Analysis of binding site similarity, small-molecule similarity and experimental binding profiles in the human cytosolic sulfotransferase family

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ABSTRACT

Motivation: In the present work we combine computational analysis and experimental data to explore the extent to which binding site similarities between members of the human cytosolic sulfotransferase family correlate with small-molecule binding profiles. Conversely, from a small-molecule point of view, we explore the extent to which structural similarities between small molecules correlate to protein binding profiles.

Results: The comparison of binding site structural similarities and small-molecule binding profiles shows that proteins with similar small-molecule binding profiles tend to have a higher degree of binding site similarity but the latter is not sufficient to predict small-molecule binding patterns, highlighting the difficulty of predicting small-molecule binding patterns from sequence or structure. Likewise, from a small-molecule perspective, small molecules with similar protein binding profiles tend to be topologically similar but topological similarity is not sufficient to predict their protein binding patterns. These observations have important consequences for function prediction and drug design.

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1 INTRODUCTION

A large fraction of cellular and biochemical processes involve the interaction between proteins (particularly enzymes) and small molecules (molecules other than nucleic acids and polypeptides). At a molecular level, these interactions are determined by the interplay of many factors including entropic effects and various molecular forces between interacting groups present in the protein and small molecule (Israelachvili, 1992; Leckband and Israelachvili, 2001). The term molecular recognition is used here as an amalgam of all these factors.

A detailed analysis of the enthalpic and entropic effects involved in molecular recognition requires extensive molecular dynamics simulations using more realistic force fields than those presently available (Becker et al., 2001). In their absence, the analysis of similarities can improve our understanding of molecular recognition by highlighting those structural elements that have the largest effect on the interactions between small molecules and proteins, thus providing a focal point for more detailed studies.

The extent to which similarities between non-homologous proteins, in terms of their overall sequence and structure, are related to similarities between their cognate ligands (substrates as well as cofactors) has been studied recently (Mitchell, 2001; Nobeli et al., 2005). Nobeli et al. (2005) report that a correlation between protein and ligand similarity can only be clearly established for very similar proteins. However, from an evolutionary point of view, while a pair of protein sequences might have diverged enough to be classified as non-homologous, the conservation of similarity between their cognate ligands constitutes evidence of homology between them.

A growing body of experimental evidence (Copley, 2003; O’Brien and Herschlag, 1999, Shears, 2004) suggests that catalytic promiscuity is integral to the function of various proteins. In recent years, the notion of binding promiscuity has also received support from computational docking studies (Koehler and Villar, 2000; Macchiaulo et al., 2004) where it has been observed that the complex formed between a given protein and its cognate ligand rarely is the most stable.

We use experimental data on the thermostability effect due to the binding of a small molecule to a protein as a measure of the strength of binding between the small molecule and the protein. The same data can be seen as (1) a set of small-molecule binding profiles to the proteins studied and (2) a set of protein binding profiles for each small molecule. In principle, unlike Nobeli et al. (2005), the set of ligands is not connected evolutionarily to any particular protein. Consequently, the binding patterns observed are not the result of any evolutionary optimization process.

The analysis of similarities in small-molecule binding profiles between members of a given family offers the possibility of comparing binding site similarities to small-molecule binding profile similarities within the evolutionary framework (i.e. overall sequence evolution) of the given family. Likewise, the comparison of small-molecule topological similarities and protein binding profiles can shed light in the relationship between structural similarity and molecular recognition from the point of view of small molecules.

The family of human cytosolic sulfotransferases (SULT) is involved in drug metabolism, detoxification and hormone regulation. The enzymatic reaction involves the transfer of a sulfuryl group from a donor molecule to a hydroxyl group in the acceptor
molecule. In general the sulfuryl donor is 3'-phosphoadenosine-5'-phosphosulfate (PAPS). Substrate promiscuity is integral to the function of SULTs, particularly in detoxification and drug metabolism where the addition of the sulfuryl group increases the water solubility of the substrate thus facilitating its excretion. SULTs belong to the Rossmann fold group of proteins within the αβ class composed of a five-strand β-sheet flanked by a series of α-helices. The cofactor and substrate binding site lie on one side of the β-sheet with several loops forming a binding site ‘trapdoor’ that interacts with both cofactor and substrate. Trapdoor loop residues have been shown to be responsible for substrate specificity (Coughtrie, 2002). Various residues in the sheet interact with the substrate and some are essential for catalytic activity (Chapman et al., 2004; Glatt and Meinl, 2004).

2 METHODS

In the present study, we analyze a subset of selected members of the human cytosolic sulfotransferase family. Table 1 gives some information about the structures used.

<table>
<thead>
<tr>
<th>Name</th>
<th>PDB code</th>
<th>Cleft Size</th>
<th>Ligands</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A1</td>
<td>1is6A</td>
<td>1.90</td>
<td>194 (51)</td>
</tr>
<tr>
<td>1A3</td>
<td>2a3rA</td>
<td>2.60</td>
<td>177 (49)</td>
</tr>
<tr>
<td>1B1</td>
<td>1xv1A</td>
<td>2.10</td>
<td>181 (51)</td>
</tr>
<tr>
<td>1C1</td>
<td>1hehA</td>
<td>2.22</td>
<td>163 (43)</td>
</tr>
<tr>
<td>1C2</td>
<td>2ad1A</td>
<td>2.00</td>
<td>21 (21)</td>
</tr>
<tr>
<td>2A1</td>
<td>1f6hA</td>
<td>2.40</td>
<td>275 (72)</td>
</tr>
<tr>
<td>4A1</td>
<td>1zd1B</td>
<td>2.24</td>
<td>106 (32)</td>
</tr>
</tbody>
</table>

*The names of SULTs follow a scheme related to their pairwise sequence identity.

*Three-letter PDB codes are followed by the one-letter chain code of the subunit used in the study followed by the resolution in Ångstroms.

*Number of atoms in cleft (number of residues contributing atoms to the cleft).

*Relevant ligands present in the binding site. A3P (adenosine-3',5'-diphosphate), pNT (p-nitrophenol), LDP (L-dopamine), PCQ (3,5,3'-tetrachloro-biphenyl-4,4'-diol), NHE (N-(3-cyclohexyl)taurine), PLO (pregnenolone) and PAP (3',5'-phosphoadenosine-5'-diphosphate).

*Values for the lower and upper bound thresholds, respectively. In what follows, maximum distance difference in defin-

*Structures purified and solved by the Structural Genomics Consortium (SGC), to be published.

*The crystal contains an unknown Escherichia coli metabolite sequenced during expression.

2.2 Binding site sequence similarity

We created a multiple sequence alignment of the sequences of the proteins used in this study using HMMer (Eddy, 1998) and the pfam (Bateman et al., 2004) Sulfotransferase_1 (PF0685.15) hidden Markov model. Sequence similarity is measured using the Tanimoto coefficient (Gasteiger and Engel, 2003). The residues defining the cleft are mapped onto the sequence alignment and used to define a local Tanimoto score of sequence identity (LTSseq):

$$LTS_{seq} = \frac{N_{comm}}{N_{total} - N_{comm}}. \quad (1)$$

where $N_{comm}$ represents the number of columns in the alignment where both sequences under comparison contain cleft residues of the same type and $N_{total}$ represents the sum of the total number of cleft residues in both proteins. The use of LTSseq is a natural choice, since there is no straightforward way to define a sequence overlap to be used as normalization factor when comparing binding site residues, as these can be spread widely in the primary sequences of the proteins under comparison. For global sequence comparison, a global Tanimoto score of sequence identity ($GTS_{seq}$) is defined using all columns in the alignment (except those containing gaps for both sequences).

2.3 Binding site structural similarity

Given the sets of atoms defining the clefts under comparison, the question that needs to be answered is what is the largest subset of atoms in both clefts in direct correspondence with each other geometrically as well as chemically. This is a combinatorial optimization problem where, in principle, each possible set of atom correspondences might be a solution and the largest such set is the global solution. Graph theory offers a means to solve this problem via the detection of the maximal (largest) clique in an association graph. Further details on graph theory can be found elsewhere (Gross and Yellen, 2004). In the present work, we use the standard algorithm of Bron and Kerbosh (1973) for the detection of cliques.

Depending on the number of atoms being compared, the size of the association graph might make it practically unfeasible to detect the largest clique when considering all non-hydrogen binding site atoms. In order to overcome this difficulty we perform the graph matching in two stages.

In the first stage, an initial superimposition is performed via the detection of the largest clique in an association graph constructed using only $C_{\alpha}$ atoms of identical residues in the two clefts. A maximum distance difference of 2.0 Å is used to create edges in the association graph, imposing an upper bound of the same magnitude in the coordinates root mean square distance (RMSD) of corresponding $C_{\alpha}$ atoms.

Once the largest $C_{\alpha}$ clique is obtained its transformation matrix and translation vector are used to superimpose all atoms in the two clefts using the least square method of Arun et al. (1987) based on the singular value decomposition of the coordinates variance-covariance matrix.

In the second graph matching stage, all non-hydrogen atoms are used. Association graph nodes are created with the requirement that two atoms, one from each cleft, be of the same atom type as well as that their spatial distance be within 1.5 Å. This spatial distance constraint is used to decrease the size of the association graph and is the reason why the initial superimposition is performed. In the present work, we use eight atoms type classes (Sobolev et al., 1996, 1999) comprising the following classes: hydrophilic, acceptor, donor, hydrophobic, aromatic, neutral, neutral-donor and neutral-acceptor. Similar to the first stage, a maximum distance difference in defining association graph edges is used. This second threshold is set to 1.5 Å and again defines an upper bound of that magnitude in the RMSD between corresponding non-hydrogen cleft atoms.
Figures and tables have been added as per the requirements. The text has been rendered in a readable format, including the conversion of tables and equations as per the guidelines.
point represents one pairwise comparison. In this respect, it is important to note that the use of a carefully curated hidden Markov model to align the sequences is far more stringent than the use of pairwise sequence alignment methods, thus adding more significance to the success in locating equivalent residues through the pairwise structural comparison. The observations that do not fall on the diagonal involve primarily SULT4A1 and are due to local backbone movements (loop movements).

Figure 2 shows the dendrograms of the clustering of the SULT proteins in terms of their binding site sequence similarity (LTSseq) or all-atom binding site structural similarity (LTS3D-All). The dendrograms are very similar but that of LTS3D-All better represents the structure within the data ($\rho = 0.992$) than LTSseq ($\rho = 0.977$). Table 3 contains a summary of the correlation between the various similarity measures of members of the SULT protein family.

### Table 3. Correlation coefficients between SULT similarity measures

<table>
<thead>
<tr>
<th></th>
<th>LTSseq</th>
<th>LTS3D-Ca</th>
<th>LTS3D-All</th>
<th>SmBP</th>
</tr>
</thead>
<tbody>
<tr>
<td>GTSseq</td>
<td>0.829</td>
<td>0.826</td>
<td>0.710</td>
<td>0.570</td>
</tr>
<tr>
<td>LTSseq</td>
<td>0.992</td>
<td>0.938</td>
<td>0.386</td>
<td></td>
</tr>
<tr>
<td>LTS3D-Ca</td>
<td>0.966</td>
<td>0.406</td>
<td></td>
<td></td>
</tr>
<tr>
<td>LTS3D-All</td>
<td>0.334</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Another question that can be inspected through the analysis of LTSseq and LTS3D-All is that of the evolution of the binding site vis-à-vis that of the overall sequence within the family. In Figure 3 we present a clustering of the SULT protein sequences according to their pairwise global sequence similarity coefficients (GTSseq).

The same plot shows the pairwise LTSseq similarity coefficients in the lower half matrix (below diagonal elements) as a heatmap. The correlation between GTSseq and LTSseq or LTS3D-All is 0.829 and 0.710, respectively (Table 3), suggesting that binding site similarities are not well represented by overall sequence relationships. The average values of GTSseq, LTSseq and LTS3D-All are 0.30, 0.24 and 0.21, respectively, showing that binding site sequences are less conserved than overall protein sequences within the SULT family as binding site diversity brings about the diversity required for the specific function of the various members of the family.

### 3.2 All-atom comparisons and the effect flexibility

Side chain and backbone movements can have a strong influence on the atomic correspondences found through structural comparisons.
While a more relaxed choice of parameters used for the structural comparison can somehow account for the effects of flexibility, a balance must be struck between the increased computational resources required (time and memory) to solve the graph matching problem and the actual resources available.

The comparison of specific LTS3D-All and LTSseq values (symmetric elements with respect to the diagonal in Fig. 3) does not show a clear tendency; in some cases, LTS3D-All is larger, in others, it is smaller. One major difference between LTS3D-All and LTSseq is seen in the pairwise comparisons involving SULT4A1, where LTS3D-All values are much smaller than LTSseq. The reason for this difference can be rationalized by the fact that the majority of the comparisons involving SULT4A1 shows large values of LTSseq.

The same ΔTagg values presented in Table 2 can be analyzed from a small-molecule perspective. That is, for a given ligand one can utilize its pattern of binding to the set of proteins as a combined measure of the effect that particular characteristics of the small molecule affect its interactions with the set of proteins. Such protein binding profiles (PBP, rows in Table 2) can be used to cluster the set of small molecules to compare the resulting dendrogram to the matrix of pairwise ligand topological fingerprint (TFP) similarities (Fig. 5).

From a small-molecule point of view, it is again difficult to rationalize the binding profiles in terms of small-molecule structural similarities (0.253 correlation). Several but not all small molecules with similar binding profiles tend to show high degree of structural similarity.

4 CONCLUSIONS

The method for the detection of local structural similarities and the Tanimoto coefficients of sequence and structural similarity developed here are able in conjunction to detect binding site sequence and structural similarities between members of the human cytosolic sulfotransferase family.

Binding site sequence and structural comparisons uncovered similarities between 2A1 and members of the SULT1 subfamily not seen through overall sequence comparisons.

Proteins with similar small-molecule binding profiles show binding site sequence and structural similarities but the opposite is not true, namely, binding site similarity is not sufficient to predict the pattern of binding of different ligands to the given protein. From a computational point of view, this fact suggests that the accurate
prediction of cognate ligands from structure might not be possible until a better understanding of the process of molecular recognition is reached. From a pharmaceutical point of view, it is assuring that even similar binding sites show significant differences in the way they bind the same molecule as this diminishes the chances that a drug developed for a specific protein will have the same effect on related proteins.

Conversely, from a small-molecule perspective, a similar situation occurs. Namely, small molecules that bind with a similar pattern to a series of proteins tend to be topologically similar but topological similarity in itself is not sufficient to predict binding patterns. This observation suggests that we do not have a proper metric with which to gauge the effect that small topological differences can have in the binding pattern of a small molecule. While even related small molecules may bind to proteins in different orientations or with otherwise different interacting parts of their scaffolds, this deficiency of topological similarity as a metric is significant given its common usage in the selection of representative molecules for the creation of libraries in drug design, potentially leading to the oversight of important molecules.

Ultimately of course, we need accurate methods to calculate ΔG values associated to the binding of small molecules to proteins, but in their absence, simple similarity-based scores provide clues that can help cluster small molecules and protein according to their binding profiles.

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**REFERENCES**


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Fig. 5. Clustering of small molecules according to their protein binding profiles (PBP, r = 0.775) compared with pairwise topological hashed fingerprint similarity (TPF). In this case, symmetric elements with respect to the diagonal are identical.