were involved in the formation of 148 different types of networks. Some of the network types, which have occurred more than two times, are listed in supplementary table 3. Among the reported 197 networks (supplementary table 3), 34 networks were formed between the polar atoms Asp-Lys-Glu, 31 networks were that of Asp-Arg-Glu combination and the networks between the Asp-Arg-Lys and Glu-Arg-Lys corresponded to 28 and 18 networks, respectively. In some cases, a common residue present in more than one network extends the network. Figure 4A depicts the three-dimensional structure of the extended network present within the subunit (chain A) of the nitrobenzene dioxygenase (PDB-id: 2BMO) (Friemann *et al.* 2005). The interactions are formed between Arg 286–Glu 169 via HOH 220, Glu 169–Lys 173 via HOH 219 and Glu 169–His 143 via HOH 143. In the above extended network, Glu 169 is common in all the three incomplete water-mediated ionic interactions and acts as a bridge to link the three individual networks, namely, (a) Arg 286–HOH 220–Glu 169–HOH 219–Lys 173, (b) Arg 286–HOH 220–Glu 169–HOH 143–His 143 and (c) Lys 173–HOH 219–Glu 169–HOH 143–His 143 to make an extended network (Pedersen *et al.* 2004). It has been reported that a total of 10% of the networks comprised extended networks. The networks within the subunits are mainly helpful in the flexibility of active site and structure stability of the protein molecules. As stated in the literature, the flexibility of the active site in Apo protein tyrosine phosphatase 1B is controlled by a network of water-mediated interaction (Pedersen *et al.* 2004). In addition, in aspartic proteinases, the water-molecule-mediated network present in the active site is conserved (Prasad and Suguna 2002).

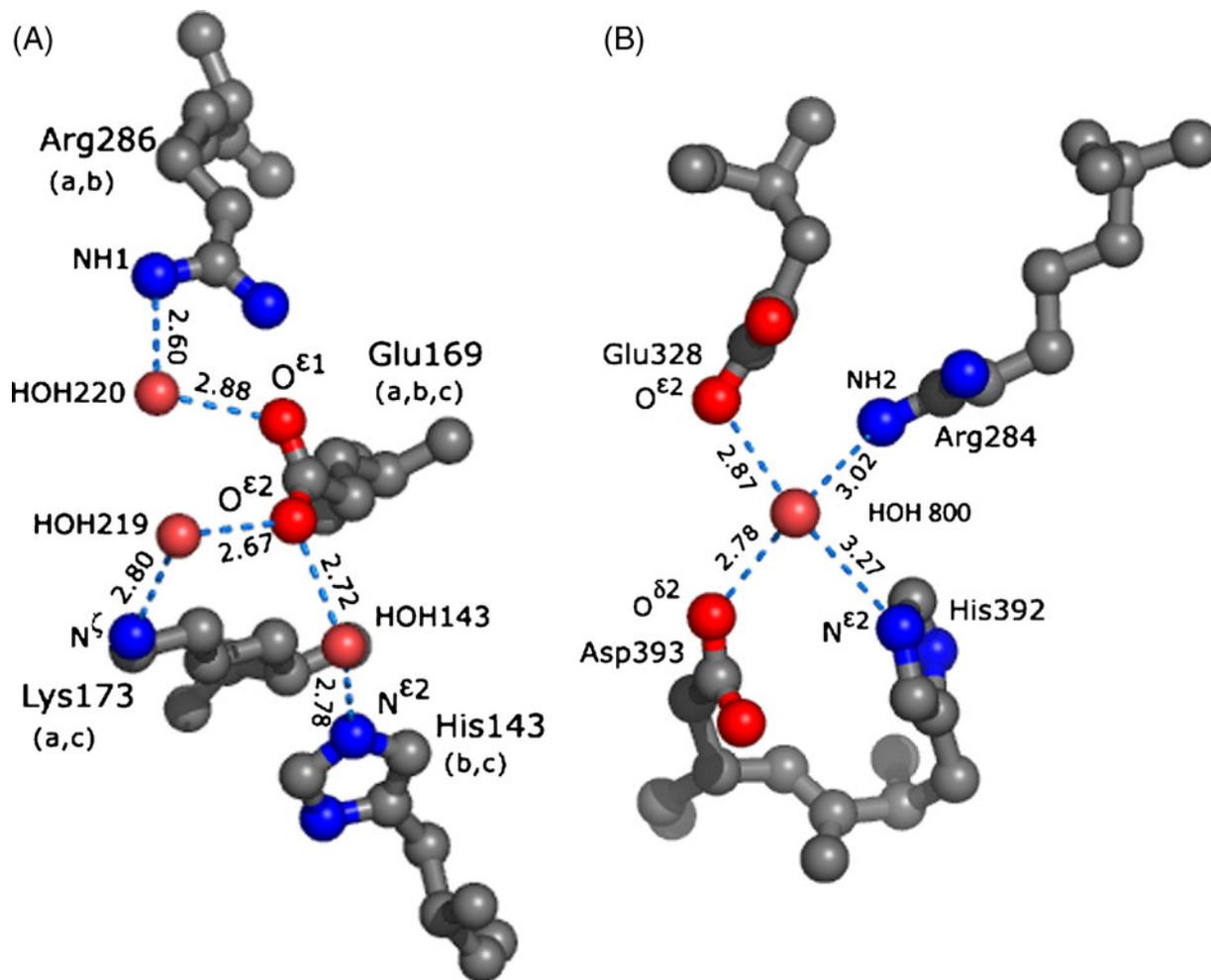


Figure 4. Networks of water-mediated ionic interactions (A) Extended network of water-mediated ionic interactions, in the crystal structure of nitrobenzene dioxygenase (PDB-id: 2BMO). The residue Glu 169 is commonly present among the three different networks (a, b and c). (B) Hubs, a water molecule HOH800 from the crystal structure of amylosucrase [PDB-id: 1G5A] act as a hub for two different water-mediated ionic interactions. The first one is between the Arg 284–Asp 393–Glu 328 via HOH 800 and the second one is between His 392–Asp 393–Glu 328 via the same water molecule HOH 800.

3.5 Hubs: A special case of networks

When two or more separated residue pairs appear to share a common (single) water molecule and when each residue pair along with that particular water molecule satisfies the criterion to form a water-mediated ionic interaction, it is called a hub (figure 4B). In the 90% non-redundant dataset, 20 different water-mediated ionic interactions are involved in the formation of four different hubs within subunits of different protein chains. A typical hub shown in figure 4B is observed in the crystal structure of amylosucrose (PDB-id: 1G5A) (Skov *et al.* 2001). Here, the water molecule HOH 800 acts as a hub for the interaction of four charged residues Glu 328, Arg 284, Asp 393 and His 392 in order to form two different water-mediated ionic interactions. Since

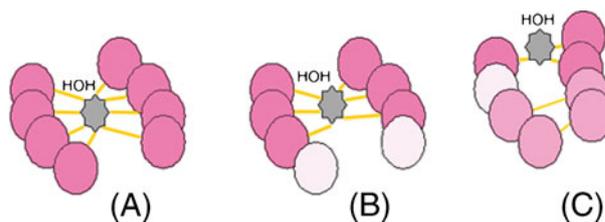


Figure 5. The diagram shows the possible movements of the residues involved in water-mediated ionic interactions. (A) Hub of water-mediated ionic interactions, (B) some interactions are broken or formed by the entry of water and (C) water molecule escapes and gives rise to ionic interaction formation (last two residue pairs). The dark coloured residues are involved in a water-mediated ionic interaction formation and the lightest ones are not involved in any interaction.

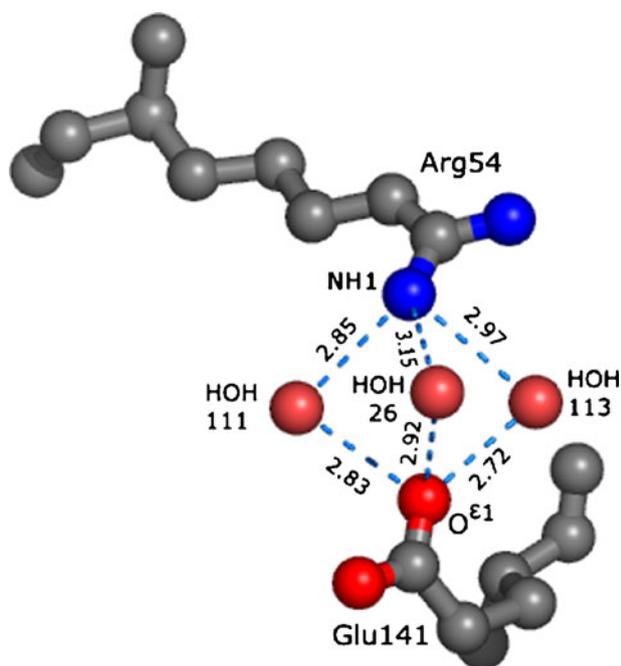


Figure 6. Cluster, a multiple water mediated ionic interaction found between the residues Arg 54 and Glu 141 found by three water molecules (HOH 111, HOH 26 and HOH 113) in the crystal structure of Alpha-tocopherol transfer protein [PDB-id: 1R5L]. The distances are in Å units.

interfacial water molecules are functionally important (Schimer and Evans 1990), inter-subunit hubs may be located near catalytic sites or may have a specific functionality (like charge redistribution). For example, the transfer of water molecules is involved in protein gating in certain enzymes (Wikstrom *et al.* 2005). The hubs observed between the dimers of Dps and its related ferritin-like structures helps in the conformational stability of the protein molecule (Ranjani *et al.* 2008).

As mentioned earlier, the water-mediated interaction between the charged residues increases the plasticity of ionic interactions. The mechanism for hydration in an

ionic interaction and its consequent plasticity has been hypothesized in figure 5. In this example, a hub comprising four residue pairs is depicted in figure 5A. Disruption of ionic interactions upon the entry of a water molecule forms the water-mediated ionic interactions. Similarly, when the water molecule exits, the water-mediated ionic interaction is broken (figure 5B). There is a reformation of the ionic interactions upon complete exit of the water molecule (figure 5C). It clearly shows the flexibility and the plasticity of the water-mediated ionic interactions.

3.6 Cluster: Multiple water-mediated ionic interactions

When two or more water molecules mediate an ionic interaction, it is called as multiple water-mediated ionic interaction and the water molecules involved are called cluster of water molecules. The definition of a multiple water-mediated ionic interaction is based on the number of water molecules involved, and so the constituent water-mediated ionic interactions can be either complete or incomplete. In the present study, a total of 57 multiple water-mediated ionic interactions were identified comprising 21 residue pairs. Most of the multiple water-mediated ionic interactions were mediated by only two water molecules (figure 3B). The rest of the interactions were mediated by three water molecules. For example, figure 6 depicts a multiple water-mediated ionic interaction between three water molecules (HOH 111, HOH 26 and HOH 113) and charged residues Arg 54 and Glu 141 in the crystal structure of alpha-tocopherol transfer protein structure (PDB-id: 1R5L) (Min *et al.* 2003). According to Royer and co-workers, a cluster of interfacial water molecules play a crucial role in the communication between the subunits (Royer *et al.* 1996). The multiple water-mediated ionic interactions are involved in the catalytic activity of the enzyme bovine pancreatic phospholipase A₂ (Shankar and Sekar 2009). Thus, it can be concluded that water-mediated ionic interactions help in maintaining the stability of the structure and facilitates inter-subunit interactions.

Table 3. Distribution of charged residues in various secondary structural elements

Secondary structure	Histidine	Arginine	Lysine	Aspartate	Glutamate	Total
α -helix (H)	120	725	481	541	762	2629
3_{10} -helix (G)	15	70	47	75	77	284
π -helix (I)	1	0	0	2	0	3
Extended conformation (E)	79	332	271	237	282	1201
Isolated bridge (B)	2	20	18	17	19	76
Turn (T)	71	241	214	499	220	1245
Coil (C)	83	216	166	295	177	937

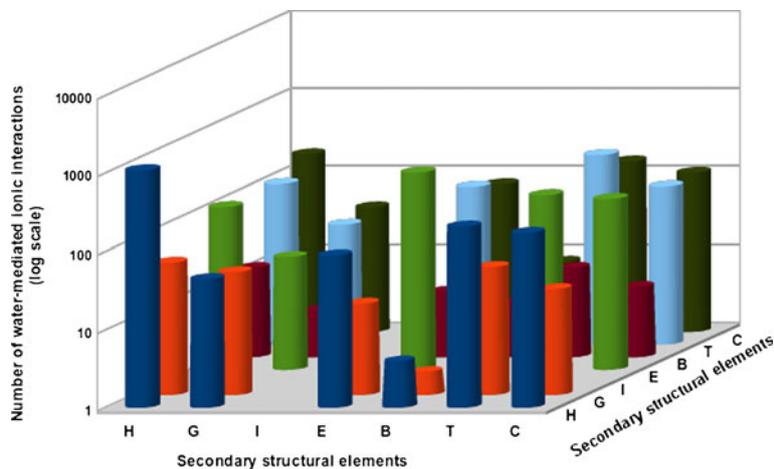


Figure 7. Distribution of water-mediated ionic interactions between the different secondary structural elements. The different secondary structural elements given in the table are H: α -helix, G: 3_{10} -helix, I: π -helix, E: Extended conformation, B: Isolated bridge, T: Turn and C: Coil.

3.7 Water-mediated ionic interactions in secondary structures

In the present study, a total of 5438 (85.30%) charged residues involved in the water-mediated ionic interactions

were found in different secondary structures (table 3). In addition, the charged residues comprised a total of 3172 (49.75%) basic and 3203 (50.24%) acidic residues. The number of residues present in α -helices was higher than any other secondary structural elements (table 3). Earlier it had

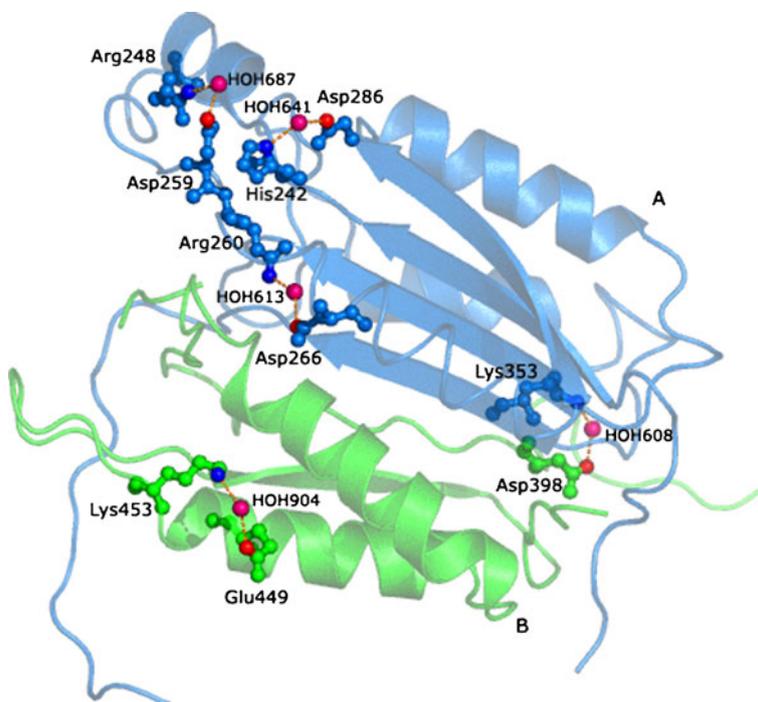


Figure 8. The water-mediated ionic interactions between the residue pairs present in different secondary structural elements of subunits A and B in the crystal structure of caspase-8 protein [PDB-id:1QTN].

been reported that direct ion pair interactions are more frequently observed in α -helices than in β -sheets (Shankar *et al.* 2007). It is a known fact that α -helices are the most abundant secondary structural element observed in proteins. Further, water-mediated ionic interactions present between different secondary structural elements were observed within and between the subunits of protein molecules (figure 7). The number of interactions between secondary structural elements like helix-helix and strand-strand corresponded to 31.84% and 23.91% (figure 7). It is evident that water-mediated ionic interactions are higher in α -helices due to intra-helical and inter-helical interactions. Inter-strand water-mediated ionic interactions effectively act as inter-sheet interactions. Due to the prevalence of hydrogen bonds, the number of water-mediated ionic interactions required to stabilize a β -sheet is low. However, α -helices are stabilized by electrostatic interaction (Vila *et al.* 1992). Thus, it can be concluded that these polar intra-helical interactions may well be treated as water-mediated ionic interactions.

The water-mediated ionic interactions observed in other secondary structural elements like helix-coil and strand-coil contributes to 15% of the total water-mediated ionic interactions. It helps to stabilize the charged residues that are present in an unstable coiled region by introducing a water-mediated ionic interaction between the corresponding residues in the stable secondary structural elements (α -helix or β -strand). Figure 8 depicts the different secondary structural combinations of water-mediated ionic interactions within and between subunits A and B of the protein molecule caspase-8 protein molecule (PDB-id: 1QTN) (Watt *et al.* 1999). As shown in the figure, the water-mediated ionic interactions among the different secondary structural elements are α -helix-coil (A-Arg 248-A-Asp 259), turn- α -helix (A-Arg 260-A-Asp 266), coil-coil (A-His 242-A-Asp 286), coil- α -helix (B-Lys 453-B-Glu 449) and turn- β -Strand (A-Lys 353-B-Asp 398). In multi-subunit proteins, the percentage of basic and acidic residues (in different secondary structures) forming water-mediated ionic interactions are 51.61% and 48.39%, respectively (data not shown). Among them, 43% of the acidic and 50% of the basic residues are present in α -helices, and the corresponding numbers are 10% and 25% in β -strands. A total of 36 different interactions are observed between the subunits. The mediated interaction contains both complete and incomplete water-mediated ionic interactions. The results of different secondary structural combinations show that water-mediated interactions are involved in the anchoring of tertiary structures. These results help to prove the earlier work of Park and Saven (2005). Similarly, the ionic interactions mediated by the water molecules at the interface region also play an important role in the formation of dimers from monomers and other biomolecular complexes.

4. Conclusion

In the present study, various patterns of water-mediated ionic interactions formed between the charged residues were classified at the atomic level. The existence of different types of water-mediated ionic interactions, charged networks, hubs and clusters are reported in the present study. Thus, water-mediated ionic interactions have a marked importance in intramolecular interaction, macromolecular association and in biomolecular recognition. Furthermore, the presence of water-mediated ionic interactions in the secondary structural elements confirms their participation in the stability of the same. Water-mediated ionic interactions also play an important role in holding the subunits in multi-subunit protein structures. Therefore, the present study is useful to understand and appreciate the role of water-mediated ionic interactions in the plasticity and flexibility of protein structures.

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